SEM LEORIGINAL

THE RIGHTS OF HEARING-IMPAIRED PERSONS

CCR 3 Meet

A Public Forum

Illinois Advisory Committee
U.S. Commission on Civil Rights

In Cooperation With

The Section of Individual Rights and Responsibilities of The American Bar Association and The State of Illinois Department of Human Rights

> Hilton Hotel and Towers Williford A Room Chicago, Illinois

Monday, June 30, 1986 8:30 a.m.

LONGORIA & GOLDSTINE
CERTIFIED REPORTERS
176 West Adams Street
Suite 2232
Chicago, Illinois 60603

(312) 236-1030

## BEFORE:

MR. HUGH J. SCHWARTZEERG Chairman

MS. THERESA F. CUMMINGS

MR. ROBERT C. SPENCER

MR. ISIDRO LUCAS

MR. JOHN LINGNER

MS. ERMA M. DAVIS

MR. THOMAS PUGH

MR. PRESTON E. EWING

J

-

CHAIRMAN SCHWARTZBERG: Good morning.

My name is Hugh J. Schwartzberg, and I am

the Chairperson of the Illinois Advisory

Committee to the United States Commission

on Civil Rights.

Today the Illinois Advisory

Committee is holding a public forum to hear presentations from experts of the rights of hearing impaired persons.

Members of the Illinois

Advisory Committee present with me today are

Ms. Theresa Cummings from Springfield,

Illinois, Mr. Preston Ewing from Cairo,

Mr. John Lingner from Chicago, and there

will be present later during the day at

various times, because some of the

Commissioners will be here at different

portions of the presentations, Ms. Erma Davis

from Peoria, Mr. Thomas Pugh, also from

Peoria, Mr. Herschel Seder from Highland Park,

Dr. Robert Spencer from Petersberg, and

Ms. Joyce Tucker from Chicago.

Also with us today are the following staff members of the Midwest Regional

Office of the United States Commission on Civil Rights in Chicago, Mr. Clark G. Roberts, who is the Regional Director; Dr. Isidro Lucas, the Deputy Regional Director, and who serves as staff to this commission; Mr. Carmella Melendez, Mr. Frank Alford, and Ada L. Williams.

At this time we want to express officially our thanks to the various organizations that have offered us their assistance, and with whose cooperation we are holding this forum.

The section of individual rights and responsibilities of the American Bar Association, the Illinois Department of Human Rights, and the Alexander Graham Bell Association for the Deaf.

This public forum is being held pursuant to the rules applicable to the state Advisory Committee and other law and regulations relating to the United States Commission on Civil Rights.

I would like to emphasize that this is an open meeting, and the cooperative

efforts involving the Illinois Advisory

Committee, the organizations already

mentioned and the experts making presentations,

written and oral, from Illinois and the rest

of the country and Canada.

In particular, this forum is also held on the explicit advice of the United States Commission on Civil Rights and its chairman.

Those participating have voluntarily agreed to appear before the Committee, and to share their expertise with us.

The meeting is open, and the media and all interested individuals are welcome to attend. We have been asked about taping these proceedings, and because this is an open meeting, taping is, of course, permitted.

The experts will share their information with the Committee, and will respond to questions and engage in discussions with Committee members. Public officials and private citizens not on the agenda are

welcome to present written statements for consideration of the Committee and for inclusion in the record. Such written materials can be given to the staff person at the registration desk outside of this room. They can also be mailed before the 20th of July, 1986 to the United States Commission on Civil Rights, 230 South Dearborn, Room 3280, Chicago, Illinois 60604.

We are concerned that no individual or specific organization be the victim of defamatory or degrading statements. In the event that statements may develop, it will be necessary for me to call this to the attention of the person making the statement and request that he or she desist in that action.

Furthermore, any person against whom such allegations are made will have ample opportunity to make a statement in response as appropriate.

There is legislation at the local state and federal level aimed at securing the civil rights of handicapped

persons. This Committee wants to know how this legislation and supplementing registrations and endorsements are being applied today to persons whose handicap is hearing impairment.

The Advisory Commission on Civil Rights of citizens of this forum will report to the Commission. A transcript of the proceedings is being made and will be available at the Commission's offices.

Civilizations are judged by their strengths and by their most powerful exemplars, but they are also judged by the way in which they treat those who are the weakest of their citizens.

In the normal course, one acquires language through one's ears, and so for millenia, even growing up within communities, the deaf child remained mute and did not acquire the power to speak, and was thereafter often treated as being less than human. The operative word, no longer appropriate, was dumb.

For the deaf, if the community,

and because of the rarity of this condition and the cost of its cure, community here means the state. If the community does not intervene with diagnosis, amplification, training and education, language itself is denied, and this process must begin very shortly after birth in order to be very effective.

There is therefore a potential claim by those physically handicapped, those hearing impaired, as against the state, a claim that the majority are bound to act so as to supply language itself in light of the probability that if the state does not so act, the potential of that human being will be destroyed by social inaction.

This claim may be grounded in revulsion against the unnecessary wasting of a potential human mind, it may be rooted in our moral queasiness at the loss of any one human's reasonable potential, it may be colored by consideration of the social costs of inaction, such as the cost

to the public of what may otherwise become a public charge.

This right may be claimed even in the absense of legislative action, although in fact much legislation does in fact exist.

Certainly none of the rights retained by the people will blossom into fruit unless they are watered with knowledge of why they are claimed.

What are the rights of the deaf? What can the hearing impaired claim as against an unthinking or even contrary majority? What claim against the state does the hearing impaired hold as a matter of right, and what should be offered by legislative grant? Where does the law stand now? We will be looking at some of these questions today.

There are some minor house keeping details. In the program, there is a misprint on the first page, and after Dr. Ling's presentation, it says that the next section will start at 11:00 o'clock a.m.

That is not true. The section initiated by Dr. Ling will continue with Drs. Northcott, McCartney and Conway, and the 11:00 o'clock section will start with Ms. Celia Warshawsky on page two.

Now I would also like to point out that the Commission has very lengthy, very lengthy papers, and many of those which we have received are of obvious quality. For the purpose of this session, we have asked their office to summarize the written presentations in ten-minute segments.

The fuller text will be available at the Commission's offices in the Federal Building here in Chicago.

At this time, and to start the forum, I have the pleasure to introduce to you Dr. William Castle, the Director of the National Technical Institute for the Deaf.

## Dr. Castle?

DR. CASTLE: Thank you, Hugh.

I would like to introduce the persons who are assisting me in the

ı

presentation of this first lecture.

We have Mickey Baron-Gerstein doing the sign-languange interpreting, we have Mary Boone doing the oral interpreting, as the day moves along others will be taking their place, and I would like to introduce each of them as well.

We have over here Lynell Weddington, who will also do some sign-language interpreting.

In front of me we have

Colin Freeman Rizla, and way to the left also

Carol Convertino, all of whom will be doing

oral interpreting, and we have two young

ladies in the front row here as well,

Jan Krefft and Rita Tacona, who will be doing

whatever reverse interpreting is required

for today's session.

It is indeed a pleasure for me to have been invited to open this today, this day-long meeting on the rights of the hearing impaired.

It is a topic that we clearly owe a great deal of attention. The subtopic

I have been asked to address, as you see by the program, is the meaning of "reasonable accommodations" and the hearing impaired.

Fortunately, the program also says that my addressing this subtopic is to be a state-of-the-art report.

tell, no easy dictionary-like definition to provide a meaning for what is referred to as reasonable accommodations.

Hopefully, therefore, my state-of-the-art report will lend meaning or means to that phrase.

As my report ensues, you will discover that, for the most part, I used the generic terms, hearing impaired or hearing impairment, which includes both the deaf and the hard of hearing. At times, however, I use the words deaf and deafness.

I wish to clarify that when

I do use the words deaf or deafness, they

are always used to refer to both severe and

profound hearing impairments, that is hearing

losses that are 70 decimals or more in the

better ear.

A great deal has happened in recent years to impact on the rights of the hearing impaired. One might say that at least in the United States hearing-impaired persons have never had it so good. This is in large part due to the fact that a great deal of legislation has been put in place in the last two decades to make a difference in their lives, especially by broadening their base of options, and hopefully their freedom of choice.

The National Technical
Institute for the Deaf Act was passed in
1965 to complement what occurs for the deaf
at Gallaudet College.

A 1968 amendment to the

Vocational Education Act provided that 10

percent or more of the funding given to any

state in the union be used for the handicapped.

The Model Secondary School for the Deaf Act was passed in 1966. The Bureau for the Education of the Handicapped, which is now known as the Office of Special

Education Programs was established in 1967.

The Captioned Films in Media for the Deaf

Program have been in existence for a long

time, and now includes considerable funding

for captioned television.

The amendments to the

Vocational Rehabilitation Act of 1973

provide for easier access for all handicapped

persons to post-secondary education, and

to a large variety of employment opportunities

through its Section 504.

The Education of All
Handicapped Children Act, otherwise known
as Public Law 94142 was passed in 1976.
The interpreter for the Deaf Training Act
was passed in 1978.

There is now also in the Education of Handicapped Act a provision for special funding for post-secondary education programs for the handicapped, four of which must be programs for the deaf.

All of these federal laws have their impact on what happens today for

3

5

4

6

8

7

9

10

11

13

14

15

16

17

18

19

20

21

22

23

24

hearing-impaired people and their rights.

Federal law also now requires that 10 percent

of all telephones in every hotel or motel

be compatible with the special needs of the

hearing impaired.

Now pending in Congress are special bills designed first to amend the Communication Act of 1934, both to expand the number of public telephones that are compatible with hearing aids and special equipment for telephone service for certain disabled persons; number two, to amend the Internal Revenue Code of 1954 to exempt the excise tax for telephone facilities designed for the hearing impaired; to provide a deduction for tv captioning equipment for the hearing impaired, and to allow extra deductions for certain disabled tax payers; three, to amend Titles 8, 18 and 19 of the Social Security Act to provide an optional part C insurance program for coverage of certain vision, hearing and dental services, and prescriptive drugs, and to treat certain sensory and communication aids as medical

or other health services; four, to amend the Court Interpreter's Act of 1978 to better provide for the hearing impaired in Court; five, to establish a pilot program of fellowships in the house of Representatives for persons who are hearing impaired and blind; and six, to provide for better quality education of deaf programs throughout the country through the newly-passed Education of the Deaf Act of 1985.

Through Friday's newsletter for this convention, we learned that the Chicago City Council unanimously approved a bill that requires all hotels and motels in the City to provide visual or vibrating fire alert systems for the safety of their hearing impaired guests.

Five of our fifty states have similar legislation, and such legislation is now pending in New York State.

18 of our states have commissions on deafness. There is no doubt that our hearing impaired population has it better than any other such population in

the world.

However, we cannot afford to rest on our laurels. Our hearing impaired people may have it good, but they could have it much better.

I will spend the major part of my time talking about how they might be even more reasonably accommodated than they currently are.

We are all acutely aware that legislation is one thing, fulfillment of that legislation is another. It seems safe to say that although our nation always has had an abiding interest in equal educational opportunity for its citizens, the fulfillment of that interest has not been something easily effected, because the makeup of our citizenry is very diverse.

The educational programming among today's 50 states is also very diverse, and among the multitude of local school districts within each of those states educational programming is very diverse.

Complications of the fulfillment

process have been the peculiar history of dealing with American Indians from the time of colonization. The uncertainty about the role of women as citizens from the time the United States Constitution was written, the awarding of citizenships to our black population, who, like their ancestors, at that time were first here as slaves.

The diversity of the nation's immigration program and the recent influx, legal and illegal of spanish-speaking Americans from Cuba, Puerto Rico, and Mexico other countries or territories, and of Asians from such places as India, Pakistan, Vietnam, Thailand, Korea and Taiwan, our inability to recognize the American Indians as real citizens persists, but so do their demands for equal rights.

Though women were assured the right to vote in 1920, they still strive today for the Equal Rights Amendment to the Constitution to give full assurance of equal educational and employment opportunities.

Though the blacks and other

minority groups were assured of citizenship in 1868 and the right to vote in 1870, they have had to demand special civil rights legislation in recent years to foster equal educational housing and employment opportunities for themselves.

Inspite of all that special legislation, they are still a far cry from where they would like to be regarding their equal rights, educational and otherwise.

We must heed this history and must assume that an important part of our interest on this very day is to have a different kind of history written for the hearing impaired, who have only recently been given their legal rights to equal education and employment opportunities.

The Education of All
Handicapped Children Act of 1976, that is
Public Law 94142 is the one piece of
legislation that has the greatest impact on
what happens for hearing impaired children
and youth today.

A second piece of legislation

that impacts significantly on hearing impaired youths and adults is the Vocational Rehabilitation Act of 1973.

In a sense, these two pieces of legislation are the civil rights bills for hearing impaired and other handicapped people.

They, therefore, deserve and receive a great deal of our attention today, and they merit a great deal of respect for what they were designed to accomplish.

It is because of them that
we have heard with frequency such things as
Section 504, Equal Educational Opportunity,
Equal Employment Opportunity, Equal Access,
Least Restrictive Alternitive, Least
Restrictive Environment, and Reasonable
Accommodations.

At the same time, these two pieces of legislation fall short of solving some of the current and future problems affecting hearing impaired people.

Public Law 94142 declares that many handicapped children do not have

successful educational experiences because their handicaps go undetected, but that law does nothing to provide for programs for early detection.

It does not make provision for handicapped children between the ages of 3 and 21, but it's mandate covers only the ages between 5 and 18.

Like other handicaps, hearing impairment in children often goes undetected. It's not a visible handicap and is too often discovered later than it should be for the undertaking of very necessary early intervention.

Mandating attention to children who are 5 through 18 years old is fine, but it is clearly not enough with respect to the hearing impaired, who, along with their parents, should receive special educational attention as soon as the hearing impairment is detected, preferably at birth or shortly thereafter.

Our country needs to put in place, as soon as it can, a universal system

Ī

of early detection for hearing impairment.

Though the United States is ahead of all other nations in what it does for the hearing impaired, it lags behind some other nations in our world in its efforts in early detection.

Audiological techniques have been developed which can do what is needed, but they are not applied universally.

Medical doctors, especially gynecologists, obstetricians and pediatricians, need to be more universally aware of these techniques and see that they are applied, because the earlier hearing impairment is known the earlier special programs and reasonable accommodations can begin.

The lack of the earliest

possible detection is the first strike

against equal educational and employment

opportunities for many hearing impaired

individuals in our country. It is therefore

the first strike against their civil rights.

Our country also needs to establish a universal program of early

education for hearing impaired children and their parents.

For some years now, several programs for hearing impaired children under the age of three and their parents, often referred to as parent-infant programs, have been in existence. Such programs are important for the adjustment on the part of both the parent and the child with respect to the problems associated with severe or profound hearing impairment.

They provide a very early opportunity for parents to become involved in the educational programming and reasonable accommodations for their children.

However, such programs are not available universally in our nation, and they should be. The absence of such programs is frequently the second strike against equal education and employment opportunities for many hearing impaired individuals, and therefore a second strike against their civil rights.

The full meaning of early

1 2 3

,

education includes what is done at the preschool for hearing impaired children who are between the ages of three and six.

Over the years, specifically since World War II, our nation has done a fair job of providing preschool educational opportunities to hearing impaired children, especially deaf children.

It is still not a universal opportunity, but it is a fairly prevalent one.

Good quality preschool education for the deaf is also not universal, because not all teachers involved are adequately trained to do the job, and because important support services may often be lacking.

For example, reasonable
accommodations may not be available from
speach pathologists, audiologists or
itinerant teachers skilled in the education
of the deaf and the use of manual communication,
if that be needed.

Further, one cannot ever say enough about the importance of the earliest

\_

possible use of amplification to optimize the use of whatever residual hearing a hearing impaired child may have.

either about the importance of providing the earliest possible opportunity for a hearing impaired child to learn how to speak the language of his native country, because in the long run of that child's life, there is probably no better basis for becoming mainstreamed as a productive adult in one's society and meeting the challenges of upward mobility in employment than having understandable language, both spoken and written, good speach reading skills, and the ability to make optimal use of residual hearing.

Development of the written and spoken language skills of hearing impaired children should be the focus of all early childhood intervention, using whatever communication modalities are useful or necessary.

It was therefore a great

pleasure to learn recently that technical amendments to Public Law 94142 were introduced to the United States Senate on April 14 of this year, which provide for an extension of that law's mandate to age three within two years, and the creation of a new discretionary grant program for early intervention of handicapped children from birth through age two.

Public Law 94142 seeks to provide appropriate educational experiences for handicapped children at the elementary and secondary levels.

has been in place for ten years, it is still a fact that regardless of whether they receive their elementary or secondary education from residential schools or from some form of mainstreaming, the deaf students among our hearing impaired in general do not have elementary and secondary attainments which equal those of their hearing peers.

Inspite of all the dollars that have been spent on education of the deaf

2 f

over the past 30 years, the average reading level of deaf graduates from secondary education programs remains bearly over the fourth grade level.

an elementary or secondary program in a school for the deaf is taught by teachers of the deaf who are not teachers of elementary and secondary subject matter or content, and the usualy deaf student in some mainstreaming format for elementary or secondary education is taught by a teacher who is qualified to teach elementary or secondary subject matters, but who is not qualified or knowledgeable about how to teach deaf students.

Thus our usual deaf student is unexposed to a great deal of what might be called fundamental knowledge.

These dynamics indeed can hardly be thought of as reasonable accommodations of the deaf childrens' needs.

A commitment to genuine implementation of Public Law 94142 for the

elementary and secondary levels of education of the deaf will require a drastic but positive revision in the teacher training programs designed to train persons to teach the deaf.

Preschool, elementary and secondary teachers of the deaf should all be certifiable by state departments of education, both as qualified teachers of the hearing and as qualified teachers of the deaf.

Unless that becomes true, we will not begin to make the headway that we need to in those levels of education of the deaf.

Public Law 94142 expresses

the fact that families of handicapped

children in youth are too often forced to

seek special educational services for them

outside the free public school system,

frequently from agencies that are a great

distance from their residence and frequently

also at their own expense, and hence most

people interpret that law to mean that all

handicapped children should and must be mainstreamed in their local public school district.

I believe this interpretation leads to restricting freedom of choice from among programs that are available.

If implementation of Public Law 94142 were to lead to the immediate or eventual demise of all residential schools for the deaf, public or private, this would be a grave error, and a severe restriction of freedom of choice, and therefore a restriction of civil rights.

Not all hearing impaired persons wish to be mainstreamed during every aspect of their education, employment and community living, and some may wish never to be mainstreamed.

Not all parents of hearing impaired children wish to have their children mainstreamed at every point of education, and some may rightfully believe that the least restrictive alternative for their child is a residential school for the

deaf, and even a residential oral school for the deaf.

Therefore today we see signs of backlash from the deaf community and from educators of the deaf against the way Public Law 94142 is being interpreted.

There are many voices of protest against what is called Manual Ten, a document which describes the principles to be used in deciding the least restrictive alternative for a given handicapped child.

That manual, like Public Law 94142, describes a continuum of alternatives which places the regular classroom on the continuum at the point of least restrictive and the special school at the point of most restrictive.

The backlash contention is that for many deaf children the special school is actually the least restrictive.

It is foolhardy to think that every school district in the United States can provide all that is needed to all handicapped children residing in that

district in order for them to be properly educated.

This cannot at any time be a cost-effective or cost-efficient way to accomplish mainstreaming.

Districts must pool their resources.

When Public Law 94142 was passed in 1976, its proponents declared that the educational needs of handicapped children were not being met adequately, and that at least 50 percent do not receive the reasonable accommodations required to assure them full equality of educational opportunity.

With respect to the hearing impaired, the percentage was higher at that time, and it continues to be so for the several reasons already cited.

When it comes to dealing with all the reasonable accommodations that are required for the hearing impaired, it's important to realize that reasonable accommodations require a greater prevalence of audiologists, special pathologists,

language specialists, school psychologists and personal and career guidance counselors at all levels of education of the deaf.

For too long it has been the practice to have the classroom teacher of the deaf, particularly in residential schools for the deaf, be all things to all deaf students.

In this day and age this is no longer appropriate, and it probably never was.

At NTID, for instance, we know that 90 percent of the deaf students who enter our programs can profit from the use of amplification, but 80 percent do not know how best to make use of amplification.

Nearly 35 percent of the students who can profit from the use of hearing aids either do not own aids or seldom wear the aids that they do own.

Even though the average age of deaf students entering NTID is 19 and-a-half, and one would expect the students to be fixed in their ways, we have discovered

at NTID, one, that with the right approaches to orientation to the hearing aids by audiologists, non-users and seldom-users alike can both be pursuaded to become users.

Two, that with the right approaches by audiologists to auditory training, poor users of the hearing aids can be taught to be good users, poor listeners can be taught to be better listeners, and poor auditory discrimination skills can be improved.

Three, that with an approach by audiologists to speach reading which concentrates on repetitious social interchanges and job-related vocabulary and language, speach-reading abilities can be greatly enhanced.

If these things can happen at the ages of 19 and 20, is it not a reasonable accommodation to see that they happen earlier?

The Vocational Rehabilitation

Act of 1973 is still another matter. It

insists that handicapped persons who are

UNITER

otherwise qualified cannot be denied employment or admission into educational programs for reasons of their handicaps, so long as the employment or educational institution is a recipient of federal subsidy.

It also suggests strongly that employment environment, education environments, and social environments, such as theaters and churches, do all that they can to provide reasonable accommodations for handicapped persons, for the deaf and the hard of hearing, therefore there are many efforts today to remove significant communication barriers inductance loop systems, fm systems and infrared systems are prevalent in theaters, churches and other public and private meeting places for those who depend on auditory amplification.

films, many captioned television programs, and even captioned live theater, including opera, for the specific accommodation of

•

hearing impaired persons.

Interpreters of the deaf, including oral, simultaneous and American sign language interpreters are prevalent in many social and educational environments and on television.

Deaf persons also enjoy great use of telephones these days, because of the multitude of teletype devices that are on the market and special techniques that have been designed for deaf persons with good speach and language to use telephones equiped with special amplifiers.

Inspite of all the good intentions of this act, it is proving to be counterproductive in some ways, especially in the realm of post-secondary education for the deaf.

There has been a net growth

from 27 post-secondary programs for the
hearing impaired to 145 in the last 13 years.

Many of those programs do not begin to
comply with the principles proposed in 1973

by the Conference of Educational Administrators

2

3

5

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

Serving the Deaf.

The first evidence of this fact is that only 63 programs, that is 43 percent, of the 145 that are listed in college and career programs for deaf students, 1976, are granted full program description, since such descriptions were provided only for those institutions which, one, have at least 15 full-time hearing impaired students enrolled in the post-secondary educational program for deaf students, two, are part of an acredited post-secondary institution, three, have a coordinator of services for deaf students who devote a minimum of 50 percent of his or her time in directing that program, and four, generally comply with the principles proposed by the Conference of Educational Administrators.

A second evidence is that the percentage of new students who are transfers to NTID has been increasing, and the admissions recruitments staff at NTID indicate that transcripts of 83 percent of these students show that no credits have

.

been earned, even though they spent an average of one and-a-half years attending another college.

that over 39 percent of the programs listed, all those with ten or fewer full-time deaf students, provide little or no special services to the hearing impaired students, other than interpreting and tutoring, and 18 percent of those programs provide no interpreters, while 24 percent provide no tutoring.

Among those programs that
have between 11 and 20 full-time hearing
impaired students, another 22 percent have
all the programs, 3 percent provide no
interpreting, 17 provide no tutoring, 29
percent provide no special counseling from
counselors who can communicate directly with
hearing impaired people, 55 percent have no
special classes for hearing impaired students,
52 percent have no paid note takers, and
45 percent have no vocational counselors
who can communicate directly with the hearing

impaired students.

or more full-time students, that is 24 percent of all the programs listed, can it be seen that 100 percent use paid interpreters, 100 percent use tutors, 90 percent have personal and vocational counselors who can communicate directly with hearing impaired persons, even then 45 percent have no special classes for hearing impaired students with teachers who can communicate directly with those students, have no paid note takers, and provide no direct speach and hearing services.

These data are considered by those who pull them together to be conservative, to say the least. That is, the situation regarding reasonable accommodations at the post-secondary level should be considered worse than these data suggest.

Also, a survey of the transfer students at NTID from 1980 through 1984 suggests that these students were disappointed

in the support services they received elsewhere, and they felt that the teachers in many cases were insensitive to their needs.

For example, the teacher talked with his or her back to the class or didn't make use of mediated materials and so on.

These data are offered because they are they are suggested and not because they are absolutely accurate, and what they suggest are the following things; one, there is indeed a proliferation of secondary programs for the hearing impaired which are not quality programs.

Section 504 of the Vocational Rehabilitation Act of 1973, with all of its good intents, has played some tricks on us.

Its insistance that hearing impaired persons who are otherwise qualified for entry to any given post-secondary program cannot be denied entry to the program has led to a very interesting irony, that is, any post-secondary program which is not

qualified for handling the special needs for the reasonable accommodations for hearing impaired students cannot at this point in time be denied the right to admit those students.

Thus many institutions are giving it a try and failing in the process, because they do not provide the reasonable accommodations that are needed.

Number two, the open or liberal admissions policies of many colleges and universities allow for many hearing impaired persons to enter their doors who are not truly qualified to be there, because lo and behold, by policy, they cannot be kept out.

This may be sheer folly in light of the fact pointed out before that inspite of at least 30 years of trying to make it different, the average reading level of secondary school leavers who are deaf is still only at fourth grade.

Three, the propensity of state agencies to force their vocational

rehabilitation counselors to insist that
their clients must remain in their home
states, and sometimes even in their near-home
community college, because it's less costly
for them to go to college, is most certainly
taking its toll on many young hearing
impaired people.

While Public Law 94142 clearly, and Section 504 more subtly preached the gospel of least restrictive environments or least restrictive alternatives, many young deaf people are unduly restricted from using quality over mediocre programs, or for having no choice, or from having any choice at all.

They are thereby denied an important civil right, and today, there is a second backlash becoming evident among educators in particular against proliferation of non-quality educational programs.

One other matter is worthy of attention before this state-of-the-art report is brought to a close.

The federally sponsored training programs designed to train

interpreters for the hearing impaired are not required to train their students to deal with the interpreting needs of oral hearing impaired adults who rely predominantly on their speach reading skills rather than on an understanding of sign language.

These programs should be mandated to do so in order that the civil rights regarding the interpreting needs of a significant portion of our hearing impaired population can be provided, since it's becoming increasingly clear that oral interpreters are often a reasonable accommodation to the severely hard of hearing as well as to the oral deaf community.

Thus we see that though the United States is the most progressive nation in the world in terms of what it does for its hearing impaired population, it still faces some significant problems that beg for solution before it can brag that it has provided for all the civil rights of that population.

As a nation we have done well,

but we certainly can do much better.

Thank you.

CHAIRMAN SCHWARTZBERG: Thank you, Dr. Castle, for that keynote presentation.

I note that I neglected to provide a detailed introduction of Dr. Castle's numerous qualifications. I would simply have that inserted in the record, since I doubt there is anyone here present in this room who is unfamiliar with him.

Let me now note that we have been joined by a member of the Commission, Dr. Robert Spencer.

Professor Spencer came in during the keynote presentation.

What follows is a somewhat different format, that is to say each of those following Dr. Castle is to be held to at least a ten-minute summary.

We recognize that will be far less useful to those of you in the audience than the presentation that you just had, the members of the Commission, however,

have these detailed papers which are available for examination, and we will in those cases be moving primarily to questions and answers.

Let me turn now to the first of these, and the first is Dr. Dennis Pappas.

I would like to ask Dr. Dennis
Pappas, and Dr. Judith Marlowe to please
come forward.

Dr. Pappas is Clinical

Professor with the Department of
otorhinolaryngology at the University of
Alabama School of Medicine, the University
of Alabama at Birmingham, and is President
of the Echo Foundation in Birmingham,
Alabama.

Dr. Pappas has provided us with a paper on the medical aspects of early identification of hearing loss and appropriate follow-up of high risks and registries.

## Dr. Pappas?

DR. PAPPAS: Thank you, Mr. Schwartzberg,

members of your committee, ladies and gentlemen.

It is indeed a pleasure to be here to give this presentation today, and this is a subject that has been of great interest of mine for some 12 years, and is greatly integrated into my practice.

Many of the diseases that primarily affect children have been controlled by legislative criteria for preventive measures augmented by the dissemination of educational information.

Hearing loss in children has not been as fortunate. Yet statistics have indicated that 1 in 700 to over 1 in 2,000 infants in a well nursery have a severe to profound hearing loss.

These figures may not be as impressive as some, but when compared with the potentially devastating adversities associated with this handicap, such as speach and language retardation, loss of education time, psychoneurological and psychosocial handicaps, behavioral problems

and employment prejudices, the significance of these statistics becomes more clearly defined.

It must also be realized that these statistics represent infants in well nurseries, the prevalence of hearing loss in neonatal intensive care units is more impressive and represents approximately 2 percent of these new births.

Nevertheless, awareness of the need for hearing screening as well as the special educational needs of the hearing impaired is still the exception rather than the rule, and the development of a philosophy of education and habilitation for hearing impaired children has been slow to progress.

The purpose of screening for hearing is to detect a hearing loss prior to the time that it becomes obvious. Screening any disorder is a task that should be accomplished rapidly, accurately, economically and will little energy or resources misspent pursuing normals.

Although no diagnostic screening device will fulfil this criteria completely, over the past decade programs and procedures for screening the hearing of newborns has been developed, modified and improved.

Nevertheless, the fact is that early identification programs continue to be absent from most hospital nurseries.

If a hearing loss is identified at birth and early habilitation education instituted, such effects may be diminished to the extent that the child can realize a more normal role in society.

Therefore it's proper to provide auditory screening early in life of all children, and the first and only opportunity to examine the entire infant population is in the newborn nursery.

Without dispute, there is a critical or sensitive learning period for language development. To prevent compounding the already complex problems inherent in a hearing loss with additional and possibly

2.16.0

l

greater social educational and psychological disadvantages, the diagnosis must be made early in the child's life.

Now, there is one point where this is very nicely done, and that's at birth, where there is a captive audience; however, there are children that develop hearing losses during the first year of life, and these, of course, will be missed with the screening.

The next screening period occurs typically at age four or five in the nursery school, and maybe in the first grade. The sensitivity then of the physician and of the parents becomes of utmost importance in identifying these hearing losses.

Such diagnosis as delayed hereditary hearing loss and progressive hearing loss of viruses, that tends to cause this problem, are the ones that I am Speaking of.

There is one high risk factor at this point, and that is of the mother who brings this child into the physician

and says I think my child has a hearing loss, we have found that she is right 25 percent of the time. That's one out of four will have a hearing loss.

The importance of identifying the cause of a hearing loss, in our eagerness placate the acute needs of the newly diagnosed hearing impaired child, we often overlook the significant repercussions of the specific cause of the hearing loss.

This knowledge should be vigorously pursued, for it should open the door to a wholeistic approach to remediation and unharness the possibilities of rehabilitative and medical intervention.

Clear examples of this would be a case of congenital hereditary hearing loss and one of hereditary delayed hearing loss.

In the latter case that I just previously mentioned, hearing acuity is lost in the first year of life, and the loss is progressive until the age of six

years.

Such children have normally heard speach for the first six months of their life. They may even have developed words by the age of 10 to 12 months, only to have their speach and language development deteriorate thereafter because of the progressive nature hearing loss.

These children benefit in speach and language development in an auditory verbal program.

Other examples of the significance of knowing the cause of the hearing loss of those due to meningitis, which may require accelerated speach language intervention and monthly audiological monitoring because of their progressive nature.

Those secondary decidua Mengo
virus or birth injury, both of which present
the possibility of brain damage and may
require specific habilitative processes,
and those associated with meningo dysplasia
in which the subsequent development of a

perilymphatic fistula should be recognized knowledge of the cause of hearing loss gives insight into the prognosis of associated problems and habilitative and medical management.

It can also be recognized that the progression of a hearing loss can be aborted in certain instances. These diagnostic identifications are very similar in the adult, it's just that the cause is different, and I might add that in reviewing and evaluating some 500 hearing impaired children over the past six years, that we have found that we can make the diagnosis in 70 percent.

Of that 70 percent, we can make the diagnosis with the history alone in 60 percent, so in 10 percent the diagnosis is made by special laboratory findings.

Of course, then in 30 percent the diagnosis cannot be made, and therefore we have to have a monitoring program for that 30 percent every three years for three times, that is up until about age 9 to 12, where

.

some, and many other tests have been introduced or repetition of tests will make the diagnosis.

