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CALIFORNIA LEGISLATURE

Senate Subcommittee on the Disabled Milton Marks, Chairman

HEARING ON MENTAL HEALTH AND THE DISABLED

STATE CAPITOL ROOM 113 SACRAMENTO, CALIFORNIA

THURSDAY, MAY 16, 1985 1:00 P.M.



HEARING Haylo BEFORE THE SENATE SUBCOMMITTEE ON THE DISABLED Hearing in re: LAW LIBRARY 5 MENTAL HEALTH AND THE GOLDEN GATE UNIVERSITY 6 DISABLED. 7 8 9 10 STATE CAPITOL 11 ROOM 113 12 SACRAMENTO, CALIFORNIA 13 14 15 16 17 18 THURSDAY, MAY 16, 1985 19 1:00 P.M. 20 21 22 23 24 25 Nadine J. Parks 26 Shorthand Reporter 27 28

INDEX	Page
Proceedings	1
Opening Remarks by Senator Milton Marks	1
Kathryn Ross, State Department of Education	2
Moss Nader, Ph.D., State Department of Mental Health	5
Collis Kimbrough, United Cerebral Palsy Association of San Francisco	16
Dexter Lane, United Cerebral Palsy Association of San Francisco	16
Ladonnis Elston, City and County of San Francisco Community Mental Health	20
Dr. Ken Brynjolfsson, Far Northern Regional Center, Redding	26, 2
Dennis Ferrell, Far Northern Regional Center, Redding	26
Wendall Fingar, State Department of Rehabilitation	38
Art Segal	46
Elizabeth Swain	54
Sharon George-Garrity	56
Adjournment	58
Reporter's Certificate	59

PROCEEDINGS

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CHAIRMAN MARKS: We're going to bring this meeting to order. Let me tell you that this is a meeting of the Subcommittee on the Disabled. I'm Senator Milton Marks, and this is Dorothy Epstein, who has been the coordinator of this committee for a number of years.

And I have a short statement I'd like to read at this time; then, we'll hear our witnesses. The other members of the subcommittee are involved, unfortunately, in committee meetings of their own. So, we'll record this testimony and they will see it.

The care of the mentally ill has been recognized as a responsibility of State government. The Legislature encouraged the creation of a system of treatment options so that mentally ill persons could obtain the proper services within their own community. The Short-Doyle Act provided a fiscal mechanism to implement this policy, and the Lanterman/Petris/Short Act, passed in 1968, has always been recognized as one of the most important pieces of mental health legislation.

This is our past, and I acknowledge we've come a long way in the last 17 years, but not far enough, because some of the problems that existed then exist now.

In the deaf community, for instance, there are not enough in-patient clinical programs and not enough therapists who understand the deaf consumer, and some additional problems exist today -- such as the large numbers of street people and

refugees needing mental health services.

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The reason that I am holding this hearing is in my discussions with the disabled community, I am constantly reminded that there are a number of critical needs not being addressed such as the quality of mental health services, patients' rights, adequate treatment for children, stigma, and more self-help services, just to mention a few.

Over the years, the changing state and federal regulations, economic distress, and funding reductions and lack of statewide standards are some of the issues that have combined to threaten the mental health system.

Today, advocates and taxpayers, who pay millions to help finance the public mental health services, are demanding more accountability, efficiency, and effectiveness in mental health services than ever before. I hope that today's hearing will address these concerns of the disabled community so that we may move on to provide the services needed.

We have a number of witnesses to testify on the subject. Let me first call on Ferd Shaw. Ferd Shaw here? I won't call him. Kathryn Ross.

MS. ROSS: My name's Kathryn Ross and I'm the administrator for special education in the State Department of Education.

CHAIRMAN MARKS: You saw the letter that I sent?

MS. ROSS: (Nodding her head). Okay. I'd like to attempt to answer the questions you asked us to address, the first one being, "Is the Department of Mental Health cooperating with the Department of Education in providing

mental health services for residential schools?"

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I surveyed our State residential schools, the two schools for the deaf -- the one in Riverside and the one in Fremont -- and discussed with their superintendents what kind of services they were getting from State Mental Health.

At this time, they indicated in the school for the deaf in Riverside that they are getting some services from St. John's Hospital. And I understand that Mental Health directly contracts for those services for deaf emotionally disturbed there.

They did indicate, though, that because of changing population -- this is true for both schools for the deaf -- they are getting many more multiple-handicapped children and have much more need for intensive services. The school for the deaf in Fremont is contracting with Dr. William Evans and he's from the University of California Center for Deafness.

And, again, they have a grant from the Department of Mental Health. And Dr. Klopping indicated also that they have about 50 children now in need of intensive services at the school.

I know they're also trying BCP to try to get some services directly, State support, too, but they aren't getting that kind of services. They are not getting very many services from Alameda County Mental Health. They have contacted them; have been working with them. Alameda County has -- well, as we know, limited amount of funds, and also, the youngsters at the school for the deaf are from many other counties, not just

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Alameda.

They do serve them, but they are not a priority. They do not come ahead of any other deaf children at their center.

The school for the blind has not used nor have they pursued giving services for mental health, and nor have the three diagnostic centers. They have not pursued it. It might have been a good idea, but that has not been in their history.

A number of private residential schools have contracts with local mental health agencies, but that's individual private schools contracting for individual children with departments of mental health.

Your other question was on the impact of
Assembly Bill 3632. Right now we don't know the fiscal
impact or really the programmatic. We're gathering data and
the data's coming in. We are trying to find out who is
actually serving which youngsters' schools could be more
properly served by Mental Health. That data is not available
yet.

CHAIRMAN MARKS: When was the bill passed?

MS. ROSS: The bill was passed last year. It was supposed to be effective -- I don't know the passage date.

We have been working with -- we're supposed to have a report into the Legislature on April 15th. We don't have that in yet on the cost. To Finance. We don't have that in yet. The cost data collecting -- the forms, I guess, didn't give us the kind of information that we really needed.

I understand that Mental Health has resubmitted

too, because there's been a lot of misunderstanding about what was going to happen. The implementation date of 3632 is July 1. It was September 30, 1984.

The -- what we have been doing with the Department of Mental Health very recently, working very diligently within our staff with members of their staff to complete draft regulations. Those regulations are almost completed. And we're looking forward to what we hope will be joint hearings on the regulations. So we're both -- we're looking at the needs together instead of separate agencies, so we can coordinate better.

The big problem appears to be adequate funding.

As you probably are aware, there's still a stalemate, I think, on 105, on the follow-up bill. That's pretty much where we're at.

CHAIRMAN MARKS: Thank you very much.

MS. EPSTEIN: Could I ask you a question?

CHAIRMAN MARKS: Put the microphone up.

MS. EPSTEIN: In terms of children receiving services, are there enough therapists now?

MS. ROSS: I don't think so.

CHAIRMAN MARKS: Okay. Thank you very much. We appreciate your being here.

Is Ferd Shaw here?

Moss Nader.

MR. NADER: Good afternoon. I'm Moss Nader from the State Department of Mental Health. I will attempt to respond to the questions that you have posed for us.

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And if there are any other questions, I'll be glad to address them.

You'd asked that -- whether the Department of Mental Health is putting its financial resources -- where we are putting our financial resources for the mentally ill population other than in the Short-Doyle county block grant and hospital bailout.

As you may be aware, the Department of Mental Health has no other financial resources besides the funds that we are spending for our community programs and the cost of the State psychiatric hospital operations.

At the present time, we spend roughly \$438 million for the local assistance, the funds that goes to counties. We have about \$4.2 million that goes to the judicially committed county programs for programs that deal with mentally ill offenders.

Then, we -- in the State hospitals, we have the Atascadero State Hospital and Patton State and other programs that deal with judicial offenders, the Penal Code patients -we spend about \$98 million for the operation of those hospitals and other State hospital programs for psychiatric Lanterman/Petris/Short Act type of patient, we spend almost \$46 million.

In addition to that in recent years we have managed to obtain \$15.7 million from the federal government which we block grant to the community mental health centers and agencies. But in a nutshell, most of the funds, except for a very small percentage that is for the administrative

are basically funnelled to the local government agencies for the administration of mental health programs.

