

MEETING MINUTES
HUMAN SERVICES SUBCOMMITTEE
FEBRUARY 4, 1987

The meeting of the human services subcommittee was called to order by Chairman Winslow at 8:07 a.m. on February 4, 1987 in room 325 of the state capitol building.

ROLL CALL: All members were present.

PUBLIC TESTIMONY - DEVELOPMENTAL DISABILITIES (DD)

(31a:000) Dave Lewis, director of Social and Rehabilitative Services (DSRS), spoke briefly on the issues that would be addressed before the committee, namely the need for expansion for the waiting list, and the problem with the level of provider support.

Chris Volinkaty, lobbyist for Developmentally Disabled, introduced the testimony that would be presented to the committee, including respite care, family training, day programs, supported work, independent living, specialized family care, and the new proposed special services support organization.

Tom Crosser, chairman of the Developmental Planning Task Force, asked the committee to consider the percentage of the developmentally disabled population that was unserved or underserved, and how to best meet those needs. He covered the final report (exhibit 2) of the task force, and described the summary of their recommendations, which he described as a roadmap to the future.

In response to a question from Chairman Winslow, Mr Crosser stated the first mile of the road map had been introduced in two (2) senate joint resolutions which reflect, in a broad way, the recommendations made by the task force. He stated the only monetary recommendation was the Special Services Support Organization (SSSO). He noted other recommendations can be accomplished during the interim without a substantial financial impact. He stated he did not see any one recommendation in particular being the first step, but noted the executive budget reflects a move into the right direction, with the Developmental Disabilities restructure under one (1) administrative system. He stated the reason the task force did not come up with a recommendation for a specific agency designation for DD was because they felt it was beyond the role and scope of the task force. He stated the team did not look at those issues, but realized there were some obvious questions that needed to be addressed. He added that one of the resolutions in the senate

calls for a legislative interim study to restructure the DD services.

In response to an inquiry from Rep Switzer, Mr Crosser defined a naive offender as a developmentally disabled individual who comes in contact with the criminal justice system and is not cognizant of the right or wrong of breaking that law.

There was discussion of the final report (exhibit 2) that was submitted by the DD task force and the general information it contained. Sen Himsel commented that the report was not specific enough and asked for more guidance and specific direction on how to proceed with the task force recommendations.

(31a:250) Brodie Mall, director of Northern Gateway Enterprises, covered the issues and concerns raised through the Priorities for People (PFP) process dealing with the DD program. He stated the primary concern was the restoration of the base at 4%. He added that it will take \$3 million to address the waiting list of 289 people identified as unserved. Other issues supported by the PFP process was a the 52 bed SSSO for the severely handicapped, supported work for the 172 graduates from special education from high school who have no services available, and upgrading respite care to the Montana minimum wage from the current wage of \$1.15 (exhibit 18).

(31a:306) Rep Jerry Nisbet, president of the board of directors for Region II Child Care Services, addressed two (2) points: (1) facilities have been at the same funding level for three (3) years while operational costs have risen and staff are paid significantly lower wages than comparable other positions and (2) a freeze or decrease in the provider reimbursement rate would result in the deterioration of services provided.

(31a:353) Peyton Terry, Glasgow, explained the services that have been provided to his daughter, who is now 32 years old and mentally retarded. He stated she had received excellent services up to the age of 18, at which time there were several unsuccessful attempts to find placement for her. She is currently in the Transitional Living Complex (TLC) and making tremendous progress. He added he supported previous statements in support of DD programs and urged continued support of the programs in Montana.

(31a:436) Rep Les Kitselman, Billings, explained that his son Brian suffered from encephalitis of unknown origin. He stressed the importance of the support his family received from the people and the programs. He added he is a strong

supporter of early intervention and supported full funding of the DD program.

(31a:466) Jim Foster, superintendent of schools, Chester, spoke on two (2) issues: (1) the problem of youth graduating from high school special education and having no services available for them, the problem of long waiting lists for current services, and the regression that takes place when there is no follow up after graduation. He stated there is no future for those youth, and encouraged expansion of funding for services for this population of unserved youth. (2) Mr Foster has a 15 year old autistic son in Billings receiving special intensive care. He said that in three (3) years, when Brian is 18, he will not be eligible for services. He stated Brian has made tremendous strides and progress, and thanked the legislature for the progress in the programs for the handicapped that are currently available. He concluded by asking the committee's consideration of program expansion for those youth like Brian who will be ineligible for services and for whom there are no other services available.

(31a:568) Jessie Schlinger, Kalispell, testified that she has been involved in foster care for 20 years. She added that she has a developmentally disabled daughter who is now 15 years of age who was not fortunate enough to receive the services that are now available. She said that two (2) years ago they took Joey into their home for care, and it took almost a year before he could receive services from the Comprehensive Development Center (CDC) in Missoula. She expressed the importance of the staff from the center in maintaining Joey in the home and providing relief for the family; their dedication and the low pay they receive for the types of services they provide. She stated it took two (2) years to get respite care, and that she could not take care of developmentally disabled children without the support of the staff and trainers and services she received from CDC.

(31a:650) Greg Olson, director of Westmont, spoke on behalf of the staff and program at Westmont. He said that since the special session in June and the loss of DD funding, there have been no raises for his staff. He noted the starting salary is \$4.30 at the facility compared to \$6.00 at the Montana Developmental Center; a position with less responsibilities than are required at Westmont. Mr Olson stated his staff are having a hard time making it; that of the 71 FTE's employed at Westmont 21 are married, 11 are single parents, 13 of which are employed outside of Westmont in other positions, 12 are eligible for AFDC, food stamps, or some type of assistance. He explained his concern that he will not be able to keep the staff he presently has unless he gets an increase in the budget, and that a 3% increase

would be wonderful, while a decrease or a freeze would result in a lay off of staff. He stated that in past contracting sessions he was told formally by the division that there was to be no layoff of direct care staff, and that any layoffs from lack of funding were to come from administrative staff. He said he was told informally that this was the case again this year. He noted of the 71 staff members, only six (6) are administrative positions, and that he couldn't maintain services without the current administrative personnel. He also supported the creation of the SSSO, and urged consideration of employee wages at the state matrix for the organization.

(31b:035) Florence and Patti Massey, mother and daughter, testified that the DD system is working well for them, but that it is just not extensive enough to be covering everyone that needs services. Mrs Massey explained that Patti had an inutero stroke three (3) weeks before birth, and she received no early intervention or support until she was eight (8) years of age. Mrs Massey stated it was a fight even then to get her into the school system. She said Patti lived at home until she was 24 years of age, at which time she entered TLC. Mrs Massey also noted that Patti has attended the Billings Workshop since she was 21. She expressed her support for a tax to cover expansion of DD services in the state, including a sales tax.

(31b:080) Randy Thomas, Billings, read his prepared testimony (exhibit 14), in support of DD services. His four (4) year old daughter, Paula, has numerous physical disorders. Mr Thomas expressed his gratitude for the services being provided in the state that enables his family to care for Paula in their home.