After identification,

amplification with hearing aids should be

provided immediately after the hearing loss

is identified and within two weeks after

identification the child and its family

should be initiated into a program of

habilitation that develops the child's

audiological speach and language skills.

The guidelines for amplification in children are different from those of adults.

It has been demonstrated that even mild hearing losses can have an impact on the development of speach, and more importantly the development of oral language.

Consequently, the approach to hearing aid assessment in children must consider the speach language and vocal factors in a systematic manner. The criteria for the needs of amplification in a child with sensorineural hearing loss are not

ь

clearly defined, therefore each patient with a mild hearing impairment must be individually evaluated according to specific defects, specific language disorders, and academic performance.

In monitoring the child with hearing aids, changes in the hearing levels are commonly seen in children wearing hearing aids. Some of the causes of these deviations are faulty functioning of the hearing aid, serositis media, additional acquired causes, such as meningitis, viral infections, etc., and the progression of the hearing loss due to any ideology.

levels more than once a year is pure inertness.

Notwithstanding the possibility of preventing an effectively -- or effectively treating the cause of the progression of the hearing loss and aborting its potential damage, a mere 15 d.b. shift in the hearing may mean the difference between hearing and not hearing speach in children with mild to severe hearing losses.

Educational programs must be in a position to detect changes in hearing thresholds if adequate amplification is to be provided and parents and educators must be able to recognize changes in aided hearing performances.

The hearing should be objectively monitored with unaided and aided audiograms every three months following identification of the hearing loss.

During periods when the hearing is unstable, weekly audiograms are recommended. If the hearing remains stable for three years, the period between monitorings may be extended to every six months.

In all cases, these children should be examined whenever a change in speach performance is noted by the parents or educators.

Often times the cause of progression in the hearing loss cannot be ascertained. Nevertheless, when one is confronted with an aided child who manifests a progression of his or her loss, the function of the middle ear must be examined

\_

)

especially for the presence of negative pressure or fluid accumulation, additional examination for medical causes of the progression, including the hereditary factor, viral origin of sensorineural hearing loss and calcification of the cochlea following meningitis, perilymphatic fistula and otomyasthenia ear disease.

Should a progression in the hearing loss be determined to be related to the hearing aid usage, it should cause a temporary threshold, and the hearing threshold should improve with the removal of the hearing aid.

In regard to cochlean implant, discussion of the use of the cochlean implant is best approached with judicious caution.

At the present time there is not enough information available to determine which device is best for which patient, and this I'm referring to adults, this is due at least in part to the nature of the procedure, which does not take the device availability for comparison studies,

14

15

16

17

18

19

20

21

22

23

24

in addition the long-range ramifications are not yet ascertained.

Nevertheless, it should be clear stated that a cochlean implant has provided limited but significant benefits in some adults, such as the ability to recognize environmental sounds, and many of the segmentals of speech.

The use of this procedure in children is still considered experimental.

In any case, the cochlea implant should be considered only when there is no measurable improvement with the use of a hearing aid or aids.

Amplification by aids still gives the patient more benefit at the present date cochlea implant, that is, when it's indicated.

There is only a small percent of children where the implant is really indicated.

So, in conclusion, I'm advocating two aspects of this presentation, and that is the early identification,

l

.

especially at birth, where there is a captive audience, and that is the best time to do that.

The second is the two monitorings, one should be medical, the other one should be that of testing of hearing and hearing aids, and the monitoring of this is based on the diagnosis.

Thank you.

CHAIRMAN SCHWARTZBERG: Judith A.

Marlowe, who will give the next presentation,
is an audiologist whose private practice is
located in Winter Park, Florida.

Her memberships and certifications include the American Speach and Hearing Association, the American Auditory Society, and the Alexander Graham Bell Association, where she serves on the Executive Committee of the International Committee for Auditory Verbal Communication.

She has developed audiology services in a variety of settings, including hospitals, medical offices, nursing homes and schools.

In 1983, she designed and implemented a unique Infant Hearing Assessment Program in cooperation with the Infant Hearing Foundation and the Telephone Pioneers of America, which has served as a model for additional screening sites.

She is the editor of a monograph The Evaluation and Mangement of Communication Disorders in Infants, 1982, a contributor to the volume Hearing Disorders, 1984, and is currently completing a text book on the auditory verbal approach to communication development which is to be published in 1987.

Dr. Judith A. Marlowe.

DR. MARLOWE: Thank you, Mr. Schwartzberg and members of the panel.

It is a privilege to appear before you this morning to discuss an issue in which I've developed a very keen interest over the 15 years that I have been practicing audiology.

As is always the case, in appearing on a panel of this nature, much

basic information is repeated again and again, and I know you have heard much of it, so with the kind permission of the panel, I will depart from merely summarizing the paper with which they were provided to offer hopefully some additional insight without the danger of repeating what has already been so well stated by my preceding speakers.

of my colleague, Dr. Pappas, on the importance of early detection, and I think that among those of us who are in the field of hearing impairment, regardless of our particular persuasions with regard to educational methodologies, we are all in favor of early detection.

However, when we discuss the populations that we should be regarding for this purpose, I think we must make a distinction.

True, there is a population considered at highest risk for hearing loss, and the first steps have been taken in at least 17 states across this country to

address their needs, early and hopefully adequately.

But I'm concerned about another group of people, I am concerned about a group of people who are perhaps not in high risk for hearing loss, but who are at highest risk for the lack of detection, and that is the 30 percent of the hearing impaired population who present to us without associated medical factors at the time of birth.

There can be little argument that the idea of testing of the newborn nurseries across this land is an idea whose time has come. A very important pioneer in the field, Marianne Downs, has recently authored a publication in which she states that for the future it is imperative that every baby be screened for hearing loss, but his presupposes a very significant change in attitude for all of us, and that is this; while there is some evidence for the fact that screening of newborns for hearing impairment meets all of the criteria

for the public mandated disease screenings that we conduct, would it not be a wise thought for us to consider for a moment moving the concept of hearing screening into the area of wellness.

Now, wellness is the buzz
word of today's age. We are very keen on
the idea of wholeness, wholeism and wellness
programs abound in hospitals around this
country.

If we are to advance in the process of human development, is it not appropriate to regard hearing because of its reciprocal role in cognative verbal and social skill development as one of the foundations for wellness, and to address in that context.

So, I would offer the modest proposal that we not restrict ourselves to mere consideration of the high risk or medically involved population in the nursery, but that we expand our horizons and expand them rapidly to include the entire population.

After all, we take

temperatures and we perform hematocrits on every baby born. We are exploring babies born for conditions such as PKU, phenylketonuria, yet the incidents of these disorders is far less common that that of hearing loss.

Now, I do not merely make an idle suggestion that we do so. My appearance before you today, I'm sure, is prompted by the fact that I know where of I speak, because for the past three years I have had the privilege of organizing and directing a program which accomplished just that, a progam which routinely screen the hearing of every baby born in a particular hospital nursery, a small beginning admittedly, but a beginning nonetheless.

In this program we have made some interesting observations, we have observed that screening in and of itself is not sufficient. There must be a multi-pronged approach.

I

\_

u

public and physician education that infants can indeed be screened. One of the most common reasons for failure to investigate hearing early in infancy is the simple and unfortunate belief that infants still cannot be tested, despite the availability of advanced technology and computerized testing techniques.

Secondly, there is a need for prenatal education, the concept of parents education is a social change which has taken over in this country.

Included in that information when we educate parents regarding the developmental milestones for which to look, they can become expert reporters, and as Dr. Pappas has pointed out quite rightfully, they can serve as a very useful piece of information for us in the period after the infant has left the newborn nursery.

Beyond parental prenatal education and physician education, public education, we need to actually conduct a

screening which will enable us first of all to target those babies who deserve to be watched.

In our program we have determined that of the more than 3,000 infants that we have successfully screened during the past three years, 15 had problems deserving of our attention.

It has long been suspected
that the incident estimates of one in every
750 or 1,000 well infants maybe in fact
conservative, because most early screening
techniques only address the severe to profound
hearing loss categories.

In our screening, we are also screening for mild to moderate hearing deficits, and therefore our yield is higher.

Moreover, we have found that
in a voluntary program such as ours, we
can achieve an 87 percent compliance rate
voluntary participation by 97 percent of
the families delivering in this hospital.

With the support of the pediatrician who made this a standing order

in the nursery, hearing testing in the nursery does not arouse parental anxiety, it is viewed as a routine part of getting a baby off to a healthy start in life, and perhaps that is the orientation that we most fervently seek in the future.

In addition to our efforts in early screening, we are able to establish a mechanism in order to follow these infants during the early stages of their life, so that we have not withdrawn our net after casting it forth to catch them during the initial period in the nursery.

We have found over the past
three years a surprising increase in referrals
among physicians in the community who,
because of their new awareness of the
possibilities of testing infants in the
nursery, are able to remember hearing when
they are conducting their evaluations and
are more willing to refer for testing since
they are now acutely aware of the fact that
newborns are successfully being tested.

Our yield in this program is

still under study. We are in the infancy of our own work, and yet as we look at what we have been able to accomplish by presenting this as a part of a wellness program, I think that it points in the direction of what can be accomplished in the future.

The idea of early identification involves in the right of every parent to know not merely whether the child is deaf, but indeed to know does my child have adequate hearing in order to complete the course of development in education required in order to become a contributing member of society.

we know from our long experience with delayed identification that the effects of late diagnosis are far reaching, that they are costly to the national economy, that they leave behind a history of devastation in terms of wasted human resources.

Therefore the concept of moving beyond early identification aimed

exclusively at the sick infant into the well population offers us a golden opportunity, an opportunity to monitor all citizens, to look out for the civil rights of all citizens in a more conscientious way than we have ever been able to accomplish before.

The prospect of discovering infants early in the nursery period when we can inform the parent and begin a program of adequate stimulation is one that has been undertaken in 17 states across the country.

It is not, however, universally undertaken. In fact, plans to offer testing even routinely to those infants who are in the level three nursery is not progressing at a rapid rate.

What is needed is quite

clear, what is needed is a national mandate

stressing the importance of hearing

detection, setting in place programs that

will enable this to be carried out, but more

importantly, following up with early

intervention services in areas where early

intervention is not mandated until the age

H

of three.

The notion of identification during the newborn period is a cruel hoax.

It simply implies that the problem has been identified and will be treated well, when in fact adequate services do not exist.

move the youngster along the road to development through a successful infancy which will result in educational success, economic independence, can be one that is a realistic goal for all of us because of innovative planning that is now available to us, the mechanized instrumentation which now allows observation by more minimally trained individuals, thus reducing the dependence upon highly specialized and highly trained personnel.

All of these developments are available to us if we will marshal our forces and move ahead in this regard.

infants, offering them the care that they need, and following them so that they may

travel through childhood on the process to becoming with good hearing is a possibility now the models exist, and it is my fervent hope that we will adopt this as a goal in the future.

## Thank you.

CHAIRMAN SCHWARTZBERG: At this point we are going to take questions if there are any, from the panel, addressed either to Dr. Pappas or Dr. Marlowe.

Are there any questions from the panel addressed to either of these two speakers?

## Yes, Mr. Ewing?

MR. EWING: Dr. Marlowe, in the program, what has been the increased cost associated with delivery and other things that the screening has added to the hospital bill?

DR. PAPPAS: The cost of the screening on the hospital bill in the design of our program is \$25, which is the same cost as an infant picture.

However, in our program, no parent has ever been denied the test because

of inability to pay, and that is because the program is designed to be staffed by volunteers.

CHAIRMAN SCHWARTZBERG: Further questions to either Dr. Pappas or Dr. Marlowe?

Dr. Spencer?

DR. SPENCER: I would like to ask a related question, I would like to ask a related question to the two panelists, what is the impact of the cost containment efforts in medical hospital services by major carriers and medical care on what you want to do for the hearing impaired as you now see it?

I have noticed a reduction in the depth of coverage and a reluctance and concern on the part of some carriers of certain kinds of very expensive care.

This would be a new dimension in infant care, would it not, in these services if it were to be universalized, what is the impact of these cost containment efforts on what you want to do?

DR. MARLOWE: I can state from our

experience in the past three years insurance companies have not reduced their cost of coverage because of this, this is only a \$25 charge, which is less than a total and direct bilirubin cost in our lab, which is quite a common procedure.

So, we have not felt an impact in terms of a reduction in insurance coverage.

CHAIRMAN SCHWARTZBERG: Dr. Pappas, would you like to comment?

DR. PAPPAS: Yes, that question is well taken, because there has been a reduction in the services that a carrier wants to cover, and this has been one of the problems with introducing a new test.

On the other hand, one cannot really figure out, Mr. Spencer, how an insurance carrier covers the cost of an individual patient.

There are certain tasks that are just thrown in, and there is not an individual charge. In other words, the carrier pays the hospital for a newborn X

,

amount and whatever is done is done.

So, on the other hand, it may not be a problem.

DR. SPENCER: Thank you.

MR. EWING: I have an additional question.

It has to do with the concern of the medical community as to malpractice and recognizing that lawsuits can be brought up to age 21 by individuals.

Have you observed any
expressions from the medical community about
the benefits of protection through the
early screening as to whether or not there
may have been malpractice, or has there been
a reluctance to gather this additional
information which may add earlier to a
malpractice claim?

DR. MARLOWE: This is an interesting question which, quite honestly, in designing this program I expected to be a major stumbling block in the establishment of early screening.

However, I found that the question arose primarily among the

obstetricians and gynecology staff who did express some concern about the identification of any condition which could implicate their behavior during the delivery process.

This was overcome by obtaining information regarding other screening programs around the country and a judicial review indicated at that time, which was 1983, that the majority of lawsuits filed involved delays in diagnosis rather than early and prompt diagnosis, and from that information it became then a more positive type of idea to the obstetricians.

DR. PAPPAS: Yes, I will agree with Judith's answer, and I just want to interject one other, because this is a very good question.

One of the problems that we are having, especially in my state, with the malpractice problem is that many of the obstetricians are not doing obstetrics, they are dropping out because of the risk of having a suit, and this is really becoming a problem for such a task as we are

considering here, because we really need the input of the obstetrician in this.

I mean he is right there,
you see, and so I think it's in that respect
it's really a problem for us.

CHAIRMAN SCHWARTZBERG: Yet I assume that failure to do the screening might well produce even greater litigation at a later point?

That may or may not be a rhetorical question. I don't know.

Would either of you like to comment on it?

DR. PAPPAS: Possibly. If it's not being done, I guess you would say well, no problem, and then it should be done it seems, and a lot of this malpractice stuff doesn't really make sense.

You may think I'm just saying that, but it really doesn't. But at any rate, it would seem that such a test would be -- that would decrease such a risk should be needed and should prevent a possible suit, and that is if it's not done.

CHAIRMAN SCHWARTZBERG: I have a final question from member Theresa Cummings.

MS. CUMMINGS: For either one of you, would the informed consent, is this being used in this position, this parent signing for the child?

DR. MARLOWE: Yes.

MS. CUMMINGS: And the parents are aware of the test?

DR. MARLOWE: Absolutely, they are presented with a detailed explanation of the test, and also given individual quotations that show the test and the results, and they are encouraged to participate.

If the parent declines the procedure, the procedure is not offered to the child.

CHAIRMAN SCHWARTZBERG: At this time, in light of our time constraints, I'm going to thank both Dr. Pappas and Dr. Marlowe.

At this time I would like to call Dr. Daniel Ling.

Dr. Ling is Dean of the faculty of Applied Health Sciences at the

University of Western Ontario, Canada, and President of the Alexander Graham Bell Association for the Deaf.

Any attempt to read a list of Dr. Ling's publications would necessarily be difficult, since he is the author of over 200 publications relating to hearing impairment, including several books on the subject.

He has worked as an educator of the hearing impaired in audiology and speach pathology, and in research programs in various universities.

His particular interests are in early intervention and in speech and spoken language development.

Dr. Ling.

DR. LING: Thank you, Mr. Schwartzberg, members of the panel, ladies and gentlemen.

The instructions for preparation of my paper were very clear, they were to descibe and briefly discuss the three modes of communication used in educational settings, the rights of children

to options involving those modes and settings, and the third thing, the extent to which current options cater sufficiently to the needs of hearing impaired children.

Certain aspects of the topic either have not been undertaken or have been carried out only to yield equivocal results, and an additional one is dealing with a field that is changing and must continue to change in response to the opportunities afforded by burgeoning technology.

An example of this is that many people a few years back would have classified any child with a hearing loss of over 70 d.b. as prfoundly deaf.

Now, many people are querying whether there is such a barrier at even 90 d.b. with current technologies.

First the mode of communication, the three principal modes of communication

I was asked to deal with were oral spoken language communication, total communication sign language, plus speech and cued speech.

There are other methods, but

j

•

they are not in wide spread use.

The methods employed to develop communication within these modes and the various settings vary widely in relation to the skills and philosophies of the educators and the perceived needs of children they serve.

Hearing impairment may be present from birth or be acquired at a later stage. It can range in severity from an auditory defect so slight that its presence can be noted only under certain adverse conditions to a deficit so great that little or no sound is ever perceived as such even through the most powerful hearing aids available, and because individual differences similar to those found among normally hearing children are also present among hearing impaired children.

A variety of communication modes and educational options are required to meet hearing impaired children's needs.

No one method or collection of methods can possibly cater for the needs

2

3

of all hearing impaired children.

4

5

6

10

11

12 13

14

15

16

17

18

19

20

21

22

23

24

Oral communication is employed in settings where the primary goal is to develop speech reception and speech production skills that will permit children to acquire the language of the country in which they live to learn through the use of spoken language in schools and to function later as independent adults through the use of speech communication in society at large.

Individual needs call for various oral options. These include auditory verbal education in which audition alone without speech reading is used for part of a child's training. Oral education in which audition is used as the primary channel in multi-sensory training and visual oral training in which emphasis is on speech reading again in multi-sensory training.

The first option in my view should be the first one. Not all children who have been placed in oral education settings have achieved intelligent

spoken language communication, and current children whose progress in oral settings has proven to be unsatisfactory are usually transferred to total communication or cued speech programs.

Total communication is taught in settings where the primary goal is to establish and develop communication by using sign, manual sign language or all other means of interacting with the child.

It has been defined by the Conference of Executives of American Schools for the Deaf as a philosophy incorporating appropriate oral manual and oral modes of communication in order to establish effective communication with and among the hearing impaired.

A secondary but major goal of TC, total communication, is that hearing impaired children should learn to use English, and in particular to read and to write it.

American sign language, ASL, is a distinct language, hence it has inherent

•

potential to conflict with the learning of English.

In attempts to avoid such conflict, several sign systems have been created. The purpose of these systems is to preserve and extend what some consider to be the essential similarities between sign and English.

The theory is that sign language presented simultaneously in combination with spoken English will be mutually reinforcing.

However, subsequent work
has not given unequivocal support to this
hypothesis, and in many cases has denied
it.

Cued speech was invented and first reported by Coronet in '67. It was designed to supplement speech reading by using four hand positions and eight finger configurations to differentiate speech patterns that look alike, such as PBN, TDN, either hand can be used to formulate the cues, all of which were made close to the mouth of the

speaker.

The hand positions are used to differentiate vowels and the finger configurations consonents.

It can be employed to provide totally deaf or near totally deaf children with a more complete pattern of spoken language than they would otherwise be able to perceive.

The system is not difficult to learn, an average person can become completely familiar with the cues but not fluent in their use in less than ten years.

It has been clearly demonstrated that cued speech can permit this reception of speech at extremely high levels of accuracy. For their research on the system is required, but probably more for logistics than for theoretical reasons cued speech is not yet in very wide spread use.

It is a viable alternative to more widely accepted oral procedures, an alternative to cochlean implant surgery, and a realistic alternative to total

communication.

Cued speech is particularly appropriate for those that cannot for some reason be fitted with or benefit from a hearing aid.

A cochlean implant or a tactile device is an aid to special reading. Educational settings, many different types of educational settings exist to cater for the wide variety of children who are hearing impaired.

of settings is determined by the age of the children, the type and degree of the hearing impairment, the levels of their communication skills and their educational achievements.

However, other factors such as the presence of additional handicaps in the population, the geographic distribution of cases, the existence or not of programs for the detection of hearing impairment fluctuations in the relatively low incidents of hearing impairment funding considerations

and prevailing philosophies of educational treatment all contribute to the provisions or lack of provision of educational settings.

Unfortunately, early
intervention options which are perhaps the
most useful way of starting a
child's career in education are not
available in all states across North America,
and early oral options are particularly
scarce.

I would now like to specify a few of the rights that are listed in my paper, just pull them out to state some of these rights relating to communication modes in educational settings.

I believe that hearing-impaired children have a right to receive the serivces and devices that are necessary to promote optimal development and use of their potential for speech reception and speech production and the use of fluent spoken language.

Next, parents have a right

\_

to receive fully documented reports on the hearing impairment of their children with complete and impartial explanation of its implications for the choice of their children's mode of communication, and most appropriate educational setting. Many parents don't get that.

Three, hearing-impaired children have a right to placement in the educational settings that offer one and only one of the three major modes of communication from the time of diagnosis throughout school life.

I want to point out that oral education is not provided as part of total communication.

Oral education requires the sole use of oral methods.

Four, children have a right to receive instruction in educational settings that most effectively develop the modes of communication chosen for the child by the parents.

Five, no child who has the

0 23344

potential to learn how to communicate fluently through speech should be denied the right to achieve that potential.

Six, each hearing-impaired child has a right to achieve educational and communicational skills at an optimal rate.

Seven, hearing-impaired children have a right to an on-going review of their performance with regard to the suitability of the chosen communication mode and educational setting.

Eight, hearing-impaired children have a right to be taught by educators who are adequately prepared for the task.

Nine, parents have a right to choose the type of education for their children that best suits their own philosophies and the children's needs as they, the parents, perceive them.

Rights relating to educational options.

Parents of your hearing-impaired

children have a right of access to parentinfant programs which they can learn to
serve as the primary agent in the habilitation
process.

Secondly, under this
heading, parents who are unable for whatever
reason to participate in early habilitation
programs as the primary agents of
intervention have a right to expect childoriented programs to be available for the
hearing-impaired children.

Next, hearing-impaired children have the right to be placed in settings that offer communication modes and education appropriate to the type and degree of their hearing impairment.

On the other hand, next, children have a right not to be assigned to a particular education setting simply on the basis of their hearing levels.

There are many, many children who are quite capable of speaking orally, learning without a great deal of intervention, who are currently in a special state school

for the deaf which is, I think, teaching potentially hearing children to be deaf adults.

Children have a right to and therefore not to be assigned to those sort of settings.

Hearing-impaired children,
lastly, attending regular schools as integrated
pupils have a right to such support services
as are necessary to maintain their optimal
performance in those settings.

One thing I want to end on, and it is a statement that I think we must bear in mind, hearing-impaired childrens' access to a full range of options, educational settings and communication modes is unduly restricted in many and probably most regions in North America.

CHAIRMAN SCHWARTZBERG: Thank you, Dr. Ling.

Questions from the panel?

MR. EWING: Do you have any information -
I'm sorry, we often hear of what services

are needed and the type of trained people

)

who should deliver those services.

Do you have any information as to the numbers of trained people currently available to deliver the kind and quality of services that you are concerned about?

DR. LING: I would like to reply,

Mr. Ewing, by reinforcing what Dr. Castle

has said, that we really need to review

thoroughly the type of training that people

receive in these areas, because in the

American Speech and Hearing Association,

there are two faculties, one speech

pathologist, speech language pathology,

the other audiology.

There is no oral habilitation

section in ASHA. There is a Council on

Education of the Deaf, who have, with the

knowledge of various of our people, produced

new guidelines for the education of people,

but there is still a great gap in the provision

of university programs that can in fact

provide the necessary training in oral

habilitation.

So, we are really short on

educational settings at the university level, that will provide the field with adequately trained personnel.

MR. EWING: As a follow-up question, would you favor this need being met through maybe a concept of -- because normally teachers come into these public school districts on a fixed salary level, but where we have shortages, and it has this impact upon the future of children, would you favor maybe a different salary schedule to bring in the qualified people that are needed to provide these skills?

DR. LING: In many other communities across the world, which has in fact been found to be essential, and I would certainly support that notion.

CHIARMAN SCHWARTZBERG: A question from Dr. Lucas.

DR. LUCAS: Dr. Ling, to what extent, the various approaches to extend communications skills to the deaf, are technical issues on dealing with the hearing-impaired person and to what extent or where is the limit of

what a social policy of the rights of the hearing-impaired person has a limit, where are the limits between those two areas, the right of the person?

Is there such a thing as a person's right to speak, how would you define for us that limit?

DR. LING: What you are asking me to do is define which side of the pair of scissors cuts the paper? I think that you have to look at the technical advances, you have to look at all of the social and legal issues and say that these somehow have to be put together.

I don't think there is a simple answer to your question.

CHAIRMAN SCHWARTZBERG: In other words, as I gather, you are suggesting that what the right is changes over time in light of technological possibilities?

DR. LING: I think so. The advent, for example, of cochlean implants of wearable devices for allowing the child to perceive speech through the skin, tactile

2

3

4

5

6

7

10

11

12

13

14

15

16

17

18 19

20

21

22

23

24

aids, these are recent developments.

a child and a parent has Now a right to choose between those type of devices, one, the cochlean implant invasive, the other, tactile aid non-invasive, I think there is a right of choice that just didn't exist before.

CHAIRMAN SCHWARTZBERG: But you place the choice itself in either parent or child rather than in the state?

DR. LING: Most certainly. I really do.

I think that the well-informed parent has to make the choice, not only an informed parent since the child can possibly make a choice then the choice has to be there.

The major thing about choice, and it is something that was spoken about last night, is that there are no choices for speech and spoken language communication unless it's developed early in life.

The only time a hearing-impaired child can choose what sort of communication they can use is when speech communication has

already been established in early infancy, because speech communication, effective speech communication probably cannot be learned after about age 11.

CHAIRMAN SCHWARTZBERG: And it was suggested in some of the papers, as I gathered, that unless certain activities occur within the first six months of life there are additional restrictions on choice which occur later, so in that sense either the state or the parent must, as I understand, under this model, choose at that point?

DR. LING: Yes, Mr. Schwartzberg, the choice has to be made early, because there is a decremental performance in terms, right away through life, it is an optimal period. It is not a critical period in that first six months of life.

I personally have taught children who have no previous experience of speech to speak very intelligently after six, seven, even eight years of age.

However, it was harder to do it, took longer to do, there is a definite

\_

decrement in ability to learn spoken language communication throughout childhood, and so the choice must be made early.

CHAIRMAN SCHWARTZBERG: Let me seek one clarification of that. You have talked about decremental ability to learn.

Is there a limitation on the eventual result which is depended upon when the choice is made?

That is to say, if the choice is not made in that early period, is the result limited by the time of the choice?

DR. LING: Yes, I think the result has to be limited by the time of choice.

I think that it's completely outside the question after about age 11 if the child hasn't got a substantial amount of learning down, but the major point I think to recognize is okay, that if when you are teaching a child or when the child has left without learning speech, it isn't that he's learning, he's not learning to communicate, he's learning to communicate by other means that may be inimical to the later development

.5

)

of good speech.

In other words, you are not learning nothing, you are learning an alternative.

CHAIRMAN SCHWARTZBERG: I'm really looking for a different question.

DR. LING: Okay. Try me again.

CHAIRMAN SCHWARTZBERG: Let's assume that a child is born who is a potential Nobel Prize winner; is it realistic to assume that that child will reach that potential if the child does not have input within the first year of life?

DR. LING: I would seriously doubt whether the child could every achieve full potential if the child didn't have good input early in his life.

MS. CUMMINGS: Dr. Ling, have you noticed any certain mode depending upon the economic level of the family?

DR. LING: Yes, most certainly. There are quite a number of demographic studies published by Gallaudet College that show that the various children certainly the ones in

lower economic socio-economic groups are in fact more likely to be placed in state schools, in sign language programs and not to achieve optimal growth, yes, and socio-economic conditions are a key determinant of educational achievement.

MS. CUMMINGS: And the mode?

DR. LING: And the mode, yes, but it is not necessarily so.

Early intervention with these children who in lower socio-economic groups could, I'm sure, result in far better achievements on their part. It's just that they are not available to them.

MS. CUMMINGS: Do you feel that with medical costs and things as they are, that the expense of some of those has a lot to do with it?

DR. LING: It may well be, but that's taking a very short-sided view, because the cost of intervention with these children and the results of getting them to be productive citizens and self-supporting citizens at a later stage is probably very much better

than looking for a whole group of adults who are either a charge on the state, unemployed, or not achieving and working at an optimal potential.

MS. CUMMINGS: What I'm saying, if all of the states would make that a requirement to test the child early, then that possibly could be avoided because apparently they would know early.

DR. LING: That's right.

MS. CUMMINGS: But if that's not included in part of the medical services early as a requirement, then that child could be excluded from testing?

DR. LING: That's right.

MS. CUMMINGS: And then we get to the later clause, this is what I'm speaking to.

DR. LING: That's right, I agree.

CHAIRMAN SCHWARTZBERG: We have a question from Dr. Lucas, and let me point out to the remainder of the panel and to Dr. Lucas I have promised that under all circumstances Dr. Ling would be allowed to leave at 10:30.

.

DR. LUCAS: Coming back to the ratio
of the civil rights of the deaf, we established
that this is a fluid, progressive thing,
what rights means in a particular case
according to technology and learning, how
would you suggest, is it a government task
to keep up-to-date of what these rights are,
how would you do that, to what extent is
government responsible at what government
level?

DR. LING: I think that what I tried to point out is that some of these rights changed as science and technology advances, some of these rights are basic like the right to learn, to learn to talk, the right of parents to make decisions, and as time passes and as technology has impact in the field, I think it is through this type of forum that we should in fact make people aware and negotiate legislation to cover the type of rights that develop and change along with technology and time, and I must congratulate you all on setting up this forum and holding this forum as a very valuable procedure.

Thank you very much.

CHAIRMAN SCHWARTZBERG: Thank you, Dr. Ling.

Our next speaker is Dr. Winifred Northcott.

Dr. Winifred Northcott is a past president of the Alexander Graham Bell Association for the Deaf and the National Council on Education of the Deaf.

She is a Fellow in the American Speech-Languange and Hearing Association.

In May of 1986 the New York

League for the Hard of Hearing presented

her the Nitchie Award in Human Communications,

citing her contributions in helping hearing
impaired children to function in the mainstream

of a hearing society and in training other

professionals to further this goal.

Dr. Northcott has international recognition as a teacher, administrator, teacher educator, author, lecturer and consultant on family-oriented preprimary auditory-verbal education and a systems

approach to the maintenance of support services required for school-age children in the mainstream.

A long-term consultant

for the deaf and hard of hearing in the

Minnesota Department of Education, she

was Director of the UNISTAPS Project for

hearing-impaired children, zero to four

years and their parents, one of five

projects validated for national dissemination

by the U.S. Office of Education under the

Handicapped Children's Early Education

Assistants Act.

Dr. Northcott is the daughter of oral deaf parents, the late Dr. and Mrs. Edwin Nies of New York City.

Her father, a graduate of
Lexington School for the Deaf, Gallaudet
College and the University of Pennsylvania
Dental School is cited in Deaf Heritage as
the first deaf dentist in the United States.

It is my great pleasure to present Dr. Winifred Northcott.

DR. NORTHCOTT: Thank you, Mr. Schwartzberg,

for your very gracious introduction.

If I could be permitted to add to Dr. Ling's presentation, I would say that freedom of speech and freedom through speech is still a fundamental human right, and no where is there an asterisk that says all but the deaf.

Blackson, a British jurist,
has said that any law reflecting the moral
sentiments of the people, and we need to look
at the climate in the mid 60's to early
70's prior to the enactment of landmark
legislation, it was a decade of civil rights
protection and action, a crescendo of
outrage by persons with disabilities.

because of lack of equal opportunity in the areas of education, health services and government benefits, and public school education was then preparing said children and youth for world citizenship.

Comfortable in dealing with the diversity, the natural diversity in the world and the pursuit of life-long learning

•

as an insurance for assimilation in the world of work and pleasure.

In the 1960's on the primary educational intervention for the hearing impaired took the form of 12 demonstration projects in a public school, hospital setting speech and hearing center of public schools and the retrieval materials which you can read in any piece of literature that the deaf children were learning to listen and to hear in family-oriented programs, support the parents and the affected as well as the instructional aspects of living with a hearing-impaired child, full-time hearing aid usage, and 11 of the 12 projects had an auditory verbal approach to natural language learning.