The second question was: Is there a Department

of Mental Health priority for doubly handicapped such as

mentally ill/hearing impaired or visually impaired, orthoped-

ically handicapped, developmentally disabled?

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Recently enacted AB 2381 of last year mandates that in developing and providing the services of a county — a county mental health program, the county shall consider and make provision for all of the following priority populations. Priority populations are then enumerated as chronically mentally ill, mentally disturbed children and adolescents, mentally ill elderly, mentally ill jail inmates, MDO's, and underserved populations.

The State Department of Mental Health has interpreted the physically and developmental handicapped to be among the underserved population.

We have communicated that to the counties. And we are doing our utmost to ensure that counties consider this group as a priority population; that is, the duly diagnosed mentally ill, the clients who have physical handicaps, as well as the hearing impaired. We have articulated that and indicated that to all the counties.

But, basically, at this moment, the identification of priorities has to come from the local county mental health program. Usually, each county through its own constituency group and mental health advisory board identify their own local needs and priorities. They submit it to us and then, of

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course, the State Department through some formal mechanism approves the county plan.

In general, the State does not identify specific priorities for specialized types of clientele. These priority groups are usually identified on a local basis based on local needs. That's the way the current laws are and we're just complying with them.

The next question was: Because the Department of Mental Health is block-granting the dollars to the counties on equity formula (sic), what standards of excellence does the Department of Mental Health expect from these counties?

If by standards of excellence you mean standards of excellence as far as programs are concerned, we have a number of formal monitoring and reviewing procedures. A special group of headquarter's staff from the Department of Mental Health on a regular basis conduct monitoring of local programs. The review protocol is designed to ensure some degree of excellence.

Furthermore, the Department has got a formal quality assurance mechanism. Quality assurance ensures that some degree of quality are adhered to in all of the 24-hour psychiatric facilities. We now have medication monitoring procedures. We have utilization monitoring and peer review for the in-service programs within the 24-hour psychiatric facilities.

As far as the individual -- delivery of individual services to a given clientele, of course, the way the mechanism is set, most of the clinicians in the field are licensed,

bona fide licensed professionals. And assurances of quality of their practice is the function of the -- another agency, the Medical Assurance Quality (sic) or the licensing agencies.

But in general, the Department does have ongoing reviewing, monitoring procedures and also quality assurance.

The last question, you have asked whether the Department of Mental Health plans to give categorical aid direction to the counties for multiple-handicapped persons.

If by categorical aid direction you mean by dollars earmarked for specific category, the answer is no.

This would be against the recently enacted mental health laws. We have no specific categorical funding per se for any specialized type of target group for specialized services. However, the Department will continue providing technical assistance to the counties for the provision of specialized services. But technical assistance that we provide is merely technical assistance.

The current trend is to rely on the local programs, local governments to identify and administer their own programs. The State has taken the lead in providing technical assistance.

At the present time we have one regional program that is funded directly from the State Department of Mental Health; that is, the small program we have at St. John's Hospital in Southern California for the hearing impaired.

It appears that that particular program may at some point be incorporated into the Los Angeles County overall mental health program.

CHAIRMAN MARKS: I've heard your testimony and I appreciated it. There are a lot of people who come to me in the constituency who are very concerned about how people are being handled. Can you give me the reason why that happens, that a lot of people are looked upon as developmentally disabled before there's concern as to whether or not they have mental problems? A lot of problems that are still going on.

It seems to me that if your Department is doing what it's supposed to be doing, I wouldn't have all those questions.

DR. NADER: We are, at the present time, working on some of these issues, Senator. Currently, there is a task force that is -- has identified some of the problem of the delivery of mental health services to the dually diagnosed. The task force is going to submit its recommendation for our Department. We'll be looking at those and working with them.

CHAIRMAN MARKS: Who composes the task force?

DR. NADER: The task force is -- the Association of California Regional Centers have created this task force. There are a number of members from mental health and developmental disability as well as constituency groups participating in that task force.

We do recognize that there are some problems with this particular target group. These are the people who are developmentally disabled and for some reason in the community they are developing emotional problems. And because of the nature of their problems, they are falling in the cracks. And

we are aware of it and in the process of developing possibly 7 some sort of a pilot project to see how we can address this 2 problem. 3 CHAIRMAN MARKS: Well, let me read you a letter. DR. NADER: Okay. CHAIPMAN MARKS: From the Marin County 6 Developmental Disabilities Council. 7

DR. NADER: Okay.

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CHAIRMAN MARKS: It's addressed to me. (Reading) The materialistic needs of the developmentally disabled, food, shelter, clothing, et cetera, are the needs which are easy to respond to.

It's the nonmaterialistic needs, the mental health problems, which are difficult to see and respond to. To know that you are different from other people creates all sorts of emotional and behavioral problems. these problems that are not being cared for at this time. The programs which serve the mental health population will not accept persons who are developmentally disabled. They say they do not fit into the system.

The few programs that will accept the disabled have a very long waiting list. The present mental health service and delivery system is not sufficient nor is it capable of meeting the needs of the developmentally disabled. More funds and more facilities are urgently needed.

The people are very much concerned. And if I may say so, with all due respect, a task force is not an answer and is oftentimes a way to delay a solution.

DR. NADER: We would like to embark on some action plans to remedy the problems of the dually diagnosed.

CHAIRMAN MARKS: Can you present anything to the Budget Committee?

DR. NADER: Not as of this moment.

CHAIRMAN MARKS: It's a little late. The budget meeting just ended today.

DR. NADER: For -- but we are planning to submit a pilot project to develop regionalized programs.

CHAIRMAN MARKS: When do you plan to submit that?

DR. NADER: This forthcoming summer.

CHAIRMAN MARKS: To whom?

DR. NADER: It will be submitted for next year's Governor's budget hopefully for the fiscal year 86-87.

CHAIRMAN MARKS: Go ahead.

MS. EPSTEIN: When you start out not knowing anything about the whole mental health picture, and you start having someone come in and you're saying to them -- who are all the groups in the mental health picture -- you start out as we started out doing. You have someone put this on a wall for you, it just starts at the top and goes all the way down to the floor (indicating). And then you go through and you say, to whom is each group responsible? Well, there are lots of different State groups that operate throughout the State, part of which we found there is no real responsibility where people are required to report back to someone. So that what you see are layers after layers of groups each doing their own thing. And what we're particularly concerned with is where

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does it get down to the DD person or the deaf person? are they affected? I mean, they're all great groups. And they all meet. But where does it all funnel down and into what?

DR. NADER: I'm afraid that was the target group that was identified as underserved. Unfortunately they've been underserved. We recognize that and we're trying to remedy that.

But, again, the identification of a target group, as the laws are today, ought to be made at the local level through the Mental Health Advisory Board. And groups need to be vocal in those places and make sure that --

> CHAIRMAN MARKS: Have you seen this? (Thereupon a schematic was shown to the assemblage and Dr. Nader.)

MS. EPSTEIN: This is the person who deals with DD people (indicating) from here to here. That's the local level.

> DR. NADER: Right.

CHAIRMAN MARKS: I'm not critical of you personally, but it seems to me that the State of California should do more in this area. And this is something I'm just not bringing to your attention for the first time. It's been brought to your attention I'm sure a number of occasions. I'm reminded of a hearing in 1980 so I presume it had your attention at least at that time. I just can't see how this can go on and on and on without any determination of what's going to be done. I'm really sort of disturbed that nothing

was done in this budget session, nothing.

I mean, you're evidently going to wait till the next budget session.

DR. NADER: Well, Senator, we have this particular year -- the Governor has allocated extra funds for mental health for next fiscal year. The existing fiscal year we have extra funds. We are encouraging the counties to pay attention to these target groups. But, as you know, our hands are tied because by law local county mental health directors have the complete jurisdiction over their administration of their programs. It's a double-bind that we are in. We would like to take leadership and tell local governments what to do, but on the other hand, we want to stay away from direct intervention.

CHAIRMAN MARKS: Why don't you write a letter telling me what your problem is. I'd like to know your problem.

DR. NADER: Okay.