(31b:099) Jerry Sherman, Billings, stated that for the past four (4) years she has managed the Pizza Hut restaurants in Billings, during which time she has worked with four (4) clients placed by Vocational Placement. She said that in 1983 Art Gunther, then president of Pizza Hut, Inc., visited the Billings businesses and met three (3) of the clients employed at that time, and was very moved by what he saw. She added that she had just received word that Pizza Hut Inc. is committing nationally to have a position available in every one of the restaurants for developmentally disabled individuals. Ms Sherman stressed the employers need for the professional services of the trainers for themselves, the staff, and the people placed in their businesses as well as a working relationship and partnership with vocational placement for the disabled.

(31b:142) Larry Sherwood explained the services he had received through the DD programs. He said he currently is

living independently, doing his own cooking and laundry, and receiving some support services. He said he was not dependent on any assistance and working at Goodwill in Billings to support himself.

(31b:164) Judy Fisher, Laurel, related the difficulties and heartbreaks her family faces as her son, now 20 years old, is unable to find employment. Mrs Fisher stated her son is a hard worker, can drive and is capable of working, but due to his handicap he has not been able to find employment. She related a very difficult episode they experienced when her son, upon arriving at her husband's employer for a part time job, was told the company did not hire people like him. Her husband has spent several weeks preparing him for this job, telling him what to expect, and it was a very traumatic experience for him and the family when he was rejected. She stated he is getting depressed and lonely, he has no social life or friends he can talk to, and she expressed the frustration of not having any avenues or programs to help him reach his potential.

Chairman Winslow expressed the committee's appreciation to those who had come to testify, and their understanding of the difficulty of talking about areas so close to them.

(31b:234) Sue Kaphammer-Myers, Stevensville, is the mother of an 11 old month son, Joshua, who is a down syndrome baby. She stated that with the services they are receiving from the Comprehensive Development Center (CDC) her son is developing and progressing. She added that they only had to wait four (4) months for services. She asked for support of funding for early intervention and an ongoing commitment to Montana's DD programs, and encouraged tax increases necessary to meet the state's responsibilities.

(31b:294) Bill Sirak, president, Easter Seal Society, Goodwill Industries of Montana, read his prepared text (exhibit 7) in support of community based services and the proposed SSSO.

(31b:353) Lura Nesmith, Special Training for Exceptional People (STEP), Billings, submitted exhibit 3, which compares the salary and benefits of the STEP direct contact employees versus school district two (2) classroom aide employees and covered the discrepancies between the programs. She stressed the need for compensation for the direct contact workers and the dissatisfaction felt by those employed as direct contact employees for the services that needed to be provided and the compensation received. She shared the responsibilities, trauma, and involvement of the workers with those they care for. Ms Nesmith stated incentives were needed to be worked into the budget, and that in six (6)

years 31 individuals have come and gone in the STEP program; most to get employment that offered benefits. She closed by thanking the members of the committee for serving on the committee and for the committee's past efforts.

(31b:455) Richard Harris, Missoula, presented his written testimony (exhibit 10) in support of services for those who are unserved. Mr Harris has a daughter who is severely brain damaged and profoundly retarded, and who is currently on a waiting list for services. He expressed his frustration and grief over not being able to find sufficient services for this daughter, and stated she is an appropriate candidate for specialized family care, but that this services has a three (3) year waiting list for services. He closed by endorsing a tax increase for funding of these services.

(31b:516) Chris Calhoun, Butte, testified that her two (2) preschool children have received services and that her son, George Christopher is developmentally delayed. She expressed the need for family services and early intervention, and asked for support for full funding of services.

(31b:542) Katharin Kelker, Parent's Let's Unite for Kids (PLUK), read her prepared testimony (exhibit 4) on the federal financial support available to states for early intervention programs for handicapped infants and toddlers. She stated that under Public Law 99-457, Montana may request a block grant for \$244,000 to plan and organize the implementation of early intervention programs for children 0-2 years of age. Ms Kelker stated Governor Schwinden has already designated the DD division as the lead agency for the state. She concluded by stating this was a window of opportunity for receiving federal support for early intervention services and asked for support of this grant.

(31b:592) Reverend John Cozby, Anaconda, with his infant daughter, who is a down syndrome baby, spoke in support of early stimulation, intervention, and home trainer services and the importance of help to meet the needs of the children and the families. He stated his family doesn't want institutionalization for their daughter, but they are unaware of all of their daughter's special needs. He stated their family needs support to maintain her in the home, and that these services and programs are cost effective in the long run in comparison to institutionalization.

(32a:016) Pat Lucero, Billings, read her prepared text (exhibit 21) concerning her two (2) year old son who has cerebral palsy. She said Tyler is receiving services through STEP in the form of trainer services, respite, a home teacher and adaptive equipment. She asked for

consideration of the children in need of special services like her son. She also thanked the committee for the services provided to her family that have made their adjustment of raising a handicapped child easier. She concluded by asking support of increased funding to maintain DD services.

(32a:054) Paul Odegaard, presented his prepared text, exhibit 5, in behalf of their 13 year old daughter who is severely mentally retarded. He said he felt they were fortunate in that they had been involved in many of the new programs implemented in the state for DD individuals, including early intervention and handicapped center services. He expressed how hard it was as a family to grow up and cope with a handicapped child. He added that respite care was the first chance they had to become a family again, and how important a couple of hours a week or two (2) weeks was to them. Mr Odegaard stated his daughter is currently in a group home, and since placement he has seen tremendous changes, and he noted his tremendous are very important little steps. He complemented the fantastic job done by dedicated group home workers and vocational placement staff. He stated he is an employer and uses DD employees, and that they are fantastic workers, with supported work a very important segment of DD that is working very well. He thanked the committee for expansion money from the last session, especially for the autistic group home which is doing very well. He said the autistic families who have had such a rough time over the years are finally feeling a little relief.

(32a:142) Sylvia Danforth, director of DEAP, Miles City, covered respite care and its importance to the families, allowing children to remain in the home. She noted DEAP provides 37.5% more services in the respite program while the budget increase for direct respite care increased only 23%, which translates into less money per family served. She stated the actual statewide average reimbursement for respite is \$1.09 an hour, and that it was difficult to find qualified individuals to provide care at this rate. Ms Danforth stated that families who must pay a much higher hourly rate for skilled care must decrease the number of hours they have available to them and/or less options. She said adequate funding would allow families to care for their DD children in their homes. She concluded by stating respite helps provide a cost effective and positive environment for most of the handicapped children being served.

(32a:168) Linda Zermeno read her prepared testimony (exhibit 16) in support of DD services. She said she was a single parent of two (2) sons aged 11 and eight (8). Her oldest son, John Paul, is autistic and currently resides in the REM

Montana group home for autistic children in Billings. She stated she strongly agreed with and supported previous testimony. She also expressed her gratitude for the funding for the services that are currently being provided. Ms Zermeno closed by endorsing a tax increase in order to maintain the current services and address the needs of those DD individuals still on waiting lists.

(32a:204) Cathy Peterson, Great Falls foster parent, representing specialized foster care, described the children served and services provided under this program. She talked about Jill, who was diagnosed as having cerebral palsy, and then rediagnosed with a deteriorating brain disease. Her health deteriorated rapidly, and with the support of home trainers, physicians, child and family care workers, nursing care and respite, Jill was able to remain in her home until the time of her death. She felt it was a big achievement to serve this child in a home environment versus two (2) years of hospitalization. She said after Jill's death last June, another child who is blind and microcephalic was placed in their home and is receiving many of the same services Jill had the opportunity to receive.