Group educational experience was in regular nursery schools.

The basis for educational programming during the elementary school years was on the basis of what does the child hear and what can he do. You literally did not know who was becoming functionally

hard of hearing or functionally deaf.

The audiogram did not indicate the prediction for success or a lesser achievement.

So, parents began taking the initiative as burgeoning numbers of hearing-impaired children with severe and profound losses were functioning as self-operative learners, could speak, although imperfectly, could express themselves and be understood.

They began putting many of those children, their sons and daughters, in regular classes. One regular class teacher I knew said I'm willing to try him, but I've never had a hearing-impaired child before, and the mother said that makes two of us. I never have either.

In more than half of 50 states prior to 94142, changes in laws governing public school operation made a commitment to educate children with special needs in settings as a part of and not separate and apart from their non-handicapped peers whenever possible.

E 2232 - CHICAGO, ILLINDIS 60

So, the landmark legislation, Congress was not acting and voting in suspended isolation, there was a history of this movement, the commitment of Congress in the end, Congress voted a law writing in not only for procedural safeguards governing the process of determining educational placement, but a philosophical text on educational programs and service and the terms were individualized educational programs, parent-school parternship all the way.

The stipulation was free and appropriate education. 023 after 1979 was an option, it was not mandated. It was an option if all others were served.

It required the state

Department of Education, the local education

agency to establish formal written procedures.

I want to read the heart of the law again, although you know it.

"To assure that to the maximum extent appropriate handicapped children, including children in public or private

educated with children who are not handicapped, and that special classes, separate schooling or other removal of handicapped children from the regular educational environment occurs only when the nature of the severity of the handicapies is such that education in regular classes with the use of supplementary aids and services cannot be achieved satisfactorily."

Section 504 was enacted,
supplementing the Rehabilitation Act of 1973
enacted in 1977 and its purpose was equal
opporutnity under the law, the insurance
of it, the heart of it, again, "No otherwise
qualified handicapped individual may be
excluded from the participation end, denied
the benefits of or subjected to discrimination
under any program or activity receiving
federal financial assistance or under any
programs or activity conducted by any
executive agency or by the United State Postal
Service."

Part of the regulations dealt

exclusively with preschool, elementary and secondary education, mandating auxiliary aids, including interpreters, or al and sign language.

So we come to the education or All Handicapped Children Act of 1975, Public Law 94142.

What is it, what is it not?

It mandates a variety of educational settings,
a variety of methods of instruction.

Assistant Secretary Madeline Well of the Office of Special Education and Rehabilitative Services in the U.S.

Department of Education talks about the continuum of alternative placements, but she speaks about saying that they are from residential environment to less restrictive environments.

Methodology in writings and in statements, she has said, "remains the possibility of the parents and the educational system."

I note dryly that Assistant Secretay Well refers to equal opportunities

•

for hearing impaired and deaf children, which is incorrect.

As someone earlier mentioned, hearing impaired is a generic term that covers all hearing, persons with a significant hearing loss. Deaf and hard of hearing is correct. Deaf and hearing impaired is incorrect.

The least restrictive environment is the regular class placement, the most restrictive is residential school institution hospital setting.

By the law, by the manual,

Manual Ten, that was referred to, the listing
is regular placement resource schools

special placement special schools, both private
oral and public residential schools.

The heart of it is the annual performance review assessment of an individual child, the development of an individual educational program for that child by parent and staff at the local school district level.

Procedural safeguards for

parents if they want to protest and move into formal hearing of various sorts, then within the school district or the courts, but the responsibility lies with the local educational agency of the child's residence to program locally, including supplementary services, academic tutors, speech therapists, interpreters, at the secondary level, or to contract with another district or agency if required.

The rules say if not in a neighborhood school in a regular classroom, then it must be a program close to home as possible approximate chronological age of the classmates provision of maximum contact with non-handicapped peers and non-adademic and extracurricular activities.

The monitoring, recently I'm told by a friend in the State Department of Education, a state department of education, monitoring by federal office of special education and rehabilitation service representatives is the interest in the process of informing parents of educational

alternatives. They are not as interested, it's not a high priority that statistics of the number of children integrated are significant.

So, what are the implications for deaf children? The teacher of the deaf no longer owns the deaf child.

The principal of the school and the state where I was the state consultant in the department of education for many years when I said are there a few children I could help you move back into local school districts, this was in the mid '60s, he said what, take our successes away from us?

The least restricted environment by law and Manual Ten is "maximum appropriate interaction with non-handicapped children."

Assistant Secretary Well in personal correspondance wrote "The goal is to see that all special education children are educated in regular classrooms with all necessary related services, including resource

rooms."

We may not realize it in this decade, but the significance of the next statement is really the heart of what I'm saying.

The least restrictive
environment for an individual child,
according to his individual educational
programs can be any setting in the continuum.
It may be a resource room. It may be a
public or private residential school.

The individual needs dictate the setting that will be competitive and motivate learning for that single child.

The mode of communication is the parents thoice.

Earlier you have heard the statement, the tragic statement so few parents know of their options.

In my judgment, in my specialized interest in an auditory verbal teacher's approach to language learning, so few parents know of their right to expect for their own child an auditory oral approach

if that is there wish.

The court decisions indicate, to date, that the needs of an oral child cannot be met in a classroom taught by the simultaneous method, finger spelling signs and speech under the philosophy of total communication.

find cooperative school districts, you find realization of services, you find two track programs, and if you find an intact good parent school organization, you find parents fighting for the right to maintain both of those options, the auditory verbal track and total communication philosophy under the simultaneous method track.

We need to look a moment at the data about total communication.

The term was coined in 1967

by Roy Holcum, a teacher and a deaf man.

It became visible in 1971, it was first

reported nationally, internationally, rather,

at the Congress on Education of the Deaf

held in Tokyo in 1975.

The earlier studies were invalid in the sense that there was no truly oral group as a control group, because they were all conducted in public residential schools.

The median meeting reading level of those today leaving school not in integrated secondary programs but in residential, from public residential schools and special classes is 3.5 years.

Dr. Hilda Schlessinger in

Luderman's book of perspective on deafness,

1986 "recent studies on tv with this myriad

of definitions and as presently carried out

have not shown the desired effects on large

schools and the distinguished research,"

Quigley & Paul in 1984 made the same statement.

The philosophy is translated into educational practice seems to be a deterrent to the active use of residual hearing.

Full-time hearing aid usage is insufficiently stressed, according to Ross & Calvert in '84, and Luderman & Chase

27 M.C.

in '81.

2

3

4

5

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

Connor & Calvert in their studies, in examination, indicate that speech development was neglected and language development was advocated.

The basic problem here is one that has been described through the extensive questionnaire given out by Gallaudet Research Institute for parents and children in a variety of education settings who use the simultaneous method that literally parents, teachers and children, by their responses, do not use additive amounts of speech and signs at the same time.

The way the monograph statement goes, when the use of speech is high, the use of signs is low. When the use of signs is high, the use of speech is low, and this is true in all interaction modes, the teacher to child, the parent to child, and vice versa.

The oralness of the environment makes the difference.

The characteristics of the

hearing-impaired children who are assimilated, not just sitting there integrated but assimilated in regular classes relates to oral communication, personally and linguistic competence according to Gale and Plaser in 1980.

Deaf children incontestably oral programs which are your private oral schools, a distinction from the residential state public school, early identification and full-time hearing aid usage is a good prediction for assimilation.

No inherent disability in deaf children says Dr. Connor, but rather a disability in their educational system in many instances, the annual survey of hearing impaired children in youth conducted by Gallaudet College, speech intelligible equates with speech usage directly, with hearing aid usage, with school achievement, with auditory oral habilitation in infancy which is one of Dr. Ling's major points.

The issues we have to face are very real in the future, first the issue

23

24

18

19

20

21

22

of methodology, what shall be the mother tongue, the native language of the child.

Shall he or she think in words or in signs?

We need to remember what

Gallaudet Research Institute's annual

survey has found, 91 percent of hearingimpaired children have two hearing parents,

six percent have one hearing-impaired parent,

only three percent have two hearing-impaired

parents, some of whom are oral.

It's not a question of which is better in the abstract. We move to the particular. It's a matter of informing parents, and that's difficult to do.

An infant program worth it's salt needs to encourage and to literally hand carry and help two parents to visit every known program in the state, in the area, to make informed judgments.

No one method is better than another, an orderly progression, the first option has to be available to all families who want it. An auditory oral approach to

.

language learning.

It's not available in most parts of the country, in the average public school setting. What can they hear, what can they do is a function of infant and preschool intervention programs.

Distortion of speech is no problem to the profoundly deaf child. He has never heard perfect speech.

We need a new data base. We critically need a source of information about how children in regular classes part and full-time are doing.

The annual surveys out of Gallaudet College served a magnificent purpose. It's purpose is somewhat diluted at the present time, because it's source are special classes and special schools. They indicate the number of children mainstreamed from the schools and special classes, public, residential schools is declining.

The number of children mainstreamed from those settings, it shows

94142 is working via the IEP to a pretty good extent.

There is need for mandated surveys within each state of the number of hearing impaired children in part or full-time regular classes, and that the information go automatically up to the U.S. Department of Education.

So, we can have the subject's potential subjects for data base studies.

Secondly, what shall be the role -- or the third issue, what shall be the role of the public, not the private, the incontestibly oral private residence schools, what about the role of the public state residence school, the CEASD, Conference of Executives of American Schools for the Deaf, now called Conference of Administrators Executives Serving the Deaf.

Their formal paper speaks

about the role of the public residence school,

that it is the least restrictive environment,

that it has highly trained personnel, it

can serve the important social emotional needs

of children.

2

3

4

5

6

7

10

11

12

13

14

15

16

17

18

19

20

21

22 23

24

It recommends that the birth to three infant programs be located in "sign language-oriented programs" in these special schools.

They recommend that the public residential school act as a statewide assessment center, the resource for consultants services to the local educational agency.

You need to be informed about all formal position papers, Alexander Graham Bell, CEASD, so that you can work in your own community and region for the direction that you want your local programs to go in.

Labels, when is a deaf child not a deaf child? It's impossible to answer today. Will the real deaf child please stand, but why are we labeling programs as programs for the deaf when those children have one, two, three, four additional educationally handicapping conditions.

CHAIRMAN SCHWARTZBERG: Let me, at the risk of treating this presentation in a sort

of warning manner, I'm going to have to call time at this point, at this particular point, because I must advise the audience, I have great difficulties worrying about exactly how we get to our planned closing time, and I am going to take the risk of cutting short at this point what is an, of course, obviously an extraordinary presentation by Dr. Northcott.

Let me start, if I may, with a question. Let me do this, I'm sorry.

DR. NORTHCOTT: I got it all in, it's all right.

CHAIRMAN SCHWARTZBERG: Let me do this, if I may. We are going to come back to questions in a moment.

I would like to suggest two things. There are one or two people standing in the rear, there are some additional seats in the front at my left, your right.

The second thing is I don't know whether Mr. Brian McCartney is in the room yet.

Is Dr. McCartney in the room?

Does anybody know?

Let me turn to the next presenter. The next presenter is Dr. David Conway.

Dr. Conway is an Assistant
Professor in the Department of Counseling
and Special Education in the University of
Nebraska, Omaha, also is Director of the
Education of the Hearing-impaired Teacher
Training Program at the University.

He received his doctorate in special education for the University of Cincinnati, Ohio in August, 1985.

Dr. Conway has spent five
years as an instructor and Assistant Professor
at Minot State College as a member of the
Education of Deaf program and as Director
of Services to parents, infants and
preschool children with impaired hearing.

He received his master's degree in Education of the Deaf from the Smith College-Clarke School for the deaf Teacher Training Program.

He taught at Clarke School for

five years.

This month, Dr. Conway

begins a three-year term as Associate
Editor for the Volta Review.

He is a member of the Nebraska Department of Education ad hoc committee on Early Childhood Special Education.

DR. CONWAY: Thank you, Mr. Schwartzberg, members of the panel, ladies and gentlemen.

The question and indeed the challenge of providing educational services to our hearing-impaired children living in rural areas presents a number of unique and complex problems for government at all levels, federal, state and local, for school administrators, for educators, for parents and for the hearing-impaired children themselves.

If the source of the problems could be summed up in one word, that word would be diversity.

First, there is diversity in the characteristics of rural areas. For as

ŀ

Doris Helge, a leading researcher in rural education, has pointed out, what constitutes a rural area is not just a matter of low population density and geographic features, but also is dependent upon a multiplicity of community and school district variances as well.

Second, there is diversity in the nature of hearing impairment and the educational needs of hearing-impaired children at all levels.

Thirdly, there is diversity in the way educational services are provided for hearing impaired children in rural areas.

The primary purpose of my presentation to this panel this morning is to look at how services are or can be provided to hearing-impaired children in rural areas.

As much as we have to deal with diversity, we also have to deal with a lack of information in this particular issue. Unfortunately, based on the

information available, it does not seem that we are doing a particularly good job of meeting the needs of hearing-impaired children in rural areas.

Just two years ago, Knox made the following observations based on his study of 33 rural school districts' programs for serving hearing-impaired children.

One, by and large, there is very little activity in the area of identification of hearing-impaired children, severely impaired students tend to surface without such identification procedures in place, and therefore most of the local services are directed towards this group of youngsters.

From this, one is led to conclude that large numbers of children with less severe losses are unidentified and thus unserved.

Two, there did not seem to be many options available for serving hearing-impaired students in rural areas.

Typically, the two choices of placement for

such a child are in the regular classroom with or without support services, or placement out of the district which, according to Knox, usually meant in a public residential school.

Three, services appear to be more effective in terms of availability and options when regionalized programs are available, or when regional cooperative arrangements between districts have been entered into.

Four, educational personnel who have been trained to work with hearing-impaired students are an endangered species in rural areas.

Five, in a typical rural program, a speech clinician is likely to be responsible for language development programs and alternative methods of communication.

Resource teachers and regular education teachers shoulder the responsibility for academic progress.

Six, one frequently has the

15

22

20

21

24

23

feeling that students with hearing impairments have a low priority in many of the districts.

It should be noted that the programs on which these observations were made were not confined to one section of the country. This was taken from school districts from the northeast to southeast upper midwest, southwest and northwest areas of the country.

The districts also operated under different administrative alignments, intermediate school districts, multi-county cooperatives, county districts and local school districts.

Despite the less than encouraging observations of Knox, the picture is not completely negative.

Effective programs that can provide quality educational opportunities for hearing-impaired children in rural areas have been developed and are in operation. Unfortunately, there is not a consistant pattern of using programs, implementing

programs across states or even within individual states.

At the infant level, zero
to three years of age, home-based models are
most prevalent. Such models often described
as parent-infant programs stress early
identification, early amplification and
early intervention.

Examples of such programs are the Sky High, developed in Utah and designed for serving specially hearing-impaired infants in rural areas, and the Portage Project, which first began in south central Wisconsin.

At a preschool level, three
to five years of age, hearing-impaired students
enter into more formal school or center-based
learning settings. These can range from
classes at state or private residential
schools or satellite classes from these
schools, college or university affiliated
clinics, hospital programs, public schools,
within the public school system or are
contracted to private agencies, multi-agency

2

3

4

5

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

centers and diagnostic centers.

Some areas continue to use home-based programming in conjunction with center-based instruction at the preschool Upon reaching kindergarten age and continuing until graduation for schoolleaving age, hearing-impaired children become a part of the more traditional schooling process.

Educational options which should exist, and I stress should because these options do not always exist and not all the options exist in any given area. educational options which should exist for these hearing-impaired students ought to include regular classroom placement with no support services if unneeded, but with monitoring of progress, regular placement with support services, including any or all of the following; consultant-teacher of the hearing impaired, speech language theory, ideological services, tutor, note taker, peer tutors, interpreters, interpreter tutors, psychological and counseling services and

team teaching, a resource room.

If categorical, the hearingimpaired students would be placed with a
teacher of the hearing impaired for part of
the day and placed in a regular classroom for
the other part of the day.

If a generic resource room, hearing-impaired students could spend part of the day with a resource room teacher who may or may not be trained in education of hearing impaired, and with students with other handicapping conditions, consulting teacher services may or may not be available on top of this.

Consultant itinerant services.

A teacher of the hearing impaired works with local personnel to design educational plans and monitor progress of the students.

The consultant or intinerant teacher may or may not work with the hearing-impaired students individually or in small groups, depending on the arrangements within the school districts.

Contained classrooms. Students

would spend the majority of the school day
in a self-contained setting with a teacher
of the hearing impaired, and with other
hearing-impaired children of similar
chronological age and abilities. Additional
support services can be added on to this.

Classes for the hearing impaired may be housed in a public school building with other regular classes, or may be located in separate facilities.

Finally, residential school.

Placement is usually considered out of district in a separate facility offering

24-hour care and programming.

Variation on any of these options is possible. Districts may enter into purchase of service agreements with other districts to secure the services.

In some locals, foster living or boarder programs have been initiated.

In these programs, the hearing-impaired children who otherwise would not be able to obtain needed services, board with host families in districts or

areas which do have those services available.

Mobil unites for consultant itinerant teachers which contain instructional materials, teacher resources, diagnostic equipment, have been designed and are being considered for use in some rural areas to provide a traveling resource center or mobil classroom for hearing-impaired students.

The keys to unlocking effective programming seem to be the ability to design master plans at state, regional and local levels, which recognize the diverse needs of hearing-impaired children and which have provisions for an array of educational options.

This does not mean that every school district must provide every option, but that there needs to be some plan for making these options available to these students based on the individual needs of those students.

Without a doubt, making such

an array of options available involves all of those individuals that I mentioned at the beginning.

At the government level there may be a need for enabling or facilitating legislation. There is a need for physical support.

The administrative level, there is a need for flexibility to be able to engineer cooperative arrangements and agreements between districts across school boundaries, and perhaps even across state boundaries.

trained teachers, both at a preservice

level and an in-service level, and not just

better training for teachers who will be

working directly with the hearing impaired,

but with all teachers who may come in

contact with handicapped children.

There needs to be continued effort on the parts of parents and students to seek the most suitable educational opportunities for themselves.

The challenges of providing effective education for hearing-impaired students in rural areas remains.

The resources and the strength to meet those challenges must come from the continued and concentrated efforts of all individuals interested in the needs of the hearing impaired.

Thank you.

CHAIRMAN SCHWARTZBERG: Is Dr. McCartney here? Is Dr. Brian McCartney here?

Let me at this time hold, if

I may, the questions for Dr. Northcott and

for Dr. Conway, and call the remainder of

those on the Right to an Education, we will

take up questions for the entire panel on

the Right to an Education.

Let me at this time call
Ms. Celia Warshawsky.

Oh, Brian McCartney is here.

Ms. Warshawsky, could I ask you to sit down
for a moment.

Dr. McCartney is here at this point.

Dr. Brian McCartney was born with a severe to profound hearing loss and attended a day school for the deaf until he was mainstreamed at the age of 10.

He attended a private school for eight years and attended Susquehanna University where he majored in Latin.

He continued his education at Teachers College, Columbia University where he earned two master's degrees and his doctorate.

He has worked with hearingimpaired individuals from infants to adults and is currently the supervisor of secondary education at the Human Resources School in Albertson, New York.

The reason we have been calling his name is the material that he had indicated that he was not going to be called until 11:00 o'clock, and we are pleased that he is with us at this time.

Dr. Brian McCartney.

DR. McCARTNEY: Thank you very much. I have never been introduced so quickly, I

have never timed my entrance so precisely.

I come in a rather awkward position, not knowing what had been said before I came, but I will delve right into my subject, if I might.

The least restrictive environment needs to be, in my opinion, viewed from the viewpoint of those directly involved with it.

It is recommended that it be viewed from four different levels, specifically, the level of the school district personnel, the level of the direct services personnel, the level of the parent and the level of the hearing-impaired individual.

On the level of the school district personnel, each individual needs to acquire a basic understanding of hearing impairment and how it may affect an individual life. They need to be updated periodically on the developments in the field, and see how they can apply to the hearing-impaired population.

They should have professional contact with an individual in the field of education of the deaf who would be able to provide explanation or an insight into hearing impairment and its ramifications.

More importantly, the school district personnel needs to see that the hearing impaired that are individuals and therefore may each require his or her own educational program suited to meet his or her own needs.

School district personnel need to have a general knowledge of the various programs their district has to offer to determine what is available to the hearing-impaired individual.

In addition, they should be aware of those programs outside of their own district which may be "the least restrictive environment."

On the level of the district services personnel, specifically teachers and the various therapists, these individuals need to possess the desire to work with the

hearing impaired. Whether they be certified teachers of the deaf or certified subject area teachers, they are presented with the challenge of educating the hearing impaired in a number of areas.

In addition to the subject at hand, they assist the hearing impaired in functioning in society to the fullest extent possible, and assist society in better understanding the hearing impaired.

Along with this willingness to teach the hearing impaired direct services personnel need to be aware and updated on the various teaching techniques which may benefit the hearing impaired.

Direct service personnel

need to be able to administer tests and

interpret test results so that the hearing

impaired may be appropriately compared to

their hearing counterparts, consideration and

explanation, to be given to further pinpoint

the strengths and/or weaknesses of the

hearing-impaired individual on a given test,

since these test results are often the

deciding factor in determining the placement of a hearing-impaired individual.

The next level is the level of the parent, and there are five stages a parent goes through when they find out that their child is hearing impaired.

These are, according to Obgen and Lipschultz; shock, recognition, denial, acknowledgement and constructive action.

As the parents go through these various stages, they learn that they must in the end take on the responsibility of educating and accepting their hearing-impaired child.

They need to be realistic about their plan for him or her.

They need to see that their child faces an unusual challenge in life, and that he or she will require a great deal of support and understanding to do the best job possible.

Being realistic also means being flexible so that changes may be made

when necessary.

2

3

4

5

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

One least restrictive environment may be perfect for a given period of time, and then another may have to be sought.

Communicating with all persons involved with their hearing-impaired child is probably one of the greatest responsibilities of the parent of the hearing impaired. They need to be constantly in communication with a direct service personnel to monitor their child's progress and to create the best possible learning environment.

They also need to be in frequent contact with the school district personnel so that they are always aware of what is being done for and with their hearing-impaired child.

They need to be in daily communication with their hearing-impaired child to see to it that he or she is doing the best he or she is able.

This communication may provide

the parent and the child with a strong common bond so that they may together face the challenge of establishing the least restrictive environment.

Lastly, the least restrictive environment needs to viewed on the level of the hearing-impaired individual.

He or she needs to be aware of his or her hearing impairment and the restrictions it may place upon his or her life.

As the hearing-impaired individual interacts more and more with the world, he must face the obstacles placed before him and decide upon ways to overcome them.

One such obstacle for myself was my inability to define many words in the English language. I know my hearing impairment prevented me from acquiring language as my hearing counterparts, and I still was determined to overcome this somehow. My spirits were somewhat dampened when I took the SAT to enter undergraduate school.

The verbal was quite low, and I honestly had no solution at the time. Fortunately I was accepted at a fine undergraduate school, and as I reviewed the course offerings, I came across a solution, Latin.

I majored in Latin, and what a boost it was in my inability to deal with the English language.

No matter what educational setting the hearing-impaired individual is placed in, he or she needs to participate as much and as fully as possible in his world. This participation will enable him to grow to a wide variety of experience, and he will be in a position to get the most out of life if he is willing.

Perhaps one of the most important areas the hearing-impaired individual must focus on is the area I mentioned with the other level, and that is communication.

The hearing-impaired individual must communicate in order to get the most out of his situation, whether it be

a given lesson with the teacher about an event at a youth rally with some friends.

In communicating, the hearingimpaired individual is providing the
teacher, the parent or the friend with
feedback regarding a given situation, and
is then opening himself up for encouragement.

According to Arthur Simon, it is this encouragement that determines the direction of the life of the hearing impaired.

In conclusion, I would like

to say that the least restrictive environment

is, as pointed out, most difficult to

define in either words or interpretation.

It needs to be viewed on four different

levels, the level of the school district

personnel, the level of the direct services

personnel, the level of the parents, and the

level of the hearing impaired individual.

On each of these levels there are a number of areas to be addressed, all of which lead one to better define the least restrictive environment.

4 5

The point which needs to

be stressed here is that the least restrictive environment is different for each hearing-impaired individual. All parties involved in the selection of the least restrictive environment should be flexible and open as to what will best serve the needs of the hearing-impaired individual at a particular time.

Thank you very much.

CHAIRMAN SCHWARTZBERG: I am insistant to our participants that they cut their formal presentations down to ten minutes.

We have provided an almost impossible task.

There are two of us on
this panel who served as adjunct professors
of journalism, and at least one of them notes
your editing was utterly masterful.

Waht would the members of the Commission like to begin with for any of these three speakers? Yes, Preston Ewing?

MR. EWING: I believe it was Mr. Conway, we talked about the reasonable time set for the delivery of special education services.

MR. CONWAY: Yes.

MR. EWING: Recognizing under the regional concept that we face a problem of time and distance, but under the concept of equal educational opportunity it may be that for many of the hearing impaired the length of the school day, in order to give equal educational opportunity, has to be as long as the length of the school day for non-handicapped children.

What is your feeling on the issue of the maximum amount of time that should be spent en route to distant facilities on a daily basis?

MR. CONWAY: I'm not quite sure how to answer that one. Having had to ride buses myself at times, a lot depends on the age of the child, and also a lot depends upon the topigraphical features that have to be crossed in order to get the child to the services available.

In some states and in some locations that do have regional arrangements where transportation is a problem, these are

the areas in which the foster living and boarder programs have been initiated to decrease that amount of travel time that's involved.

The children come into the district sometimes within 10 or 15 minutes of the actual school that will be providing the services, and board with a family in that area, that is one option that's been approached.

As to a maximum time limit,

I really can't say. 45 minutes to an hour
is a long time for a five or six-year old
to be on a bus.

It's also a long time for a high school student to be on a bus.

MR. EWING: Have you found under the regional concept that some school days have been shortened because of the length of transportation?

MR. CONWAY: To my knowledge, that has happened in some programs, and in some regional arrangements that has been developed; however, there has been a backlash

of response to that based on the introductory statements you made to your question that needs to provide a school day that's as long for the hearing-impaired student as it is for regular students.

CHAIRMAN SCHWARTZBERG: Dr. Lucas?

DR. LUCAS: This question is for any member of the panel, the civil rights of the hearing-impaired person, we seem to have established a change as the knowledge about this handicap and the means to treat it change.

Whose responsibility, and specifically what governmental unit's responsibility is it to continue to the interpretation of the meaning of civil rights for the hearing impaired in the face of these changes?

DR. NORTHCOTT; Well, I think it
depends upon the monitoring team within
a state education agency, whether they are
brought into the U.S. Department of Education
periodically for an examination of the
drafts of Manual Ten which relate to the

implementation of Public Law 94142, the monitoring at the state level is supplemented, of course, by visits periodically from OSERS, the Office of Special Education and Rehabilitation Services, which means again that the interpretation of the rules.

advocates for a change related to something that everybody can relate and have information about, then it's fair game for saying that the program that's initiated within a district or coop or region is or is not living up to the spirit of the law and the spirit of the changes.

DR. LUCAS: If I may follow-up what we would have here is somewhat of a conflict or at least an interfacing of two areas of responsibility of the government, one is the civil rights, the securing of the basics as defined, basic rights of the individual, and the other the provision of services that may or may not be a right to provide.

Am I hearing you say that in this particular case these two elements of

law have to be cooperating because you seem to assign to the group the rehabilitation services, the task of defining not only the services, but also the rights, how would you limit the tasks of both groups of services?

DR. NORTHCOTT: Well, in my judgment, since it's education, the primary responsibility is within the state education agency, the district receiving the service.

The adjunct and the supplemental area of monitoring of civil rights comes in the due process procedures.

of a formal registration of objection to something that is being done locally or regionally, but I'm not an expert, and I don't think I can give you a better answer than that.

CHAIRMAN SCHWARTZBERG: Further questions?

Any other member of the panel, or any one of the three speakers who would like to comment further on Dr. Lucas's question?

MR. CONWAY: If I could follow-up what Dr. Northcott has said, the notion of the

interface between education and civil
rights has already been established and is
there when you look at Public Law 94142,
which deals with education, and in Section
504 of the Rehabilitation Act, which is
more civil rights legislation, yes, there
does have to be an interface and cooperation.

CHAIRMAN SCHWARTZBERG: Thank you.

## Dr. Spencer?

DR. SPENCER: This question is addressed to the teachers, I'm concerned, this might be just off the wall for you, on the impact of what is taught the child by the priorities set in the classroom environment, and elsewhere by the disability itself by hearing impairment.

My own observation has been that children who teach the handicapped, teachers who teach the handicapped are better educated and more dedicated to children and their real needs than the usual run-of-the-mill school teacher.

Does this also affect what is taught and the quality of what is taught,

because I think at least some of these children are denied the opportunity of watching television seven hours a day, which is what most American children have thrown at them, but there should be an advantage then in not having that exposure to the tube.

You might even be able to major in Latin in elementary school if you could be denied those things, but what about the quality of education they really get because of this?

MR. CONWAY: Being both a former teacher and still a teacher, and a teacher trainer, there is some truth to what you are saying, some valid points made there that not everyone can be a teacher of the hearing impaired or wants to be, nor should they be.

The teachers that do enter into the profession do seem to have some special qualities which make them want to be teachers of the hearing impaired.

Being a teacher trainer, I

also feel that we don't always do the best we can do in preparing our teachers to go out into the field and serve hearing-impaired children.

Hopefully that self-doubt is something that continues to make us evaluate what we are doing, and hopefully continue to improve the quality of teachers who are able to provide to the profession and thus the quality of education that's provided to hearing-impaired children.

CHAIRMAN SCHWARTZBERG: At this point,

I'm going to suggest that we break off
questioning of the panel, since there may
be some overlap between this panel and the
panel which is to follow, if the three of
you would continue to make yourselves
available, it may be that we will want to
have you participate in some of the questions
addressed to the board following.

Thank you very much. I'm sorry to the five who are to follow. Actually, that may be six.

For everyone's general

background, I'm not at all certain that we are going to complete the Right to an Education this morning, and that may well throw the Right to Education panels over to the afternoon.

For that purpose, is Mr. J.B. Davis here? Does anyone recognize Mr. J.B. Davis, and is he here?

Mr. Davis is supposed to be on at 3:00 o'clock. He had originally been invited to give a paper on technology in the work place.

The paper that has been submitted by Mr. Davis would appear to more clearly fall under the Right to an Education category, and it may be that we will carry that over first thing in the afternoon.

At this point I would like to call on Ms. Celia Warshawsky, and at the same time, Ms. Ellen Rhoades, Ms. Inez Janger, Mr. Virginia Stern, and Mr. Bruce Goldstein, would you come forward?

Celia Warshawsky has been hearing impaired since birth, and she notes

that she has "more or less dedicated her life to problems of hearing impairment."

Ms. Warshawsky received her

BA degree from Gallaudet College and undertook

additional course work at Northwestern

University and the University of Nebraska
Lincoln.

She was the first hearingimpaired teacher hired to teach in the
public schools in Illinois, and has been
honored many times for her devotion and
dedication to the disabled.

In 1983, she was named
"International Deaf Woman of the Year" by
Quota International, Inc. and "Frater of the
Year" by the National Fraternal Society
of the Deaf.

She was elected to the NFSD Hall of Fame.

In addition, she was honored by Illinois Parents of Hearing Impaired at their 1985 State Day as "an outstanding advocate for the educational rights of all hearing-impaired children in Illinois."

In 1986, she was inducted into the Hall of Fame of the National Congress of Jewish Deaf.

She has served on the Task

Force for the Illinois HB 1814 and was

instrumental in the passage of that recent

Illinois Act which provides free

telecommunication devices for severely

hearing-impaired people in Illinois.

She is currently on the Board of Directors of Chicago Hearing Society,
Advisory Board of the Department of
Rehabilitation Services, co-chairs the
Skokie Advisory Council on Disabilities,
the Advisory Council for Chicagoland Advocates
for Signed Theater, and is a member of the
Governers Executive Committee, Decade of
Persons with Disabilities.

She also serves as Educational Specialist for the Illinois Association of the Deaf.

Ms. Warshawsky.

MS. WARSHAWSKY: Thank you, Chairman, members of the panel and ladies and gentlemen.

a

The topic assigned to me -- can you hear me?

The topic assigned to my portion of the program is the rights of the hearing-impaired child at the earliest stages, including children from birth to five years of age.