CHAIRMAN MARKS: Why don't you write me a letter.

DR. NADER: Okay.

CHAIRMAN MARKS: Let me see what your problem is.

CHAIRMAN MARKS: Well, I'd like you to be specific.

DR. NADER: We'll send you a letter, Senator.

CHAIRMAN MARKS: And what you are doing to eradicate those problems.

DR. NADER: In regard to the dually diagnosed and developmentally disabled, we have to be specific because --

Tell me what your problem is and what you're doing.

DR. NADER: Okay.

MS. EPSTEIN: In the past couple of years there have been some very large community meetings. One in particular was put together by Independent Living in San Francisco. It was enormous.

At that time the theme that we seemed to hear out of that talked about types of medication, self-help groups, stigma.

My question is when the Department is making its decisions about how to help these people, is it talking to people from any of these consumer groups as well as the government bodies?

DR. NADER: We are indeed. We now have a representative of these consumer groups in a number of our committees, action, working committees. Things are changing tremendously. The consumer groups are becoming very vocal through a program that we have had. We've managed to bring in an awful bt of consumer groups to our planning, organizational committees, a number of our action committees. I just don't want to mention the task force, but they are expressing their voices.

CHAIRMAN MARKS: One of the reasons we formed the subcommittee on the disabled was to try to get a group together that would hopefully be the ones that the disabled community in every aspect of society would be able to turn to. And when I tell you I'm not satisfied, I'm not satisfied. I'd like an expression from you as to what you think the problem is and what you're doing to resolve this.

16 DR. NADER: Okay, Senator. I appreciate your -We'll work on it and I will be sending you a letter. concern. 2 CHAIRMAN MARKS: Thank you. 3 DR. NADER: Thank you. 4 CHAIRMAN MARKS: Collis Kimbrough here? Dexter 5 Lane? Do you want to testify together or separately? 6 Why don't you both come up? 7 MR. KIMBROUGH: My name is Collis Kimbrough. 8 coordinator of program development for United Cerebral Palsy 9 Association of San Francisco. I only have a few brief 10 comments to make. 11 Developmentally disabled people in California, 12 like everywhere else in the world, present a cross-section 13 of our culture in our society. They run the gamut of people 14 racially, ethnically, vocationally, et cetera. 15 As California, like other states, moves towards 16 serving these people in the community, increasing problems 17 such as mental health needs are going to be exposed, expressed, 18 and hopefully dealt with. 19 I don't have any prepared testimony for you today. 20 Dexter Lane does. 21 CHAIRMAN MARKS: Okay. 22 MR. LANE: I'm going to read from a letter that I 23 prepared and I also would like to follow it up with some other 24 comments. 25 A 1981 study by the State Department of Mental

one-third of the developmentally disabled were . . .

Health among California Regional Centers concluded that ". . .

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handicapped to a significant degree by disturbances in thought

emotion, or behavior."

If this percentage holds true for Golden Gate Regional Center clients, well over 1,000 persons in its caseload have significant mental health needs.

In San Francisco, one person in the Department of Health is currently responsible for coordinating services to an estimated 550 persons with mental health and developmental disability dual diagnoses. One person, however well qualified, is insufficient to provide comprehensive coordination of services for these clients. Simply stated, the vast majority of these persons are underserved for their mental health needs if they are being served at all.

Compounding the inadequate level of professional coordination of services for these clients is the under-developed state of counseling methods for the dually diagnosed.

Again, from the 1981 study, quote, ". . .Research into the causes, incidence, prevalence, and treatment of mental illness in those without developmental disabilities has produced much information and has contributed to the development of widely distributed and sophisticated programs of restorative care. More modest gains have been made in understanding the same issues for the developmentally handicapped; consequently, treatment resources are in a primitive state compared with what is available for others."

Treatment for psychiatric needs of dually diagnosed clients currently available through the Golden Gate Regional

Center case management coordination are behavioral evaluations and treatment, private psychotherapy funded by Medi-Cal, and a behavioral treatment program at the recreational center for the handicapped.

The behavioral evaluations are reactive rather than preventive, and are generally provided in response to severe behaviors such as violence or destructive acting out.

The recreation center program has 28 clients and a waiting list. For reasons which will be described shortly, a private psychiatrist, funded by Medi-Cal, is the treatment of choice usually only for the very highest functioning clients.

Roughly speaking, it is usually the dually diagnosed person who most severely disrupts the lives of others or the very highest functioning who receive specific treatment for their mental health needs.

Those clients whose major diagnosis the Regional Center has determined is primarily psychiatric in nature have been shuffled back and forth between the developmental disability system and the mental health system, neither system wanting to assume primary responsibility for treatment.

Fortunately, a very recent agreement between the Regional Center and the De partment of Health has been reached which clarifies responsibility for the coordination of services to these clients.

For most mental health service providers, clients with developmental disabilities are not as attractive to work with as are clients with a single mental health diagnosis.

and potentially amenable to insight as are clients without retardation. Few mental health service providers are aware of issues raised by the developmental disability diagnosis. The client with the dual diagnosis is simply short-changed between two separate service provision systems, one for mental health and one for developmental disabilities, neither of which has sufficient expertise to work in the other's domain.

As the social worker for United Cerebral Palsy of San Francisco, I see a very small percentage of the many dually diagnosed clients and their families who fall between the large cracks in the system. The population I see is even further narrowed by the requirement that my clients have a physical disability, generally cerebral palsy.

For some clients I have devised behavioral treatment programs, some I work with in individual psychodynamic-based counseling and some in family therapy.

In two cases I do not even see the identified dually diagnosed client, but provide supportive counseling to the single mothers of the clients with the goal of stabilizing their chaotic family environments.

My education and seven years of hospital and halfway house work experience did not address the developmental disability issues which are important to my current work.

I have had to integrate information from two largely separate service methodologies to provide treatment to my clients.

Clients dually diagnosed with developmental and psychiatric disabilities need more flexible and informed responses to address their treatment needs than is currently available.

The developmental disabilities and the mental health service systems need much greater coordination and much greater information sharing before the majority of dually diagnosed persons will be adequately served.

CHAIRMAN MARKS: Thank you very much. We appreciate your both being here. Thank you.

MR. LANE: I have copies.

CHAIRMAN MARKS: Would you, please.

Ladonnis Elston.

MS. ELSTON: Okay. My name is Ladonnis Elston.

I am the coordinator for services to people who are disabled for the San Francisco Community Health Services.

My report is going to be very brief today, but it has to do primarily with some of the statements that was previously made by the speakers, such as lack of continuum of services for the disabled people who have recently been discharged from State Hospitals without developing programs in the community prior to discharging them in the community. So, we find that we are experiencing a lot of disabled people coming back to the community without having adequate programs.

A lot of times we find that we are having to keep people in acute places, such as hospitals, San Francisco General, Mount Zion Hospitals, longer than necessary because

there are not places to put them such as halfway houses or 1 programs where people who are developmentally disabled cannot 2 go into halfway houses with people who are higher functioning 3 and those who are psychiatrically disabled. 4 They often need some kind of specialized treatment, 5 so we find that we're spending a large sum of money on people 6 in acute services where, if they were able to have some kind 7 of community programs, it would better meet the needs of the 8 client in addition to less cost for the community. 9 CHAIRMAN MARKS: Let me ask you a question about 10 11 spending money. Is all the money coming from the State or does the city --12 MS. ELSTON: The city puts in money. 13 CHAIRMAN MARKS: What percentage would you say 14 comes from the city approximately? 15 MS. ELSTON: Oh, I really -- about 20 percent from 16 the city and county. I mean --17 CHAIRMAN MARKS: And some comes from the federal 18 government? 19 MS. ELSTON: Some come from the federal government 20 and some comes from the State. We have ad valorum, which is 21 money that comes from the city and county that is not State 22 money. 23

Thank you. CHAIRMAN MARKS:

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MS. ELSTON: This is approximate. I don't have the --

> CHAIRMAN MARKS: I won't hold you to it. MS. ELSTON: Okay. Don't hold me to it.