(32a:265) Chuck Super, Billings, asked for support for funding for providers and their staff, endorsed the SSSO, and the proposed reorganization of SRS.

(32a:330) Chairman Winslow then addressed the audience and expressed appreciation for everyone's efforts and that the committee understood the emotions that were felt when dealing with these issues. He noted, however, that the committee deals with these issues across the board in a lot of other areas as well. He stated the unfortunate situation is that to date, the financial situation has not changed, and that the public does not seem to be in support of any kind of tax increases. He said as some have testified in favor of the sales tax proposals, but at the present time sales tax proposals will not provide any general fund money, but are in fact replacement taxes for property tax relief. He noted a 20% property tax reduction translates into a \$400 million deficit. He said these facts are to bring everyone back to where the state is at. He stated the need to look at the priorities, reorganize, and in some cases, make people more accountable in the human services areas. He pledged efforts to do the best job possible in the next few days under very difficult situations.

The meeting was adjourned to room 108.

Eligibility Determination

Peter Blouke, LFA, covered the budget and issue sheets concerning the eligibility determination program.

(32a:403) Dave Lewis stated one of the mandates of putting the new department together, from the governor's office, was that this was to be a no increase cost situation. A problem arose with the salaries for county directors and clerical support staff, who are funded 20% from social services and 80% from eligibility determination. The eligibility determination program absorbed the 20% of the salaries from social services to make the transfer to the new department on a revenue neutral budget basis. He said the difference on the budget issues is that the 9.24 FTE in the executive needed to have 15.8 FTE added on to have the existing positions funded 100% from the eligibility determination program with no net increase in cost. He stated the current level FTE for eligibility determination was 385.6 FTE, and that he had agreed to absorb \$300,000 in vacancy savings to comply with the concerns of keeping the new department at a no cost increase level. After a budget review by the committee and Gene Huntington, it was determined that all of the money taken out for funding the new department was not needed, and Mr Lewis asked that the money cut from the budget be returned.

Lee Tickell, administrator of the Economic Assistance Division (EA), SRS, stated this budget covered staff and travel for both the state and non state administered counties, i.e. allowing directors and staff to travel for training purposes, home visits and investigations.

In response to an inquiry from Sen Himsl, Mr Lewis clarified that \$7.3 million is raised by the 12 mill levy from the counties and the actual cost of those services is \$20 million, with the state putting in \$13 more than they are receiving in reimbursement. He noted that for ease of accounting, the 12 mill levy is placed in the medicaid program.

(32a:650) Norm Waterman, director, Lewis and Clark County Welfare, covered some of the problems in the county that cause people to complain about service delivery and to give his support to the proposed FAMIS automation system. He stated the agency is a paper oriented agency with dozens of forms requiring handling by four (4) or more individuals for completion. If there are any errors in the process and a client utilized medicaid services without the proper information being in the system, payments for the bills submitted by the providers are rejected. He noted the stress this creates on staff, providers, and the clients needing services. Mr Waterman stated the agency could not handle an increase in the caseload with the current delivery system,

and that automation would alleviate the multiple handling and diminish the error rate. He stated he would never say automation would reduce the need for staff in future years, he said it might cut the need, but as caseloads grow there will probably be a need to increase staff to handle those caseloads, but that with automation paperwork processing would be faster and could be done more effectively. He concluded by stating his staff are at the saturation point, and without the pilot automation program currently in place, they would be buried in the paperwork.

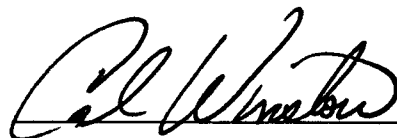
(32b:052) Wanda Stout, director, Jefferson County Human Services, read from her prepared statement (exhibit 11) in support of the FAMIS automation system.

In response to a question from Sen Hims1, Ms Stout clarified that on the pilot project their agency has inquiry access only into the state system, which they pay by the length of time on-line.

(32b:106) Chairman Winslow asked if there would be any opportunity to reduce staffing when computers are implemented, and, as testimony has indicated, it will take less time to process the claims and paperwork.

Ms Stout stated that tasks are prioritized and those at the bottom of the list are not getting done due to the workload, and with automation those tasks could be completed.

The meeting was adjourned at 10:45 a.m. (32b:170)

A handwritten signature in cursive script, appearing to read "Cal Winslow", written over a horizontal line.

Cal Winslow, Chairman

HUMAN SERVICES SUB COMMITTEE

Date

February 4, 1987

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Name: F. Lee and Marianne Swartz
Address: 2517 Briegleb St
Missoula, Montana 59803

People:

As the parent of a mentally handicapped child I feel I must express to you my concerns over the upcoming budget changes concerning Social Services for the disabled. As a former recipient of some services rendered by the Western Montana Comprehensive Development Center (CDC) I am acutely aware of the necessity of early education for the developmentally disabled.

My daughter Rhannon is a five and one half year old developmentally delayed child. At age twenty-six months, with the recommendation of our family physician, we enlisted the services of CDC to help determine the best possible course of action to take in order to give Rhannon the opportunity to become her fullest and most productive self. At age thirty months, we began a process called Home Training, to teach we the parent and the child herself, how to care for her, meet her everyday needs, and give her the best possible start to an absolutely necessary early education.

At this age (30 months) Rhannon had no intelligible speech, her gross and fine motor skills were vastly underdeveloped. She had only begun to develop toilet skills. She could not hold a spoon, was barely able to grasp a cup and it was extremely tedious to dress her because she had neither the physical capabilities to dress herself nor the mental or verbal capabilities to explain a way to help us help her. She was withdrawn and mechanical showing little emotion other than occasional outbursts of frustrated anger. If she was sick she had no way of telling us and there were no outward signs. If she was hungry she did not express it. All these things, you and I as "normal" people, take so very much for granted. We assume growth as a matter of fact. Rhannon was a

part of our lives, but lived in her own world and her father and I had no idea how to enter that world or how to invite her into ours.

Rhiannon is now five years, eight months old. She is a bright, beautiful child with huge, dark, eyes that light up with life, hope and best of all curiosity. She has received speech therapy physical therapy and occupational therapy all included as part of her special education. Her speech, though not equal to that of her chronological peers, is colorful and understandable. She walks, runs, jumps, plays, sing Sesame Street songs at 135 decibels, and causes mischievous trouble capable only from a child with intelligence enough to know she's garnishing a reaction! This mind has been unlocked to a world she may have never realized without the help from C.D.C.'s most important services.

Rhiannon is still developmentally delayed, but the gap between her and her peers is slowly closing. Without early intervention by the skilled and concerned disability providers of C.D.C., we most assuredly would have had to make a painful decision concerning the future of our child. An institution would

surely have been more expensive not only in financial terms but in the terms of the loss of dignity of a human being.

Rhianon will most probably need the continued services of State disability services. In order for her to become a productive member of society instead of another burden on the welfare system she will need many more years of special education. Through careful planning and work she will be an asset versus a liability to herself and her community.

