

Maine Developmental Disabilities Council

February 14, 2023

Testimony in favor of LD 258 An Act Making Unified Appropriations and Allocations from the General Fund and Other Funds for the Expenditures of State Government and Changing Certain Provisions of the Law Necessary to the Proper Operations of State Government for the Fiscal Years Ending June 30, 2023, June 30, 2024 and June 30, 2025

Senator Rotundo, Representative Sachs and the distinguished members of the Joint Committee on Appropriations and Financial Affairs:

My name is Nancy Cronin and I am the Executive Director of the Maine Developmental Disabilities Council (DD Council). I am here to testify in support of PART A Section Z185.

Who We Are:

Councils on Developmental Disabilities were created through the Developmental Disabilities Assistance and Bill of Rights Act (DD Act) in 1970. Maine's DD Council has been advocating for individuals with Developmental Disabilities (DD) for over 40 years. The DD Council is a federally-funded, independent organization with members from across the state, including persons with disabilities, family members, and representatives of public and private agencies which provide services and/or funding for services for individuals with developmental disabilities.

As required in federal law¹, our purpose is to promote systems change to ensure that all individuals with developmental disabilities are able to live and fully participate in their communities of choice. Working in partnership with people with disabilities, parents, advocates, and policy makers, Maine's DD Council works to promote the independence,

¹ The Council's work is governed by the federal Developmental Disabilities Assistance and Bill of Rights Act, which defines a "*developmental disability*" as a severe chronic disability of an individual that is attributable to a mental or physical impairment or combination of mental and physical impairments, manifested before the individual attains age 22 and likely to continue indefinitely, which results in substantial functional limitations in 3 or more of the following areas of major life activity: self-care; receptive and expressive language; learning; mobility; self-direction; capacity for independent living and economic self-sufficiency.

integration, and inclusion of all people with disabilities through advocacy, capacity building, and systems change activities throughout the state of Maine and on the national level. The DD Council's mission is **to create a Maine in which all people are valued and respected because we believe communities are stronger when everyone is included.** The Maine DD Council employs 3.5 full-time staff who are responsible for all of the internal projects, external grants, and business of the Council.

What We Do:

- Promote and provide grants for activities that expand the capacity of communities to provide opportunities for individuals with developmental disabilities to actively participate in all aspects of community life.
- Advocate for systemic changes that allow individuals with developmental disabilities to achieve full integration and pursue meaningful and productive lives.
- Increase public awareness and work to eliminate barriers that impact the independence, productivity and inclusion of people with developmental disabilities.
- Foster and support coalitions and other advocacy and community groups.
- Support close working relationships among the various public and private service providers.

With our Federal Funds and the required State Match, the Maine DD Council will continue our novel work to improve the lives of Maine people with DD and the systems they utilize. The DD Council works with State Agencies to identify potential solutions to systemic problems and funds projects to test whether or not the solutions are effective. We are a unique agency in that it is our job to try out new ideas, evaluate their effectiveness, and apply the lessons learned for systemic improvements. While more can be read about our programs in the 2023 legislative report, I would like to highlight a few projects that we have recently worked on to improve the system, for example the Maine DD Council:

- Provides funding and technical assistance to Speaking Up For Us (SUFU) to support individuals with DD to have real, self-determined lives in the community. In just the last couple of years, in the midst of a pandemic, hundreds of individuals with DD either received training or achieved greater independence as a result of SUFU's work. SUFU's work was especially important during COVID preventing complete isolation by providing peer contact and opportunities to be involved and share what concerns in the middle of a global pandemic.