Okay. So this is what we have been finding. The recent trend is to keep people who are possibly able from — in the community without putting them into the hospitals. What has happened is that there has been a decrease in the number of beds of people in hospitals. However, when the people are in the community, the money that was allocated for the hospitals did not trickle down or follow the client into the community. So we have it very difficult, because the traditional mental health models was not set up to deal with clients who have special needs. The primary focus was on people who could benefit from the traditional psychotherapy.

In San Francisco we have recently done some surveys and found that some people -- some groups of clients can be mainstreamed with the existing mental health programs.

The survey identified the groups having the most difficulty are those clients who are violently mentally ill. They are violent. They have fights and so forth. And people who are substance abusers and they are also mentally ill, and people who are developmentally disabled and they are mentally ill. This is the -- these are the high user groups identified.

of the three groups, the developmentally disabled who are mentally ill continues to be the most difficult group to provide mental health services. With the increase in the number of developmentally disabled who are mentally ill returning to the community from State hospitals without funds for follow-up programs, they continue to go between the systems of the Regional Center system, and the community mental health

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However, we have tried to address this with the State Interagency Agreement. And I think that that is beginning to address some of the issues that is between clients who are Regional Center clients and clients who are Mental Health clients. That seems to be starting to work better than it had in the past.

Funding to the developed community programs should be allocated prior to discharge of persons into the community. The mental health system has been and is focused on the short-term treatment orientation. And many of these clients if they were -- are to benefit from the Medi-Cal system have to be in a short term kind of program. In other words, we have to have them in treatment for six weeks at a maximum or so many visits. And a lot of these clients need to have extended visits. They might need maybe three more visits, but due to the Medi-Cal system and the funding, the way it's set up now, we are not able to just automatically extend their treatment without having quality assurance review, which means we have to go through some people to say, well, this person needs to extend treatment; therefore, would you approve so many days. And this gets to be a real issue a lot of times in terms of how you develop a treatment plan for people who need extensive treatment. And because the mental health system has been focused on the short-term crisis or short-term treatment program, I think one of the areas that needs to be addressed is in terms of disabled people who need mental health services should be looked at with maybe a

different point of view in terms of extension or amount of time that one should be able to receive treatment.

And in conclusion, I think the major efforts that the San Francisco Community Mental Health Services is providing at this time is to do some training with our staff in terms of trying to sensitize our staff about the needs of people who are disabled and develop programs that would be modified to work with disabled people.

Again, I would like to ask that legislation and funding sources should be considered in terms of allowing for some more flexibility with funding for the mental health system or, if that's not possible, create some kind of special funding whereby people who need — disabled people who need specialized treatment can benefit from this kind of services.

I believe that the State policies must demonstrate and reinforce the fact that disabled-mentally ill persons have personal value and that they will be provided with the necessary resources, social, medical, and vocational, to help the disabled develop to their potential.

The other thing that we discovered is that public education promoted on a statewide basis providing accurate information through the media and to the general public about mental health and disability issues will help to improve the services to disabled people.

MS. EPSTEIN: So, what you're saying is that we need to go back to some of the things we did before, which is a lot of public awareness.

25 1 MS. ELSTON: Absolutely. 2 MS. EPSTEIN: I'm not sure in terms of -- the Senator's office, many years ago, was able to do public 3 awareness in terms of physical disabilities by everybody 5 taking on -- in terms of people being allowed to work who had 6 disabilities. MS. ELSTON: Right. 8 MS. EPSTEIN: And in mental health, where people 9 also have disabilities but still need to work to get back 10 into the system, that's still true. 11 MS. ELSTON: This is true. And the thing about 12 people who have emotional problems and disabilities, they have so many stigmas in addition to having the mental illness 13 stigma, they're also --14 15 MS. EPSTEIN: Disabled. 16 MS. ELSTON: -- there are also disabilities. There's 17 also a double disability. 18 MS. EPSTEIN: That's right. 19 MS. ELSTON: That really makes it very difficult 20 for families, our social agencies who are traditionally accustomed to dealing in -- with this only kind of disability, 21 having to deal with a double disability, sometimes more than 22 one, multiple disabilities. 23 MS. EPSTEIN: Would it be wise, then, that perhaps 24 the Senator's office, perhaps if they can take on some kind 25 of project that might make the public more aware? 26

And I think for people who have these kinds of stigmas it is --

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MS. ELSTON: I think that would be very helpful.

26 1 a lot of them it's kind of hidden and it becomes even more 2 difficult to. And I think it would be very helpful if people 3 started to understand or to start to even look at and 4 realize that there are these disabilities. 5 CHAIRMAN MARKS: How many people are in your 6 office? 7 MS. EPSTEIN: Her. 8 MS. ELSTON: There are several offices. 9

have about 40 people working in the community mental health system.

CHAIRMAN MARKS: Thank you very much.

MS. ELSTON: Okay.

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CHAIRMAN MARKS: We appreciate you being here.

Is Dr. Ken Brynjolfsson here?

DR. BRYNJOLFSSON: It's an Icelandic name.

CHAIRMAN MARKS: Well, that was pretty close.

DR. BRYNJOLFSSON: Yes, indeed. I'm going to defer to Mr. Dennis Ferrell who is the project director of our project regarding the dually diagnosed client in the area of Far Northern Regional Center, which is nine counties in the northeastern corner of the State of California.

CHAIRMAN MARKS: Okay.

MR. FERRELL: My name is Dennis Ferrell. staff psychologist for the Far Northern Regional Center serving the developmentally disabled.

Our testimony today represents the Association of Regional Center Agencies and their preliminary report on the mental health needs of the developmentally disabled. And our

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testimony will also represent the activities of our pilot project at Far Northern Regional Center, which is an outgrowth of the mental health task force.

The problems of the mentally -- mental disorders in the mentally retarded and otherwise developmentally disabled population, as other speakers have discussed before, are problems which have only begun to come to light in the last ten to fifteen years.

Services to each of these populations by themselves have been developing, especially in the State of California, at a fairly rapid pace much — quite a bit before that. The reasons for this particular population, dually diagnosed population being underserved I think fall into two primary categories. One has to do with the clinical state of the art. Diagnosis and treatment of the dually diagnosed individual has lagged far behind clinical advances in other areas of mental health and other areas of developmental disability.

It's an emerging field. There are beginning to be national experts and hard empirical data which are showing that there can be effective clinical diagnosis and there are effective clinical treatments for this population and it's very recent that these developments have occurred.

The second major reason specific to California are the -- what I see as the fundamental philosophical differences between the mental health system and the regional center system.

The mental health system, as I see it, was conceived

of as a system which was to deliver in the most efficient way possible direct services to the consumer at the local level. This is the way the local mental -- the county mental health agencies were organized. They consist primarily of direct services personnel with less personnel at the top, at the administrative level.

However, the regional center system was conceived of to be a broker of services at the community level to integrate the mentally retarded and otherwise developmentally disabled persons into the existing service system of the community, part of which from our point of view at the regional centers is the community mental health agency.

We fully -- the system was created fully expecting that we could refer our clients with mental health needs to county mental health agencies and they would be served. It's a wonderful idea. It has not worked.

These people have not been served. By and large, the county mental health agencies are not aware of the procedures for how to diagnose mental disorders in a mentally retarded individual or any other disabled individual who cannot speak or otherwise communicate clearly. There are biases and prejudices which were existing in graduate school when I was training which lead a lot of mental health professionals to the point of view that a mentally retarded person is not amenable to treatment. If they have a mental disorder, it is directly related to the mental retardation or developmental disability. And, therefore, that person is not a good candidate for treatment by traditional psychotherapeutic means.

As a result of this, there have been accusations flying back and forth between the two systems at the State level and the local level for the past 15 years regarding inappropriate referrals and a lack of responsiveness and a lack of cooperativeness between the two systems and this has led directly to this population being inadequately served, and in many, many cases in these individuals being forced to be institutionalized.

CHAIRMAN MARKS: Do you think the State's doing enough?

MR. FERRELL: I think only now has the State at the State level begun to recognize this problem by the fact that the Department of Developmental Services was willing to fund a \$40,000 six-month pilot project at Far Northern Regional Center. I have to say I believe that in my view the leadership has come from the Regional Center side of the system to the agitation, if you will -- equate that with leadership -- for funds to be directed for this purpose and for staff time to be directed for this purpose also.