I want my daughter to be able to provide for herself, not depend on the government for her desires and needs. I want my child to be proud to a part of this state and to be able to continue to say "All by myself! I did it, Mom! All by myself!" We need these services in order to see this hope realized. The money you save now will be nothing compared to the money it will cost to support the uneducated children of today, tomorrow. Please reconsider the absolute necessity of services for the disabled. It is important to the pride and dignity of a community of people who because

2.4.87

they are labeled different or disabled, they are not considered viable parts of society. They are a part of us and with our help and support they will be a valuable part of our world as is the life of any human being.

We reap what we sow. Let us nourish beautiful flowers, not thorny weeds.

Thank you.

Sincerely
Marian G. Swartz

EXHIBIT 22
DATE 2-4-87
HB _____

Final Report
Developmental Planning Task Force

December 1986
Helena, Montana

JANUARY 1987 SALARY AND BENEFIT COMPARISON:

STEP Direct Contact Employees Versus School District #2 Classroom Aide Employees

	School District #2 Special Education Classroom Aides	STEP Group Home Direct Contact Employees	Recommendations
1. Training Requirements	None	A. SRS Medication Administration Certification B. Montana Chauffeur's License C. CPR Training D. American Red Cross Certification	
2. Pay Scale			
0-3 months	\$4.69	\$4.30	
3-6 months	5.16	4.73	
6-9 months	5.16	4.85	
9-12 months	5.16	4.96	
12-18 months	5.27	5.08	
18-24 months	5.27	5.08	
24-30 months	5.40	5.40	
30-36 months	5.40	5.61	
36-48 months	5.51	5.82	
48-60 months	5.60	6.03	
60-72 months	5.73	6.24	
72-84 months	5.84	6.45	
3. Holiday Pay	All Holidays off without pay.	Double time on holidays worked.	
4. Overtime Pay	None	One and one half times base salary for hours worked over 40 hours per week.	
5. Vacation Time	None	Paid vacation after 18 months continous employment equal to the average number of hours worked in the previous 6 months, not to exceed 32 hours. (Averages to 2.6 days per year maximum)	

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Name: KATHARIN A. KELKERAddress: 2210 FAIRVIEW PL.
BILLINGS, MT 59102
252-7596 or 657-2035

The Developmental Disabilities Division has an exciting opportunity to participate in the benefits of P.L. 99-457, federal legislation that provides financial support to states with early intervention programs for handicapped infants & toddlers. Under P.L. 99-457 states may request a block grant (\$244,000) has been allotted to Montana) to plan for the implementation of early intervention programs for children 0-2. To receive these monies, a state must designate a lead agency, constitute a coordinating council of 15 members and, by 1990, have in place mechanisms for identifying, evaluating and providing case management to handicapped infants & toddlers.

Governor Schwendener has already designated the Developmental Disabilities Division as our lead agency. Now the legislature needs to authorize the DDD Division to request the federal funds and begin the planning for early intervention programs. Participation of Montana in this planning effort will require no general fund money during the next biennium.

We have a "window of opportunity" for receiving federal support for our early intervention services. It makes good sense for us to take advantage of this valuable assistance. I hope you will support the DDD Division in this worthwhile activity.

EXHIBIT 5
DATE 2.4.87
HB _____

To: Honorable Members of Joint Human Services Committee of
the Appropriation Committee
Re: Hearing on the DD Budget
From: Paul and Ande Odegaard, Parents and Taxpaying Voters

We are the parents of a thirteen-year-old girl who is severely mentally retarded. After many years of trying to cope with her on a 24-hour basis and using the services available to us, we had her screened for a group home and she was accepted in March of 1986. Her progress has been steady, she is beginning to become more independent in self-help skills and her behavior is under much more control. Personally, we as a family are more productive citizens because the stress level has decreased measurably.

Today we are writing about the possible cuts in funding of the DD budget for the future. We realize that we have economic problems within this state, but please don't take it out on our DD population who have gained so much in individual growth in the past few years since deinstitutionalization was begun. In fact, in order to maintain the services we have for those persons who are not on the waiting list, we ask for a reinstatement of the 3% increase for '88-89 fiscal year.

A decrease in funding for DD programs would mean a cut in direct care staff for the clients. In the past two years there has already been a cutback in the amount of funds for the DD division. We cannot handle any more additional trimming without seriously jeopardizing the programs in existence. In our own situation, the cutback will undoubtedly result in the loss of stable employees in our group home. These people have not received a pay raise for two years. Their knowledge and experience of how to cope with our DD individuals is invaluable. You cannot put untrained, inexperienced persons immediately into a group home and expect them to understand how to control behaviors, and teach basic skills such as brushing teeth, combing hair, toileting, walking, and talking. Could you do it? Could you do it without training? Would you have the knowledge and patience to break down the training of a simple task such as brushing teeth into 13 steps and repeat the training of this task daily for unknown years? It is because of the trained staff that our individuals have made positive gains toward becoming independent. Our people need to be in constant training- maybe at a lower level on certain skills that they have "learned", but there are new skills being taught all the time.

High on the priority list for elimination in several group homes is transportation because of the high cost of liability insurance for their vans. If this budget item gets higher and your budget cut happens, it will be necessary to cut out transportation - then our people become prisoners of

EXHIBIT 5
DATE 2.4.87
HB _____

their own group home. We know of families with a DD person in their home that experience enormous 24-hour stress, consequently making the adults and children unproductive in their day to day activities. We were one of these families. Respite care and specialized family care were important help for our family and is a critical need for many families. Please don't let these programs be cut back.

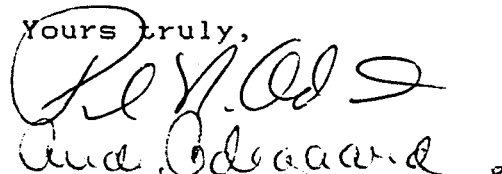
We appreciate the support you have given DD individuals in the past. We must take pride in the deinstitutionalization of Boulder and giving these people a chance to live in their own communities. Everywhere you look in Billings you have a DD individual working or participating in "normal" activities. Their life is so much more meaningful and it is certainly a lot cheaper then institutionalizing them.

We urge you to oppose the proposed budget cut and ask that you fund 3% above the executive budgeted level. Both the families and our DD individuals have a very tough time making it in this world- the proposed cut is going to make it that much harder. Do you want to take responsibility for their inability to become productive citizens of this state? Through programs such as supportive employment they can earn money, spend it, and pay taxes as well. They will return to society what they receive from society.

We also ask that you please work toward restructuring the tax structure of this state so that we don't have to go through this struggle every biennium. The services for DD clients are a necessary part of the total committment of our state.

If you need any help with convincing other legislators to understand your positive attitude towards funding of the DD system, please call on us to help you. As the Yellowstone Boys and Girls Ranch says, "It really makes a difference when you know somebody cares."

Yours truly,


Paul and Ande Odegaard

3238 Parkhill
Billings, MT 59102

NAME: Stephen J. Wong DATE: 2.4.87

ADDRESS: 1302 391 *Washington*

PHONE: _____

REPRESENTING WHOM? *[Signature]*

APPEARING ON WHICH PROPOSAL: Human Service Appropriation

DO YOU: SUPPORT? X AMEND? _____ OPPOSE? _____

COMMENTS: _____

PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.

EXHIBIT

DATE

7
2.4.87

Comments Supporting

A SPECIALIZED SERVICE AND SUPPORT ORGANIZATION

William N. Sirak

President

Easter Seal Society/Goodwill Industries of Montana

Members of the Appropriations for Human Services Subcommittee.