- Provide expertise in public policy about existing systems and how systems work together. Forever grounded and directed by our Council members with lived experience the Maine DD Council takes the long view in systems change with a keen eye towards outcomes as opposed just to outputs. A couple examples:
 - Maine DD Council recognizes that the earlier a child is identified as having a disability and receiving intervention the better. Over the last 15 years we have worked to identify children with delays earlier and advocated for barriers to be removed to access necessary educational and medical interventions. We have done this through multiple advocacy projects to increase screening, identification, and decrease barriers to children accessing effective early intervention. Our hope is through our efforts the system will be improved and we have seen some progress. While we recognize early identification still needs to be even earlier the needle has moved from an average identification ten years ago of age 8 to age 4 in Maine. We look forward to reporting one day that the average age has reached 18 months – until that time we will continue to push the system to do better.

We continue to be a prominent voice in Maine for early, comprehensive, and effective early intervention through our advocacy, but we also provide funding to test novel approaches. Currently we are funding a pediatric center to house an enhanced model of care navigation aimed at advocating and guiding families of young children after diagnosis until they are able to fully access services. We hope that this project not only helps some children and families but that we learn more about current gaps and gain insight on how to close those gaps so that all children with disabilities are able to access the services they need faster. The ultimate goal is to ensure all young children with delays are able to maximize early intervention.

- Another long term change that the Maine DD Council has focused on is to advocate for a self-directed service delivery model to be available here in Maine as an alternative to traditionally delivered and managed services, such as an agency delivery model. Self-direction of services allows participants to have the responsibility for managing all aspects of service delivery in a person-centered planning process. After many years of advocacy and work we are thrilled to report that a self-directed option is available to recipients of one of the adult DD waivers.

- Provide funding and technical assistance to improve healthcare for people of all ages with DD. Over the last few years the Maine DD Council has partnered with the Maine Primary Care Association through a multipronged approach, including increasing understanding of adverse occurrences experienced by individuals with DD, planning and implementing a response to address adverse occurrences, increasing clinical capacity to address the needs of individuals with DD through evidence-based practices (project ECHO), and modifying typical clinical processes/practices to address specific needs of individuals with DD through evidence-based practices (project ECHO). Part of that work included the creation of a “practice readiness document”, outlining how to create a safe and supportive environment for these patients. ECRI, a nationally renowned institute for safe medical practices, became a partner in testing that document. As a result of this work MDDC has been recognized by HRSA for its role in innovation in utilizing a patient safety organization to partner with a local agency on a shared need (in this case, how to best provide care for patients with developmental disabilities). An impact study highlighting our work as a promising practice can be found on The Administration for Research and Health Quality (AHRQ) website.

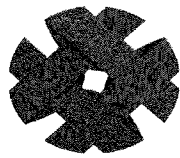
All of the above work products are high-quality and are highly beneficial to the systems that serve people with DD. The State Match Funds go and will continue to go to funding projects throughout Maine to develop and test models of services of supports that meet the needs of Maine communities and directly improve people’s lives.

I really could go on and on about the wonderful work that the DD Council does.

How We Are Funded:

Every State and Territory in the United States of America has its own Developmental Disabilities Council. The Administration for Intellectual and Developmental Disabilities provides each state with an allotment of federal funding to support Council work. In return, the federal government requires that each state provide funds to their DD Council via a State Match. The funding proposed in this bill would do just that by providing a State Match for the current fiscal year. The Maine DD Council works on a five-year State Plan system, in which as soon as staff completes a project they move on to the next.

Thank you for this opportunity to speak. I am happy to take any questions.



**Maine Developmental
Disabilities Council**

Annual Report

**Submitted to the 131st Legislature
State of Maine, 1st session**

Pursuant to 34B: 17001

January 1-December 31, 2022

This information has been compiled by Maine Developmental Disabilities Council, pursuant to its duties under the federal Developmental Disabilities Assistance and Bill of Rights Act of 2000, to complete and update a comprehensive review and analysis of services, supports, and other assistance for individuals with developmental disabilities (as specified in 42 United States Code, Chapter 144, Section 15024(c)(3)).