DR. BRYNJOLFSSON: If I could, Senator, it is to the credit, however, of the task force, the joint task force under ARCA's leadership between Mental Health and Developmental Disabilities — it is to their credit that they did endorse and support this pilot project and have been fully behind it. And Mr. Nader, in his capacity, has been entirely supportive of this effort.

CHAIRMAN MARKS: I'm not specifically opposed to task forces, but oftentimes a task force is a way to avoid

an issue.

DR. BRYNJOLFSSON: Yes, sir. In this case -CHAIRMAN MARKS: It would not be the case?

DR. BRYNJOLFSSON: Yes, sir. In this case, the
task force has seen fit to see that something is implemented
at the grass roots level and has seen that a project has been
developed that results in direct services to clients.

MR. FERRELL: I'd like to echo Mrs, Epstein's statements that there exists no clear responsibility. I have read the memorandum of understanding between the Department of Mental Health and the Department of Developmental Services. I have read the memorandum of understanding between my agency and each of our county mental health agencies as well as others around the State. All of those on paper -- when I ask people who does it work, nobody knows. Very few people actually follow what has been written down on these pieces of paper.

CHAIRMAN MARKS: I wonder if you'll write me a letter and tell me what you think your problem is.

MR. FERRELL: We have submitted a letter that I primarily wrote which comes from ARCA, which summarizes our position as well as the preliminary report from ARCA. And those have been submitted. I will also give you extra copies today.

I believe that the primary needs at this point four months into our project -- almost five months into our project -- we must come up with a jointly acceptable clinical definition of dual diagnosis, a pragmatic definition of

somebody who needs services from both systems.

However, one of the problems is that there are clinicians in both systems who make these decisions and they are used to using empirical clinical data to make these decisions on the primacy of the disability, on treatment decisions as well.

MS. EPSTEIN: Up to very recently part of the problem that we seem to hear about is that in making a clinical decision for a person, very often the doctor making that decision really looks at the person as being -- in the case of DD people, developmentally disabled, and couldn't look beyond that to understand that there was a second problem.

MR. FERRELL: Absolutely. Clinical research literature has shown. The concept is called overshadowing in the literature.

There have been clinical studies where professionals, psychiatrists, et cetera, looked at case studies, written-up studies as the only difference between two cases was the I.Q. score, which indicated one person was mentally retarded and the very same description of symptoms in another vignette. Without that I.Q. score indicating that that person was mentally retarded, the diagnosis was entirely different. The professionals by and large could not see beyond the definition of the diagnosis of mental retardation the clinical symptoms. And the symptoms do present differently in many cases. I want to emphasize it is an emerging field and the clinical skills are not out there.

In my time with the pilot project, I have been

absolutely appalled at the lack of information that each service system has about the other. One of the things that we have done in our pilot project is to initiate interagency training, getting agencies together to talk about the services and make contacts at the line level, the people who see the clients, the mental health line workers and the Regional Center line workers to establish lines of communication directly between them and not necessarily having to talk through the administration and director to talk to director.

That's not the most effective way to get administrative decisions made and it's not the most efficient way that clinical decisions are made. And it's important to have the agencies work together in any kind of efficient way to meet the human needs of these folks.

Additional clinical training is needed as well.

Administrative flexibility must be there as well as adequate funding. We're identifying a population whose needs are substantially different from solely mental health needs and solely needs based on developmental disabilities. This is a new population of different needs. There must be more funding in order to adequately serve these folks.

I would say the two biggest problems -- the two biggest programmatic problems, at least in our region, and I think all over the State lie in the prescription and monitoring of psychotropic medication for the developmentally disabled who also have mental disorders and in-patient crisis services for the developmentally disabled persons who, as a

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result of a mental disorder, act in ways that render them a danger to themselves and others in the community.

Monitoring psychotropic prescription medications for our clients -- Regional Centers are not staffed with psychiatrists. These are the people who have appropriate clinical training to monitor these things. Mental health agencies in our area are also by and large not staffed to handle these extra people. We refer to psychiatrists whenever we can. Psychiatrists in our area have Medi-Cal quotas. They don't take -- they take only so many persons who are funded by Medi-Cal. Quite often by the 5th of each month they have reached their quota. Therefore, we cannot get these folks seen.

What has resulted is that general practitioners, family practitioners have ended up prescribing and monitoring psychotropic meds because they know they are needed, but they also are aware that it's outside their clinical expertise.

CHAIRMAN MARKS: I'm a member of the Budget Committee and we have the ability to increase or decrease the budget. And I would be glad to do something about it if I'm given the information at the appropriate time.

MR. FERRELL: As a result of our pilot project, sir, you'll get some very specific recommendations from our point of view about what can be done to solve these problems.

CHAIRMAN MARKS: Okay.

MR. FERRELL: And our final report will be generated conjointly with the directors of the --

CHAIRMAN MARKS: When will this report be ready?

MR. FERRELL: We'll be forwarding it to the Department of Developmental Services no later than August 31st.

CHAIRMAN MARKS: Why don't you send me a copy?

MR. FERRELL: Yes, sir, certainly. I certainly will.

Our pilot project has several pages of objectives which I won't go through. But I would like to group them into four main categories which relfect, I hope, the comments that I've made thus far.

One of our primary activities is in the area of training, the interagency training I mentioned, the clinical training I mentioned. We've contracted with an expert in the field from Northern Illinois University to come to our area and to do two two-day workshops in our area on diagnosis and treatment of dually diagnosed individuals. We also are in the midst of setting up what I'm calling administrative training which boils down to being the directors of the four county mental health agencies we're working with, our director, Walter Baldo, and the directors of the Department of Mental Health and the Department of Developmental Services, all in the same room at the same time with these issues laid out on the table in front of them and let them discuss it on an administrative level.

CHAIRMAN MARKS: When and where is the hearing?

MR. FERRELL: The where is in Sacramento. The date

I am not sure has been set yet.

DR. BRYNJOLFFSON: We'll inform your office of the

time.

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MR. FERRELL: The second primary activity is data collection regarding the mental health needs of the developmentally disabled. Only very recently -- as recently as March, '85 issue of the Journal of American Society of Mental Deficiency has there been instruments designed to assess the mental health needs of the mentally retarded.

We are involved right now in a project assessing in a systematic way exactly what are the mental health needs of our clients at Far Northern.

made myself available on a 24-hour basis to each of the in-patient units in our four county mental health agencies. When one of our clients shows up in an in-patient crisis, they call me. And I go. And I see what's going on. And I interview that client. If that client is in such a state, I interview the family, the care provider where that client lives, and I interview the sheriff's officers wherever possible who brought that client in in a crisis and --

CHAIRMAN MARKS: What county is this?

MR. FERRELL: We serve nine counties. The four counties that are involved in our mental health project with us are Butte, Glenn, Tehama, and Shasta.

CHAIRMAN MARKS: I'm glad you're in Northern California.

MR. FERRELL: Thank you. The fourth primary mode of activity is the administrative mode. We are reviewing the memorandum of understanding between ourselves and each of

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these four counties, as well as the rest of our nine counties We will revise those at the outcome of this in our area. project to reflect the plans and procedures we want to put into place.

We are reviewing the memorandum of understanding between the Department of Mental Health and the Department of Developmental Services at the State level and will make recommendations in our final report as to changes we feel need to be made.

We will also make recommendations regarding legislative changes that need to be made.

And I'd like to quickly point out one change -one legislative problem that hamstrings Regional Centers in dealing with our clients.

Welfare and Institutions Code Section 6500 gives Regional Centers the power to petition Superior Court to involuntarily detain a mentally retarded individual who has shown themselves to be a danger to themselves or others.

The law says specifically mentally retarded individual. This excludes a great many developmentally disabled individuals for whom the Regional Centers have primary case management responsibility -- the cerebral palsy individual, the epileptic individual, the autistic individual.