Before addressing the specific age of the child, I would like to share with you a list of rights of hearing-impaired children of all ages.

The right to be myself, the right to know his own name, the right to communicate with his parents and receive love and understanding from them, the right to express myself in the manner in which I feel, which he feels most secure, the right to freedom from physical abuse, the right to special care so my condition does not become a handicap, the right to an appropriate education for his individual needs, the right to make his own decisions, the right to choose his own friends, the right to reach his full potential as a human

being and not equate the tension of a "normal child."

I'm going to have to condense my report, so I will just pick out selections.

The hearing-impaired child in a hearing family is at a high risk for communication deprivation and subsequent problems which attach.

With proper counseling and education with the family members, the young hearing-impaired child is afforded the opportunity to learn about the same aspects of his environment that a hearing child would pick up automatically.

More important, however,
is the opportunity provided through
communication to establish relationships
within the family. The implication of
this outline for today's program on rights
during early childhood is that the access
to the hearing-impaired baby's rights is
through a formal program, although a
formal program isn't necessarily equal.
In most cases the actual access to the young

Į

.

hearing-impaired child's education is not the formal program per se.

It is the child's development of communication since the child's access to education is through the development of communication, competence, it is essential for an early childhood program for the hearing impaired to have a thorough understanding of how the communication process is established, reinforced, maintained, expanded in all children.

Since the parent and child play an equal role in developing the communication, the parents' acceptance of the child and comfort in relating to the child is critical.

The first concern in the effective parent-infant intervention program is to help the family deal with the crisis of having a hearing-impaired child.

Effective parent counseling with other parents is a way to facilitate exploration of feelings and strategies for coping.

Much has been written about
the wording process that an association for
the child who is not meeting parental
expectations, it is important that the
process be recognized, acknowledged, allowed,
facilitated and/or understood.

A critical component in the acceptance process is a growing understanding of hearing impairment and the needs of hearing-impaired people.

A way to promote this understanding is through topics related to parent education programs prior to preschool years.

Parent counseling groups

may help remove parent feelings of

incompetence and loss of self-esteem upon

learning that the child is handicapped.

In a short period of time, needs are introduced to family-infant concepts. A formal education program will enable the parents over a period of time to integrate all this new information.

Up to this point I have

N

emphasized the parent avenues of parentinfant intervention process, a comprehensive
educational program for children is
essential to a quality program.

In order for families to work through the necessary emotional issues that lead to acceptance of the child's hearing impairment, the family members must have confidence in this child's educational program.

Optimal organization of parent-infant programs with regard to the children's needs would include individual sessions with parents and children together, as well as groups of children working with a teacher without the parents.

While the children are together the parents can have their counseling and/or educational program with the social worker and through a parent educator.

Since peer interaction plays an important part in the child's social development and subsequent communication

development, peer facilitation and reinforcements of communication is a critical component of a quality early intervention program for the hearing-impaired child.

In Illinois the biggest issue today regarding the rights of the parent, infant and preschool hearing-impaired children is what is currently happening politically at the state and federal levels.

a standard ingredient in public school and private programs for hearing-impaired children, however, public school programs have not been mandated and are therefore subject to severe budget cuts.

The programs that have survived have proven themselves effective inspite of the rigid budget cuts in the profession of comprehensive services for family and hearing-impaired children.

The state is currently studying comprehensive programming for all handicapped

children and their families from birth to five years of age. By January, 1989 the information gathered from the early childhood state plan grant and the pilot project model sites will be compiled and the State Board of Education, in a report to the General Assembly, the intent of the current effort to mandate birth to five-year programs in the State of Illinois is potentially beneficial to the hearing-impaired child and his family.

The potential danger to the hearing-impaired child with regard to the upcoming mandate is that the program from birth to five years will become noncategorical, which means all handicapped children will be placed together without regard for their physical impairment.

Most other groups of handicapped children speak and hear English and therefore are part of the community and culture of the hearing majority.

The mainstreaming of these handicapped groups of children is chiefly

done by creating some physical access for them to enter regular programs.

Physical access is not the deaf child's problem, his problem is understanding of the basic language communication when one is placed in the group.

The family of the hearingimpaired infant and preschool child, as well
as the child himself, has the right to have
verbalized and comprehensive services
funding at an adequate level to meet the
specific needs of the population.

I would like to make some recommendations, one, mandate for free public and appropriate education from birth to five years of age for hearing-impaired children.

Two, representation from the deaf community on a state-wide task force in decision making.

Three, insure through certification requirements in-depth education with all the components, early childhood education, counseling and administration of

early childhood and so forth.

Four, discplinary team

effort where all staff work together at

agencies such as medicals, social, emotional

and educational also work together and

on-going research and education in the

field of preschool education with focus on

which models work best and why, and how

deaf babies learn language.

CHAIRMAN SCHWARTZBERG: Now, turning to the next speaker, who is Ms. Ellen Rhoades out of Atlanta, California -- I'm sorry, Atlanta, Georgia. I think really that might well be a sign for us to stand up for about ten seconds so at least the court reporter can rest her fingers for that long.

(Whereupon a short recess was had.)

CHAIRMAN SCHWARTZBERG: Ms. Ellen A.

Rhoades is the Founder and Executive Director
of Auditory Educational Clinic in Atlanta,
Georgia and has been such since 1977 to the
present.

She is Project writer/director

to the UNIsensory Project for HCEEP from

1980 to 1983, a member of the Board of

Directors of the A.G. Bell Association,

a member of the Executive Board of the

Auditory Verbal International, a member of
the Citizens Advisory Board for Southern

Bell Telephone.

She is Children's Rights
Coordinator of Georgia for A.G. Bell.

She has had public and private teaching experience with oral deaf children from infancy to 16 years of age, and has congenital bilateral severe deafness and a defined d.b.

At this point I would like to call on Ellen A. Rhoades.

MS. RHOADES: Thank you, Honorable Chairman and distinguished members of the panel.

My topic is the rights of hearing impaired children to early intervention.

The research basis for early intervention is ample. There is no doubt whatsoever that our efforts to improve

the lives of imperiled children and their families is affected.

We have well known this since the pioneering work of Skiles, Bloom, Hunt, Kirk, Riekirk, Gray and Klauss.

It was these very pioneers who in the 1930's and 1960's gave us such data as the Iowa study and the working project and the Prairie Preschool Project.

Because of them, early intervention has been substantiated, we need no longer defend nor should we the right of early intervention.

In fact, it's no longer debatable that the most successful federal program of our generation has been and is the Handicapped Children's Early Education Program, or first-chance network.

To such federally funded programs we know that handicapped children reap immediate and long-term gains as does our society.

Today the right question to ask is how, how shall we better provide

early intervention and with what tools, and focusing specifically on the hearing-impaired child, the prenatal deaf child, if you will. We need to review gathered research.

The 1973 survey data, it indicates that the typical deaf adolescent has spent ten years within a self-contained cell, be it a school or a class for the hearing impaired, that this deaf person is either functionally illiterate or had a third-grade reading level, that he uses sign language as a primary means of communication, and does not have intelligent speech, and finally, this costs the tax payer between \$6,000 to \$15,000 per nine months school year.

This survey data is based on oral and manual deaf students identified by schools for the deaf.

If we then review our survey data of 20 years ago, specifically in 1964, we shockingly observed no all-over improvements for deaf children, yet we know early

intervention programs have been implemented in the last 20 years.

Does this then mean that the deaf child does not significantly profit from early intervention? We all know better than to ask such a question.

However, perhaps we need to delve more deeply into the how of early intervention. In 1947, most deaf children did not use amplification, and when the few did, it was for only a portion of each day, however, a handful of clinicians or teachers believed that the deaf could somehow learn to hear.

One of them as a doctor who put hearing aids on two and three-year old deaf children. This experimental program, federally funded, was first initiated the practice of reverse mainstreaming, it consisted of early amplification.

My brother and I were a part of this three-year program. Short-term results of that project were published in a book entitled Deaf Children in a Hearing

World. We were two of twelve case studies.

20

21

22

23

24

Around the same time, also in the '40s, in Pennsylvania and in New York City and also in Denver, this approach was being implemented to teach deaf children basic listening skills. For a year these teachers struggled along with the belief that the deaf can hear. It had been considered ridiculous in the '50s to, number one, try putting hearing aids on babies, two, to try and buy oral amplification, three, to deprive deaf children of visual means of communication, therefore dooming them to a life of frustration, and four, to mainstream their children into a hearing a world, and therefore denying their deafness.

But their efforts were not in vain. Ever so slowly, one by one, a few more deaf children overcame their handicap and learned to function at a hearing college.

A few more families were helped, and many costly difficulties were avoided, but more importantly, a slight ripple effect began to occur among our

)

profession.

It began to include auditory training as part of the curriculum in the '60s and in the '70s, we began to see a change in the terminology of teaching methods.

Manual communication advocates recognized their limitation, and the total communication aspect was born.

Oral communication advocates also recognized their own limitations, and the auditory oral concept was born.

Auditory training had come of age, but this was not enough. The survey coming out of Gallaudet told us so.

The auditory verbal pioneers recognized the dangers of auditory verbal approaches being misinterpreted, so joined together a network to advocate teaching and hopefully to gain strength in numbers.

They founded the auditory special committee, which is the special committee of the A.G. Bell Association for the Deaf.

.

This committee was and remains committed to the right of every hearing-impaired child to become a functional member of our hearing society.

We are committed to the provision of equal opportunity to the first option of learning to listen. We do not claim that every deaf child can learn to listen well enough to use the telephone without needing special education services.

We do claim, however, that at least, at the very least, 50 percent of our deaf children, if given the opportunity, can become functionally part of hearing and productive tax paying citizens.

We espouse the very basic principles of one early consistent and effectively powered amplification, two, one-on-one individualized auditory verbal communication training sessions, three, parents are fully informed, on-going and active partners, four, forming screening for every preschooler, and five, a realistically high expectation level that the young deaf

can hear.

hour.

These principles mean. that the deaf infant is trained every working

We focus on the infant learning to listen, not reading. The pioneers realized that auditory training cannot be maximally effective if it is merely a 30-minute exercise.

In addition, the expectation of hearing must be a way of life. Listening schools must be integrated into the total personality development of the deaf infant if the deaf infant is to overcome his deafness and become assimilated into our regular education programs.

The studies we have on the auditory verbal approach are unequivocal in demonstrating that deaf children indeed can learn to hear and understand.

We have normal auditory
educational scales, therefore we assess this
approach with the use of evaluation instruments
which have been normal and standardized with

the normal hearing population.

We feel the use of such assessment is more likely to give an accurate portrayal of achievement. We already know that the deaf tend to fail rather badly.

Panek's study had demonstrated that in a given year on 49 subjects, 95 percent of those children developed auditory verbal communication skills. 50 percent of these children were successfully mainstreamed at that time with those who were in special education classes, also considered to be multi-handicapped.

In a follow-up study of 422, they found 85 percent still used speech as a primary means of communication. 76 percent were mainstream, 41 percent were multiply handicapped. Of the 10 adults in this study, 7 were college graduates, and all but one were gainfully employed.

When the auditory educational clinic was part of the first transnetwork we saw 29 children on a weekly basis in three years as typical of a small sample

size were too many assumptions being violated, the resultant data was necessarily descriptive in nature. Case studies did indicate that the degree of deafness did not determine the success or appropriateness of the auditory verbal approach to the children.

We also were able to determine that a cost \$3,000 to \$4,000 per a 12-month school year over three to five preschool year period to fully assimilate a deaf child into a hearing environment. The tax savings are tremendous.

Unfortunately, the number of children being trained through the auditory verbal approach are not yet legion.

We have small samples, we are not accepted by the teaching profession at large. We need to support higher education programs which provide for the professional training of auditory verbal clinicians, be they teachers, speech technicians or audiologists, because unfortunately we no longer have any such appropriate clinicians and training programs in this country.

~

Well, with our support and our belief that the deaf children need not go in silence, we will get larger samples, and with the auditory verbal approach we can indeed fulfil our primary reasons for early intervention of deaf children. Once we can enhance the child's development so that he hears, we can provide support and assistance to each preschool child's parents, and we can enhance the child's and family's benefit to society, and finally, we can avoid some or all of the costly difficulties which accrue when intervention is delayed.

Thank you.

CHAIRMAN SCHWARTZBERG: Thank you. The next speaker is Ms. Inez Janger.

From 1961 to 1969, Inex Janger, as a parent, chose to stay at home to raise a family.

However, various activities, such as the League of Women Voters and civic advisory groups enhanced her skills in community organization, public speaking and lobbying.

6

8

10

11

12

13

14

16

17

18

19

15

20

21

22

23

24

From 1959 to 1961 she was personnel supvervisor for Crowell-Collier, MacMillan, Inc. and Hayden Publishing Company, and she secured her BS degree from Simmons College, and MS from Cornell University as a Candidate in Personnel Administration, and additional work at Columbia University, New York, graduate work in law, finance, education and administration.

With Citibank, she completed courses in compensation, staff relations, EEO, Position Evaluation, Management Process, Consulting Skills and Targeted Selection.

Therefore it is perhaps not surprising that she became President of the Board of Education, for Hastings-on-Hudson, New York, where she served as a member from 1979 through 1985, as President from 1983 to 1985.

She is a member of the Executive Committee of the International Parents Organization of Alexander Graham Bell Association for the Deaf and is Past President of the New York State Parents of

Hearing-Impaired Children.

Ms. Inez Janger.

MS. JANGER: I have been asked today to address specifically the elementary and secondary school years.

I want to focus on the educational decisions made for deaf and hearing impaired or hearing-impaired children, and how they could but often do not comply with the current laws and the rights of hearing-impaired children to grow, communicate, live and work in the hearing world.

You have heard much earlier today about how research proves that it can be done, and during the past few days here at this convention we have seen the products of education that allows this to happen.

A young, hard-of-hearing child who has already begun to develop speech at home is placed in a public school kindergarten program out of district, self-contained except for recess, using total communication. Her parents are told that

nothing else is available, despite their clear desire to have her develop speech and liberating skills.

This child quickly begins socializing with her deaf classmates and drops all attempts to communicate orally with the rest of the kids in school.

Oral language development

begins to decline noticeably at home. Another

child who lives, reads and speaks well is

mainstreamed in a regular class through

sixth grade, functions quite well at grade

level, and upon graduation to junior high

is placed in a self-contained class for

the deaf.

The school says he won't be able to keep up because the school program is now departmentalized and there is no support for the regular academic teachers.

Or another version of the same story, and I have heard this repeated numerous times these past few days, a similar child oral functions extremely well in the mainstream, is placed in junior high school

in regular classes with a sign interpreter and grouped with several other non-oral students in academic classes, although there has been no request for this nor was it in his IEP, which is the prescriptive vehicle for special services, supposedly, according to the child's needs.

In each of these situations, and I could go on and list many others, the parents, however, were able to apply common sense and political pressure and use the IEP to bring about program changes and services more consistent with their child's needs.

They wouldn't accept established patterns. They wouldn't accept what the bureaucracy was telling them.

In the first case, the parent convinced her local school board to place the child in a regular kindergarten in her home school, and to find an itinerant tutor to work with the child.

The parent was able to show that aid was available to this service, the

child did not have to be in a special class in order to save the school district money.

Aid was available as long as the IEP indicated it as a need for the child.

In addition, this parent hires a private therapist to work with the child.

In the next situation, the parent continued to browbeat the local placement committee to change the IEP to allow regular classroom attendance with appropriate subject matter tutoring and an hour a day with special language therapy.

The IEP now indicates that all academic subjects will be taught in regular classes using regular school curriculum and support will be provided by the regular classroom teacher if requested.

Realistically, however, much of this support has turned out to be provided mainly by this practical, educated and articulate parent, not by the school. The child, who had a sign interpreter, succeeded in getting placed in different sections of

the same class, and now uses voluntary note takers and extra tutoring after school.

These in fact are all success stories. These parents have succeeded in breaking through the red tape, have been highly articulate and quite sure of their objectives.

They have been pushy and bothersome, they have pulled heart strings, they have cried, they pulled political strings, and they have been well prepared, much more knowledgeable than anyone else involved about what their child really needed.

One can wonder, in the face of this, what the law, the new laws have really changed. These kinds of parents have always been able to beat the system with or without IEPs, with personal and often financial resources.

They have insured the rights of their children, and we have seen many examples again of these children in the last few days here at the conference, but

,

what about the parents who can't do it, who lack the energy, the money, the knowledge and time to move bureaucracies they come up against.

People who cannot educate the uniformed bureacrats who cannot out maneuver them or threaten legal action and then deliver it when necessary. For parents without resources, these same stories might have ended like this, the five-year old hard of hearing child who has been speaking at home begins to perform as if she were functionally deaf, unaccustomed to using her considerable residual hearing and speech, placed in an environment with limited oral communication and lowered expectations for language development.

The child performs as expected, a star among the deafer kids, but far behind her hearing peers.

The seventh grader who had been successfully mainstreamed through sixth grade, following the simplified curriculum of self-contained class and does not have the

background to move back into the mainstream eighth grade class, so he's mainstreamed into a slower-moving class for kids with learning problems, with non-categorical classes, I guess the term is now, and unfortunately does not get a chance to move back into the mainstream.

bans together with the non-oral kids, learns sign and a subsection of the hearing class if formed. All communication between the deaf kids and the teacher is through the interpreter. The teaching ends up being done by the interpreter, not the subject matter teacher.

In all these cases, the schools use the individual education plan, the IEP, to describe what was available, not what the individual student needed.

The IEP became a uniform curriculum and placement tool for any deaf child rather than a plan for the unique needs of the child at a particular point in time.

Much to my surprise, I have

learned these last few days that these endings are in fact happening with increased frequency to parents who one would have thought, because of their determination, could have moved the system, the bureaucracies are becoming more entrenched in many places in the United States.

What Mr. Regan had to say about the hungry and the homeless in the United States was also true for the hearing impaired and their parents. The services are out there to be had, but it takes a fair amount of knowledge, drive and luck to make them available for our children, more than it should be, more than it should take in a reasonably civilized society.

Given this situation, what should we, as parents, be asking for?

There are no easy answers.

The task can be left to legal redress, which is in fact where we are today, witness the number of cases I have heard about the last few days.

The laws exist, and it's up

to the parents to make them work. Of course, that takes knowledge, resources, great determination, and extracts an enormous emotional toll on the whole family unit, including the hearing-impaired child.

Moreover, it limits the benefits of the law and services to the very few, or there can be substantial improvement in parent-child advocacy, I believe, utterly professional and nonpartisian.

All professionals having contact with the child and parents must be aware, must be mandated to help parents become aware of options for oral skills development for mainstreaming, and for a variety of services which can and have been used in the past to help children develop their full potential.

There must be a public awareness about options, services, about the fact that hearing-impaired children can learn to speak, live and read in the real world, can go to hearing colleges, work in

-

o

major corporations, be lawyers and doctors, be elected to public office.

Public information targeted at the medical communities is also crucial. It is, after all, the medical communities where the first diagnostic contact is often made.

The lack of knowledge for the potential for speech, hearing and language development often leads to careless referrals and provision of limited information or worse, misinformation, to parents.

When a pregnant woman goes
to her obstetrician, she is bombarded with
literature about babies, child rearing,
and preschool education, information about
services, options, potential information
about services, options and the potential
for speech reading, speech and lip reading,
and useful references should be readily
available at pediatricians, in the audiology
clinics, and hearing aid dealers offices,
parents should be bombarded with the
necessary information for deciding on methods,

[4

programs, assistive devices, etc.

Who should provide this public information, where are the advocates, where is the funding, that's the question I don't have the answer for, and I will leave that to you.

Non-profit voluntary advocacy agencies like A.G. Bell simply do not have the resources, government-funded agencies very often tend to have a bias and a nitch or geographic base only.

They may be only interested in career training and finding out anything from that agency about elementary education may be impossible.

Our concern is for all parents, all locations, all ages of children with the laws, knowledge and the will, they can get what they need.

Without the knowledge, the laws and the will, the advancements in services haven't changed very much for many of our children, their right to learn to speak intelligibly, to communicate and live

in the hearing world, these rights are still unfortunately being denied.

CHAIRMAN SCHWARTZBERG: Because some of the members of the audience are scheduled to appear at a luncheon that begins at noon, we are going to take our break at this point.

We will, however, attempt to start a little bit earlier, at 1:15, so if you will please be back here at 1:15, we will break at this point.

(Whereupon the proceedings were adjourned until June 30, 1986, at 1:15 p.m.)

STATE OF ILLINOIS )
) SS:
COUNTY OF COOK )

COLETTE M. KUEMMETH, being first duly sworn, says that she is a court reporter doing business in the City of Chicago, and that she reported in shorthand the proceedings had at the hearing of said cause, and the foregoing is a true and correct transcript of her shorthand notes, so taken as aforesaid.

Colette M. Kummeth

to before me this day of August, 1986.

Notary Public

1	THE RIGHTS OF HEARING-IMPAIRED PERSONS
2	A Public Forum
3	
4	Illinois Advisory Committee U.S. Commission on Civil Rights
5	
6	In Cooperation With
7	The Section of Individual Rights and
8	Responsibilities of The American Bar Association and The State of Illinois
9	Department of Human Rights
10	
11	Hilton Hotel and Towers
12	Williford A Room Chicago, Illinois
13	
14	Monday, June 30, 1986
15	
16	
17	
18	
19	
20	
21	
22	
23	
2.4	

1	BEFORE:	
2		MR. HUGH J. SCHWARTZBERG Chairman
3		MS. THERESA F. CUMMINGS
4		
5		MR. ROBERT C. SPENCER
6		MR. ISIDRO LUCAS
7		MR. JOHN LINGNER
8		MS. ERMA M. DAVIS
9		MR. THOMAS PUGH
10		MR. PRESTON E. EWING
11		
12		
13		
14		
15		
16		
17		
18		
19		
20		
21		
22		
23		
24		

1	MR. SCHWARTZBERG: Let me call the
2	session back to order, and at this time I
3	would like the following people to come
4	forward: Virginia Stern, Bruce Goldstein, and
5	Mr. J.B. Davis.
6	For those of you who may not have
7	been present at the morning session, in nearly
8	all cases extensive papers have been presented
9	to the Commission and to the Advisory
10	Committee, and summaries are being given here
11	of not to exceed 10 minutes.
12	I am sorry to have to limit it to
13	10 minutes because of inconvenience to the
14	speakers, but also because it may give those
15	who are here much less of a flavor of what
16	are, in many cases, just very fine and highly
17	educated submissions.
18	You will find in your program
19	that Mr. Davis appears on the right to
20	employment technology in the work place.
21	He has submitted a paper
22	marked closely right to an education; and
23	we have therefore added him in at this time.
24	I would like to let me call on

1 Mr. Davis at this point. Mr. Davis was elected Vice 2 3 President of the Chicago Chapter, NFSD at 19; 4 for 25 years he has served as Grand Trustee, 5 NFSD; and for 18 years he has been Law Chairman of GLDBA; President of the Illinois 6 7 Association of the Deaf for 11 years, and is a 8 charter member of the Chicago Club for the 9 Deaf. 10 He has been active in legislation 11 on deaf driver rights, on TDDs being required 12 in county sheriff offices, and in the recent securing of legislation to provide TDDs for 13 14 deaf people in Illinois. 15 He serves as Chairman of the 16 Board of the Silent Coperative Apts. and has served on many committees in education, and 17 18 has for three years been Chairman of the State 19 Board of Rehabilitation. 20 He has been a delegate to the 21 National Association for the Deaf three times. 22 Mr. J.B. Davis. 23 MR. DAVIS: Thank you.

My name is John B. Davis.

1	I was born deaf in Evanston.
2	As a result of my background, it
3	has been real jumbled.
4	I have gone to different
5	education experiences.
6	I spent ll years in a strict oral
7	environment in Chicago in the public schools,
8	four years in mainstreaming, two hearing high
9	schools with no interpreters and no
10	notetakers, one year in a residential school
11	and three years at Gallaudet College, both
12	with the use of total communication.
13	But most of my education stems
14	from the Evanston library, as I was an avid
15	reader of books at an early age.
16	At age 18 I entered a deaf
17	entered the deaf community and started using
18	total communication through sign language.
19	When my school days ended I
20	became involved in sports and club activities.
21	I also joined the NFSD and the
22	IAD, the Illinois Association of the Deaf.
23	I am speaking today about the
24	establishment of my own school experiences,

1	what I have learned from the experiences in
2	the Association of the Deaf education, and on
3	my own perspective of meeting deaf people in
4	all walks of life, and talking with them about
5	their school experiences.
6	As a long team leader of the IAD,
7	I began to be concerned about the slow
8	deterioration of the Illinois School for the
9	Deaf; that deterioration is going on in other
10	state schools also; that is the effects of
11	PL42142 that supports mainstreaming of
12	handicapped children in regular school
13	classes; that bill really ruined IAD's
14	enrollment.
15	Madaline Will (phonetic),
16	Assistant Secretary of Education and a mother
17	of a Down's syndrome child, is responsible for
18	this legislation that caused the state schools
19	to deteriorate throughout the United States.
20	She slowed the process of deaf
21	education starting in 1813 with the Hopkins
22	with Thomas Hopkins Gallaudet establishing the
23	first state school at Hartford, Connecticut.
24	Since 1975 and PL94142 the

1	percentage of deaf children attending state
2	schools has dropped to 30 percent, while 70
3	are attending day programs in their home
4	communities.
5	Each special education director
6	operates his or her area as a one out of
7	about 600 special education directors has a
8	degree in deaf education in Illinois.
9	There are about 25 coordinators
10	of hearing impaired programs compared to about
11	2500 administrators in special education,
12	including the coordinators.
13	The expertise of people in deaf
14	education just isn't being applied to the
15	education of deaf children, yet the special
16	education expect all deaf children to make
17	complete and rapid progress side by side with
18	their hearing peers.
19	Valuable communication time is
20	lost in bus trips from home to school twice a
21	day.
22	At home after school, the child
23	often has no one to play with.

State schools offer a deaf child

1	more hours of learning by encouraging
2	communication 12 to 15 hours a day in the
3	cafeteria, dormitory, classroom, vocational
4	shop, and playground.
5	This compares to a day school
6	with only seven hours and often none at home.
7	In most cases the child is even
8	isolated in his own family.
9	It is a fantasy to expect a deaf
10	child after mainstreaming in school to become
11	normal and be swallowed in the hearing world
12	as an adult, to marry a hearing spouse and
13	live happily thereafter.
14	No traditional school can match a
15	state school in the services offered deaf
16	children.
17	These services include psychology
18	services, audiology, speech therapy,
19	counseling, apparent education support groups,
20	language skills development, occupational
21	therapy, physical therapy, and otolaryngology
22	services.
23	What is particularly disturbing

is the concept of least restrictive

1	environment that has been applied to the local
2	neighborhood school.
3	While the state school is seen as
4	the most restrictive environment local
5	educators have joyfully accepted LRE, least
6	restricted environment, and have expanded
7	their programs to absorb deaf children at the
8	expense of state schools.
9	Local directors play the numbers
10	game, the more deaf children in a program the
11	more money they receive; thus an administrator
12	is very reluctant to release a child to
13	Illinois State School for the Deaf.
14	While a provision exists for
15	parents to seek redress in the courts to
16	overrule the decision of a director, very few
17	parents have the determination, money, and the
18	guts to outlast a director in the courts where
19	cases can be postponed time and again.
20	To me this is a gross civil
21	rights abuse and is at the expense of all deaf
22	children now and in the future.
23	Parents as taxpayers whose money

help support state schools are denied the

1	right to enroll their deaf children in those
2	schools.
3	This is another civil rights
4	abuse in taxation without representation.
5	We deaf people and concerned
6	parents must work together to be more forceful
7	in demanding that Congress and state
8	governments change this law by removing the
9	deaf from PL94142; growing number of deaf
10	educational professionals are ready and eager
11	to serve in administrative positions and to be
12	available in consultive capacities.
13	It is a waste of time to talk to
14	special education directors who know nothing
15	about deafness and are experts in double-talk.
16	Legislation is the best source of
17	redress.
18	That's where the power is; it is
19	our civil right to do so.
20	MR. SCHWARTZBERG: Thank you, Mr.
21	Davis.
22	The next speaker is Miss Virginia
23	Stern, the American Association for the

Advancement of Science.

1	Miss Stern, let me first
2	introduce you.
3	Virginia Stern is Co-Director of
4	the Project on Science, Technology, and
5	Disability of the American Association for the
6	Advancement of Science with headquarters in
7	Washington, D.C.
8	The Project is a national center
9	for information about education, careers, and
10	technology for disabled people in science,
11	math, and engineering.
12	She has conducted workshops and
13	conferences throughout the country on access
14	to education at the pre-college and
15	post-secondary levels, and has published in
16	these areas.
17	Before coming to AAAS, Stern was
18	editor of "Ideas for Families", a national
19	publication for parents of the Lexington
20	School for the Deaf in New York.
21	She is a former board member of
22	the A.G. Bell Assocation for the Deaf, and
23	currently serves on the Executive Board of the

International Parents Association of the A.G.

1	Bell Association.
2	She holds an M.A. degree in Deaf
3	Education from Gallaudet College 1971 and a
4	B.A. degree in History from Stanford
5	University 1955.
6	Mrs. Stern is the mother of four
7	children, three of whom have completed
8	undergraduate degrees, including a profoundly
9	deaf son who has just been awarded a Bachelor
10	of Science from Stanford University.
11	Virginia Stern.
12	MS. STERN: Thank you, Mr.
13	Schwartzberg.
14	The simple fact is that a hearing
15	impaired child born in this country before
16	1960 had limited prospects of higher
17	education, and then primarily in a special,
18	segregated setting.
19	It is true that a small number of
20	talented deaf individuals set their goals
21	independent of prevailing expectations and
22	succeeded in getting admission in and
23	graduating from the college of their choice

24

Some continued to graduate school

1 and professional credentials. 2 But if these hearing impaired individuals did not have a superior academic 3 4 record and an extraordinary family, there was 5 almost no one in the education community at large who would say that their goal was 6 7 possible. 8 A hearing impaired child born after 1960, that is, turning 18 after 1978, 9 10 when the 504 regulations were first 11 implemented, lives in a different world. 12 Today's world, under that 504 13 law, offers a hearing impaired student the 14 same opportunity as his or her hearing peers, 15 the same right to a higher education, the same right to individual choice on where and in 16 17 what setting that education might take place. 18 It is now considered right and 19 reasonable for hearing impaired persons to 20 compete with those not hearing impaired. 21 However, that right cannot be 22 exercised without information, and the right 23 to that information must be protected.

Under the law, as it is written

1	and as it is being upheld in a startling
2	number and variety of settings, disability,
3	and that included hearing impairment, can't be
4	used to discriminate against a student in the
5	college admissions process.
6	Colleges and universities are
7	required to provide any reasonable
В	accommodation that may be necessary for a
9	disabled student to have equal access to
LO	programs and services available to nondisabled
11	students, if the disabled students request
12	them.
13	Please note that the college is
14	under no obligation to seek out disabled
15	students who do not identify themselves to see
16	if there is something they might need.
17	And I might add here that there
1.8	are some very unreasonable expectations in
1.9	interpretation of the law where individuals
20	feel that they will go up to the college of
21	their choice and that they will be greated at
22	the door by six interpretors in black ties who
23	say "Ready to serve you," doesn't really work
2.4	that want the corpicat have to be developed

7	and worked out as a cooperative venture.
2	A broad range of options now
3	exists for hearing impaired students to attend
4	and complete courses of studies at colleges
5	and universities throughout the country, in
6	any field and at any level of achievement.
7	However, for hearing impaired
8	students public information and counsel given
9	to students making choices has not caught up
10	with reality; thus the choice is not there for
11	all hearing impaired students because they are
12	not being told about alternate paths.
13	We can get some idea of the
14	impact of the law from the statistics of the
15	higher education community, and some other
16	ideas from anecdotes of direct experience.
17	I am quoting first from the
18	American Freshman National Norms which are an
19	annual national longitudinal study of freshmen
20	in college.
21	1978 was the first year in which
22	any question was asked about disability.
23	In 1978 2.7 % of the entering
24	freshman classified themselves as being

1 disabled. 2 In fall 1985 there was a three-fold increase where 7.7 % of the 3 entering freshman who were disabled. 4 5 Now, these are the freshman all 6 over the country so you can see the impact of 7 the law that has trippled the number of freshman who are entering colleges and 8 9 universities. Among disabled freshman the 10 11 percentage of hearing impaired students 12 remained relatively stable: in 1978 it was 11 13 percent and in 1985 11.7. 14 Well, where do these hearing 15 impaired students get their education? 16 The most popularly known sites 17 are the federally-funded programs designed 18 specifically for hearing impaired students, 19 Gallaudet College in Washington, D.C., which 20 has a student population of about 1500, and 21 NTID, the National Technical Institute for the 22 Deaf, within the Rochester Institute of 23 Technology, with enrollment of about 1300.