This project was supported, in part by grant number 2001MESD-02, from the U.S. Administration for Community Living, Department of Health and Human Services, Washington, D.C. 20201. Grantees undertaking projects with government sponsorship are encouraged to express freely their findings and conclusions. Points of view or opinions do not, therefore, necessarily represent official ACL policy.

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Maryann Harakall
CSHN Representative

Throughout FY22, as the Maine Developmental Disability Council initiated its 2022-26 State Plan, we heard from members, stakeholders, and community members about their “ending the pandemic” experience. As restrictions eased, persons at high risk, including many persons with developmental disabilities, continued to experience challenges in accessing their communities and needed supports. Families report gaps and long waitlists to access both community and education services. Concern from advocacy groups and providers regarding young children failing to access needed services increased. Adult’s report continued un and underemployment, even with historically low unemployment. This stress has been exacerbated by financial challenges related to inflation and by ongoing workforce challenges.

We appreciate the leadership of Governor Janet Mills and her administration as over the past several years the system continues to adjust to meet the needs of individuals with DD in a post-pandemic world.

We thank our members and staff for their flexibility and commitment throughout 2022.

Finally, we wish the 131st Legislature all our best as it undertakes a difficult job.



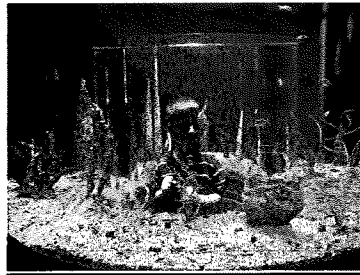
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I. Maine Developmental Disabilities Council Overview

Who We Are

Councils on Developmental Disabilities were created through the Developmental Disabilities Assistance and Bill of Rights Act (DD Act) in 1970 to "engage in advocacy, capacity building, and systemic change activities that are consistent with the purpose of the DD Act; and contribute to a coordinated, consumer and family-centered, consumer and family-directed, comprehensive system of community services, individualized supports and other forms of assistance that enable individuals with developmental disabilities to exercise self-determination, be independent, be productive and be integrated and included in all facets of community life."

Maine Developmental Disabilities Council (MDDC) is a partnership of people with disabilities, their families, and agencies which identifies barriers to community inclusion, self-determination, and independence, and acts to effect positive change. The Council has a vision that all people are included, supported, and valued in communities that provide opportunities to participate and succeed as they choose. MDDC acts to affect positive change through advocacy, capacity building, training, demonstration projects, and support for other inclusive and collaborative systems change activities.

Mission

Maine Developmental Disabilities Council is committed to creating a Maine in which all people are valued and respected because we believe communities are stronger when everyone is included.

Purpose

To **promote systems change** to ensure that all individuals with developmental and other disabilities are able to **live and fully participate** in their communities of choice. Working in partnership with people with disabilities, parents, advocates, and policy makers, MDDC works to **promote the independence, integration, and inclusion** of all people with disabilities through advocacy, capacity building, and systems change activities throughout the state of Maine and on the national level.

Enabling Federal Legislation

Under the federal Developmental Disabilities Assistance and Bill of Rights Act of 2000, MDDC is required to ensure that individuals with developmental disabilities and their families participate in the design of, and have access to, needed community services, individualized supports, and other forms of assistance that promote self-determination, independence, productivity, integration, and inclusion in all facets of family and community life.

Council Membership

Maine Developmental Disabilities Council's specific priorities are shaped by its governing council. The Council consists of 27 members, 19 of whom are appointed by the Governor. 60% of MDDC's membership is made up of people with developmental disabilities or their representative family members. The remaining 40% of MDDC's membership is made up of representatives of State agencies administering federally-funded programs related to individuals with developmental disabilities, local non-governmental agencies/organizations that serve people with developmental disabilities, and MDDC's "sister agencies", so-called because they are also funded through the DD Act: Disability Rights Maine, Maine's Protection and Advocacy agency for people with disabilities,

and the Center for Community Inclusion and Disability Studies at the University of Maine, Maine's University Center for Excellence in Developmental Disabilities.