My name is Bill Sirak. I am the President of the Northern Rocky Mountain Easter Seal Society/Goodwill Industries of Montana. My organization has been involved in providing direct services to people with disabilities for more than 40 years. I have personally been involved with the organization as its President for the past 15 years. It is an organization that currently provides services in Montana, Wyoming and Idaho. We have observed that the State of Montana has consistently provided a leadership role in encouraging, developing and coordinating an outstanding community based service delivery system for people with disabilities. The Governor's office, his administration and the Montana legislature must be commended for their leadership role in this area.

The Montana Easter Seal Society has been a partner with the State of Montana since the beginning of the deinstitutionalization

process. We are now the largest provider of adult DD services through our adult training facility in Great Falls and our Goodwill stores and plants located in Missoula, Billings and Great Falls.

The Governor's office and Social and Rehabilitative Services have recommended the development of a specialized service and support organization to serve an additional 52 severely disabled individuals in a community based setting. We compliment the SRS staff for bringing to you an extremely well-thought-out plan of action to serve people in their communities. We have found this plan to reflect national trends in a very unique, innovative and cost-effective manner. The SRS staff, and specifically the DD Division, should be commended for their research, study and recommendations in this area.

As a provider of community based services, we share the concerns of many other providers, as well as the legislative body regarding the current financial crisis in the State of Montana and the future funding base for current services. Although we feel that adequate funding for current services must remain a priority, the adequate funding, development and implementation of this proposal must be taken under consideration at this time.

Services to the most severely disabled population is a problem that simply will not go away. If it is not addressed immediately, the problem may become more profound and more costly. We are, therefore, encouraging the development of this

EXHIBIT

DATE

NO

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2.4.87

concept at this time.

One cannot deal with this issue without taking into consideration the practical and political issues centered around the future of the Montana Developmental Center, formerly known as Boulder. Past study groups such as the 909 Commission and subsequent task forces have recommended very specific and appropriate functions for the Montana Developmental Center and East Mont. The development of this project, therefore, should not be viewed as a competing or threatening project to those legislators and residents of Boulder and Glendive.

The fact remains, however, that national accrediting standards and national trends pose a very real threat to the way these two institutions currently provide services. It is important to note that the threat of loss of certification is not a reflection of the quality of care provided at those institutions, which I personally feel is excellent and beyond reproach. Rather, it is a reflection of a national trend toward community based services.

The proposal presented to you regarding the development of a specialized service and support organization is timely, reasonable, cost effective and deserves your immediate and serious consideration. If it is not dealt with now, we will deal with it at some future time with a greater dollar expenditure.

PARENTS OF SPECIAL KIDS

EXHIBIT

DATE

8
2-4-87

OFFICE BOX 6760

GREAT FALLS, MONTANA 59406

February 2, 1987

The Honorable Cal Winslow
House of Representatives
Capital Station
Helena, Montana 59620

Dear Representative Winslow and Members of the Committee,

The Committee on Human Services will be considering ^{Expansion of} ~~to mandate~~ Developmental Disability services. I strongly support this service and feel this service is in great need to continue the future service of this program.

My daughter Cathy, who is 2½ has been with Region II Child and Family Services for 2 years. Cathy was without oxygen at birth and has cerebral palsy. Cathy is my only child and I wish each of you could meet this darling bundle of life. If you don't have a handicapped child, it's hard to understand what happens when a handicapped child enters your life. In the two years we have been with Child and Family Services I have seen Cathy make some excellent progress by their service they have given us. You don't know how thankful I was to find there was such a service. The hardest part of having a handicapped child is being told you have one. By no means is the rest easy, but with the help of my Home Trainer, Terri Jones, we have been able to learn so much on how to help not only Cathy but ourselves as well in living with her disability. I thank you for the support you have given his program, as without it I would be lost in a world I know nothing about.

As you know the cost of raising a child is expensive, but the cost associated with a handicapped child is greater. There is extra medical costs and adaptive equipment that are part of the increase. As with everything the cost goes up every year and it becomes harder for this organization to continue their service. I would greatly appreciate you supporting a rate increase for this agency so they continue to help other families as they have help me.

Thank you for your time and consideration of my views.

Sincerely,

Della Sandrock

Della Sandrock - Great Falls, Montana

My name is Susan Kaphammer-Myers; my husband, child and I live in the Stevensville area. I am a high school librarian, my husband is a chemical dependency counselor, and we are both life-long Montana citizens. I am asking for support of reinstatement of full funding for programs for the developmentally disabled.

Our eleven month old son, Joshua, is a Down's Syndrome baby. A generation ago he would have been institutionalized for life. But today he is a healthy, happy child who, thanks in large part to services we receive from the Comprehensive Development Center in Missoula, is progressing in motor, language, and cognitive skills. Joshua has received two detailed evaluations by a speech pathologist, a physical therapist and a psychologist. A home trainer, through weekly visits, provides a developmental program for Joshua. My husband and I could not alone provide for our child these services that are so crucial for his growth.

I understand that developmental disability services suffered a funding freeze as did most state programs. A continued freeze, let alone a cut-back of funding, could well mean staff reductions as increases in fixed expenses such as liability insurance and worker compensation insurance cut into their budgets. An increase over the next two years will be needed to maintain current services, even without salary increases. Staff reductions could mean, for example, longer waiting lists for services (we waited about four months for services for Joshua), and bi-weekly or monthly training sessions rather than the weekly sessions that keep Joshua's development progressing.

I am sure all legislators receive pleas for funding that combine to amounts far beyond the resources available. Knowing this, I don't know how to stress enough how services for the developmentally disabled make sense, not only for humanitarian reasons, but also for financial considerations.

It costs far less to provide home training such as Joshua receives than it costs to maintain a person in an institution. In addition, because of such services, the developmentally disabled can become self-sufficient to varying degrees, and even become tax payers, rather than tax burdens. Joshua's future holds the possibility of independent living and a productive lifestyle, because he is now receiving special help.

Money is tight; but programs for the developmentally disabled save state monies in the long run. Services cost money, and I am willing to pay. I would actively encourage support for a state tax structure, including increases, necessary to meet the state's responsibilities to all Montana citizens.

Thank you for your time and attention.

Susan Kaphammer-Myers
344 Sky Pilot Lane

NAME: Richard B Harris DATE: February 4 1987

ADDRESS: 808 Woolford Missoula MT 59801

PHONE: (406) 728-9156

REPRESENTING WHOM? self

APPEARING ON WHICH PROPOSAL: _____

DO YOU: SUPPORT? / AMEND? _____ OPPOSE? _____

COMMENTS: Montana has a group of qualified, dedicated
service providers who are presently underfunded. Our daughter,
Ellen B. Harris, profoundly retarded, currently sits on a long
waiting list for services. My wife and I are currently considering
moving to another state, simply to obtain necessary services for
our daughter. The state's services are good but inadequate.

PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.

EXHIBIT 14
DATE 2.4.87
HB

P. O. Box 836
Boulder, Montana 59632

I am here to speak in favor of the FAMIS automation system.

In the Jefferson County office, we have been a part of the S.R.S. pilot automation project. In this project, we have utilized a personal computer with various software and have had on-line inquiry to the Employment Service Wage Match and the S.R.S. mainframe.