These are both national programs.

1	They were established before the
2	504 legislation with the specific mission of
3	educating deaf students, in addition to
4	providing a post-secondary education.
5	They do a lot of outreach
6	service, train interpreters, develop technical
7	knowledge and specialized curriculum.
8	The Federal Government also gives
9	direct support to four post-secondary programs
10	specifically for deaf students.
11	These are located in different
12	regions of the country, and have a combined
13	enrollment of about 500.
14	Earlier this morning you heard
15	quoted from a resource directory, College and
16	Career Programs for Deaf Students, which was
17	quoted in the first presentation by Doctor
18	Castle, the new 1986 edition.
19	This directory, in addition to
20	the six federally-funded programs lists 136
21	other post-secondary institutions which
22	currently offer what is called "programs" for
23	deaf students.
24	They have more than 15 students

They have more than 15 students

1 and they meet certain guidelines. This is an excellent document, 2 3 however the six federally-funded programs and the 136 other programs listed in it represent 4 5 only slightly more than 10 percent of the 6 sites in the United States at which hearing 7 impaired students are now receiving a higher 8 education. 9 The second edition of the Directory of College Facilities and Services 10 11 for the Disabled, which has also been 12 published in 1986 by the Oryx Press, surveys 2300 colleges and universities throughout the 13 14 country on how they serve disabled students. 15 This volume lists more than 1300 16 post-secondary institutions that are currently serving hearing impaired students. 17 18 Now, they do not call themselves programs, you understand, but they are 19 20 offering a rage of support services which is 21 comparable to that offered by schools 22 considered to have programs. 23 These services include oral and

manual interpreters, notetakers, paid or

1	volunteer, visual alarm systems, TDDs,
2	captioned television, speech therapy, free or
3	at a fee, tutoring, counseling, career
4	services, and accomodation for all activities
5	on campus.
6	To put this statistic in context,
7	you have to realize that the all the
8	post-secondary schools in the United States
9	which are certified by the Department of
10	Education that total number is 3300, now
11	1300 of these, one third, are serving in some
12	way the nation's hearing impaired students,
13	not two, not six, not 136, but 1300.
14	In order to underline this
15	statistic with some direct communication and
16	bring it alive in an absolutely up-to-date
17	fashion, I contacted about 100 colleges and
18	universities throughout the country.
19	I wrote them a letter on the 4th
20	of June and told them that I would speak to
21	you today and asked them what they were doing
22	in providing services with hearing impaired
23	students on their campus; and I got 56 people
2.4	have answered me up to this morning

1	You may be interested to know
2	almost all of them are providing notetakers, a
3	large percentage are providing sign
4	interpreters, somewhat less but still very
5	many are providing oral interpreters.
6	Almost all have TDDs or visual
7	alarm systems.
8	Many have captioned televisions
9	and offer other services such as speech
10	therapy, tutoring, counseling.
11	These are not obscure
12	institutions.
13	To give you an example of the
14	sort of response our office received, this is
15	right now in this year, NYU has 30 hearing
16	impaired students; Boston University has 10;
17	Minnesota has 40; Emory has 4; Purdue has 11;
18	Oberlin has 5; Notre Dame has 12; Northeastern
19	in Boston has 35; and UC/Berkeley has 20.
20	Now, to put a little more direct
21	experience in this survey, I made a site visit
22	to one of the institutions, the University of
23	Minnesota in Minneapolis, St. Paul, less than
24	a week ago on my way to Chicago.

7	The University of Minnesota
2	serves 30,000 students in total during the
3	semesters, 600 disabled students on campus, 40
4	of these students are hearing impaired, 20 of
5	them do not require interpreters in classes,
6	20 of them do.
7	Both oral and sign interpreters
8	are available upon request.
9	Bringing this information about
10	available options back to focus on the
11	individual hearing impaired students of
12	precollege age, I would like to quote Rhona
13	Hartman, Director of the HEATH Resource
14	Center, the National Clearinghouse on
15	Postsecondary Education for Handicapped
16	Persons.
17	She reports that every week the
18	Clearinghouse receives a call from a counselor
19	or perhaps a parent saying "I have a student
20	who is deaf, where can he go to school?"
21	And I would propose that this is
22	an inappropriate question.
23	It can't be answered because it
24	makes the assumption that the design to

1	attend a given institution should be based on
2	one's disability; that puts the emphasis on
3	the status of being hearing impaired rather
4	than the status of being an academically
5	qualified student who happens to have a
6	hearing impairment.
7	If you are interested in denistry
8	and you are qualified to be a dentist, you go
9	to a program that teaches denistry, not a
10	program for the deaf.
11	If you are interested in
12	anthropology and have the skills to pursue
13	that science, you might choose to go right
14	here to the University of Chicago; that is
15	what you choose first is the excellence of the
16	program.
17	If you are interested in
18	journalism, you might choose to go to the
19	University of Missouri because of the
20	excellence of their program, not because of
21	their services for the deaf.
22	And then you work with the
23	service providers to receive the services that
24	you need that belong to the student under the

1	law.
2	The question remains how widely
3	known is this information?
4	When a hearing impaired student
5	meets with high school guidance counselor or
6	vocational rehab counselor, do they have
7	access to this information?
8	Do the people who guide the
9	hearing impaired students understand the law?
10	Do they understand that they
11	that the students can exercise their
12	educational choices?
13	A school does not have to have a
14	special program, it can develop services; and,
15	of course, it makes only it only makes
16	sense that the hearing impaired student work
17	with the college to develop those services.
18	We must ask the question: Does
19	the public relations network channel many
20	students into a limited number of choices when
21	in fact they may have a very wide number of
22	choices?
23	Does the vocational rehab system

in certain states favor a very narrow range of

1	options?
2	Even within colleges that offer
3	support systems, is there a channeling of
4	hearing impaired students to take certain
5	courses so that all hearing impaired students
6	will be conveniently grouped rather than
7	following their intellectual interests?
8	Are students who request oral
9	interpreters told that they are not available
10	because the counselor who is asked the
11	question is not aware that this is a support
12	option?
13	We must educate all our children,
14	including our disabled children, early on to
15	make choices.
16	We must teach them, as they grow,
17	to be responsible for their education and not
18	expect others to prepare every aspect of
19	education for them.
20	And so information about
21	responsibilities and about rights must reach
22	down to those who need it and who will need it
23	in the future.

This information is not reaching

1	far enough today.
2	MR. SCHWARTZBERG: Our next speaker is
3	Mr. Bruce Goldstein of Edwardsville, New York.
4	Bruce Goldstein is Special
5	Counsel to the law firm of Bouvier, O'Connor,
6	Cegielski & Levine.
7	The firm's practice includes
8	advising and representing school districts and
9	other education agencies, with particular
10	emphasis in the areas of education law and
11	handicapped law.
12	In private practice for seven
13	years, Mr. Goldstein was previously an
14	Assistant District Attorney and Assistant
15	County Attorney.
16	Mr. Goldstein taught for one year
17	in the City of Buffalo School District, and
18	was an undergraduate instructor in
19	Constitutional Law at the University of
20	Michigan and an instructor at Bryant and
21	Stratton Business Institute.
22	He was formerly a member of the

Association Conference on Legal Rights of the

faculties at the New York State Bar

23

1	Handicapped in 1983 and at the Fifth National
2	Institute on Legal Problems of Educating the
3	Handicapped, the 117th American Association of
4	School Administrators Conference and the 1985
5	National Association of Pupil Personnel
6	Administrators Conference.
7	In addition, he has been a guest
8	lecturer at the State University of New York
9	at Buffalo Law School in the areas of
10	Disability Law and Trial Techniques, and a
11	lecturer in the Lay Advocacy Training Program
12	at the University which he co-founded.
13	Mr. Goldstein holds the degree of
14	Juris Doctor from the University of Michigan
15	Law School where he graduated Cum Laude.
16	While in law school he was a
17	research assistant for Arthur R. Miller in the
18	preparation of the treatise, "Federal Practice
19	and Procedure."
20	He has been lead counsel in the
21	class action lawsuits <u>Gebhardt, et al. v.</u>
22	Ambach, et al., Bushey, et al. v. City of
23	Buffalo Board of Education, et al., Andres,
24	et al. v. Reville, et al., and Bonar v.

1 Ambach, as well as counsel for the Handicapped Intervenors in Arthur, et al. v. Nyquist, et 2 3 al. 4 Bruce Goldstein is Vice-Chairman 5 of the International Parents Organization, 6 Alexander Graham Bell Association, member of 7 the Executive Committee of the New York State 8 Developmental Disabilities Planning Council, past president of the Western New York Parents 9 10 Association of the Hearing Impaired, founder 11 of the Erie Coordinating Council of 12 Handicapped Organizations, a member of the 13 Board of Directors of the Western New York 14 Independent Living Project, member of the Board of Directors of the Association for 15 16 Retarded Children, Erie County Chapter, and a former member of the Boards of Directors of 17 18 Language Development Program of Western New 19 York, Inc., Autistic Services, Inc., and 20 Buffalo Speech and Hearing Clinic. 21 Finally, Bruce Goldstein is a 22 member of the New York State Assembly Task 23 Force on the Disabled, the New York State Bar

Association Committee on Mental and Physical

1	Disability, the Education Committee and Early
2	Intervention Committee of the New York State
3	Advocate for the Disabled, the Erie County Bar
4	Association Committee on Disability, and the
5	New York State Trial Lawyer's Association, and
6	the Western New York Disability Coalition.
7	Mr. Goldstein, for all these
8	reasons and others, is also a recipient of the
9	Erie County Bar Association Special Service
10	award.
11	Bruce A. Goldstein.
12	MR. GOLDSTEIN: As a parent of two deaf
13	daughters and someone who has raised those
14	daughters under 94142, I am of the generation
15	of higher expectations.
16	To paraphrase Doctor Martin
17	Luther King, "I have been to the top of the
18	mountain and I can see the promised land, but
19	we are not there yet."
20	94142 makes parents a partner
21	with the development of their children's
22	educational program.
23	It is a very sophisticated law
24	that guarantees a right to a free, appropriate

1	public education, but that promise of 94142
2	goes unfulfilled and will continue to be
3	unfulfilled unless and until we realize that
4	the 94142 is but a tool, an excellent tool,
5	but like all tools it is only effective if the
6	person who implements it has the skill to use
7	it properly.
8	Most parents, unfortunately, lack
9	the essential knowledge to effectively
10	implement 94142.
11	School districts share this
12	burden because they are often deficient in the
13	information and knowledge that is necessary in
14	order to reach the promise of 94142.
15	This holds true for all
16	handicapped children.
17	A unassisted parent is often
18	viewed by the school district as an uneducated
19	layman.
20	The parent who has some knowledge
21	and attempts to utilize that is often fearful
22	to speak up in the face of the experts or
23	alternatively is made to feel unreasonable or
24	ill-informed.

1	Many areas lack the assistance
2	that is necessary to help parents to implement
3	the law.
4	There are no trained advocates in
5	many areas in this country, and there are no
6	trained lawyers who know anything about 94142;
7	and those urban areas fortunate enough to have
8	some lawyers who are knowledgeable, it is
9	often too expensive for most parents to access
10	that assistance; the attorney's involvement in
11	special education is very labor intensive, and
12	the cost is often prohibitive.
13	Under Federal law there are no
14	protection advocacies that are required to
15	exist in every state, but their obligations
16	are to represent all of the developmentally
17	disabled, and there are a numerous number of
18	people that that requires is much too
19	burdensome for the staffing that is permitted.
20	In addition, their
21	responsibilities are not just for education
22	but for all areas of handicapped and
23	violations of civil rights.
24	And finally the quality varies on

1	a state-by-state basis, so as a practical
2	matter protection advocacies are not available
3	for parents to assist implementing 94142.
4	There is a if not a complete
5	absence of training for parents to allow them
6	to take advantage of the benefits of 94142.
7	When we address the needs of the
8	hearing impaired, this program is all the more
9	exacerbated.
10	Unfortunately, using the poor
11	term, it is the blind leading the blind
12	because the school districts don't have the
13	knowledge, and the parents are not given the
14	assistance of the knowledge to join with the
15	school districts.
16	The success of oral, oral hearing
17	impaired persons is in fact part of the
18	problem.
19	It is the very fact of the
20	integration and assimulation of those
21	successful oral hearing impaired persons that
22	leads to a lack of publicity about their
23	success which allows the continuation of
24	misperceptions and stereotypes.

1	Even a knowledgeable parent can
2	become caught in the system.
3	94142 in spirit calls for a
4	particular of services available to meet the
5	individual needs of all persons, and that
6	includes deaf persons.
7	Every school district should have
8	available a continuum of services which
9	provides total communication and an oral, oral
10	approach; however, all too often only total
11	communication is offered as an alternative in
12	many school districts.
13	This leaves the parent with a
14	Hopkins choice, either total communication or
15	placement of the child in the mainstream
16	setting with no support services.
17	Even more insidious is the recent
18	development of the so-called mixed classes
19	whereby children using sign language are
20	placed side by side with children following an
21	oral approach.
22	The school district will respond
23	that's no problem, we won't make the oral
24	child sign; he or she need only read the lips

1	or the translator.
2	Little thought as to how that
3	works in actuality on a day-by-day basis when
4	children are dependent on one another and
5	when the translator ends up being the aide for
6	all of the hearing impaired children, actually
7	happens, undercuts the ability of the
8	effectiveness of the oral program.
9	94142 establishes the concept of
10	least restrictive environment, which says we
11	will not segregate our disabled population;
12	however, the interpretation of least
13	restrictive environment as pursued by the $\mathtt{US}$
14	Government and by school districts and states
15	pursuant to 94142 is of no help to oral/oral
16	children.
17	The concept has been interpreted
18	as a simplistic interpretation and definition
19	which is based solely on physical setting.
20	A child who is in a segregated
21	building is in a more restricted setting than
22	the child in the public school setting.
23	The child who is in the
24	self-contained class is in a more restricted

1	setting than the child in the mainstream
2	class.
3	But when we talk about hearing
4	impaired children, we must however take a step
5	back.
6	We must look at the life plan of
7	the child.
8	Where do we foresee that child
9	being as an adult?
10	Because that is what an
11	appropriate program is about.
12	An appropriate program is to meet
13	the needs of the child, to let him gain those
14	skills and abilities that will allow him to
15	function as best as possible in society as an
16	adult.
17	Accordingly, we must enhance the
18	ability of the child to function in the
19	mainstream to the best of his or her ability.
20	Thus, if an oral/oral child has
21	the potential to function in the mainstream of
22	society as an adult then we have a
23	responsibility to provide that oral/oral
24	option in order to allow him to do so.

1	Unfortunately, parents don't know
2	what is available to them under the law.
3	If they know how to utilize that
4	tool and the U.S. Department of Education
5	refuses to recognize the significance of the
6	least restrictive environment that will permit
7	a child to sometimes be in a temporarily more
8	restrictive setting in order to gain the
9	foundation to allow him or her to be in a
10	therefore less restrictive setting and later
11	as an adult.
12	Finally, the promise of 94142
13	goes unfulfilled because the cornerstone,
14	which has been called the individual education
15	program, is either not used or only
16	superficially utilized in implementation of
17	the law.
18	School districts and parents both
19	lack of the training to draft true IEP's or
20	individual education programs.
21	Every IEP must be specific; it
22	must be measurable.
23	However, almost all schools in

this country in setting forth annual goals

1	have for handicapped children, will increase
2	their functional vocabulary.
3	Johnny will increase his reading
4	level.
5	If one takes a step back and
6	thinks about that, Mary will increase her
7	functional vocabulary will apply to virtually
8	every child, how can that be an individual
9	educational program if it can apply to every
10	child?
11	Johnny will increase his reading
12	level will apply to virtually every child from
13	the first grade to the 12th grade, how can
14	that truly be an annual goal if it applies to
15	every year through the child's educational
16	career?
17	IEP's must be drafted to be
18	specific and measurable.
19	Mary will use or identify the
20	following 30 words in spontaneous conversation
21	80 percent of the time while tested three
22	times over a two-week span.
23	Johnny will increase his reading
24	level from X level at the present time to Y

1	level one year from now, then the IEP becomes
2	the real tool of 94142, then it acts as a
3	yardstick or a red flag so that when we all
4	analyze at the end of the year, has Johnny's
5	annual goal been met, and we find that it has
6	not then we can go back and try to determine
7	what went wrong.
8	It doesn't mean the teacher was
9	wrong; it doesn't mean the child was wrong; it
10	doesn't mean nobody worked hard, but it may
11	mean that the goal was too high.
12	It may mean we have to take a
13	step back and breakdown the annual goal into
14	smaller steps, take those smaller steps bit by
15	bit, make the child feel good about himself or
16	herself and progress until we eventually reach
17	that goal.
18	The nature of the program or the
19	mode of instruction may be inappropriate,
20	maybe we need to change that, but until we
21	provide the assistance to parents and until we
22	provide the training to educators and parents,
23	and until the Federal Government through the

states monitors the meaning of 94142 in its

1 true spirit, it will only remain a dream. 2 MR. SCHWARTZBERG: I have a problem in 3 terms of those attempting to use the 4 audioloop. 5 I gather that there is no 6 audioloop established in this room at this 7 time; is that correct? I am advised that this can be 8 9 done within a five-minute period; is that also 10 correct? 11 I am going to take a five-minute 12 break for the purpose of attempting to do 13 that. 14 If it can't be done within five 15 minutes, I have lost any ability to hold my 16 interpreters, so it is going to be a 17 five-minute break. 18 (WHEREUPON, a short recess was had.) 19 MR. SCHWARTZBERG: The court reporter 20 will indicate to me at the point which she is 21 ready to proceed. 22 At this time I would like to ask 23 whether there are any questions addressed to

the last panel or actually the last two

1 panels. 2 Any members of the Commission? Yes, Mr. Ewing. 3 To the members of the 4 MR. EWING: 5 panel, I had a lot of questions. 6 The Chairman would never allow me 7 to ask this many questions. 8 I wanted to sort of summarize my 9 questions into one. 10 It has been pointed out that 11 parent involvement is one of the most 12 important aspects of securing equal education 13 opportunity. 14 And when you read public law 15 94142, you come away with a sense that there 16 is no other document anywhere in State, 17 Federal Laws and Regulations that gives a kind 18 of power to parents than does public law 19 94142. 20 So I think my question is: 21 you believe that parents really recognize the 22 power that they have whereby a local school 23 district needs their signature of consent in

order to implement an educational program?

1	And my other question is that in
2	view of the fact that there is massive
3	undereducation of parents for the
4	responsibility that they have to make
5	educational decisions, and inasmuch as this is
6	required by law that parents be trained, and
7	since there is a crisis in that parents are
8	not trained, how can this be brought about?
9	MS. JANGER: Inez Janger.
10	I think I am speaking for
11	first of all, the three of us up here are
12	parents.
13	Your first question was do
14	parents recognize the right they have to sign
15	the IEP and the power of the IEP?
16	Absolutely not.
17	Why don't they recognize it?
18	I think what I was trying to say
19	in my presentation is to a great extent it
20	is a combination of the awe we all hold of
21	professionals, the expertise or supposed
22	expertise that they present us with; our own
23	fear and sort of grasping for hope at any
24	particular point in time, the numerous

1	situations we have heard of where the IEP is
2	what Bruce mentioned, Mary will read better or
3	Mary will have the services of a tutor period
4	without anybody saying what the tutor will do,
5	and then the statement of the school districts
6	but that's what an IEP is.
7	Parents do not know what an IEP
8	is; parents are not educated, and I don't
9	exactly know the answer.
10	What I tried to talk about, what
11	public information I don't have an answer.
12	I think the problem is parents do
13	not recognize their right.
14	They are undereducated, and one
15	cannot count on the educational system, I am
16	afraid, to educate the parent properly, but
17	they sure as heck haven't since the law was
18	passed.
19	MR. GOLDSTEIN: Just to add, to bring
20	that change about would require some mandated
21	programing upon school districts to require
22	the training of those involved in the special
23	education system and to require that they
24	provide the information to the parents or

1	arrange for its provision, which they don't
2	đo.
3	Obviously, if you view yourself
4	in an adversarial relationship with parents or
5	sharing power, and you are not used to sharing
6	power, those you can keep out of information
7	keeps you in the power of position.
8	Mass education in this country
9	has always been to teach the public
10	education teaches the masses; this law runs
11	contrary to at least a century of public
12	education in this country because we talk
13	about individualism.
14	School districts don't like that;
15	that is administratively difficult, if not
16	inconvenient, so we must mandate training for
17	parents so that they know what to do and how
18	to effectuate; that it won't hurt to educate
19	the educators at the same time.
20	MR. EWING: I would like to point
21	out it is already written into Public Law
22	94142 in three different places within the law
23	that parents must be trained; it is even
24	written under the personnel development

1	section which mandates that funds be spent to
2	train teachers.
3	In the same line is written that
4	parents must be trained, but I wanted to point
5	out is that this is not happening, and that's
6	why we are seeking some ideas on how it could
7	happen in view of the fact that parents don't
8	seem to be effectively organized, but I wanted
9	to point this out since you used the term
10	"mandate," that it also exists in the law.
11	MR. SCHWARTZBERG: Further questions
12	from the panel?
13	There are no further questions
14	then we will thank the participants, and we
15	move next to Doctor Richard Stoker.
16	While this panel is moving down,
17	I also ask that Doctor June Grant of Trinity
18	University come forward.
19	Doctor Richard Stoker is an
20	Associate Professor in the School of Human
21	Communication Disorders at McGill University.
22	He is the author of over 50 book
23	chapters, research reports and published
24	articles on hearing impairment.

1	He is the Editor of <u>The Volta</u>
2	Review, an academic journal in the field of
3	Education for the Deaf.
4	He is a member of several
5	national and international standards
6	committees concerned with technical standards
7	for electronic devices for the hearing
8	impaired.
9	At this point I would like to
10	call on Doctor Richard Stoker of McGill
11	University.
12	DR. STOKER: Mr. Chairman, Members of
13	the Committee, it is a pleasure to be here
14	this afternoon to talk about technology and
15	its impact on the educational process for
16	hearing impaired children.
17	I think that it would be fair to
18	say that technology represents an unfulfilled,
19	as of yet, trump card in our effort to provide
20	educational and other opportunities to hearing
21	impaired children and their parents.
22	Technology has a very special
23	role to play in the education of hearing
24	impaired children.

1	It is in this field that many of
2	the technological advances that have come from
3	the laboratories and research workers have
4	found truth and have been applied and have
5	been proven successful over the years.
6	For example, the use of the
7	electronic hearing aid which was the first
8	commercially successful use of the transistor
9	invented in Bell Telephone laboratories in
10	1951.
11	This is not to say, however, that
12	technology is or can be or necessarily will be
13	an unqualified success in the hearing impaired
14	children.
15	For example, again with hearing
16	aids, several studies have shown that in
17	programs for the hearing impaired between 40
18	and 60 percent of hearing aids being worn by
19	hearing impaired children are not functioning
20	on any given day.
21	That is to say that approximately
22	half of the children in educational programs
23	in the United States today are being educated
24	with ear plugs instead of technological

1 assistance. This, I am afraid, does not well 2 to use to advance technology such as some of 3 those I would like to briefly discuss this 4 morning; however, I do believe that with 5 6 increased sophistication of teachers and 7 parents and a realization that we do in fact 8 live in a technological world, that there is 9 great promise for such devices in education of 10 hearing impaired children. 11 As for myself, make no doubt of 12 this, I am a child of the technological world. 13 I personally owe a great deal of 14 my own existence to the use of technology. 15 I realize this very strongly 16 every morning when I awake to the bright light 17 flashing in the room waking me up, and reach 18 over to switch off the sound switch which 19 turns on the flood lights in my bedroom. 20 As I then reach for my glasses, 21 which allow me to locate my amplifier hearing 22 aids, and as I put them on and switch them 23 into being, I am not so pleasantly reminded of

the auditory world that I have escaped from

for an evening's rest.

I have been brought over to the

dresser to turn off a noise sensitive monitor

which warns me of a fire alarm or another loud

noise, of course, in my apartment while I

6 sleep.

15

16

17

18

19

20

21

I am then able to check my

answering machine to see if any messages have

been left on my TDD or telephone device for

the deaf, as the messages tumble out on the

little letter brightly lit.

I often marvel at the fact that
this wonderful piece of equipment was not
available only a few short years ago.

I frequently give a silent thanks to the deaf inventor, Doctor Robert, who pioneered TDD communication along with other individuals some of which -- some of whom are sitting in this very room; what a positive affirmation of the potential for the deaf to lead the deaf.

I often muse about how wonderful
it will be when I will be able to hook my
telephone up and understand what a person is

1	speaking on the other end, without them having
2	to have a similar TDD-type device.
3	There are four basic areas where
4	technology stands to impact very greatly on
5	education of hearing impaired children.
6	The first of those is century
7	aids, century aids such as hearing aids, such
8	as devices which transfer the spoken word into
9	sensation that can be picked up on the skin,
10	such as devices which present visual signals
11	in a pair of glasses where a person can
12	actually read what the other individual is
13	saying.
14	All of these marvelous things are
15	on the drawing boards and in the research labs
16	and are going to be things that will confront
17	us in the very near future.
18	The second area is that of
19	teaching, teaching hearing impaired children
20	as with all other children is being and will

LONGORIA & GOLDSTINE, CHICAGO, ILLINOIS (312) 236-1030

teach many things to children under the

continue to be revolutionized by the use of

programmed to individualize and patiently

technology such as microcomputers which can be

21

22

23

1	guidance	of	a	teacher
---	----------	----	---	---------

- I would like to emphasize that

  because technology cannot just be turned on

  and worked, it needs to be cultivated in the

  context of an intelligent educational plan to
- 6 utilize these things.

11

The third major area is that of
telecommunications, of using the
telecommunications network to create advanced
potential for hearing and deaf people to

communicate on an equal basis.

- We are all aware, I think, of the
  increasing use of telephone communication in
  our society; and one of the most promising
  aspect of this increasing use of
  telecommunications is that it will allow
  individuals to communicate with hearing people
  on a daily basis without having to overcome
- the entire phase of the communication

  difficulty experienced by the hearing

  impaired.
- The last area then is the medical professions, the ability to accumulate and quickly diagnose hearing loss in infancy is

1 now a reality. 2 It is not -- this equipment is not used as effectively and as consistently 3 4 and as often as it should be, however it is available or does allow the hearing impaired 5 child to be diagnosed virtually at birth, and 6 for educational programing to begin 7 8 immediately with all of the attendant 9 benefits. 10 What do we need to bring about a 11 heightened use of technology in this field? 12 First of all, we need that 13 universal bomb that we call money. 14 It is not inexpensive to 15 implement many technological aids to the 16 hearing impaired; and there will need be a 17 commitment among various government agencies, 18 parents, and schools to providing these 19 technological devices that will make the lives 20 of hearing impaired individuals more full, 21 more capable, and able to utilize the various educational services available to them. 22 23 We will require more than just

this however; it also will require training.

1	It will require the hearing
2	impaired individuals themselves become expert
3	consumers; that they become knowledgeable;
4	that they become demanding; that they do not
5	accept poor quality or poorly designed
6	technological assistance.
7	It will also require that
8	professionals and parents become more
9	knowledgeable; that they learn to accept the
10	limitations and work with the strengths of
11	various pieces of technology.
12	This leads to what I believe will
13	be an entirely new field and that is the
14	assessment of technology for the disabled,
15	where centers and professionals will be
16	trained to evaluate the technology needs of
17	people with disabilities.
18	Not everyone needs the same
19	packet of goodies; not everyone needs the same
20	assistance.
21	It is very individualized, very
22	unique situation for every individual; and
23	therefore it will be necessary to develop
24	specialized assessment procedures and

1	practices to make sure that the technology is
2	available to the hearing impaired.
3	With this heightened individual
4	need, awareness of individual need, I am
5	confident that we can work together to bring
6	about the technological revolution that has
7	thus far evaded our grasp, even though many of
8	us have lived long enough to see an enormous
9	increase in use of technology in society at
10	large, and the hearing impaired have lagged
11	behind in their ability to make use of these
12	excellent, unique, and creative devices.
13	I am sure that with the
14	cooperation of all involved in this process,
15	this day will come sooner rather than later.
16	Thank you very much.
17	MR. SCHWARTZBERG: Doctor June Grant.
18	June Grant earned a Bachelor's
19	Degree from Central Institute for the Deaf,
20	Washington University, Saint Louis in 1941, a
21	Master's from Trinity University in San
22	Antonio, Texas, and a Ph. D. from the
23	University of Texas in Austin.
24	She has taught hearing impaired

1	children at the Central Institute for the Deaf
2	and at Sunshine Cottage in San Antonio where
3	she served as Academic Head.
4	In addition, she has taught
5	learning and language disabled children.
6	Currently she is a Professor of
7	Education at Trinity University in San Antonio
8	where she is Director of Special Education.
9	The Department has programs to
10	prepare teachers for hearing impaired,
11	language disabled, emotionally disturbed,
12	physically disabled, and mentally retarded
13	children.
14	Doctor June Grant.
15	MS. GRANT: My assignment was to speak
16	on education and the hearing impaired student,
17	but the mandate and what are the practices.
18	And when we talk about bilingual
19	hearing impaired children, we are talking
20	about a minority within a minority.
21	In other words, the number we
22	all know that hearing impairment is a very low
23	incident handicap, we talk within that
24	handicap, the number of children who come from

1	the bilingual homes very often not bilingual,
2	they are simply nonEnglish speaking homes
3	we are talking about a very small percentage
4	of children, but yet a large number of
5	children.
6	Some of the predicted figures of
7	things to come are really frightening.
8	It has been predicted that by the
9	year 2000 there will be 40 million nonEnglish
10	speaking in the United States, so if we think
11	we have a little problem now, we have one that
12	is going to grow bigger and bigger as the
13	years go on.
14	Right now there are four and a
15	half million school children from homes where
16	English is not the spoken language in the
17	home, so this even though that is a small
18	percentage of the number of school children in
19	the United States, that is a large hunk of
20	children.
21	The demographic studies put out
22	by the by Gallaudet College every year,
23	last year accounted 57,731 hearing impaired

24

children.

1	Now, these are the children who
2	are in programs for the hearing impaired; this
3	does not count the hearing impaired children
4	who are mainstreamed or those who are out in
5	schools and receiving no services whatever.
6	But of that number over 7,000 of
7	those children or 15 percent of them come from
8	homes where English is not the language of the
9	home.
10	So we have a large percentage and
11	yet a minority of children who come from homes
12	where English is not the language spoken; and
13	to talk about those homes or these children as
14	bilingual is a real misnomer because very
15	often it is very monolingual as far as the
16	homes go and the children themselves are
17	monolingual in many cases rather than
18	bilingual.
19	But we do have legislation to
20	provide for each of these conditions; we have
21	Title 7 of the Elementary and Secondary School
22	Act which was enacted in 1968, the public law
23	95561; and we have public law 94142, which has

been mentioned so many times here, to provide

1	for all the education for all handicapped
2	children.
3	Now, what would it take to have
4	the kind of program that we need for children
5	from nonEnglish-speaking homes who are hearing
6	impaired?
7	I think there are four factors
8	that need to be attended to.
9	Those are: Adequately trained
10	personnel and enough of them, appropriate
11	assessment instruments, adequate and
12	appropriate materials for the children, and
13	parental and community support.
14	I think those are four main
15	topics we need to address.
16	I haven't prioritized these, and
17	I don't know how to prioritize them.
18	I think they are all equally
19	important, but I think in order to talk to
20	them at all we will put them in an order of
21	what would have to come first; and I think the
22	thing that has to come first would probably be
23	parental and community support, and here we

have a dichotomy again because within each

1	handicapping condition and I speak of
2	bilingual, if it were true it wouldn't be
3	handicapped but I speak of the child from
4	the nonEnglish-speaking home, and within this
5	condition there are options and within
6	education of the hearing impaired there are
7	options also which have been amply covered,
8	and I won't talk about that but within
9	bilingual education there are several options.
10	Two main categories are bilingual
11	education as opposed to ESL, English as a
12	second language; and we will take ESL first.
13	In ESL the idea is to immerse the
14	students in English-speaking programs and have
15	them pulled out that is a term that is used
16	for intensive English instruction for a period
17	a day, usually an hour, something like that,
18	and the kind of program that is usually is a
19	program that is behaviorly oriented where
20	there is constant drilling on using English
21	without too much concern, according to a lot
22	of the experts, on what kind of comprehension
23	the children are having, but this is a
24	drill-type material.