The breakdown of membership includes:

- 5 seats designated for individuals with developmental disabilities
- 5 seats designated for parents, guardians, or family members of individuals with developmental disabilities
- 6 seats designated for *either* individuals with developmental disabilities *or* parents, guardians, or family members of individuals with developmental disabilities
- 1 seat designated for a local non-governmental agency/organization that serves people with developmental disabilities
- 1 seat for the University Center for Excellence in Developmental Disabilities
- 1 seat for the Protection and Advocacy agency for people with disabilities
- 1 seat for a statewide self-advocacy organization
- 1 seat for each of the following State Agencies:
 - Department of Health and Human Services, Office of Aging and Disability Services
 - Department of Health and Human Services, Office of Child and Family Services
 - Department of Health and Human Services, Children with Special Health Needs
 - Department of Education, Special Services
 - Department of Education, Child Development Services
 - Department of Labor, Vocational Rehabilitation

Due to its diverse membership, the Council provides opportunities for disparate perspectives to come together to work toward systems that support individuals with developmental disabilities to be fully included in their communities.

What We Do

Maine Developmental Disabilities Council is required to engage in advocacy, capacity building, and systems change activities that address:

- | | |
|--------------|--|
| ❖ Employment | ❖ Transportation |
| ❖ Health | ❖ Community Supports |
| ❖ Child Care | ❖ Quality Assurance/Self-Determination |
| ❖ Recreation | ❖ Education and Early Intervention |
| ❖ Housing | |

MDDC develops a five-year plan that best suits the needs of the community. Current strategies include:

- Promote and provide small grants for activities that expand the capacity of communities to provide opportunities for individuals with developmental disabilities to actively participate in community life
- Advocate for systems change that allow individuals with developmental disabilities to achieve full integration and to pursue meaningful and productive lives
- Increase public awareness and work to eliminate barriers that negatively impact independence, productivity, and inclusion of people with developmental disabilities
- Foster and support coalitions and other advocacy and community groups
- Support close working relationships among the various public and private service providers

Five-Year State Plan – See Appendix A.

II. Defining Developmental Disability

The US Department of Health and Human Services' Administration on Intellectual and Developmental Disabilities states that:

"A *developmental disability* is a severe chronic disability of an individual that:

- Is attributable to a mental or physical impairment or combination of mental and physical impairments
- Is manifested before the individual attains age 22
- Is likely to continue indefinitely
- Results in substantial functional limitations in 3 or more of the following areas of major life activity: self-care; receptive and expressive language; learning; mobility; self-direction; capacity for independent living; and economic self-sufficiency
- Reflects the individual's need for a combination and sequence of special, interdisciplinary, or generic services, individualized supports, or other forms of assistance that are of lifelong or extended duration and are individually planned and coordinated

An individual from birth to age 9, who has a substantial developmental delay or specific congenital or acquired condition, may be considered to have a developmental disability without meeting 3 or more of the criteria described above if the individual, without services and supports, has a high probability of meeting those criteria later in life." (Developmental Disabilities Assistance and Bill of Rights Act of 2000)

The federal definition is *functional*; that is, it is intended to describe the nature and scope of limitations without reference to medical diagnosis.

How Many People Have Developmental Disabilities?

The Administration on Intellectual and Developmental Disabilities estimates the population prevalence of developmental disabilities at 1.58%.

Approximately 21,000 Mainers of all ages have developmental disabilities.