Just during the course of this pilot project, we have run across several cases in which we, as a Regional Center, had absolutely no standing to intervene in spite of the most gross assaultive or self-injurious behavior on the part of these individuals. We have been very fortunate with our four county mental health agencies because they have begun to work with us to pursue conservatorships on these individuals. But it's a very cumbersome process. By and large, some of these folks really do not get immediately into their system either. We believe — at least I believe and I'm going to recommend in my final report — that legislative changes need to be made so that Regional Centers can have this standing also with all of our clients and not just the mentally retarded.

This section of the W & I Code has been very effective in protecting the rights of the developmentally disabled. Unfortunately, it has also prevented them from getting certain services which they need.

Thank you.

DR. BRYNJOLFSSON: Thank you for your patience in hearing us out, Senator.

CHAIRMAN MARKS: Thank you. We appreciate you coming before us and I hope you'll furnish us with the information I have requested. Maybe hopefully we can do something in the area of training and improve the system.

DR. BRYNJOLFSSON: Thank you, Senator.

CHAIRMAN MARKS: Okay. Is Wendall Fingar here?
MR. FINGAR: Yes, sir.

CHAIRMAN MARKS: You're representing Mr. Shaw?
MR. FINGAR: Yes, sir, I am.

CHAIRMAN MARKS: Let me say this. This completes the testimony of those who are on the agenda. There are a couple of other people who wish to testify and we will give

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you that opportunity. Go ahead.

MR. FINGAR: My name is Wendall Fingar. I'm a staff assistant with the State Department of Rehabilitation here in Sacramento testifying for Ferd Shaw, our Deputy Director for field operations.

Fortunately I prepared the notes for Mr. Shaw not being aware at the time that I would be using them personally.

But I think having listened to your presentations, that it might be a better use of your time if I were to share a copy of these background notes with you and to concentrate on the one problem area that seems to be most related to your concerns.

CHAIRMAN MARKS: You'll give us a copy?
MR. FINGAR: Yes, I will.

CHAIRMAN MARKS: Give it to the Sergeant.

MR. FINGAR: Right there.

Our program serves about 60,000 people statewide at any one time through about a system of 600 vocational rehab counselors.

In all the categories that I looked at, about one-third of the total represents the mentally ill, whether it's a referral or writing a plan to resolve a person's problem, or the end result of our system, of course, is employment in the labor market. But the figure of 33 to 35 percent seems to come up over and over when we're looking at those represented by the mentally ill, not including the DD and MR. That's a separate category in our system.

I asked our program consultant who works with the mentally ill, alcoholism, drug abuse, and who has experience of about 25 years in that system to review your announcement and the three questions you had.

And we kept coming back to the same thing. The problem in his perspective and our receiving fewer numbers of mentally ill in recent years and a more severely disabled group, that we have a harder time succeeding with seems to, he feels, resulted from the shortage of funds in the community mental health service system statewide in the sense that the funds that are available have had to be geared more into the acute and continuing care patients as just heard by our earlier witnesses, and more attention, therefore, away from the outpatient service area. And dovetail more directly with the outpatient services. That's where the referrals to our system tend to come from other than those that are self-referred to our field offices.

The reason we identified so closely with the outpatient services component of the system throughout the State is that we depend very heavily u pon the continuation of counseling, therapy, and medication for the mentally ill client that we're serving while our vocational rehab counselor is working with them to develop a plan to get them back to work.

So often our vocational client falls apart if the ongoing underpinning of therapy and medication is not kept available for maybe the 12 to 15 months that we're serving him.

CHAIRMAN MARKS: Let me ask you a question. I'm on

one of the subcommittees but not on that particular subcommittee that handles your funding. But did you make requests for additional funds to those subcommittees?

MR. FINGAR: To serve mentally ill specifically? CHAIRMAN MARKS: Yes.

MR. FINGAR: I do not know. I'm not -- I'm new in this assignment. I don't really know.

CHAIRMAN MARKS: It's hard to do anything unless we get requests.

MR. FINGAR: We are working, for example, with L. A. County to try to increase the number of specialist counselors in a special co-op contract.

CHAIRMAN MARKS: I'd like to find out if you made requests to a subcommittee to get additional funds. If you didn't make the request, I cannot conceive of how you espect to get them.

MR. FINGAR: Right. But that relationship to the outpatient services and the difficulty of receiving referrals who are continuing to receive therapeutic support services have been a problem in recent years for sure. That was the one pint I wanted to make most urgently.

I think our success because of that in recent years has tended to be with those mentally ill referrals who are less severely disabled and does not need the ongoing supportive services and medication after they come into our system. We tend to be able to serve them better because of the facts I mentioned earlier.

Other than that, I did take a look at the

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double handicaps that was mentioned in your announcement.

That's a little hard to get a handle on in our statistical system, but I found that — I talked about one-third of our cases earlier. If we look at mental illness combined with any other physical disability, such as the hearing, visual, or orthopedic problem, that percentage goes up to about 38 percent; you add another five percent to that group that we're serving if we try to identify them in that way and that's one way of getting a handle on that.

It was an interesting thing that I have never looked at before. We do code both the primary and the secondary disability and do have retrieval of that information.

CHAIRMAN MARKS: Well, what I don't quite understand is why they're so hard to find. You just said a moment ago that it's hard. Why should it be hard? Yours is is a department which is designed to help people presumably. Why should it be hard to find?

MR. FINGAR: My understanding from talking with the consultants that I referred to is that the people are there but the ease of identification referral to our field offices is what has been more a problem in recent years. They aren't as identifiable in outpatient services programs in many of the counties as they were in the past. We don't have the same volume and support services provided by the community mental health services programs that we had.

And, therefore, even though we may get them into our system, it tends not to be as successful as they fall apart once we

try to develop a written rehab plan to get them back to work. That's my understanding.

CHAIRMAN MARKS: I would presume that the local agencies must know about the problem and must notify you about these problems. I cannot conceive of how they don't.

MR. FINGAR: I think what happens more is they tend to continue to refer the same volume of people but the severity of the disability is greater and the supportive services from the mental health system that we rely on heavily tends to not be as thorough as it was in the past. And that's what causes that success ratio.

MS. EPSTEIN: What's happening with some of these people who are the dual diagnosed in terms of the Department putting them into either workshops or places where they would go and start to be rehabilitated and go back into the community?

MR. FINGAR: I do not know what happens in terms of outcomes because that would take quite a bit more of investigation to really tell what has been happening. All I can tell is within our active caseload there is this percentage and these numbers by disability groups and you will see the variation in the size of the group there. So, I don't really know if our success rate has been more successful with one of these subgroups or another. I really don't know.

CHAIRMAN MARKS: It would seem to me that your

Department can come up with more information about this

particular problem and should do so. I'm very concerned

that you did not. You obviously don't have the information.

It's inconceivable to me that the local agencies that are handling these problems are not informed of these problems. I find it very difficult to believe.

MR. FINGAR: Are you referring specifically to the persons with a double disability or --

CHAIRMAN MARKS: I'm referring to any or all the problems that are relating to people in this category.

We've heard witnesses. We heard the gentleman right down there talk about the problems that he has. I'm sure he must have notified you.

MR. FINGAR: We have the counseling staff available to serve the persons referred. What I was referring to is that the change in the mental health system in funding has caused more of a shift toward emphasis to the acute and away from outpatient. And that kind of causes a dislocation on those being referred to us. The volume may be the same, you know, as in the past, but the results I think are not as good.

MS. EPSTEIN: I think the questions that keep coming back to us is that there are a number of people who have the double diagnosis and who just seem to float from one place to the next place and don't seem to get help.

MR. FINGAR: We also have, apart from our mainstream program, we also have an habilitation services program with many DD persons identified there. It's a State funded separate program. I don't know that for sure, but I'm certain it must be a large block of MI/DD combinations represented in those other programs, too. So that shouldn't

be overlooked. But I was thinking more of our mainstream vocational rehab program which is what I was referring to.

CHAIRMAN MARKS: Does that complete your testimony?

MR. FINGAR: Right.

CHAIRMAN MARKS: I would like the people who represent local agencies to indicate -- to send to me copies of the information that you've sent to the State Departments. I'd like to see what they received. I'd like to see whether they acted upon it because I'm very much concerned that -- I've heard a lot of the problems that the people are having and I'm very much concerned that a lot of the problems are not being resolved.