Even though our capabilities through this project are considerably more limited than the FAMIS system would offer, we have saved considerable staff time in various areas. For example; monthly we are required to issue food stamp authorization cards. A duty that manually took at the least a full 8 hours of staff time. This task now takes 20 minutes. This is only one example of the many staff saving areas that can be utilized, reducing the need to increase staff.

With on-line direct input by workers, days of turnaround time can be saved and errors can be edited and corrected immediately. By having instant statewide search available, double-dipping and fraudulent reporting can be squelched at the time of application. Something that can go on for months before discovery under the current system.

In addition to saving staff hours, discovering the fraudulent applicant, and reducing errors, we would be able to provide a much faster service to the needy. We would no longer require days of mail turnaround time to acquire verifications, etc. Medical providers would also find their payments expedited as they could bill immediately rather than wait two weeks or more for our information to clear the current input system.

Thank you for hearing this testimony. I appreciate the opportunity to speak to you and hope that you will contact me if I can be of any further service to you.

Respectfully submitted,


Wanda Stout, Director
Jefferson County Human Services

TESTIMONY SHEET:

EXHIBIT 12
DATE 2.4.87
HB _____Name: Irene Olsen
Address: 318 Windsor Circle
Billings mt. 59105

Feb 3/87

I Irene Olsen am writing in behalf of Jane A. Wanner who resided in Boulder 23 yrs.

Since leaving Boulder Jane was in several Foster homes in Billings. Resided with Mrs. Sarah Tipps for 10 1/2 yrs.

Jane has been living with me, Irene Olsen, Jane's Natural Mother since May 26th 1983.

Jane has really improved due to the Programs. She does beautiful Latch Hook Rug. She worked at The Billings Sheltered work shop for a number of years, but for some reason got bored and didn't like working there anymore.

Jane worked at The Pizza Hut on Billings Heights with Steve Abdallah for 2 1/2 weeks, was doing real good and enjoyed her work, but developed a serious allergy to Soap, she was very disappointed about this and maybe that's why she didn't want to work at the work shop anymore.

She was transferred to The Senior Activity Program and is very happy there and is progressing.

Jane is now 46 yrs old and will be needing programs for a lot of years. One of my wishes is to have a Sr. Group Home in Billings.

There are a lot of Older people in her Sr. Activity program and they need a Sr. Group home also.

The Sr. Program is more leisurely which is better for them as they are no different than we are, except that their aging process comes earlier in life.

They're enjoying their retirement or near retire-

ment.

We appreciate her programs and her progress
but every program she has there must be
enough staff and suitable wages to cover
the staff.

Sincerely
Irene Olsen

Name: DIANE SAYLER
Address: Box 261
Red Lodge, MT 59068

Mr Chairman, members of the Committee. For the record my name is Diane Saylor. I live in Red Lodge. I am the mother of a 5 year old developmentally delayed girl Sarah has been diagnosed as severely mental retardation and without Special Training for Exceptional People, she or I, would not be nearly as far as we are at this point in time. Step has helped me not only in teaching basic self-help skills, but also helped me to cope and deal with behavioral problems special children sometimes develop. Without Step and their support and guidance I would have felt alone and isolated, in trying to help my child grow to her full potential.

Sarah has a speech impediment, seizure disorder, left frontal temporal lobe damage resulting from placenta deficiency. Because of our child's handicaps we face more emotional and marital stress and difficulty finding quality child care.

A proposed 4% increase will help cut the inflation rate and help us to at least maintain programs. To freeze the budget will mean cutting our home trainer services and respite even further. Our home trainer from Billings has a wider area to cover and less time in order to do her job.

We need the services of Step and Respite in our community. My daughter and all special children deserve equal opportunity to develop to their full potential. Without your support our kids may never know the feelings of self-respect obtained through personal accomplishments.

In order to have a progressive state we must have justice for all peoples.

For the past services that have helped me and other parents of handicapped children, I wish to sincerely, Thank you.

TESTIMONY SHEET:

14
2.4.87

Name: Randy Thomas
Address: 3142 Yaleth St. S.W. R.R1
Billings, Mt. 59106

Mr. Chairman, members of the committee for the record my name is Randy Thomas I live in Billings.

The DD system is important to me because my daughter Hollie Ann is handicapped. She is 4 years old. She has a cleft lip & palate, cerebral palsey, hydrocephalus, a seizure disorder and chronic aspiration pneumonia. We receive respite care, family training, specialized family care and preschool. With Hollies many serious physical problems and with the severity of her handicap without these services we would be forced to institutionalize our little girl. We are grateful for all the services we have received and hope they may continue and maybe expanded to help other families that have problems similar to ours.

Thank you

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2.4.87

Name: Bruce Owens
 address: 8025 N.E. Bliss Blvd. #70
Billings, Montana 59101

Mr. Chairman for the record my name is Bruce Owens from Billings.

I'm the mother of a 24 year old Down's Syndrome man, I'm also the Pres. of the Montana Assoc. for Retarded Citizens.

Mark lives at home and is employed at the Billings Workshop.

Any more freezes or decreases in the D.D. budget will affect him greatly. With out an increase more staff will have to be laid off and that will mean more idle or down time because there wouldn't be enough trainers to work with him or job productivity increases.

I have talked with other parents around the state that have had their children on waiting lists for 2 years after getting out of Spec Ed. More will be graduating this spring with some or little job training. Those that have had a lot of training are unable to find jobs.

The need is so great I have more funding to keep the programs going & I create new ones.

I'm the handling director for 72 special persons in Bliss. Let handling is a big light for them.

Many live in group homes. Until recently they arrived at the Bowling alley in vans provided by the Group Home Corp.

When the budgets were frozen and the cost of insurance went up the insurance co. limited the driving to certain trainers.

And some group homes must use special transportation & some handling. After handling they go home. At that time stay until Monday AM when it's time to go to work unless a trained person from another home can take time to take a group in on meeting.

Not many years ago when the funding ^{was} ~~got~~ low the ~~week~~ staff was 1 trainer for 8 clients.

Can you imagine what you would do if you were in charge & 1 person had a bad seizure - no one to call for help and a client would panic & run out the door.

The staffing got better. Two trainers & most group home & some have 3 trainers for the more intensive training homes.

If there isn't an increase there may be some staff that would have to be laid off or have a wage cut.

I greatly appreciate the programs & help that Mont. has given our son and others. Your investments have paid off but he will still need a group home in the future.

I realize we have made great strides in DD programs but we must not go backwards.

I would be willing to pay more taxes for the programs.

All of Montana's D.D. persons are looking to you for their future.

Representative Cal Winslow and members of the Committee,
My name is Linda Zermeno and I'm from Billings 16
DATE 2-9-89
ID

I am a single parent of two sons age 11 and 8. My oldest son John Paul ~~has~~ has the problem of Autism and currently resides in the REM MT Group Home for Autistic Children in Billings. I thank God that you recognized the needs of Autistic Children and supported our cause by appropriating funding for this Group Home.

Prior to his move into the Group Home in August, John Paul had received services from Respite Care, Family Training and Specialized Family Care. We've been very fortunate.