1	Most educators of bilingual
2	education, but they are very prevalent because
3	they are easy to do, and it doesn't take as
4	much training for personnel.
5	Now, even within bilingual
6	programs there are choices.
7	The choices are a maintenance
8	program as opposed to a transitional program;
9	and in a transitional program the idea is to
10	give these children as much English as give
11	them the idea is to get them into a totally
12	English speaking programs as soon as possible.
13	So, in other words, they are
14	given programs in the native language as long
15	as necessary, but the goal is to get them into
16	English only programs just as soon as
17	possible; whereas the maintenance programs are
18	supposed to maintain the home language plus
19	the home culture throughout the program.
20	However, even in these programs
21	it usually is for a limited time, ordinarily
22	three years.
23	If it starts at the preschool
24	level, generally it is while the children are

```
1
         ages 3, 4, and 5; and they tend to put them in
2
         English-speaking programs at 6 years or
         first -- or it is the first and second and
3
         third grades, but by the fourt grade most of
4
5
         these programs are finished, but there are
6
         exceptions of course.
7
                       So as I said, within each
8
         category we have options, and strong opinions.
9
                      We have had some strong opinions
10
         already addressed as far as education of
11
         hearing impaired children go so I won't
12
         address that now.
13
                       To get back to the topics, the
14
         community.
15
                       What can we do to get community
16
         support -- for the community I mean the total
17
         community; I mean the legislatures, the school
18
         districts, the local governments, the religous
19
         organization, the health services, the news
         media, the parents' group, all of these
20
21
         categories.
22
                       We have the legislation; it is on
23
         the books.
```

It is not always implemented, and

Ţ	it is certainly not implemented to the extent
2	that we would like for it to be, as we've had
3	evidence here already.
4	School districts, most of them
5	comply to the letter of the law, if not within
6	the spirit of the law.
7	Local governments, many of them
8	give lip service without giving actual
9	support, and some of them don't even do that.
10	Religious organizations, and I
11	speak of this because I come from an area
12	where most of the nonEnglish-speaking children
13	represent Spanish-speaking homes, and most of
14	the Spanish-speaking population is catholic;
15	and we could get and do get in some great
16	support from the Catholic church, but I think
17	all of the religous organizations need to be
18	able to lend support to this sort of thing.
19	The news media do in San
20	Antonio we have good news coverage and
21	positive in bilingual education but not for
22	hearing impaired but and the parents!
23	groups, it is a sad situation.
24	In San Antonio in particular we

have found the Mexican American parents very 1 willing and very eager to do what we want them 2 3 to do and what they can, but these people often are nonEnglish-speaking completely; many 4 5 of them illiterate who cannot even read or 6 write, and so we have a group of really 7 intimidated people who are not going to be very assertive or at least as assertive as we 8 9 would like them to be, in which case, I think, it is up to the professional to assert the 10 rights of the parents or get some assertive 11 12 parents to assert their rights because I think 13 that's what is going to take before we get the 14 kind of programs we need for these children. 15 ' And -- of the presentation that 16 just preceded this about parents' rights and that sort of thing, in Texas as far as IEP and 17 18 what is called the ARD -- it is the meeting 19 that happens once a year. 20 It is admission review of 21 dismissal of all handicapped children; the 22 parents must be notified of this meeting if 23 they are only Spanish-speaking they must be 24 notified.

1	If they illiterate, they must be
2	notified either by telephone or by personal
3	call at the home, but there is nothing that
4	says they must come; and in some cases we get
5	cooperation and sometimes not so good.
6	But so much for the community and
7	the parental support.
8	I think this is going to have to
9	be the first thing, that there is going to
10	have to be a great deal of pressure from this
11	group what about personnel, now?
12	We have requirements and
13	certification or endorsement requirements for
14	teachers of the hearing impaired; and we have
15	requirements, certification, and endorsement
16	requirements for teachers of children in
17	bilingual programs, but we have nothing that
18	combines the two.
19	It is as if these two entities
20	never interface, and they do very, very
21	frequently as we know.
22	And what is the answer to this?
23	I don't know if we ask teachers
24	to get additional certification to be a

1	teacher of the hearing impaired, to be a
2	bilingual teacher and then to get
3	certification in education of the bilingual
4	hearing impaired.
5	I think we are asking for the
6	moon.
7	I think that it's take extra
8	time; it will take extra money, and we don't
9	have that kind of personnel.
10	And what does it take of these
11	personnel?
12	These people have to be should
13	be affluent in Spanish.
14	We have in the University I
15	represent, we have worked very hard to recruit
16	Mexican-American students into the university
17	in general; and we have very high entrance
18	requirements, and those Mexican-American
19	students who meet these entrance requirements
20	very often are middle class students and very
21	frequently don't even speak Spanish.
22	I mean these are people who have
23	been raised in the American milieu, as it
24	were, and they have lost tract of their

1 culture. 2 All right. So we have a great deal of trouble getting the right kind of 3 4 personnel. 5 The only thing that I know of 6 that has been rather successful that we have 7 been able to get a number of bilingual fluent Spanish speakers for teacher aids, but these 8 9 are not professionals, they are 10 paraprofessionals. 11 Now, the standard -- all teacher 12 preparation programs meet certain standards 13 put out by National Colleges of American 14 Teachers of teacher education programs; and 15 two of the standards within the requirement 16 are that any teachers who are certified must 17 be sensitive to the plurality of American 18 culture. 19 They must realize there are

They must realize there are
students who come from different kinds of
cultures, and the students must be sensitized
to this; also they must be sensitized to the
special needs of children who have needs -need special programs for learning, in other

words, Public Law 94142 and the program -- the 1 2 Title 7 of the Secondary Elementary and Secondary School Act. 3 But there is, of course, that 4 5 would say that they have to know and have some б experience in the education of handicapped 7 hearing -- of hearing impaired children from 8 nonEnglish-speaking homes. 9 So we have a long road here. 10 We have a very small of personnel 11 who are qualified to teach these children in 12 their native language, who are capable of 13 dealing with the parents in their native 14 language; and I just don't know what the answer is going to be in order to make that --15 16 to increase the size of that group of 17 personnel. 18 Materials, the publishers, 19 publishers of textbooks and things of that 20 nature have been very slow to address the 21 needs of bilingual programs in general, much 22 less the programs for hearing impaired 23 children.

Maybe this isn't so bad for

1	teachers of hearing impaired children because
2	many of the materials that they use are
3	teacher made anyway; however, we need to have
4	something from which to start, and there is
5	very, very little available.
6	The Skyhigh program was mentioned
7	here this morning.
8	It is a program that is used in
9	Texas for parent-infant programs.
10	Now, parts of that have been
11	translated into Spanish so that parent-infant
12	facilitators can deal with parents in their
13	native language and help them out with that;
14	and there is a what is called a Hispanic
15	Parent Outreach Program in Texas that is
16	teaching parents who do not speak English sign
17	language, and instead of using English
18	translation are using Spanish translation of
19	American sign language.
20	Now, the problem with this is
21	that American sign language in general is not
22	the sign system used even in the schools that
23	do not have oral/oral programs; most of these
24	use some sort of manual English, some sort of

l signs of English.

- 2 Texas has its own particular sign
- 3 system called briefed signs.
- 4 It is very much like seeing
- 5 essential English, and parts of that have been
- 6 translated into Spanish, but there really are
- 7 no -- there is no volume of materials for
- 8 hearing-impaired kids in Spanish.
- 9 This same program, this Spanish
- 10 Outreach Program has listed 248 volumes on
- 11 sign language in Spanish that is in all of the
- 12 United States.
- 13 123 of them are published by the
- Lexington School of the Deaf, well, they have
- a large population of nonEnglish-speaking, and
- 16 60 of them are from Texas, which is the
- 17 Texas-Mexican border, and 15 of them from the
- 18 school in California, and the rest of them in
- every school will have one or two books for
- 20 parents in Spanish.
- 21 Another problem that I mentioned
- is that of an assessment and children who do
- not speak English, like children who come from
- Spanish in the home traditionally have

l suffered bias.

2	And all of us who are educators
3	of hearing impaired children know how they
4	suffer from a bias if they are measured by a
5	verbal test for intelligence or anything else
6	because their verbal skills are not generally
7	what they are hearing.

So just to start talking about an assessment for these children brings us at a really disadvantage, a terrible disadvantage; and as far as I know there are no real solutions to this problem.

There are tests that have been translated into Spanish, but these have been highly criticized by the leaders in bilingual education; they say — that existed in English version are simply translated into the Spanish version, and the students are not much better off.

What has been suggested that not -- that educators not rely on these standardized tests as the only measures of placing students, but instead develop some observational techniques that would help them

1 place children in appropriate programs. 2 So in summary, I would like to 3 mention two programs that, I think, are really 4 doing a wonderful job and hope that maybe they 5 can lead the way into better things: 6 Rhode Island School for the Deaf has a project 7 called Project Opportunity, and they provide services for children of nonEnglish-speaking 8 9 homes, and they have more than one language, 10 they have Spanish, children from 11 Spanish-speaking homes, Portuguese, and even 12 some others; and they are doing an admirable 13 job. 14 Their -- is utilize the language 15 and the culture of the home because even after 16 our children learn English, and that is our 17 main objective, is that they should be 18 functional and fluent in English, but even 19 after that, their home culture is always going 20 to be a part of their total life, so don't 21 ignore it. 22 And the literature in bilingual 23 education emphasizes to the greatest extent 24 that the home language should be the one to be

1	the primary one in the language of the
2	starting point because this is where the
3	children are going to get the input.
4	MR. SCHWARTZBERG: Doctor June Grant, I
5	think at this point I am going to have to call
6	time because we are really running quite a bit
7	over.
8	DR. GRANT: The other program is
9	Lexington School for the Deaf; and those two
10	programs need to be exploited really.
11	MR. SCHWARTZBERG: At this point I
12	would like to ask if there are any questions
13	to the advisory committee?
14	I see no questions.
15	I have one question from Doctor
16	Lucas.
17	DR. LUCAS: A quick question for Doctor
18	Stoker.
19	When does a technological advance
20	become so commonplace that it would be needed
21	or would be mandated by the regular civil
22	rights legislation and regulations?
23	MR. SCHWARTZBERG: Doctor Stoker?
24	DR. STOKER: Let me make sure that I

1	understand the question.
2	When will they become so
3	commonplace
4	MR. SCHWARTZBERG: Let me try to
5	rephrase it.
6	Technology changes over time.
7	What is the rule that helps
8	determine when a particular technological
9	change becomes reasonable accommodation?
10	Is there something other than a
11	rule of reason that one can apply or is it
12	just all wrapped up in this wonderful word
13	"reasonable"?
14	DOCTOR STOKER: I am afraid I am going
15	to have to go through your later suggestion, I
16	don't think I do not think there is any
17	general rule that I could give you there.
18	What is reasonable to one person,
19	however, may seem very unreasonable to
20	another; and I would like to add that very
21	often even very expensive technology is
22	less expensive than the consequences of
23	ignoring the ability of the hearing impaired
24	individual.

1	MR. SCHWARTZBERG: Thank you, Doctor
2	Stoker.
3	Any additional questions?
4	None. Let's turn from the Right
5	to An Education to the Right to Employment.
6	And as this panel steps down, I
7	will ask that Robert Menchel of the National
8	Technical Institute for the Deaf, Miss Bonnie
9	Tucker of Phoeniz, Arizona, Mr. Sy DuBow,
10	National Center for Law and the Deaf, and Mr.
11	Rocky Stone of S.H.H.H. for the hard of
12	hearing, Mr. Alan Post, a lawyer of
13	Springfield, Illinois, if those five would
14	please come forward.
15	I will begin with Mr. Stone is
16	going to be the first to leave, although he is
17	going to be able to share some of the panel,
18	because he has a plane to catch and he will be
19	leaving here at approximately 4.
20	Mr. Rocky Stone lost his hearing
21	at age 19 while serving in the United States
22	Army.
23	He graduated from the University

of Southern California with honors in 1949.

1	His combined scholarship and
2	fellowship at the school at John Hopkins
3	University in 1949 and 1950 which led to his
4	Master of Arts Degree, and then spent 25 years
5	in a period during which he was severely
6	deafened with the Central Intelligence Agency
7	of the United States, the CIA, including 17
8	years abroad.
9	His last position in Washington,
10	he was responsible for all Soviet operations
11	worldwide.
12	He returned to the United States
13	at age 50, and in the year 1979 he founded the
14	organization known as S.H.H.H. Self-help For
15	the Hard of Hearing, in November of that year,
16	and started membership solicitation for that
17	organization in the summer of 1980.
18	Various observations have
19	suggested that the growth of that organization
20	since that time has been one of the most
21	remarkable organizing feats that we have seen
22	in recent years.
23	Mr. Rocky Stone, President of
24	Self-help For the Hard of Hearing.

1	MR. STONE: Thank you, Mr.
2	Chairman.
3	I don't usually read speaches,
4	but testimony before a distinguished
5	Commission such as this, I try to be a little
6	bit more precise.
7	Most of my work is motivating
8	work in which we try to stimulate people to
9	understand their problem and try to do
10	something about it, but civil rights
11	safeguarded by the Constitution and by the
12	Bill of Rights, but that is just the
13	principal.
14	But because of the fact that I
15	have been assigned the task of speaking about
16	practical aspects of access legislation and
17	policy changes implies that there must be a
18	considerable gap between the principal and the
19	reality of what occurs today, and that is
20	certainly true.
21	If I were to attend the very
22	highly advertised Statue of Liberty
23	celebration next week in New York, I would be
24	unable to participate in any of those

1 ceremonies for the simple reason that they are 2 communicatively inaccessible. 3 There will be millions of people 4 like me in the United States who, if they had 5 the ability to receive the benefit of the 6 technology that Doctor Stoker and others have 7 described to you, we could be in the mainstream of life. 8 9 I think there is a very definite 10 communication here implicit in the existence 11 barriers is the denial of fundamental rights, 12 the right to assembly, the right to 13 employment, as we are discussing here, and 14 even the right to due process because many 15 courtrooms are among public facilities that 16 remain communicatively inaccessible; certainly 17 denial of access, be it physical or communications is in violation of both the 18 19 spirit and the intent of civil rights laws, 20 particularly as they have evolved through 21 clarifying legislation and court tests over

LONGORIA & GOLDSTINE, CHICAGO, ILLINOIS (312) 236-1030

resource and sometimes the course of last

The law represents a major

the last 20 years.

22

23

1	resort in which handicapped people must turn
2	if their rights are to be affirmed.
3	Did you hear that worried
4	"affirmed"? That is the key word because that
5	brings us to the practical aspects of access.
6	We, those of us who are hearing
7	impaired, must do the affirming.
8	Our record in this regard is very
9	weak.
10	Since 1973 Congress has sought to
11	forbid discrimination against all handicapped
12	individuals; and yet just four months ago in
13	1986: the National Counsil on the Handicapped
14	issued a report to the Congress and to the
15	President in which the Council noted that
16	existing civil rights laws do not cover
17	discrimination on the basis of handicapped.
18	NCH recommended that Congress
19	enact a quote, "new comprehensive law"
20	requiring equal opportunity for individuals
21	with disabilities with broad coverage in
22	setting clear, consistent, and enforceable
23	standards prohibiting discrimination.
24	The law remains a tool which we

1	hearing impaired people must be prepared to
2	use in asserting our rights within its
3	framework, without this existing or improved
4	legislation will mean nothing.
5	Access is imposed by law in two
6	ways: By the element of equal opportunity
7	and/or by the legal provision guaranteeing
8	equal access to services and to programs.
9	Where can hearing impaired people
10	who need help in finding and holding a job go
11	for help?
12	Many turn to state vocational
13	rehabilitation agencies. Even the employed
14	person may need vocationally-oriented advice
15	or counseling to individual in advancement, to
16	determine more personally satisfying
17	employment or to avoid further injury to his
18	or her hearing in the present employment
19	context.
20	Those too young to work need to
21	help coordinate their education with
22	vocational goals, and older persons who desire
23	employment to supplement retirement income

should also have advice and assistance

1	available, but there are many problems
2	inherent in the nature of the population to be
3	served.
4	The term hearing impaired
5	includes deaf and hard of hearing people;
6	typically however services and benefits from
7	legislation and/or regulations have focused on
8	deaf people.
9	Recently a trend has developed to
10	include to some extent that population which
11	we describe as hard of hearing.
12	In terms of ability effectively
13	to receive spoken communication through the
14	auditory process, there is a thin line between
15	profoundly deaf and severely hard of hearing
16	people.
17	Intent to provide services to
18	both immediately triples the constituency; in
19	other words, with two million deaf people, we
20	add over four million who are severely hard of
21	hearing to total over six million possible
22	recipients of services.
23	There is talk too of giving more

recognition to the functional aspect of

1	hearing disability in addition to the more
2	diagnostic and measurement of decibel loss.
3	If you have a 60 decibel loss,
4	and I have a 110 decibel loss, and neither of
5	us can understand what the other person is
6	saying to us, it really doesn't make much
7	difference diagnostically.
8	The key word is function, and the
9	degree to which we cannot only direct the
10	attention of our medical and professional
11	society in order the functioning aspect of
12	hearing loss and the government as well that
13	will determine how far we are able to advance.
14	Interpreting and implementing
15	existing regulation becomes increasingly
16	difficult, as we move away from something that
17	is very clearly defined, it is at the far end
18	of the spectrum of need, profound deafness;
19	people understand that a little better anyway
20	than the ambiguity which often occurs in
21	moderate to severely hearing impaired persons
22	in the work place.
23	Recent trends in government
24	and a recognition of the problems and

1	perils that deaf persons and a company's
2	willingness to provide services and to make
3	accommodation for this large population are a
4	beginning, but it is slow going and it is
5	going to continue to be slowing going for
6	sometime.
7	The hard of hearing person often
8	has disability in using the telephone to set
9	up an interview.
10	Let me give you an example, what
11	I think is a pretty representative type of
12	conversation that goes on among people that
13	are in our particular the phone rings and a
14	female voice says "Mr. Jones, this is Mrs.
15	Shift. We received your resume, and we would
16	like to have you come in tomorrow morning at
17	10:00 o'clock; our address is 3347 Street,
18	have you got that?"
19	"Is this Mrs. Smith?"
20	"Yes, yes."
21	"Mrs. Smith, we have got your
22	resume.
23	We would like you to come in at
24	10:00 o'clock tomorrow."

1	The tenativeness, the hesitation
2	immediately conveys a picture in the eye of
3	the interviewer of somebody who is not too
4	sharp.
5	The question of whether they even
6	go through with the interview at that point is
7	a very significant one, but you usually
8	struggle through to get the time and the
9	address, and you get there the next morning
10	if you can get by that next interview by some
11	miracle of a combination of empathy and
12	assertiveness and perhaps even admitting that
13	you have got a problem which is Catch 22
14	because as soon as you admit that you have got
15	a problem, they form, the supervisor can and
16	frequently does form in his or her mind the
17	degree of limitations that you have in the job
18	because they don't know a thing about you or
19	your problem.
20	They simply react to the fact
21	that if you are severely hard of hearing, you
22	cannot do certain things.
23	I was fortunate in the CIA; they
24	didn't treat me as a disabled person.

1	They related to me on the basis
2	of my abilities, and it is because of that
3	reason that I was able to rise to the top of
4	that organization, plus my own drive and
5	competence, but if I didn't have the chance,
6	the openness of the personnel and the
7	administration of that organization permit me
8	to do that, I would have been dead in the
9	beginning.
10	Most of the people who interview
11	us have make these decisions in the past
12	and continue will continue to the degree
13	that they remain uneducated.
14	And we have to do a lot of
15	educating;
16	they cannot decide what our limitations are.
17	In relating to the supervisor, if
18	we do get a job, we have a serious problem.
19	Again the question is do you
20	admit the hearing loss or do you try to hide
21	it?
22	And if you do admit it and try to
23	work with it, to what degree is that person
24	going to accept those conditions and those

1	moderate differences of approach which make
2	the the difference of life and death to you
3	as a hearing impaired person.
4	Again it is a matter of
5	education.
6	We have to make some modification
7	of the environment on the job, not much, but
8	some, but again there has to be an educated
9	willingness to accept these modifications so
10	that they don't they are not construed as
11	simply giving somebody something special as
12	opposed to equal treatment of all other
13	employees.
14	What they are actually doing is
15	giving us the access that we need to perform
16	at whatever level we are capable in that work
17	place by making these accommodations; and as
18	Doctor Stoker said, most of the time they are
19	not very expensive.
20	We have trouble in acquiring the
21	necessary tools with which to do the job,
22	environment considerations.
23	We have trouble in maintaining
24	the job after acquiring it, and after we put

1	on a visible hearing aid.
2	Throughout the United States I
3	have talked to people who, when they put a
4	hearing aid on in a sensible effort to
5	maintain effective communication in the work
6	place and in society, they are fired.
7	The people who are fired most
8	frequently are teachers, school teachers.
9	There are faces in this audience
10	that I recognize who have lost their jobs
11	because of the onset of hearing loss.
12	One, I believe, is a
13	stenographer, another was a research person,
14	but right here in Chicago.
15	You have very specific examples
16	of what it is we are talking about; job
17	maintenance becomes a very important aspect of
18	civil rights in this context.
19	If you travel, the barriers of
20	the use of the telephone, inability to hear
21	announcements at airports and train stations,
22	difficulty in hearing a door lock, smoke
23	alarm, or a phone ring in a hotel room, and
24	the overall strain of attending leaves us limp

<u> </u>	at best, and at worse costs as our job.
2	Legislation effecting all of
3	these situations already exists, but few hard
4	of hearing people are aware of it, and even
5	fewer employers take the initiative in
6	addressing the problem.
7	The size of the gap between
8	legislation and compliance, in my opinion, is
9	the sum of ignorance on the part of both
10	employer and employee; neither can educate the
11	other, which is the beginning of compliance,
12	if they have no knowledge of the subject.
13	In those organizations where
14	serious efforts are being made to understand
15	and comply with laws regarding people who are
16	handicapped, inclusion of the visible
17	condition in their consideration has been
18	slowed to and that is understandable.
19	There is no point in simply
20	waving our arms and getting angry about it.
21	Many of these people sincerely
22	are interested in trying to do something in
23	the way of accommodation, but they don't
24	really look at a hard of hearing person as

1	having a problem; and it is only when the hard
2	of hearing person is willing to explain that
3	in a nonemotional way, that we can make any
4	real progress.
5	Oh the laws on the books aren't
6	going to help too much unless the employer is
7	sensitized to the fact that a communication
8	access is on a bar with physical access; by
9	and large they gravitate toward what they can
10	see.
11	They gravitate toward what they
12	think is obvious; clearly education is the
13	major problem.
14	Employers must be made aware that
15	hard of hearing people are covered under the
16	same legislation that relates to people with
17	physical disabilities; communication access is
18	on a bar.
19	It is accepted in the standards
20	of the architectural there are specific
21	definitions in answer to the question that was
22	raised here before about what constitutes
23	reasonable accommodation in terms of
24	communications access; and they are described

1 as an annex to this paper, which I have submitted to this Board. 2 The employer must be taught what 3 problems arise in the work place due to 4 5 hearing impairment and what can be done to 6 resolve them. 7 Hard of hearing people must be informed of the rights, very difficult due to 8 9 the size of the population, its diversities, and the cloaks of anonymity worn by so many of 10 11 its members. 12 Hard of hearing people must 13 assert their rights; no one is going to do it 14 for them. 15 In spite of the goodwill of many 16 of our hearing people, there is no better advocate in this issue than we who have the 17 18 problem. 19 In our society there is no 20 automatic implementation of law, people have 21 to insure compliance, that requires deep 22 involvement in their own cause; and until 23 recently hard of hearing people did not even

perceive a cause, but they are learning.

1	In our economic and political
2	system social issues usually face an uphill
3	fight, and it is not just in this
4	administration, I do not foresee any
5	administration coming to Washington that is
6	going to be significantly more responsive in
7	our economic constraints over the next 20, 10
8	years.
9	The issues must be made highly
10	visible and presented in a way so that the
11	person having no experience with the problem
12	will understand it and perceive that action is
13	required along the lines recommended by
14	proponents of the issue.
15	This has always been difficult
16	for those conditions which are invisible,
17	first and foremost among that group are those
18	who are hearing impaired.
19	Sign language makes deafness
20	visible; it tells the world that
21	communications are being sent and received
22	manually.
23	For 100 years, organizations of
24	and for deaf people have worked to providing

T	support structures disseminating information
2	about the civil rights of deaf people, and
3	with varying degrees of success have organized
4	constituencies to participate in the process
5	of securing those rights.
6	Just six years ago this month the
7	only successful national organization of hard
8	of hearing people began its struggle to
9	organize and inform a constituency which
10	should soon play a major role in improving the
11	lot of all hearing impaired people.
12	The reason I use that word "all"
13	is that we have the numbers.
14	If you have two million deaf
15	people struggling for years with the
16	assistance of parents and professionals to
17	meet the kind of demands that are placed on
18	by our society, it is difficult.
19	If you add the other 18 million
20	hard of hearing people in this country to that
21	2 million then the numbers become very
22	significant; that last statement implies a
23	commitment on our part to work closely with
21	deaf moonlo and their organizations in nursuit

of our mutual objectives. 1 2 Self-help for Hard of Hearing 3 people is only six years old. We have 12,000 due-paying 4 5 members. 6 We are embarked on an educational program through our publications, foremost of 7 8 which is our Journal S. H. H. H, which has an estimated 200,000 readers. 9 10 We have negotiated a cooperative 11 action plan with the Counsil of State 12 Administrators of Vocational Rehabilitation, 13 Rehabilitation Services Administration, and 14 the National Institute of Handicapped 15 Research, to include specifically services for 16 hard of hearing people. 17 We work closely with and on an 18 equal basis with -- through a joint task force 19 set up by President Gary Lee and myself -- we 20 have been successful in enlisting the support 21 of professional groups in the hearing health 22 delivery field; and we have excellent 23 relationships with professionals throughout

24

the country.

1	Unlike the nice lady who said
2	that the parents were in awe of professionals,
3	I must say we are not.
4	We work with professionals on an
5	equal basis; and we appreciate very much their
6	input, but in awe, we are not.
7	Our 192 chapters and groups in 43
8	states; we have members of all 50 states, are
9	embarked upon a national advocacy program
10	requiring communications access for hearing
11	impaired people in all buildings supported by
12	government funds and all programs supported by
13	government funds, such as the National
14	Endowment for the Arts.
15	Our members are engaged at state
16	and local levels at a variety of legislative
17	activities.
18	In a national survey to determine
19	priorities for services and research during
20	the next 10 years, the problem of hard of
21	hearing and deaf person who face
22	discrimination in the labor market is being
23	examined.

We note that employers have been

Ţ	reluctant to depend upon the worker with
2	impaired hearing, despite evidence that their
3	fears are unfounded, and that such an employee
4	can be as good or better than any other.
5	The prejudices is often deep, and
6	it is reinforced when the hard of hearing
7	person or deaf employee is placed in
8	situations that exacerbate the difficulties in
9	communication.
10	The problem remains large, but
11	the framework for action is in place by
12	including the hard of hearing person in the
13	interpretation of existing legislation and
14	regulation, which has come about through
15	policy changes in the last six years.
16	We are opening a new area.
17	The large numbers of hard of
18	hearing people can reinforce efforts by deaf
19	people to obtain practical means of access in
20	the labor market, through national programs of
21	employer education, grassroot supports for
22	appropriate legislation, and increased
23	representation on Federal and State
24	Commissions, Boards, and organizations which

1	frequently affect us directly.
2	While our problems require
3	different solutions in many cases, deaf and
4	hard of hearing people in combination can
5	begin to realize a world of equal access in
6	by the Rehabilitation Act of 1973.
7	Thank you very much.
8	MR. SCHWARTZBERG: Our next speaker is
9	Miss Bonnie Tucker.
10	She is an attorney specializing
11	in litigation for large corporate clients,
12	particularly antitrust, trade secrets,
13	business torts, and contract disputes.
14	At this point I would like to
15	call just about a minute break for the benefit
16	of the court reporter.
17	(WHEREUPON, a short break was had.)
18	MR. SCHWARTZBERG: I had begun to
19	introduce Miss Bonnie Tucker.
20	Let me, at the risk of repeating
21	myself, go back to the beginning of that
22	introduction.
23	Bonnie Tucker is an attorney with
24	the firm of Brown and Bain in Phoenix,

1	Arizona.
2	She specializes in litigation for
3	large corporate clients, particularly
4	antitrust, trade secrets, business torts, and
5	contract disputes.
6	She received her Doctorate in
7	Jurisprudence from the University of Colorado
8	in 1980 when she was Editor in Chief of the
9	Law Review let me repeat, she was Editor in
10	Chief of the Law Review at the University of
11	Colorado Law School; and she graduated Order
12	of the the highest honor society.
13	She clerked for one year for the
14	honorable William E. Doyle, the 10th Circuit,
15	United States Court of Appeals.
16	She is a member of the Arizona
17	Bar, the Colorado Bar, and the California Bar.
18	She is chairperson of the Arizona
19	Counsil for the Hearing Impaired.
20	She served for the past four
21	years as member of the National Advisory Group
22	to the National Technical Institute for the

She is a member of the Board of

23 Deaf.

1	Directors of the Arizona School for the Dear
2	and Blind.
3	She is the author of several
4	articles on rights of the hearing impaired
5	people.
6	She will teach in 1986/1987 a
7	three-credit course at the Arizona State
8	University College of Law on the rights of
9	physically handicapped persons.
10	She is profoundly deaf, and she
11	has been profoundly deaf since infancy.
12	Miss Bonnie Tucker.
13	MS. TUCKER: Mr. Schwartberg, Members
14	of the Commission.
15	I am very happy to have the
16	opportunity to address you today with respect
17	to employment, with respect to the employment
18	the rights of hearing impaired people have
19	their rights in three courses, the United
20	States Constitution, Federal Statutes and
21	State Statutes and Regulations.
22	Due to the impossibility of
23	covering this very broad I will limit my
24	summary this afternoon to a discussion of

1	Federal.
2	There are three primarily Federal
3	Statutes that are intended to assist
4	handicapped people with respect to employment,
5	Section 501, 503, and 504.
6	The three primary Federal
7	Statutes that I intended to are we all set
8	here?
9	Okay, thank you.
10	I will start from the beginning,
11	okay?
12	With respect to employment, there
13	are three primary Federal Statutes that are
14	intended to assist the handicapped people:
15	Section 501, 503, and 504 of the
16	Rehabilitation Act.
17	The Section 501 of the Act
18	prohibits the Federal Government from
19	discriminating in matters of employment on the
20	basis of handicapped; and it requires federal
21	employers to take affirmative action to hire
22	qualified handicapped people.
23	A qualified hearing impaired
24	person in section 501 is one who with or

1	without reasonable accommodations of and if
2	in terms of education and experience for the
3	particular job.
4	Under this Section federal
5	employers are required to make reasonable
6	accommodations on an individual's hearing
7	impairment.
8	With this statute we have a
9	little it is reasonable if it does not have
10	undue hardship on the operation of a federal
11	agency program.
12	The term reasonable accommodation
13	is specially defined to include interpreters
14	that require job restriction to eliminate that
15	barrier for hearing impaired persons that a
16	medical technology answer the telephone
17	when the secretary is away from the desk.
18	The second statute, Section 503
19	of the Rehab Act requires any company with the
20	federal contract or subcontract with more than
21	\$2,500 to take affirmative action to employ
22	qualified handicapped people.
23	Several hundred thousand private
24	businesses are affected by this statute which

1	requires federal contracts to make reasonable
2	accommodations for employees hearing
3	impairment.
4	Reasonable accommodation per
5	Section 503, but those are not cause undue
6	hardship for the employers when looking into
7	these as the business of necessity and the
8	expense of the accommodation.
9	There are two significant factors
10	to understand with respect to Section 501 and
11	503.
12	First, they both require
13	affirmative action. No discrimination is not
14	sufficient.
15	Therefore, employer is subject to
16	Section 501 and 503 are advocated to actively
17	recruit qualified handicapped people.
18	Second, they both require the
19	basis of reasonable accommodations.
20	Now, if that represents
21	accommodation to allow hearing impaired person
22	to perform the function of a particular job, a
23	reasonable of that must be the nature and

cost of the accommodation.