You have plenty of money. If you don't have enough money, you should ask the budget committee for more money. If the Governor or the administration doesn't like the appropriation, you should still ask for it.

MR. FINGAR: That's on a policy level that I'm not familiar with. I assure you.

CHAIRMAN MARKS: Talk to your chief. Talk to the chief of your Department if there's not enough money. We may not approve it. We may or may not approve it. We'd have to look at it. We can't do anything at all unless you ask for additional funding. If a specific problem exists, you're to tell it to us.

MR. FINGAR: Well, our Department I don't think is usually that very bashful about asking for money. But in this case, the money is outside of our system that would, I think,

45 1 cause the change that we would benefit from. It would be 2 money in outpatient services in the community mental health 3 system. 4 CHAIRMAN MARKS: You could make some suggestions 5 to us. 6 MR. FINGAR: Sure. 7 CHAIRMAN MARKS: And we in the budget committee 8 cannot do anything unless we know what you want. 9 MR. FINGAR: Certainly. 10 CHAIRMAN MARKS: Okay. 11 MR. FINGAR: Thank you, 12 CHAIRMAN MARKS: Thank you. 13 MS. EPSTEIN: I have just one question for you. 14 MR. FINGAR: Certainly. 15 MS. EPSTEIN: Do you know if people within the 16 Department in the local levels who deal with dual diagnosed 17 people are trained sufficiently to be able to understand 18 their problems when dealing with the individuals? 19 I would -- my personal opinion is MR. FINGAR: 20 that only in part. And I'm thinking of the example of 21 the co-op contract we have with the Department of Mental Hygiene. Those 14 specialists counselors -- excuse me -- 15 22 23 in 14 different counties I think receive some good specialized 24 training in working not only with the mentally ill but 25 other disability groups that also are mentally ill. 26 There I feel more secure that they would have it. 27 Generally speaking, I don't know. I doubt that that's the

case throughout the whole Department.

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MS. EPSTEIN: I ask the question specifically because in the San Francisco office where the Senator's subcommittee is located very often people come into the office seeking advice. We, in turn, send them down to the Department. And, in turn, we get the people back seeking the same advice.

MR. FINGAR: Right.

I'm familiar with that problem. I just do not know, but I think it's generally only the special programs that have been identified, specialty training would be better.

MS. EPSTEIN: Is there a way that the Senator's office could help the Department see that people on a local level are more attuned to the needs of these people that come in?

MR. FINGAR: The way it would seem to help would be to increase the coordination between those agencies and yet I don't -- I'm not that familiar to know what the best vehicle to do that would be at the actual service level, you know, how to accomplish that. I'm not sure.

CHAIRMAN MARKS: Okay. Thank you.

MR. FINGAR: Thank you.

CHAIRMAN MARKS: We have a couple more witnesses I believe. Mr. Art Segal and Elizabeth Swain. Is there anybody else who wishes to testify at this hearing? Okay.

I'll call on you.

MR. SEGAL: I have a prepared statement which I'll read. I also have a number of comments which I've taken in the

past ten minutes which I'd like to --

CHAIRMAN MARKS: How long is your prepared statement?

MR. SEGAL: Three or four minutes at most. Then I'd like to tell you a little bit about Marin County.

CHAIRMAN MARKS: I think I know a little bit about it.

MR. SEGAL: I hope so.

I think I need to say, first of all, that Marin

Community Mental Health has a full contingent of staff that

serves disabled people and always has since its inception in

the mid-sixties. I need to say this because I need to be able

to point out that the problem is not limited to mental health

centers not having the staff or having different attitudes.

It's much deeper than that. And I want to get into that.

It does also include in some parts of the State at least that mental health centers do not want to serve disabled people.

I also need to say for the record that I have served as a psychotherapist with disabled people for almost 35 years now. And I can support anyone who says that disabled people respond to psychotherapy in just about the same way as a nondisabled person does.

Just a little bit of information about what happens when a person has a disability. What happens is that it places all members of the family in a high risk category for serious depression. In taking a look at what happens to families, we find that disability is prevalent in divorce

situations, in suicide. A recent study in San Mateo County of suicide in adolescents found that more than half the adolescents who committed suicide that one year had learning disabilities. There is a high prevalence of disability in criminality and in mental illness. This does not suggest a cause and effect relationship; that disability creates these problems. However, it does suggest that the findings provide evidence that disability creates situations that are beyond the disabled person's coping abilities. This is true.

The mental health system today lacks the resources to prevent the deterioration of the disabled person. We don't have them. Furthermore, and for the same reason, we are unable to help disabled persons gain the emotional strength requisite for independent living. We can't even do the prevention work that needs to be done.

Since Proposition 13 went through, I have witnessed a slow yearly deterioration of my ability in Marin County and we have a strong focus on certain disabled people and people in high risk categories —— I have witnessed services fall apart because of lack of funds, not because of a negative attitude, because of lack of funds.

And as a consequence of lack of funds, I have been part of I guess a thrust from the State I would think to serve the most severely mentally ill person or the chronically ill person. This has resulted in decrease in prevention, has resulted in pretty much wiping out our outpatient department, not just for the disabled, but for everybody.

49 1 Outpatient services are available if you're 2 seriously mentally ill. If you are a parient who has an 3 autistic child and need supportive help, five years ago 4 that was available through community mental health. That is 5 no longer available. 6 If you are an adult who has a brain injury and you 7 have a spouse who needs some help -- what do I do next? -that used to be available. That is no longer available. It's 8 not only attitude. It's priority. The priority comes from 9 10 the State. We cannot simply close down the crisis --11 CHAIRMAN MARKS: Proposition 13 didn't come from 12 the State. It came from the people who voted for Proposition 13. 13 14 MR. SEGAL: Well, it came from the people, but there's no money available. 15 16 CHAIRMAN MARKS: What I'm saying is that Proposition 17 13 was adopted by the people. MR. SEGAL: Okay. True. Yeah. Okay. 18 19 I have some recommendations that I want -- before 20 I get to those, I want to read some notes that I took while people were talking. I find myself in some disagreement as 21

well as in agreement.

CHAIRMAN MARKS: You're not required to agree.

MR. SEGAL: Good. Thank you.

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I agree with Mr. Nader that localities set priorities rather than the State in many instances. This is true. And it's also true that I work in Marin County where the locality has its priorities and we are serving disabled

people. So, that needs to be said.

We also need to remember that we talk about dually diagnosed and we're not limiting ourselves to persons who are served by Regional Centers. And that sometimes we seem to limit our thinking to that category.

The problem for the brain injured and the learning disabled and the physically disabled people who are not served by Regional Centers are manyfold greater than the persons served by the Regional Centers. And there is no commitment anywhere in the State or Federal Government to serve persons with these other disabilities. Medi-Cal funds are decreasing. Private agencies don't have the resources to do it.

The other problem that is exists exists within the State Department of Developmental Services. I think that needs to be looked at as well. As a -- what I like to think as forward looking mental health system -- several years ago, I received permission from Beverly Abbott, who was then our mental health director, to work with the Golden Gate Regional Center to develop a group home to serve dually diagnosed where we would provide the mental health service. And we tried to encourage agency people in the county to do this. We went to the group homes and the residential programs who serve the developmentally disabled and they didn't want anything to do with developmentally disabled who have behavior problems because they were not getting enough reimbursement from the State to serve that population.

I went to professional agencies that serve the

mentally ill. I found one that said, yeah, it really makes sense and we really ought to have a group home for the developmentally disabled. That person sat down with me, the Executive Director of the Regional Center and talked about reimbursement and said, "I can't afford to do it. I'll go broke."

That is part of the problem.

CHAIRMAN MARKS: Well, last year I authored legislation to give a lot more money to local governments. Maybe you don't know about that. And it provided additional funding for each agency in local government, more money than they had anticipated for many years. You ought to go to the Board of Supervisors and ask them for some of that.

MR. SEGAL: Okay. Marin County Board of Supervisors does contribute far greater than the 10 percent than they're required to.

CHAIRMAN MARKS: Each city, and the supervisors, and the county, and each agency in each county and city throughout the State of California, each county in the State of California has additional funding so maybe some of it can come from them.