I agree with what others have asked for during this hearing. I especially ask your support of ~~the~~ ^{the} ~~in~~ ⁱⁿ ~~stat~~ ⁱⁿ ~~ment~~ ^{ment} in fiscal years 1988 and 1989.

I would support tax increases in order to maintain current services and address the needs of those DD individuals ~~and~~ still on waiting lists.

Our kids have truly benefited from services provided ~~the~~ and it would ~~be~~ ^{have} be a tragedy, indeed, for them to lose ground.

I thank you for your time and concern.

Summary of Recommendations
Developmental Planning Task Force
December 1986

- #1 The Task Force recommends the consolidation of all services for persons with developmental disabilities under the same administrative authority.
- #2 The Task Force recommends consideration by the Legislature of a joint resolution of support for and policy commitment to supported work services for Montana's citizens with severe disabilities.
- #3 The Task Force recommends that programs be established to fill gaps that exist in the array of DD services. To meet the needs of Montana's unserved and underserved citizens with developmental disabilities, the following programs must be developed: 1) specialized service and support organizations, 2) supported living, 3) adult congregate living and 4) group homes. In addition, new and perfected programs must be established to serve geriatric citizens with developmental disabilities and those with intense medical and behavioral needs.

The Task Force views Montana Developmental Center as an integral part of the array of services and recommends that it serve the following specific missions: 1) residential service provider for persons with severe behavior problems, 2) residential service provider for naive offenders, 3) residential service provider for persons with severe medical and/or care needs, and 4) professional resource for community-based DD service programs.

The Task Force views Eastmont Human Services Center as an integral part of the array of services to individuals with developmental disabilities and recommends that it specifically serve as an exemplary geriatric program for Montana's senior citizens with developmental disabilities.

- #4 The Task Force recommends improvements in community services in the areas of case management, respite care, and staff training, and the establishment of an additional service component--independent reviews of placement and treatment.
- #5 The Task Force recommends that the Developmental Disabilities Division of the Department of Social and Rehabilitation Services be designated as the lead agency for a new state grant program for handicapped infants and toddlers.
- #6 The Task Force recommends passage by the Legislature of a joint resolution supporting increased public awareness of naive offenders and the establishment of policies and procedures to identify and treat them.

Joseph L. Carter

EXHIBIT 11
DATE 2-4-87

NAME: Nannie Koenig DATE: 2/4/87

ADDRESS: 430 Church Dr. Kalspeel

PHONE: 752-3370

REPRESENTING WHOM? Parent 7 Lakeland Industries for the Handicapped, DD PAC,
Region V DD Council, 7 Lakeland ARC

APPEARING ON WHICH PROPOSAL: _____

DO YOU: SUPPORT? yes AMEND? _____ OPPOSE? _____

COMMENTS: Thank you for your continuing concern & support
for services for the developmentally disabled.

PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.

18
2487

PHONE: 1-18-9658

APPEARING ON WHICH PROPOSAL: Human Services

COMMENTS: YFP international reading

PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.

DATE: 2/4/01

PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.

NAME: Dorothy A. Schack DATE: 2/14/87ADDRESS: 14501 Alexander Rd Billings MT. 59105PHONE: 259-6666REPRESENTING WHOM? Developed Disabled Montell's people at MontAPPEARING ON WHICH PROPOSAL: Funding D. D. ServicesDO YOU: SUPPORT? X AMEND? _____ OPPOSE? _____

COMMENTS: My heart went out to the parents & others who
so bravely testified this morning. I totally relate the heartbreak
each have endured in raising a handicapped child. I have lived
through it all, I have a multiple handicapped son, now 37 years old
He was an R. H. baby which caused Cerebral Palsy, deaf-
ness & retardation. After going from the C. P. Center at Eastern
Mont. College to Montana School for Deaf Inc. Falls to
Special Ed. in Billings etc etc etc. my son is also termed a
Diase offender & at present is at the Montana Developmental
Center at Boulder. I can tell you how much good they are doing
at the center - their academic training, behavioral training,
physical ed plus my son has 2 exceptionally fine teachers
who teach Signing for the Deaf. However the center needs
extra staffing & a remedy to help the over crowding in cottages
 PLEASE LEAVE ANY PREPARED STATEMENTS WITH THE COMMITTEE SECRETARY.
Boulder is no longer looked at in the degrading way it once
was, however unless funding is continued the rate of upgrading
the center will definitely be delayed.

Sincerely-

Dorothy A. Schack

Name: Pat Lucero
Address: 744 Petrified Tree Pass
Billings, MT 59111

2-4-87

MR CHAIRMAN, MEMBERS OF THE COMMITTEE FOR THE RECORD, MY NAME IS PAT LUCERO. I LIVE IN BILLINGS. I AM THE PARENT OF A TWO YEAR OLD BOY NAMED TYLER WHO DUE TO A PREMATURE BIRTH HAS CEREBRAL PALSEY. TYLER RECEIVES SERVICES FROM THE DEVELOPMENTAL DISABILITIES BUDGET THROUGH S.T.E.P. (Special Training ~~from~~ for Exceptional People). STEP PROVIDES A HOME TEACHER AND AN EDUCATIONAL AIDE FOR TYLER WHO VISITS TYLER WEEKLY HELPING ME SET APPROPRIATE GOALS FOR MY SON. THIS SERVICE IS EXTREMELY IMPORTANT SINCE OUR SCHOOL DISTRICT HAS CHOSEN NOT TO OFFER A PRESCHOOL FOR HANDICAPPED CHILDREN.

RESPIRE CARE IS ANOTHER SERVICE WE RECEIVE. RESPIRE HAS ALLOWED ME TO KEEP MY SANITY AND HAS GIVEN ME TIME TO SPEND WITH MY HUSBAND AND DAUGHTER.

MY HOME TEACHER HAS HELPED US ACQUIRE SPECIALIZED ADAPTIVE EQUIPMENT THAT IS ESSENTIAL FOR OUR SON'S DAILY LIFE.

IF YOU PUT A FREEZE ON THE DD BUDGET AND DO NOT GRANT A 4% INCREASE WHICH IS NEEDED TO ALLOW AN EXPANSION OF SERVICES, IT IS MY UNDERSTANDING THAT THE END RESULT WILL BE A CUT IN PERSONNEL WHICH WILL EVENTUALLY CUT SERVICES.

THESE SERVICES HAVE MADE OUR ADJUSTMENT OF RAISING A HANDICAPPED CHILD EASIER. FOR THESE SERVICES WE WISH TO THANK YOU.

PLEASE CONSIDER ALL THE CHILDREN OUT THERE LIKE TYLER IN NEED OF SPECIAL SERVICES. I FEEL WE MUST PROVIDE FOR OUR HANDICAPPED, ESPECIALLY CHILDREN WHO CANNOT PROVIDE FOR THEMSELVES.

AS A PARENT I WILL TRY TO DO ALL I CAN FOR MY HANDICAPPED CHILD. MY SON IS LUCKY! BUT WILL THE NEXT CHILD BE SO LUCKY? THE STATE OF MONTANA SHOULD ASSURE THAT HE IS.