1	So the ultimate decision will
2	rest on a large part upon the state of
3	technology, and often what services are
4	available to assist the hearing impaired
5	person and how much it will cost the employer
6	to purchase such a system.
7	As technology advances, therefore
8	more and more necessary accommodations will be
9	held to be reasonable.
10	The third Federal Section of
11	Section 504, the Rehab Act, prohibits all
12	recipients of federal from discriminating
13	against otherwise handicapped qualified people
14	because of the term "Federal Finance
15	Assistance" to encompass anyone as well as
16	to this section applies to a vast number of
17	businesses.
18	Moreover, the Supreme Court
19	recently ruled contrary to the ruling of
20	separate lower courts that when a handicapped
21	person seeks relief for employment
22	discrimination on Section 504, it does not
23	have to be shown of the primary objective the
24	federal financial assistance was to promote

7	embrol ment.
2	This ruling will expand the
3	number of situations for Section 504 as held
4	to apply.
5	Unfortunately, however, the
6	recent Supreme Court under a recent Supreme
7	Court ruling, Section 504 will probably be
8	limited to specific department between the
9	business or program that receives the federal
10	financial assistance.
11	For example, if only one branch
12	of a company receives federal financial
13	assistance, that branch will have to comply
14	with the mandate of Section 504 but the other
15	branches of the company will not.
16	Now, this is a serious detriment
17	to the implementation of Section 504 unless
18	and this really needs to be amended.
19	The need for that amendment was
20	recognized and the amendment was proposed in
21	Congress in 1984, that amendment passed the
22	House; as far as I know nothing is happening
23	about that right now.

The goal of Section 504 is to

1	allow a hearing impaired person to reach the
2	same level of achievement in his skill of
3	employment as that reached by others employed,
4	the most integrated setting appropriate to the
5	person's needs.
6	Unlike Section 501 and 503,
7	Section 504 does not contain affirmative
8	action component so employers subject to
9	Section 504 do not have to take steps to
10	affirmively create handicapped people.
11	They do, however, have to provide
12	reasonable accommodations, the same as they
13	are subject to Section 501 and Section 503.
14	With respect to Section 504 an
15	otherwise reasonable accommodation can perform
16	the essential function of the task in
17	question; accommodation is held to be
18	reasonable, it does not cause the employer
19	undue hardship when looking at the nature of
20	the player's program, the nature of the
21	program operation, including the composition
22	and structure and the type and cost of the
23	accommodation need.
24	Now, the issue of reasonable

T	accommodation is directly related to the
2	question of whether a handicapped individual
3	is otherwise qualified for the Supreme
4	Court has ruled to be otherwise qualified for
5	a job, a handicapped person must show that he
6	is qualified inspite of his handicap.
7	The court interpreted this as
8	meaning that the individual must show that he
9	meets the criteria for the job apart from the
10	handicap, and that the physical standpoint
11	relating to his handicap is either not
12	job-related or cannot the court have been
13	very confusing and conflicting decisions with
14	respect to the which will render a
15	handicapped person otherwise qualified for a
16	particular job.
17	Each case is decided in light of
18	the, and unfortunately in accord with the view
19	of the particular judge hearing the case.
20	No one has come up with a clear
21	cut formula for the uniform application of
22	Section 504.
23	It will be helpful if we can to
24	evaluate whether a handicapped individual is

1 otherwise qualified. 2 Even if we succeed in that -- I don't think we will ever eliminate the 3 4 subject. 5 From the things that I am going to assign to my class that I will teach in the 6 University for the first time in my life, I am 7 8 going to give them all the exercise to bring 9 to class one day a summary of how they think 10 we can establish objective criteria for 11 deciding if the person is otherwise qualified. 12 Maybe they are just starting law 13 students, but maybe they will come up with 14 something the rest of us haven't been able to 15 come up with.

16

17

18

19

20

21

22

23

24

There are two other issues in relationship to 504, they need to be mentioned because in my opinion the amendment to this section deals, for example, with the programs specifically of the Federal Financial Assistance or a prospective criteria for determining whether a person is otherwise qualified should also address these two issues.

1	First, Section 504 does not state
2	who has the ability of proving that a person
3	is not otherwise qualified or is otherwise
4	qualified.
5	I think that the statute should
6	be amended to require the employer to bear the
7	burden of proving that the handicapped is not
8	qualified to perform the job in question, even
9	with his assistance of reasonable
10	accommodations rather than with the
11	Handicapped Act to prove that he or she is
12	qualified.
13	Section 504 is intended to
14	eliminate discrimination based on stereotype
15	judgments concerning the capabilities of
16	handicapped people.
17	Thus, the burden should lie with
18	the employer to show that he has conducted a
19	careful and objective inquiry into the actual
20	capabilities of a handicapped person and has
21	concluded that there are real and not imagined
22	reasons why the handicapped person is not able
23	to perform in a job or that there are real

reasons unrelated to the handicapped but for

not hiring the handicapped person.

Absent this requirement, the goal

of Section 504 is easily defeated and has been

4 easily defeated in many published court cases.

5 The second issue involves a

6 question of whether to be given to the

7 decision of administrators of federally-funded

8 programs of the -- that it does not have to be

9 shown that an employer intentionally

10 discriminated on the basis of handicapped in

order to prove that the employer violated

12 Section 504.

19

20

21

22

23

Some courts upheld a considerable

deference should be given to decisions made by

administrators in Federally-funded programs,

as long as there is no evidence that -- intent

or as long as that is not shown that the

purpose of the employer's requirements was to

deny benefits for handicapped people.

Section 504 should be amended to preclude such difference since administrators or employer's decisions may be based upon the very type of erroneous -- about the

24 handicapped being a condition or paternalistic

attitudes that Section 504 was intended to 1 2 remedy. 3 These three statutes, Section 4 501, 503, and 504 of the Rehabilitation Act 5 apply only to federal employers or contractor 6 and to recipients of federal financial 7 assistance. 8 Many states, however, have 9 intended laws that provide educated people 10 with some relief from employment and 11 discrimination in the private sector; 12 additionally almost all states have statutes 13 for prohibiting discrimination. 14 These statutes may be broadened 15 and offer more protection than the federal 16 statutes. 17 Analysis of these laws, many of 18 which are relatively new and untested is 19 beyond the scope of my talk today. 20 The difficulty of acquiring 21 private employers in a free enterprise system 22 to hire a handicapped person and to make 23 reasonable accommodations for the disabilities for these employees is obvious. 24

1	Private employers are concerned
2	with safety, and the maximization of profits.
3	There is little uniformity among
4	the states with respect to questions of how
5	far they are willing to go in telling private
6	employers who they must hire and how they must
7	spend their money.
8	A hearing impaired person who has
9	been discriminated against by a private
10	employer should look to the laws of the
11	applicable state to determine, first, whether
12	this state's antidiscrimination laws apply to
13	private as well as public employers.
14	Second, whether the hearing
15	impaired person falls within the state's
16	definition of a handicapped person.
17	Some states, for example, define
18	that term "handicapped" as a condition
19	unrelated to the ability to perform a job.
20	So in this case all an employer
21	has to do to avoid compliance with this
22	statute is to show that an applicant's hearing
23	is somewhat related to the ability to perform
24	the iob. which can always be shown.

1	Third, the hearing impaired
2	person has to find out the number of employees
3	that the employer must have in order to be
4	subject to this state law.
5	Fourth, if the state law requires
6	that the employer to take steps to take
7	affirmative action to hire and promote
8	qualified handicapped people, and we need to
9	look at the scope and breath of the state
10	statute.
11	On the positive side, more and
12	more states are making substantial efforts to
13	prohibit discrimination against handicapped
14	people, and every year the existence of
15	technology improves, making it easier for
16	employers and administrators or administrators
17	of service to make accommodations for hearing
18	impaired people.
19	On the negative side, the Supreme
20	Court is currently reading the federal law on
21	a very restrictive manner which, in
22	combination with the Reagan Administration,
23	the negative attitude about the civil rights
24	has not been encouraging.

1	Obviously, however, things are
2	looking up for hearing impaired people.
3	We need to insure that this
4	positive trend continues; and I hope that the
5	Commission will take an active role in that
6	part.
7	Thank you very much.
8	MR. SCHWARTZBERG: Sy DuBow has been
9	Legal Director of the National Center for Law
10	and the Deaf since 1975.
11	He has worked on federal and
12	state legislation to advance the rights of
13	hearing-impaired people.
14	Mr. DuBow has also been counsel
15	in several federal court cases involving
16	disability rights.
17	These cases have included issues
18	related to interpreters in colleges and
19	universities, special education, and
20	employment.
21	Two recent Supreme Court cases he
22	has been involved with are Consolidated Rail
23	Corporation versus Darrone supporting full

coverage against employment discrimination in

1	any federal program, and Irving Independent
2	School District versus Tatro finding clean
3	intermitten catherization as a necessary
4	related service for a handicapped child.
5	Finally, he has lectured and
6	written extensively including co-authoring the
7	paperback book <u>Legal Rights of Hearing</u>
8	Impaired People.
9	MR. DUBOW: My job is to wake you up,
10	the end of the day here.
11	Employment opportunities for deaf
12	people, there are real severe restrictions
13	because of negative employer attitudes.
14	When deaf and hard of hearing
15	persons are hired, it is usually at low level
16	jobs, at low pay with few job protections.
17	Even if an employer hires a deaf
18	or hard of hearing person, they show
19	resistance to promote them.
20	Both blue collar and white collar
21	employees who are hearing impaired are
22	continually passed over, ignored because they
23	are deaf from various kinds of promotions.
24	So these employer attitudes are

1	the number one barrier to employment
2	opportunities for deaf and hard of hearing
3	people.
4	Employers often make stereotyped
5	wrong assumptions about the capabilities of
6	hearing impaired people.
7	One study indicates that
8	handicapped people must show generally that
9	they are more qualified than the hearing
10	people that are hired.
11	For deaf people and hearing
12	impaired people the most frequent problem with
13	attitudes relates to that is a sign meaning
14	safety safety.
15	Because of fears of safety they
16	have refused to hire or promote qualified
17	handicapped people, especially deaf and
18	hearing impaired people.
19	One example here in Chicago
20	involved the Chicago Transit Authority; there
21	was one deaf man who graduated from the
22	Illinois Rehab Department, number one, top of
23	his class in training to become a bus

24 mechanic.

1	He was a bus mechanic or a bus
2	depending on the size; and he was turned down.
3	Because they said there would be
4	a danger for all of these buses going into the
5	maintenance garage, and because of that they
б	couldn't hire him.
7	Now, that deaf man himself
8	suggested a very simple reasonable
9	accommodation.
10	Why not just move my worksite to
11	an area that I can see all of those buses
12	coming in?
13	No problem, it is not going to
14	cost you anything.
15	They still refused until we
16	threatened a lawsuit, and then they agreed.
17	But that reasonable accommodation
18	was very simple, and again was suggested by
19	the deaf person themselves, who frequently are
20	in the best position to recognize what is a
21	reasonable accommodation.
22	A little off to the side, an
23	interesting example involves professional
24	architects.

1	There was a library, and it was
2	being constructed, a beautiful library.
3	The problem is they needed a
4	ramp, a ramp, okay.
5	Well, the architect had a
6	beautiful idea; and he had this ramp going
7	like that, a very steep angle with three steps
8	at the bottom.
9	Now, that was a ramp for Evil
10	Kanevil (phonetic) but not for a handicapped
11	person, but not for a handicapped person, but
12	they do that because they do not ask for input
13	from the handicapped person themself.
14	We see that again and again.
15	We see sometimes an employer will
16	call a meeting and not invite the deaf person
17	or the hearing impaired person.
18	For example, the post office has
19	safety meetings, important to discuss what
20	they should protect themselves for; and they
21	refuse to invite the deaf person, and there
22	are many deaf people that work for the post
23	office.
24	Instead they tell the deaf person

1 it is okay, it is all right; we will give you 2 a summary later. 3 We will tell you what is important or what is not. 4 5 What if they had a question that 6 could cause or prevent an injury? 7 Another recent example was from 8 Florida that we were involved representing, 9 where an agency refused custody to two deaf 10 parents. 11 In Florida recently, a deaf 12 couple were denied custody of their children 13 simply because they were deaf, you hear that 14 better? 15 Again, it showed an attitude, an 16 attitude that deaf people or hearing impaired 17 people could not make decisions themselves. 18 You are saying, well, that is 19 kind of crazy in this day and age, but it 20 happens again and again. 21 There is the problem of hearing 22 people never meeting a deaf person; never 23 meeting a hearing impaired person so they

don't know how to handle that situation.

1	Common practice with
2	professionals or with lawyers is the deaf
3	person comes to see the lawyer, and an
4	interpreter comes along or a hearing friend,
5	and the lawyer says pointing the lawyer
6	says to directly to the interpreter or to
7	the friend what is his problem?
8	Tell me about his problem, never
9	looking at him in the eye, the deaf person.
10	How many times has that happened
11	where they ignore you?
12	You are isolated.
13	We see that again and again
14	because of hearing people's insensitivity and
15	inability to deal with a person who has a
16	disability.
17	Recently in Washington I was in
18	court, and they appointed a lawyer who only
19	knows one sign; the sign is "Not to worry."
20	The deaf person walked into the
21	courtroom, and the lawyer said "Not to worry,"
22	and that was it; and the deaf person never
23	knew what happened in the proceeding.
2.4	Thou fust work out prother door

1	and had to pay some money; and they never were
2	explained the proceedings, but we had seen
3	that again and again where deaf people are
4	left in the dark in employment situations and
5	they are frequently, frequently isolated.
6	We have here a situation where
7	courts, employers, legislators are making
8	basic decisions about the rights of hearing
9	impaired people but without involving them in
10	those decisions.
11	Too often in the past hearing
12	people, maybe good intentioned, have spoken
13	about the needs of hearing impaired people;
14	and as Rocky pointed out he just left it
15	is for the deaf people, the hearing impaired
16	people themselves that must affirm their
17	rights, but they have to be given an
18	opportunity to affirm their rights.
19	It is with that in mind that I
20	must say, with a bit of shock and chagrin,
21	that this Advisory Committee failed in meeting
22	their responsibilities.
23	It seems to me that when we talk
24	about an attitude problem, we have one right

1	here on this Committee, when they refused to
2	permit the largest consumer organization of
3	hearing impaired deaf people, the National
4	Association of the Deaf, ten minutes to speak
5	at this forum when they allowed 10 minutes to
6	speak on the qualifications of one lawyer
7	here.
8	I found it totally shocking
9	because when I was informed of this conference
10	to substitute for Mr I noticed that they
11	were all representives of the Alexander Graham
12	Bell Assocation, that's fine; there should be
13	your presence here, but I noticed no members
14	from Illinois.
15	I noticed no members from the
16	frat I notice no members from the National
17	Association of the Deaf.
18	Now, these are groups and no, not
19	the President of the Illinois Association of
20	the Deaf; and I recommended them to Mr. Lucas,
21	and he said fine.
22	And then on my urging J.B. Davis
23	and were here but no others; and I find
2.4	that a hit chocking in this time if we are to

1	have a balanced forum, not to include the
2	major organizations involving deaf people.
3	A. G. Bell should be represented
4	and should be involved, but we should be all
5	of us to discuss all of our problems and not
6	have just half a loaf.
7	It is with that that I close; and
8	I have submitted my paper and my
9	recommendations, but I would hope that the
10	proceedings here will be open for addition
11	from other groups that have a contribution to
12	make so that the future of hearing impaired
13	people will not be determined by hearing
14	people without full input.
15	Thank you.
16	MR. SCHWARTZBERG: I am sorry that Mr.
17	DuBow did not look at the here at the
18	out at the time first that additional
19	materials can be submitted in writing at any
20	time through July.
21	Second, we regret that the time
22	that made any discussions we include
23	three of his discussions; and we further

24 regret that Mr. DuBow has seen fit to take the

1 attitude which he takes. 2 Nevertheless, this Commission 3 will as it always does be open to people who 4 wish to appear before it. 5 We have often opened up to hear 6 those who wish to appear before us. 7 We meet on a regular basis in the federal building. 8 9 And if there are additional 10 people who wish to appear before us, we will, 11 of course as we always have, hear them. 12 Now, let me turn to the next 13 person on the agenda, and that is Mr. Alan 14 Post. 15 Alan R. Post is a graduate of the 16 University of Wisconsin, Madison, with a law 17 degree in 1972. 18 He is admitted to the Bars of six 19 states, including Wisconsin, Maryland, 20 District of Columbia, Nebraska, Minnesota and

He is currently an attorney with Sorling, Northrup, Hanna, Cullen and Cochran, Ltd., Springfield, Illinois.

21

Illinois.

1	He has been an attorney with the
2	Interstate Commerce Commission Washington,
3	D.C., 1972-76, the Union Pacific Railroad
4	Company, Omaha, Nebraska, 1976-78, the
5	Burlington Northern Railroad Company, St.
6	Paul, Minnesota, 1978-83, Illinois Bell
7	Telephone Company, Chicago, Illinois, 1983-86.
8	Activities involving persons with
9	handicaps include membership in the State of
10	Minnesota Counsil for the Handicapped, St.
11 -	Paul, 1982-83, Legal Review Chair of the Spina
12	Bifida Association of America, Chicago, 1983
13	to present, and President of the Spina Bifida
14	Association of Minnesota, Minneapolis, 1982.
15	His office address is P.O. Box
16	5131, 800 Illinois Building, Springfield,
17	Illinois, 62705.
18	It says here and I am not
19	really here to give his phone number I am
20	here to provide you with the information that
21	we welcome.
22	Alan R. Post.
23	MR. POST: I suppose that makes me
24	sound like a redneck conservative utility

T	lawyer that doesn't necessarily mean that I
2	I share a lot of frustration, and I have felt
3	them personally, and I know exactly what he is
4	talking about, where he is coming from.
5	The correct pronunciation is
6	Spina Bifida.
7	I appear before you also as a
8	parent of a 10-year-old daughter who cannot
9	walk; that young lady was put in the hospital
10	this morning for some major surgery, so if I
11	seem a little less connected, she is at
12	Children's Hospital undergoing the knife right
13	now.
14	The Commission did ask some very
15	pertinent questions, and rather than burdening
16	them, I would like to address some of the
17	questions.
18	One of the questions was how can
19	you help parents carry out their legal rights?
20	One way you can do it is to pay
21	their legal bills when they win.
22	There was a brief mention of the
23	Tatro case, they ran up a quarter of a million
24	legal bill to require that their 7-year-old

1	daughter have some very personal private
2	administration through plumbing so that she
3	could go to the bathroom, otherwise they are
4	going to keep her out of school.
5	Now, every other civil right,
6	whether you are a female, black, anything but
7	the handicapped, we can get legal bills paid.
8	Handicapped folks, sorry, you
9	cannot get your legal bills paid when you win;
10	that's assuming you are willing to put up with
11	the trauma of the litigation, the hassles of
12	having your daughter's toliet matters made
13	public.
14	I understand that, my daughter
15	has the same procedure, that's one suggestion,
16	cover the legal bills, go along with Senator
17	Witer's (phonetic) bill to amend 503, 504.
18	When the parents win, to cover
19	the legal bills; the parents will then sit up
20	and listen because they will have some
21	incentive to hire I am not in that area of
22	practice; I have been a salaried employee
23	worried about utility price increases, you all
24	love that when your electricity that has

1	been my fun and games for the last 15 years.
2	Technology, I agree
3	wholeheartedly with Doctor Stoker that I am a
4	child of technology.
5	Our sights are not set high
6	enough here.
7	I have been fooling around with
8	my briefcases, kind of inbetween speakers,
9	with all of the gadgets and garbage that I
10	carry around to cope with courtrooms, noisy
11	meetings, but our sights are nowhere near high
12	enough.
13	I started with headsets from
14	crystal radio and my electrical engineer
15	father said "Here, you can hear better with
16	this."
17	And we graduated to a headset for
18	the family tv set, and all of this junk back
19	here, and basically radio and headsets, jam
20	the sound into your ears.
21	That is Rocky's philosophy,
22	self-help for the hard of hearing; don't buy
23	the thing from the hearing aid dealer, buy it
24	from Radio Shack or the local fix-it store,

1	but jam that sound into your ears in some
2	self-help manner.
3	I will give you a crazy example,
4	but the next time you see a Federal Express
5	truck, look inside the front seat and see if
6	you don't see a little tv screen; that thing
7	is a little TDD, and somebody can write
8	messages across that, and that's exactly what
9	they do in that Federal Express truck.
10	You can be deaf, and they can
11	write you messages; stop the truck before you
12	do that.
13	And you can see the messages;
14	very interesting because of office automation
15	you have a tv set.
16	You have secretaries, you have
17	people pounding things out.
18	Pretty soon we will all be
19	writing messages back and forth; we come into
20	the mainstream.
21	I have been into the mainstream;
22	it is difficult, but we are starting to get
23	somewhere.

How much should it cost?

1	I know people who are paralyzed
2	from the neck down, they breathe on oxygen.
3	They live in an electrofied
4	wheelchair; it costs the employer \$20,000 to
5	modify the work place so they can modify
6	our sights are nowhere near high enough.
7	None of this stuff behind me
8	costs I am a \$45,000 a year taxpayer
9	isn't that a reasonable invention?
10	It is cheap. Please wake up to
11	it; it is cheap for us.
12	One of the speakers emphasized
13	early diagnosis, absolutely essential and
14	critical.
15	Start jamming the normal sounds
16	into the kid's head as soon as possible.
17	People look at me and say "How
18	did you ever learn to talk normal?"
19	My mom and dad jammed that sound
20	into my ears, \$20 headset for the tv set.
21	Again, what is reasonable
22	accommodation?
23	\$20,000, you make 50,000 a year,
24	that is cheap.

1	Over a course of a lifetime you
2	pay quite a few taxes, why not blow your money
3	some more on cellular technology or any kind
4	of technology.
5	That was but it is basically a
6	telephone, and all you do is use it like an
7	ordinary telephone.
8	Why worry about the carry your
9	own phone with you; put it up to your ear and
10	bing it out and get a volume control.
11	Now, that is 3,000, but you will
12	find them running over in the Chicago
13	Mercantile Exchange, the traders make enough
14	money, and they live by the telephone.
15	The doctors start doing the same
16	thing not so crazy.
17	I have a phone like this in my
18	car because it was a phone that I know I can
19	depend on.
20	I come to the Chicago Hilton, I
21	can't use the phone in the room; my wife says
22	"My God, I can't even hear."
23	The old trusty Radio Shack

device, \$30.

1	It happens to be made by Walker,
2	and it is pretty good; plug it in, and it
3	works if everything is compatible.
4	I have been accused of wanting to
5	be a big shot; I asked for a speaker phone.
6	It costs you \$70, but someplaces
7	have very rigid regulations how many square
8	feet you can have, depending on what your
9	grade level is.
10	But if you are hearing-impaired,
11	you deserve it.
12	How much does it cost?
13	Go down to the store and buy one
14	if you have to, but buy a good one; buy a
15	cheap one, it is no good.
16	I think I have got everybody's
17	attention.
18	I got started on this thanks to
19	Mother Bell, and I don't have that many good
20	things to say about Mother Bell, but one of
21	the good things they did in 1974, they came
22	out with one of the first hearing aid
23	compatible adaptors that cost all about \$7.
24	How did I find out about it?

1	Not through a program.
2	It was buried in a dark fine
3	print of the Wall Street Journal, but you
4	cannot get this today.
5	It costs about 90 to \$100.
6	When we had Mother Bell as a
7	group, they provided them to everybody for
8	about \$7.
9	The basic purpose is to make any
10	phone hearing aid compatible, that's what this
11	gadget does.
12	And you thought that I was going
13	to talk about the law.
14	So I decided to bring the
15	technology with me; that was a small tv set.
16	If you are in the yuppy you
17	see people carrying these things and wearing
18	the fancy headset along with it, yuppy shoes
19	and their fancy clothes and everything else,
20	but it is a damn good hearing aid.
21	A transistor radio from 1957,
22	what would we do without it?
23	It was invented in 1947, just
24	think how far we have come.

1	And, of course, last but not
2	least, it is a tape recorder that you can also
3	use with the headset.
4	And if you look at all of the
5	Japanese tv sets, they are two steps ahead of
6	us.
7	They have two jacks on virtually
8	every tv.
9	You can plug in one of these in,
10	and you can listen at your own pleasure or you
11	can kill the sound completely.
12	How much does this cost?
13	\$20, cheap.
14	I got lots of batteries back
15	there.
16	But this is a transistor radio
17	that I started out with in 1958, probably one
18	of the first ones made.
19	You might laugh if you see the
20	ads from the Chicago area or other major
21	metropolitan areas.
22	I think if you want to be an
23	outside salesperson some day that has a lot of
24	value because like the Federal Express truck,

1	that machine has the of shipping and
2	receiving written messages, which is what a
3	TDD does.
4	And if you can turn the sound up
5	in a speaker phone box in the sort of a car,
6	it is amazing how well you can control your
7	sound environment.
8	All of you who are
9	hearing-impaired, you have to reach out and
10	try it.
11	If you sit around and wait for
12	the government to do it all for you, you are
13	going to be waiting an awful long time in
14	today's environment.
15	These people are
16	well-intentioned, but they are living proof
17	how they aren't going to do it for you.
18	\$3,000 for this, go ahead and
19	laugh.
20	Try it, it is hearing aid
21	compatible; and for those of you who have
22	money or who have access to it or if you
23	want to go to work.

Hey, I have a hearing impairment,

1	but provide me with one of those things that
2	you give to the brokers, I want to be able to
3	use the phone.
4	And you become a taxpaying
5	citizen, and you can raise hell when you feel
6	like it when you have to.
7	Now, this is a long way from my
8	speech I am open for questions.
9	I know we are running way behind,
10	so I will just stop there.
11	MR. SCHWARTZBERG: Robert Mencehl.
12	Robert S. Menchel is a Senior
13	Career Opportunities Advisor and Assistant
14	Professor at the National Technical Institute
15	for the Deaf at the Rogers Institute for
16	Technology.
17	He received his BS in Physics
18	from Clarkson University and his MBA with
19	honors from the Rochester Institute of
20	Technology.
21	An expert in the employment of
22	handicapped people, he served as the National
23	Role Model under the AAAS in 1977-78.

He is on numerous Boards of

1	Directors and served as the chairperson of the
2	New York State Governor's Committee on
3	Employment of the Handicapped.
4	He has published and presented
5	more than 60 papers related to the employment
6	of hearing-impaired people.
7	Mr. Robert S. Menchel.
8	MR. MENCEHL: I think that I can speak
9	my experience inside myself when I
10	interviewed once I interviewed for 63 jobs
11	after I graduated from college, and so I think
12	I have a little bit of background; and I feel
13	very strongly that all people have the right
14	to employment and job satisfaction.
15	Work is more than a job.
16	Work offers us independence,
17	securities, to earning an income.
18	It also gives us a social
19	mobility.
20	There are so many conflicting
21	things right now about the underemployment and
22	unemployment of the hearing-impaired, and this
23	is very difficult to really know where we
24	stand.

1	For example, in 1974 it was said
2	there was a very high percentage of
3	hearing-impaired people who were employed,
4	something like 97 percent were working, oh
5	that's wonderful.
6	In 1985 they reported that males,
7	20 years old or over, had an employment of
8	73.9 percent, and that was a little bit lower
9	than national average of 73.3, again that is
10	wonderful.
11	But I recently received a report
12	that said not only are many deaf Americans
13	underemployed, but underemployment among deaf
14	Americans is very high, compared with a
15	general public and even other disability
16	groups.
17	52 percent 71 percent in
18	California.
19	It is very hard to believe
20	that I don't know if we have on what it
21	is and what employment is of the
22	hearing-impaired, but I am seeing that it is
23	higher for the average for some and probably
24	not for others, but I will explain as I get to

the end of my paper.

I have a question now that I am

very much afraid of, one that I'm -- one that

I have -- I have seen becoming -- probably one

that we have overlooked, and, that is, are we

6 preparing the hearing-impaired for the job

7 market?

What I am scared of, and what

people are overlooking, is that those who are

not hearing-impaired, those who have a college

education or those who have even one or two

years of college education are taking away the

entry level jobs that the hearing-impaired

people used to go into.

They are no longer going to be available for them.

A second thing that many of us
may not be thinking about is hearing-impaired
people are not competing with other

20 hearing-impaired people for jobs.

We have to remember that they are competing with the hearing people.

23 If hearing people have better and 24 better education than we are providing to the

1	hearing-impaired, then we are going to see
2	them employed.
3	It is very clear that the
4	connection between higher unemployment rates
5	and low-level education shows the importance
6	of education in the job market; and we will
7	need more of it as we progress.
8	We also have begun to see that if
9	you look back on it, that the deaf people were
10	put into jobs what were called where they
11	are working with things.
12	Some of them were working with
13	technically a very, very small percent were
14	working with people.
15	But as we look down the road, we
16	are going to see more and more jobs be the
17	ones where we are working with people.
18	Employers are looking for
19	qualified deaf people who have the skills that
20	are in demand in the marketplace.
21	' These people will be found in the
22	educational system.
23	The fact that the educated and

qualified deaf people can successfully compete

```
1
         in the labor market is borne out by many
2
         records.
                      NTID has had a very high
3
4
         percentage of employment of 95 percent in
5
         1983, that is just an example.
6
                      So if we are providing direct --
7
         the right type of education, then we are
8
         seeing that we can meet the labor market okay.
9
                       I think that we will take --
10
         excuse me just a minute -- I think that
11
         parents and educators and vocational people
12
         should take pride, and also I think that we
13
         must give a lot of credit to young deaf people
14
         themselves that have entered the work force,
15
         and have composed what they have done, but we
16
         still have a long way to go.
17
                       Let me give you quickly what we
18
         have already spoken about, some of the
19
         barriers of employment for the
20
         hearing-impaired.
21
                       The employers again and again,
         through all of the studies that I have looked
22
23
         over, going way back into time right up to
24
         today, is that they worry about communication.
```

1	They worry about safety,
2	insurance rates, both real insurance rights
3	and unreal insurance rates, and what kind of
4	attitudes do employers have towards the
5	hearing-impaired?
6	But probably the most important
7	thing is that they do not have the
8	information.
9	There is a great amount of
10	information available, but it is not getting
11	down to the people who need it.
12	They are worried about safety.
13	There are many papers that have
14	proved that hearing-impaired employees are
15	safe.
16	I, for one, already have one
17	handicap; I don't want another one.
18	They have had negative
19	experiences with one deaf person so they go
20	and lump all of them into the same category;
21	and they stereotype them, and they will not
22	hire another one.
23	They are worried about
24	performance on the job.

1	They will use every reason in the
2	book they can not to hire one.
3	They will use medical exams, and
4	say you did not pass that because of your
5	hearing impairment.
6	They will use tests with
7	vocabulary that can't be understood by a
8	hearing-impaired person; and they will use
9	that as an excuse for not hiring.
10	They will say they cannot afford
11	the time and afford the requirements that are
12	necessary to train them.
13	They think it would take more
14	time, but there are the mistakes there is
15	information; there are training programs.
16	The problem is that that
17	information is not reaching the employers, or
18	if it is reaching the employers, it is
19	reaching a very small number of them.
20	We do not have the resources to
21	reach out all across the United States.
22	We do not have the resources to
23	supply all of the employers with information.
24	NTID does have some excellent

1	training programs, but the numbers of
2	employers that we can reach are so small
3	compared to the need.
4	They also think that there are
5	only some jobs for deaf people.
6	They feel that deaf people cannot
7	do this, this, this or this, but they can do
8	this.
9	Sometimes they think that the
10	jobs that are very noisy are best suited for
11	deaf people, and that is a mistake.
12	We know that, but they don't know
13	that.
14	It is becoming very clear that
15	they do not have the information.
16	They do not know what kind of
17	support there is.
18	They are not aware that there are
19	resources available for them.
20	We have found from our
21	experiences that we will supply them with
22	backup, backup support, they are more than

Also we have told our employers

willing to take a chance on it.

23

1	that if you have a problem five years down the
2	road, call us up; we are not going to place
3	someone in a job and then close the door on
4	them.
5	We will be able to help them any
6	time.
7	We are working with a company in
8	New York right now who hired one of our
9	students about 8 years ago, and they called us
10	up a month ago; they are having some problems
11	with him.
12	Also we have people working in
13	affirmative action with people who are working
14	with hearing-impaired people.
15	We have placement people in the
16	Federal Government who cannot communicate with
17	the hearing-impaired employees; they don't
18	have TDDs in their office, so the people can
19	communicate with them.
20	We must provide this information.
21	We must develop some type of
22	system that will provide this information to
23	the people who so badly need it.
24	It is so important.