MR. SEGAL: Two or three other notes I have here, one has to do with resources. You know, community mental health, at least in Marin, works under the philosophy that we would like private agencies to provide the service and we will contract with the private agencies. You know, private agencies do not want to serve this population from either system. So, when we have on our crisis unit a developmentally

disabled person and we try to move that person through the next step, a day treatment program or a residential program, or an outpatient program, there's no place for that person in Marin County.

When we have the money and say we'll pay for it, there's no place to send that person, no agency wants to provide the service.

The other problem is that the State Hospital system -- if we put a developmentally disabled person in a State Hospital for the mentally ill, the State Hospital for the mentally ill tries as quickly as possible to get that person out to a State Hospital for the developmentally disabled.

The State Hospital for the mentally ill by and large does not have the resources, they claim, to serve people who are developmentally disabled.

I think the problem is a systems problem. And I think it needs to be looked at that way.

CHAIRMAN MARKS: It's possible that the task force could look at the last problem you're talking about.

MR. SEGAL: The task force went on for six years. I was on it for two of them.

CHAIRMAN MARKS: Maybe I was right about the delays. It delays things.

MR. SEGAL: I have five recommendations. One, that there be a health and welfare agency commitment to serve the dually diagnosed.

Two, that State Mental Health priorities include

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serving high risk populations without the requirement that the applicant be mentally ill and that additional funding for this service be made available.

Three, that State Mental Health actively promote the development of mental health services for the dually diagnosed.

Four, that State Mental Health promote in-service training for mental health service to developmentally disabled and other disabled individuals.

Someone said earlier that a lot of people don't know what to do. They're generally correct. Mental Health people tend to look at the psychiatric side of people. don't see disability or any physical problem.

Five, that State Mental Health monitor local community mental health services to the dually diagnosed. I don't think it's correct that the State Department of Mental Health has nothing to do with services on the local level or that priorities are set on the local level is fully responsible.

The State Department of Mental Health does come around periodically and does conduct program evaluations; they do have coordinators for children's services, for adult services, for disability services. These people do come around. They do make demands on local mental health centers, and I think those demands can be increased perhaps.

Thank you.

CHAIRMAN MARKS: Do you have a copy you're going to give us?

> MR. SEGAL: Yes.

CHAIRMAN MARKS: Thank you.

MR. FINGAR: Thank you.

CHAIRMAN MARKS: You want to testify?

MS. SWAIN: Yes. I'm Elizabeth Swain and I'm speaking as a member of the Board of the Marin Brain Injury Network.

And my concern -- I just have a couple of notes on mental health needs for the traumatically brain injured.

Currently, there is no State or Federal funding for psychotherapy for families of traumatically brain injured individuals. It's been shown that the effect of such an injury for family members is devastating to the family often resulting in divorce, suicide of a family member, severe psychological disturbances, and disabling depression.

The real cost to society over time is very severe. We need to heal this problem at the point of trauma rather than later after years of suffering where the resulting problems — medical, psychological, and social — which do fall under government funding become too severe for society to ignore.

For the families also I believe that respite care is very much needed for traumatically brain injured individuals who are taken care of at home by the family.

There are many programs in existence, a growing number right now, for Alzheimer victims, but it seems to me that both the facilities and the staff could work in a combined way that could be economically a good thing.

It should be, however, I think, productive respite

allowing the family to rest and heal and having some therapy for the brain injured person during that stay.

Also, psychotherapy is needed for the brain injured individual. Psychotherapy -- also done by a psychotherapist who is knowledgeable about brain injury.

There is research showing that brain injured patients given psychotherapy alone made a better recovery over time than those given speech and physical rehabilitation therapy with no psychotherapy. Dr. Roberts down in Orange County (sic).

Also, it's shown that after a head injury, at the point in recovery when the patient is beginning to have self-awareness, to understand the nature of the damage to themselves, stress, depression, and anxiety increase and performance and I.Q. drop. And this is in spite of continued improvement in cognitive and physical functioning.

Also, research by Marshall Levy at the Los

Angeles Department of Mental Health -- he was doing research
on the criminal justice system -- turned up that 30 percent
of the inmates in the State Prisons in California are brain
damaged. It seems to me that it would perhaps help the
prison system also to treat the brain damaged population
in a different way. These are behavioral byproducts of
lack of psychotherapy.

And I had a thought -- a final thought -- that it took a long time for us to come to understand the benefits of returning disabled people to a productive life in our society. It took a long time for creating small changes that

make a big difference, like wheelchair ramps, curbsides, and into buildings. And it seems to me that now we need ramps for the mind.

CHAIRMAN MARKS: I like that. Thank you very much. We appreciate your both being here.

MS. SWAIN: Thank you very much.

CHAIRMAN MARKS: Sharon George-Perry.

Good afternoon.

Ms. Perry: Thank you. I originally had not planned to say anything, but I heard something that I think needs some comment on. I made some notes, so I don't have anything to give you in writing.

The first thing I want to say is in response to something Mr. Ferrell said. The first thing he did that I want to expand on was -- comes out of the experience that those of us who have been involved with 3632 have found. That has to do with the lack of horizontal linkages between systems. And a part of the problem that we see with this -- I remember meeting back with Dorothy as long as two years ago trying to understand the communication systems as they affect the disabled, and couldn't at that time understand, have since come to understand that system linkages follow the flow of funds.

We have found during this struggle over the implementation of 3632 that since funds have to flow parallel, horizontally. Horizontal linkages are being developed. And I'm not sure how that's going to be done in the field of the disabled. But it's clear that some specific system --

attention is going to have to be given to building those linkages.

The second thing had to do with the prejudice that Mr. Ferrell alluded to on the part of mental health clinicians towards disabled people. In the mental health field for a very long time mental health monies were held to be wasted on people who weren't amenable to treatment. Basically, that meant we didn't know how to cure them. They were hard to deal with. Unlike other disability systems, mental health put its money on the people with the best chance of getting better. People then who are — come from physical disability systems have a different orientation. The disability is classified by the degree of dysfunction and the amount of public benefit available bears directly with the degree of dysfunction (sic).

So, the physically handicapped person comes to the mental health clinician looking for help and the clinician's orientation is that we can't change the basic position.

They've got a hearing problem or a seeing problem or whatever.

It can't be changed, so why should we put money into it?

They're not amenable to treatment. That's a system issue. And I don't think any individuals have to be blamed for it. It has to do with treatment philosophies that have existed.

The implication in the mental health field has been that the most seriously mentally ill people were not cared for by the counties at all. They were shunted out to board and care homes. And, of course, the pressure of

mentally ill people on the streets is causing the mental health system to reevaluate that some, but in my judgment not enough.

But clearly, when you look at systems and design systems that will address the mental health needs of physically disabled people, there are going to have to be specific provisions to address that prejudice on the part of the mental health people.

And those two things just stood out to me, something that needed to go into your thinking and your process.

CHAIRMAN MARKS: Thank you very much. We appreciate your being here. And your suggestions are very good.

Ms. Perry: Thank you.

CHAIRMAN MARKS: Does anybody else wish to testify?

Let me say that this hearing I think has been a good one because we've got some idea of the problems and also we had some idea of what the State should do to try to resolve some of these problems and hopefully go forward to try help, because I think this is a group that definitely does need help.

And I would like to offer office anyway that I can to assist in suggestions you made for legislation, for budgeting, anything that can be of help will be beneficial.

And I'll do my very best to see this is done. Thank you very, very much.

(The hearing was adjourned at 2:55 p.m.)

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CERTIFCATE OF SHORTHAND REPORTER

I, Nadine J. Parks, a shorthand reporter of the State of California, do hereby certify:

That I am a disinterested person herein; that the foregoing hearing of the Subcommittee on the Disabled was reported in shorthand by me and thereafter transcribed into typewriting.

I further certify that I am not of counsel or attorney for any of the parties to said hearing nor in any way interested in the outcome of said hearing.

IN WITNESS WHEREOF, I have hereunto set my hand this 11th day of June, 1985.

Nadine J. Parks Shorthand Reporter