VISITORS' REGISTER

HUMAN SERVICES SUB COMMITTEE

BILL NO. _____

DATE

2/4/87

SPONSOR _____

DEPT _____

NAME (please print)	Representing	SUPPORT	OPPOSE
Dave Lenn	S R J	X	
Bob	ADORE, ADPAC	X	
Jim Foster	Chester Schools	X	
Reslie Foster	Chester	X	
Linda Zernano	Billings Hunt	X	
Aude Edgaard	Billings Parent	X	
Wayne & Donna	Billings Stateless Hunt	X	
Cheryl Pully	Billings	X	
Paul M. Olds	Billings	X	
JERRY NISBET	HD 35 Great Falls	X	
Jim Smith	HRDE / MAR	X	
Major Waterman	Mont. Assoc. of Churches		
Sue Fifeild	MCIC	X	
Jo Ann Russell	SELF	X	
Mrs. Mrs. Calhoun	Self	X	
John A. Long	SELF	X	
Myrtle & Cagle	"	X	
Chuck Supers	RSD	X	

IF YOU CARE TO WRITE COMMENTS, ASK SECRETARY FOR WITNESS STATEMENT FORM.

PLEASE LEAVE PREPARED STATEMENT WITH SECRETARY.

VISITORS' REGISTER

HUMAN SERVICES SUB COMMITTEE

BILL NO. _____

DATE

February 4, 1987

SPONSOR _____

DEPT _____

NAME (please print)	Representing	SUPPORT	OPPOSE
Rene Wheeler	Special Training for Exceptional People	✓	
Vonnie Koenig	Parent, Halted Industries, DDPK	✓	
Joan Obermatt	Foster parent CDC	✓	
Jessie Schlegel	foster parent CDC	✓	
Beverly Owens	parent Mont ARC	✓	
Florine Massey	Parent	✓	
Patti Massey		✓	
Tom SEEKINS	Family Outreach, Helena	✓	
Wallace Melcher	Reg II C&FS - Gt. Falls	✓	
Connie O'Seen	Big Sandy Activities	✓	
Sylvia Danforth	DEAP, Miles City	✓	
Cheryl Andersen	Devlop. Planning Task Force, Helena	✓	
LARRY ASHERWOOD	VOCATIONAL PLACEMENT Job Training	✓	
Steve Abdallah	" " " "	✓	
Charles Bantz	Governor's Office		
Dakota Jean Curtis	Sp Foster Care	✓	
Betsy Stalling	Voc. Placement	✓	
John W. Vesscher	Counterpoint - Livingston	✓	

IF YOU CARE TO WRITE COMMENTS, ASK SECRETARY FOR WITNESS STATEMENT FORM.

PLEASE LEAVE PREPARED STATEMENT WITH SECRETARY.

HUMAN SERVICES SUB COMMITTEE

DATE 2/4/87

DEPT _____

NAME (please print)	SUPPORT	OPPOSE
---------------------	---------	--------

Representing

SUPPORT

OPPOSE

August B. Herzog

Rem Mt. two

✓

W. H. Blatt

REM, MT, Inc.

✓

Chas. S. Huss

Ennis, MT parent

✓

Mrs. Anne D. Boyer

Parent, Missions

✓

James S. McSparrin

parent, Missoula

✓

Wine Days

W. Lohmeyer Fund

Thomas H. Johnson

SRS

Wanda Stout

S. R. S. - Jefferson Co.

4

Sandy Chaney

WLF

DATE

2/4/82

COMMITTEE ON

SubCommittee on Human Services
Appropriations

VISITORS' REGISTER

NAME	REPRESENTING	BILL #	Check One	
			Support	Oppose
JENNIS H. Taylor	DD D/SRS		✓	
Nellie James	Self		✓	
James & Harris	Self		✓	
Susan Kaphammer-Myers	Self and family		✓	
LURA NESMITH	STEP		✓	
Paul Nesmith	Self		✓	
Hatharine A. Kelker	Parents, Let's Unite for Kids		✓	
Patricia A. Lucero	DD / S.T.E.P.		✓	
Beyton Perry	DD Reg I		✓	
Ann Thomas	DD		✓	
Gandy W. Thomas	DD		✓	
Jacky Rogers	Self		✓	
Glenn Sawyer	DD		✓	
Richard B. Hain	Self		✓	
Vol J. H.	DD / RSC		✓	
Barbara M.	DD / NGE		✓	
Bill Sirak	Easter Seals / Goodwill		✓	
Judy Hainsmith	family		✓	
Tom Crosser	DD Planning Task Force		✓	
Chew Wanner				
Jane Olsen				
Donna Schack	DD Council Reg III		✓	
John C. Olsen	WEST MONT		✓	
Jane Swaster	Regional Living Service		✓	
John Hain	SRS		✓	
Jill Sherman	Vocational Placement		✓	

(Please leave prepared statement with Secretary)

DATE February 4COMMITTEE ON Subcommittee on Human Services Appropriations

VISITORS' REGISTER

NAME	REPRESENTING	BILL #	Check One	
			Support	Oppose
Rob Tallon	Bozeman	DD Approp.	X	
Jean Brackford	Billings	DD Approp.	X	
Dick Nordberg	MATTA	DD Approp.	X	
Margie Baker	Harlem	DD Approp.	X	
Laura Brockway	Miles City	DD Approp.	X	
Rock Thompson	Missoula	DD Approp.	X	
Janice Frisch	Helena	DD Approp.	X	
Sue Means	Helena	DD Approp.	X	
Clyde Muirhead	DDPAC	DD Approp.	X	
Mike Baker	Westmont H.	DD Approp.	X	
Pat A. Mulligan	Missoula	DD Approp.	X	
Linda Woodell	Helena	DD Approp.	X	
Rev. Bob Craver	Butte	DD Approp.	X	
Nana Craver	Butte	DD Approp.	X	
Ken Jackson	Helena	DD Approp.	X	
Larry Noonan	" "	DD Approp.	X	
Joseph L. Curtin	Choteau	DD Approp.	X	
Pharage Leher	Bozeman	DD Approp.	X	
Karen Oyler	Manhattan	DD Approp.	X	
Mune Mayhew	Montana Cath Conf- MAC	DD Approp.	X	
James R. Wolf	MARK	DD Approp.	X	
Carl M. Neal	Mont. Asian Pacific J. & T. C.	DD Approp.	X	
Bob Kessel	Butte Shelter & Training	DD Approp.	X	
John P. Kuyper	Montana Advocacy Program	DD Approp.	X	
Ricco Fichette	Helena -	DD Approp.	X	
Barrie F. Green	Great Falls	DD Approp.	X	

(Please leave prepared statement with Secretary)

2/4/87

DATE 2/9/
Sub Committee on Human Services

VISITORS' REGISTER

NAME	REPRESENTING	BILL #	Check One	
			Support	Oppose
Judy Fisher	Vocational Placement		✓	
Cathy Peterson	Specialized foster care		✓	
Terri Jones	Child + Family Region II Great Falls		✓	
Ida May	DDIS		✓	
Ben Jones	SRS		✓	
Barbara Prichard	Billing Parent		✓	
Maureen Battistol.	Billing		✓	
Betty Huff	Butter			
Sally Adams	Butter			
Forrest	Blas (REM)		✓	

(Please leave prepared statement with Secretary)

DATE February 4

COMMITTEE ON Subcommittee on Human Services Appropriations

VISITORS' REGISTER

[illegible]

(Please leave prepared statement blank)