1	I can quote from one person, "A
2	deaf employee's successes are related to his
3	or her supervisor's awareness and
4	understanding of deafness."
5	Another problem of I guess it
6	is upward mobility.
7	It has been a serious problem for
8	years, but again we are not really sure what
9	the problem is about upward mobility.
10	Some people have said that it is
11	very, very big problem.
12	Other people have said that they
13	don't really know if it is there.
14	I feel presently that it is a
15	serious problem.
16	We find hearing-impaired people
17	who have been working wonderful on their job,
18	who are receiving very good job ratings, but
19	we do not see them being promoted.
20	Why is that not so?
21	If we really think about it for a
22	minute, if the deaf workers are underemployed,
23	it is not because it is not the deaf person
24	who is losing out, it is society itself

1	because we are wasting a valuable resource by
2	keeping these people underemployed.
3	It is a waste of our natural
4	resources that could be better utilized.
5	And if we commit them part of
6	the problem with underemployment is due to the
7	employers' attitudes, but I also think that
8	part of the problem is due to communication.
9	And for many of them the use of a
10	telephone, very often an employer will say "As
11	you move up, you will have to communicate more
12	and more with other people, $^{n}$ and the employer
13	is afraid that the hearing-impaired person
14	cannot do it.
15	They don't know or they may not
16	be willing to give the support service needed
17	for that; maybe they are not willing to bring
18	in an interpreter, maybe they are not willing
19	to provide the technology that would make
20	those things available to them.
21	Yes, I will agree that the
22	telephone has been a problem, but what we have
23	heard already today about technology and what
24	it is and I myself use the telephone

1 everyday -- my computer is becoming more and 2 more my telephone. 3 And I think that we are seeing, a little ways down the road, when we'll have an 4 audiovisual voice that people can communicate 5 б on the computer what you are hearing on the 7 telephone. 8 But right now, again to quote, "To a deaf person the telephone is a constant 9 10 remainder of his handicap and of his 11 dependence on others for its use. " 12 It also stands as an invisible 13 barrier to his vocational advancement for he 14 has found, from sad experiences, an 15 employment -- he is considered for employment 16 only to positions which do not require the use 17 of the telephone. And I think another thing that 18 19 employers are not really aware of is that they 20 are not really aware of the fact that most, if 21 not all people, have -- hearing; they do not 22 know that many, many deaf people can use the

We have to educate them on that.

23

24

telephone.

1	There is so many things that have
2	already been brought up; I am not going to
3	repeat the things that are on the market.
4	We must also look at the barriers
5	of training that we give for upward mobility;
6	and we must look at the problem of retraining
7	for those people who may lose their jobs or
8	who may changing technology that will
9	become very, very serious problem if we do not
10	think right now about retraining and preparing
11	deaf Americans, we will be facing an even
12	higher unemployment.
13	But I am not going to put all of
14	the blame on the employers.
15	I am going to state right now
16	that educators, parents, and vocational people
17	must take part of the blame.
18	If you do not get these children
19	or give these children the encouragement that
20	they and the information that they need,
21	they are not going to reach the potential that
22	they can.
23	Creative programs are important
24	to these deaf children.

1	We, as teachers, we, as parents,
2	and we, as educators, must be aware of what
3	deaf people have done.
4	We must show them that if they
5	want to try something, they can do it.
6	You must not limit them; you must
7	not limit them in any way.
8	You must give them all of the
9	opportunity you have.
10	You must never, never say to a
11	deaf person you cannot.
12	Let them try. How do we know
13	what they can do if we don't let them try?
14	That is your responsibility as
15	teachers, as parents, and as vocational
16	people.
17	MR. SCHWARTZBERG: I think it is on
18	that note that we are going to have to call
19	time.
20	And we thank you very much.
21	As you know, the Commission has
22	the full draft, and this is certainly one of
23	the papers where the full draft is of
24	extraordinary strength.

1	Thank you very much, Professor
2	Menchel.
3	The first one is Joyce Tucker who
4	is the Director of the Illinois Department of
5	Human Rights.
6	I have been carefully garnering
7	biographies of each of the participants, and I
8	looked in front of me, and I find that I do
9	not have a biography of Miss Tucker.
10	I suppose the reason for that is
11	that Miss Tucker has been an extremely active
12	member of the Committee which planned this and
13	serves as sponsor of the Illinois State
14	Advisory Committee of the Illinois rights, but
15	she does that over and beyond her work as the
16	Executive Director of the Illinois the
17	State of Illinois Department of Human Rights.
18	And it is my very great pleasure
19	to introduce to you at this time Miss Joyce
20	Tucker.
21	MS. TUCKER: I think that there are
22	those who are in the audience that thank that
23	bio was not there.
24	It is enough to say that I am the

1	Director of the Department of Human Rights,
2	and I have been since July 1 of 1980 when the
3	Department was created.
4	I am also an attorney.
5	Miss Tucker, the other Miss
6	Tucker, shared with you the Federal Law
7	governing the rights of hearing-impaired; and
8	she suggested that it was important that you
9	take a look at the state law in order to
10	understand the full protections available to
11	the hearing-impaired in any particular state.
12	In Illinois the state law
13	governing the rights of the hearing-impaired
14	is the Illinois Human Rights Act as well as
15	the fact that the State Constitution itself in
16	Section 19R001 prohibits discrimination
17	against handicapped people.
18	Illinois is one of the few states
19	with explicit constitutional protection for
20	disabled people.
21	The Illinois Human Rights Act,
22	the Act which the Department of Human Rights
23	administers prohibits discrimination in
24	employment, housing, access to financial

1	credit, and public accommodations, and
2	employment on the basis of race, sex, color,
3	religion, age, ancestry, marital status,
4	unfavorable discharge from the military
5	service, and handicapped, physical and mental.
6	This Act establishes civil rights
7	for hearing-impaired people in the following
8	four areas: employment, housing transactions,
9	access to public accommodations, and financial
10	credit.
11	The Act also expressly prohibits
12	discrimination against people who use hearing
13	dogs.
14	The landlord or manager of a
15	housing unit is restricted from adding a
16	surcharge to a lease because of the presence
17	of a hearing dog.
18	The employment coverage of the
19	Human Rights Act is probably one of the
20	broadest of any states civil rights law.
21	All employers, with the exception
22	of the Federal Government in the State of
23	Illinois, is covered by the handicapped
24	provisions of the Human Rights Act.

1	The prohibitions in the areas
2	other than handicapped require an employer to
3	have 15 or more employees, but with regard to
4	handicapped the law says you have one or more
5	employees and that covers everyone.
6	In addition, an employment agency
7	cannot really refuse to hire or refer an
8	individual that has a hearing impairment; and
9	no employer in the state can lawfully deny
10	employment for a person that has a hearing
11	impairment.
12	Any labor organization is
13	prohibited from refusing to represent a person
14	because of a hearing impairment.
15	Offering further protection for
16	hearing-impaired people, the Act, through its
17	interpretive rules, requires all to provide
18	reasonable accommodations.
19	Accommodations can include job
20	restructuring, reassignment of duties of a
21	hearing-impaired employee, acquisition of
22	equipment such as a TDD, and the provision of
23	sign language interpreters and other similar
24	actions.

1	Miss Tucker, I think, explained
2	in quite detail what a reasonable
3	accommodation is, and the Illinois law tracts
4	the federal law.
5	Job applicants in Illinois have
6	the right to request reasonable accommodations
7	to the test procedure or other personnel
8	processes, such as the interview.
9	An applicant who would like a
10	sign language interpreter for an interview can
11	request that from the employer.
12	Labor organizations have to
13	provide accommodations to the hearing-impaired
14	members which may mean providing an
15	interpreter at union meetings.
16	In some cases reasonable
17	accommodation can make the difference between
18	the hearing-impaired person being able to do a
19	job at all.
20	The Human Rights Commission, the
21	adjudicatory body under the Human Rights Act,
22	has recently ruled that an employer must
23	reinstate a hearing-impaired man to an
24	alternative position and provide accommodation

1 of a pager that has a flashing light instead of an audiosignal. 2 3 The employer, a hardware store, 4 had argued this, a deaf person could not work 5 in such a capacity because when he was in the warehouse he could not be reached for work 6 7 assignments. Two things that should be 8 stressed in regards to employment cases are 9 that in the State of Illinois a charge must be 10 11 filed with the Department within 180 days of 12 the alleged discriminatory event, and that 13 necessary accommodation must have been 14 requested by the employee or the job 15 applicant. 16 Various forms of relief are 17 available under the Human Rights Act, 18 including reinstatement, back pay, restored 19 benefits, cease and desist orders as well as 20 attorney's fees. 21 The Human Rights Act is a very 22 comprehensive bit of legislation, but no one 23 law can address all of the concerns of the 24 hearing-impaired.

1	There are two other recent
2	Illinois laws affecting hearing-impaired
3	people which you should know about.
4	The first requires the state to
5	provide TDDs for all deaf people in the state
6	free of charge; this law must be fully
7	implemented by 1997.
8	The second law requires
9	transportation centers such as Am Track,
10	airports, and bus stations to provide TDDs for
11	its deaf customers.
12	We believe that Illinois has a
13	good foundation in law for insuring the rights
14	of the hearing-impaired; more is needed.
15	The Department of Human Rights is
16	willing to work with others to not only
17	broaden but strengthen the laws governing the
18	rights of the hearing-impaired.
19	You probably have this flyer that
20	my staff prepared on the rights of the
21	hearing-impaired.
22	We have a separate flyer on the
23	variety of handicapped issues which the
24	Department governs in terms of the Human

- We are willing to listen to your

  suggestions and receive your input in terms of

  what the State of Illinois should be doing and

  what the Department of Human Rights can be

  doing with respect to the rights of the
- 7 hearing-impaired.

13

21

22

- 8 Thank you.
- 9 MR. SCHWARTZBERG: At this time I would 10 like to call Ms. Jill Wine Banks.
- Many of us first became aware of

  Miss Jill Wine Banks at the time of the
- 14 At that time she was a member of
- the prosecutorial staff.

co-conspirator.

Watergate matter.

- I think to some extent some

  17 newspaper photographers particularly enjoy

  18 singling out Miss Banks at that time, but

  19 subsequently some of us were less aware

  20 perhaps that she tried the coverup conspiracy
- 23 After that point she next 24 achieved visibility as General Counsel of the

case in which President Richard M. Nixon was a

I.	United States Army.
2	She served at the Pentagon in
3	1977 thru 1980, and her appointment by
4	President Jimmy Carter.
5	After that she became a partner
6	of the Chicago firm of Jenner & Block.
7	When the Attorney General of the
8	State of Illinois indicated that he was going
9	to restructure his office to establish a
10	Solicitor General with primary concern over
11	all of the appeals within the state, he chose
12	as the first person to hold that office Miss
13	Jill Wine Banks.
14	She presently serves as Deputy
15	Attorney of Illinois; and it is my pleasure at
16	this point to present her to you.
17	Miss Jill Wine Banks.
18	Thank you very much.
19	MS. BANKS: Personal regards to you
20	from the Attorney General.
21	Attorney General Hartigan indeed
22	sends regrets to everyone.
23	He is very sorry that he could
24	not be here with you today personally,

1	unfortunately the responsibilities of his
2	office have forced him to be out of town.
3	It is my pleasure to represent
4	him here today; and I would also like to point
5	out that many members of our staff are here
6	today, the head of our Disabled Advocacy
7	Division, Susanna Smith, and I think Karen
8	Myer was on a program this morning, and Leanne
9	Marshal Cohen, I see sitting in the back who
10	works for our division.
11	We also have in the audience, I
12	think, many members of our Advocacy Division's
13	Advisory Counsels, and although I am not going
14	to take the time to introduce them all, they
15	are all here and can be helpful to you if you
16	have any questions later on.
17	The concern that has brought all
18	of you to this forum, the rights of the
19	hearing-impaired is a subject about which
20	Attorney General Hartigan feels very strongly.
21	As a result of his concern the
22	office has provided vigorous new
23	representation for hearing-impaired persons
24	since the time he became Attorney General in

1 1983. 2 This representation has been --3 new legislation, changes of policy, and negotiated agreements that make an important 4 5 difference in the lives of the 6 hearing-impaired citizens of Illinois. Before I go over some of the 7 8 office's accomplishments concerning the 9 hearing-impaired, let me explain briefly the 10 new direction of the office and the focus of 11 the office under Attorney General Hartigan. 12 Attorney General Hartigan, more 13 than any other person who has held that office 14 in history, has emphasized the Attorney 15 General's role as the lawyer for all of the 16 people of this state. 17 By Illinois Statute the Attorney 18 General represents other offices and agencies 19 of the state government, but state law also 20 empowers the Attorney General to act as the 21 lawyer for all of the citizens whenever their 22 rights as a group are at sake. 23 Our common law provides a broad

interpretation of this power.

1	In addition to continuing and
2	strengthening the advocacy programs begun by
3	Attorney General Hartigan's predecessor, which
4	included programs for consumers, utility
5	customers, crime victims, and also the right
6	of people to have a safe and clean
7	environment, Attorney General Hartigan has
8	created new divisions to expand this role.
9	He has created divisions to help
10	the interests of senior citizens, farmers,
11	veterans, and disabled persons.
12	These new divisions have been
13	greatly aided by individual councils which the
14	Attorney General formed to give guidance to
15	these divisions.
16	Representatives of other state
17	and local agencies, businesses, and the
18	professions, as well as members of the
19	academic community and citizen groups serve on
20	these counsils.
21	They serve voluntarily and
22	without pay and give the various divisions to
23	the office invaluable insight and expertise.
24	The new division of the office

1	which most directly concerns all of you here
2	today is the Disabled Persons Advocacy
3	Division, which the Attorney General created
4	in the spring of 1983.
5	Ours is the first and only state
6	Attorney General's Office in the country to
7	have a full-fledged division devoted solely to
8	protecting the rights of the disabled persons
9	and investigating the legal and societal
10	problems they face.
11	The division, which is guided by
12	two advisory counsils, one composed of lawyers
13	and one composed of consumers, brings issues
14	concerning the rights of disabled persons to
15	the attention of the courts, the state and
16	federal legislatures, and the public.
17	I am going to concentrate here
18	this afternoon specifically on the division's
19	advocacy for hearing-impaired persons, but
20	first I would like to point out that the
21	division's efforts over the past three years
22	have resulted in important victories for
23	citizens with all types of disabilities.
24	To cite just one example, at the

1	recommendation of the division and its
2	advisory council Attorney General Hartigan
3	represented 30,000 disabled Illinois residents
4	in a successful lawsuit against the Federal
5	Social Security Administration.
6	As a result, these citizens are
7	now receiving millions of dollars in benefits
8	which they had previously been denied.
9	They had been wrongfully
10	terminated, and as a result of this lawsuit
11	they were reinstated.
12	In order to be of the greatest
13	possible help to hearing-impaired citizens on
14	a daily basis, the office has acquired
15	necessary specialized equipment and staff
16	members have necessary specialized skills.
17	The main office in Springfield
18	and Chicago and regional offices located
19	throughout the state are equipped with TDDs.
20	Division staff in Chicago are
21	fluent in total communication; other staff
22	members are aware of the obligation to provide
23	interpreter services and are learning total
24	communication through an inhouse program or

1	through qualified organizations at the local
2	level.
3	Staff from the Disabled Advocacy
4	Division in Chicago are travelling statewide
5	this summer to provide regional office staff
6	members with training concerning the rights of
7	the hearing-impaired.
8	Karen Myer, who I mentioned
9	earlier was on a program this morning or at
10	some other time during this conference, has
11	organized this entire training program
12	throughout the state for our office.
13	The division is working with
14	other divisions to provide appropriate
15	referral of hearing-impaired citizens to the
16	various programs that the office offers.
17	For example, a hearing-impaired
18	citizen with a consumer complaint will be
19	referred to the consumer division, but a
20	qualified member of the Disabled Persons
21	Advocay Division will remain available for
22	interpretation and necessary assistance.
23	With this system of linkage
24	within the office and a statewide training

1	program, Attorney General Hartigan is
2	determined to improve on the record of service
3	to the hearing-impaired, which the office has
4	already established.
5	Over the past three years the
6	office has negotiated and successfully
7	resolved numerous individual cases involving
8	the rights of the hearing-impaired; many of
9	these cases have centered on the right of the
10	hearing-impaired to have interpreter service
11	when necessary.
12	For example, the division upheld
13	the right to interpreter services for
14	hearing-impaired persons involved in courtroom
15	proceedings and in dealings with the Social
16	Security Administration.
17	The division also successfully
18	defended the right of the hearing-impaired
19	parent to have interpreter services provided
20	by a child's school so that the parent could
21	fully participate in meetings concerning the
22	child's education.
23	The division also upheld the
24	right to interpreter services for holding

1	employment in either private sector or the
2	public sector.
3	In one case the division
4	negotiated with a suburban park district to
5	provide interpreter services for a young girl
6	who wanted to play on a softball team.
7	The division successfully
8	resolved a case in which a citizen was denied
9	auto insurance on the basis of hearing
10	impairment, and an out of court I am
11	sorry an out of date and unfair hearing
12	impairment testing procedure which was being
13	used by a transit authority for employment
14	applicants was changed as a result of the
15	division's negotiations.
16	The procedure the transit
17	authority had been using failed to measure
18	improvement of hearing through the use of
19	hearing aids; job opportunities were being
20	lost as a result of that unfair test and are
21	now opened because of the results of the
22	office.

Chicago O'Hare had only one operable TDD in

After receiving a complaint that

23

1	the entire airport and that it was available
2	only during limited hours, the division
3	entered negotiations with O'Hare officials,
4	and as a result there are now TDDs in all
5	three O'Hare terminals; and they are available
6	at all hours.
7	When a major health insurer
8	refused to provide coverage for an implant for
9	a severely hearing-impaired child, the
10	division became involved, and the matter was
11	resolved; the operation was performed, and the
12	insurance companies are now doing this on a
13	routine basis.
14	In addition to the litigation
15	efforts and the successes we have had in that
16	area, the Attorney General and the Disabled
17	Persons Advocacy Division have actively
18	supported legislation benefiting the
19	hearing-impaired.
20	During the past three years we
21	have seen important new bills become law in
22	Illinois: The Hearing Aid Consumer Protection
23	Act created standards and enforcement

24 procedures to prevent fraud and

1	misinterpretation in the marketing and sale of
2	hearing aids.
3	Prior to the enactment of this
4	law, Illinois was a mecca for unscrupulous
5	hearing aid salespeople.
6	Bill 984 created a public
7	requiring transporation facilities such was
8	train stations, bus terminals, and airports to
9	provide TDDs.
10	The Universal Telephone Service
11	Protection Law of 1985 requires in part the
12	provision of TDDs at no cost to eligible deaf
13	and severely hearing-impaired consumers.
14	The Attorney General's Office
15	concerned about the Illinois Commerce
16	Commission proposed rule to implement this new
17	law is working with hearing-impaired
18	organizations statewide to propose changes in
19	the rule.
20	We believe that there should be a
21	provision for an advisory council representing
22	the hearing-impaired community to work with
23	the ICC staff and the state's telephone
24	companies in developing the TDD program.

1	We also believe there should be a
2	voucher system so that hearing-impaired
3	consumers can select the equipment they need
4	and rather than having to accept what is
5	provided under a central distribution system.
6	Finally, we want to assure that
7	responsibility for the program will rest with
8	local carriers and that existing
9	administrative and procedures are
10	utilized.
11	The Illinois Environmental
12	Barriers Act, which was drafted by improves
13	and expands upon the State's previous
14	accessibility standards.
15	It applies to both publically and
16	privately owned buildings used by the public.
17	Specific features of the new law
18	which relate to the hearing-impaired include,
19	one, a requirement of specialized emergency
20	signals such as visual smoke alarms and
21	elevator signals; two, a requirement that
22	public telephones must provide amplification
23	devices for the hearing-impaired; three, a
24	requirement that new multi-story housing must

1	be adaptable to the needs of environmentally-
2	limited residents.
3	The Attorney General's Disabled
4	Division has made extensive and detailed
5	recommendations to the Illinois Capital
6	Development Board, the State Agency which is
7	developing standards to implement this
8	requirement.
9	Attorney General Hartigan, as a
10	State's chief legal enforcement officer, will
11	have the responsibility of enforcing Illinois
12	new Environment Barrier Act.
13	Considering the Attorney
14	General's record in advocating for the
15	disabled for the past three years, it can
16	fairly be said that the responsibility for
17	enforcement is in good hands.
18	He actively supported a city of
19	Chicago ordinance requiring visual smoke
20	alarms in one out of every 50 hotels rooms in
21	all Chicago hotels; under the ordinance which
22	was passed by the Chicago City Council last
23	week and which becomes effective January 1 of
24	next year, hearing-impaired hotel guests will

1	be able to request rooms with visual alarms.
2	Attorney General Hartigan has
3	already testified in support of a pending
4	Chicago ordinance that would require buildings
5	with elevators to maintain a registry of
6	environmentally-limited persons on the
7	premises, whether living there, working, or
8	visiting.
9	This means that in an emergency,
10	fire department personnel would know the
11	locations of persons requiring special
12	assistance.
13	The ordinance would also require
14	preparation and posting of an evacuation plan,
15	which provides for the needs of
16	environmentally-limited persons.
17	Also actively supporting another
18	bill which has been introduced in the Illinois
19	Senate and which will be voted on next fall,
20	this bill would create the Illinois Insurance
21	Act.
22	It would provide an insurance
23	pool for persons who have been considered high
24	risk by the insurance industry and who have,

1	therefore, been denied coverage or offered
2	only limited coverage or very costly coverage.
3	As many of you here know persons
4	with disabilities can experience extreme
5	difficulties in obtaining adequate and
6	affordable coverage.
7	The passage of the Comprehensive
8	Health Insurance Program Act would be a long
9	overdue solution to this problem.
10	Everything Attorney General
11	Hartigan and the Disabled Persons Advocacy
12	Division has done and I have discussed only
13	the highlights of the past three years
14	stems from a Committee of the citizens of this
15	state with disabilities, including hearing
16	impairment, are entitled to work, to travel,
17	and to communicate just like anybody else.
18	They are entitled to feel safe;
19	and they are entitled to enjoy the public
20	facilities and other offers in their
21	communities like anyone else.
22	Over the past three years the
23	office has achieved meaningful gains for the
24	hearing-impaired and for citizens with other

1	disability.
2	In every case what has been
3	gained is no more than what other citizens
4	already have.
5	As a division builds on what it
6	has accomplished so far, equal opportunity
7	will remain the central motivating factor.
8	Working with the two advisory
9	councils to develop new programs and
10	legislation, making certain that laws and
11	programs we already have are working,
12	responding to complaints and requests for help
13	from individual citizens, the division will
14	continue making Illinois a place where
15	citizens with disabilities have vigorous
16	representation in government and equal
17	opportunity under the law.
18	Thank you very much for letting
19	me be here today.
20	MR. SCHWARTZBERG: Let's turn now to
21	the members of the Committee.
22	Questions addressed to this
23	panel?

24

I do not have questions addressed

1	to this panel so we will turn then directly
2	to Doctor Breunig, Doctor Latham Breunig.
3	It might be possible to begin to
4	introduce Doctor Latham Breunig's biography
5	which begins with the time when he first
6	became an Eagle Scout.
7	I do not propose to do that
8	today.
9	Doctor Breunig is retired from
10	Eli Lilly and Company.
11	At age 3 Doctor Breunig had a 25
12	percent hearing loss, and by age 7 his loss of
13	hearing was 95 percent.
14	He attended Wabash College,
15	Indiana, earned his Ph.D. in Chemistry from
16	John Hopkins University, and also studied at
17	Purdue University, Indiana, in the field of
18	statistics and quality control.
19	Among the many organizations in
20	which he has been active, the President's
21	Committee on Employment of the Handicapped,
22	the American Statistical Association, the
23	Clarke School for the Deaf, the Alexander
24	Graham Bell Assocation for the Deaf, where he

1 served on the Board of Directors for 24 years, 2 two of them as president. In 1968 he founded 3 Telecommunications for the Deaf and was Chief 4 5 Executive Officer of Telecommunications for 6 the Deaf until the year 1978. 7 He has been particularly active 8 in the development of oral interpreters for 9 deaf people. 10 Even to list the special awards 11 presented to Doctor Breunig would take more 12 time than we have, but they include those by 13 the President's Committee on Employment of the 14 Handicapped Commendation by President Johnson 15 in 1967, Service to Mankind in 1968, and the 16 Governor Branigin's Award in 1969. 17 The list goes on and on, it 18 certainly -- the presidents who have chosen to 19 turn to Doctor Breunig include the present 20 incumbent who appointed him to the National 21 Council on the Handicapped in 1981 and 22 reappointed him in the year 1983. It is my pleasure to introduce 23

for our wrapup Doctor H. Latham Breunig.

1	DR. BREUNIG: Thank you very much, Mr.
2	Chairman, and members of the Illinois
3	Committee on the U.S. Commission of Civil
4	Rights.
5	It is a honor to be here to
6	attempt to put closure on this important
7	forum, with so many distinguished advocates,
8	that we have a very important responsibility
9	to provide for the hearing-impaired citizens
10	in the State of Illinois.
11	You have heard the right to
12	knowledge and education and employment and
13	focused on the state of the art.
14	My charge is to look to the
15	future of the hearing-impaired people, and
16	that it reminds me of my fraternity initiation
17	based upon a book of Issiah, Terrors of the
18	Night, the recently published book
19	perspectives in which was edited by David
20	gives us an excellent overview of the field,
21	although my personal opinion is that my name
22	was misspelled in the book.
23	Time permits only excerpts from
24	the chapters.

1	In his chapter on audiology
2	states for most of the children that we work
3	with amplified sound can provide the brain for
4	the sum of the raw material it needs to
5	naturally evolve in order to enhance a
6	communication system, whatever the degree of
7	residual hearing.
8	It is a biological birth right
9	that should be used and depended upon whenever
10	possible.
11	Rationalization of the
12	methodology dispute so that it does not
13	dominate the field is one of the charges of
14	the author; it is indeed a civil right of
15	every hearing-impaired child to have the
16	fullest opportunity to develop inherent verbal
17	communication skills.
18	It is unfortunate that an
19	otherwise excellent publication, Legal Rights
20	For Hearing-impaired People, which is
21	published by Gallaudet College propogates a
22	myth about lip and speech reading.
23	President Johnson, the greatest
24	enemy of truth and very often said and that

1	was a contrived lie continued the myth.
2	The myth persisted and is
3	unrealistic.
4	The perfection of the speech is
5	designed is secondary to the act to
6	generate meaningful communication, but it is
7	about time when we say we have to stop
8	worrying about what hearing-impaired children
9	can't do or have not learned about English;
10	and we need to pay attention to what they know
11	about communication.
12	And we know now for 400 years the
13	teaching of speech has been pretty much the
14	same.
15	There have been breakthroughs
16	over the last 20 years.
17	And the stage is set for
18	progress.
19	My crystal ball tells me that the
20	spoken language will always be the most
21	respective key for the entry of
22	hearing-impaired people into the educational,
23	economic, cultural and social affairs of the

world about them; whether their speech be --

1	or smooth, their capability to use it is still
2	the and the chapter in the book on academic
3	placement has raised some difficulties with
4	this concept of the least restrictive
5	environment.
6	The National Counsil on the
7	Handicapped has developed a book called
8	Independence that the United States Department
9	of Education promulgate for the application of
10	the least restrictive environment requirement
11	should clarify that the primary determinant of
12	which educational setting the least
13	restrictive is the educational appropriateness
14	of the program.
15	Now, after discussing the
16	characteristics of the mainstreaming programs
17	of the deaf child, it concluded that the least
18	restrictive environment for deafness is one
19	that is populated by people who are especially
20	trained to understand deafness; and it
21	concludes we must keep in mind that the
22	educational placement should not be
23	competitive in that the final act should be
2.4	designed and evaluated constantly in the light

1	of its effects on the whole life of the child.
2	I was asked to look at trends in
3	the general population, and they will continue
4	to exist.
5	There will continue to exist the
6	stereotype of deaf persons; that more deaf
7	youngsters having acquired spoken language and
8	will grow up and interact with the world
9	around them.
10	And I can tell you that in 1975
11	my wife and I moved to Washington; as we have
12	gone about the shops and the stores in the
13	city and we talked with the people, oh, I
14	didn't know deaf people could talk in
15	Washington D.C., so there is a lot that needs
16	to be done.
17	For services and sources, it
18	seems to me these will be enhanced if Congress
19	and the Executive Branch of Government could
20	implement the 45 recommendations of the
21	National Counsil for the Handicapped that are
22	spelled out in the book Independence.
23	Incidentally, this book as a
24	reference to the Civil Title 2 in the Civil

1	Rights Act of 1964 entitled 8 of the Civil
2	Rights Act of 1968 so there is a lot of
3	meat in that book that this Committee could
4	look at with respect to government and
5	private the role of the Federal Government,
6	what may well diminish through the monetary
7	and social constraints well those of state and
8	local governments become important, and
9	instead of running to Washington
10	hearing-impaired people need to cultivate
11	their local resources and their participation
12	in independent groups.
13	Hearing-impaired people need to
14	reach out and become active in their community
15	life, and this way they will be able to make
16	contact with people in local governments and
17	the projects within industry.
18	Needs to be directed to
19	diminishing the stereotypes on the whole; it
20	would seem that the signs of promise are
21	there.
22	This morning I believe Doctor
23	Lucas asked one of the people if there were
24	any laws requiring speech; there is one basic

1	right guaranteed by Article l of the Bill of
2	Rights of the United States of America, the
3	freedom of speech.
4	MR. SCHWARTZBERG: I would like to
5	thank Doctor Breunig.
6	I would like to, at this point,
7	point out some things which I pointed out
8	earlier but which I would like to repeat at
9	this time.
10	We are open to additional
11	materials on the subject that we considered by
12	this forum today.
13	Written statements may be
14	submitted for consideration of the Committee
15	and for inclusion in the record, et cetera.
16	The materials can be given to the
17	staff person at the registration desk at the
18	outside of this room.
19	They can also be mailed before
20	the 20th of July, 1986, to the United States
21	Commission on Civil Rights, 230 South
22	Dearborn, Room 3280, Chicago, Illinois, 60604.
23	This public forum of the Illinois
24	Advisory Committee to the United States

1	Commission on Civil Rights has focused on the
2	civil rights of hearing-impaired persons.
3	Under Federal Law this Advisory
4	Committee is to report to the U.S. Commission
5	on the information gathered here today, so
6	that after considering this information, the
7	Commission may make recommendations to the
8	President and to the Congress.
9	We again thank the organizations
10	in cooperation with which we have held this
11	forum today: The Section on Individual Rights
12	and Responsibilities of the American Bar
13	Association, the Illinois Department of Human
14	Rights, and the Alexander G. Bell Association
15	for the Deaf.
16	We also offer our most cordial
17	thanks to all of the presentors, the experts
18	that shared information with us today.
19	We face the extraordinary
20	experience of a great many more people wishing
21	to give testimony at this forum than there was
22	possibly time within which to do it.
23	We regret that there are not more
24	hours in the day, but we stand ready to hear

1	additional information and to learn more from
2	anyone who wishes to enlighten us on the
3	problems of the deaf, and on the ways in which
4	this society may deal with those problems, not
5	only simply as a matter of right against the
6	state but as a matter of legislation to deal
7	with these problems in that more general
8	context that we have often labeled civil
9	rights.
10	And so this meeting stands
11	adjourned.
12	Thank you very much.
13	
14	
15	
16	
17	
18	
19	
20	
21	
22	
23	
24	

I	STATE OF ILLINOIS )
2	COUNTY OF C O O K )
3	
4	MARYJO JORGENSEN-BARNES, being first
5	duly sworn, says that she is a court reporter
6	doing business in the City of Chicago, and
7	that she reported in shorthand the proceedings
8	had at the hearing of said cause, and the
9	foregoing is a true and correct transcript of
10	her shorthand notes, so taken as aforesaid.
11	
12	
13	Many formenson-Bann
14	
15	SUBSCRIBED AND SWORN TO before me this /300 day
16	of (3 118) 1 986 .
17	Notary Public
18 19	Notary Public
20	
21	
22	
23	