The overall prevalence of disability among all people in the United States is 12.6%¹; among people in Maine it is 16.3%² Approximately 217,000 Mainers report having a disability

¹ <https://www.disabilitystatistics.org/reports/acs.cfm?statistic=1>

² <https://www.disabilitystatistics.org/reports/acs.cfm?statistic=1>

III. Concerns of Maine Families Affected by Developmental Disability

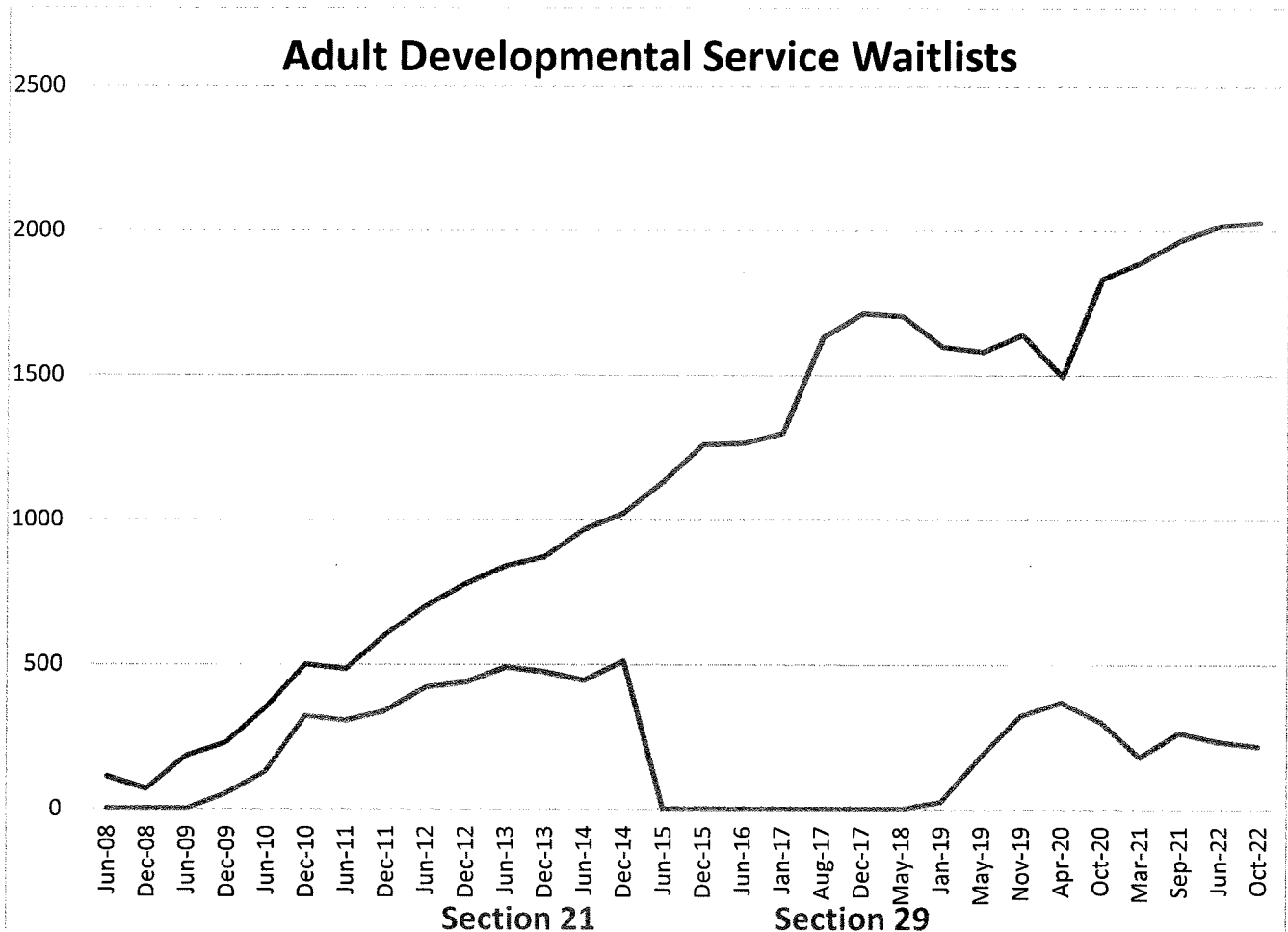
Access to Services and Supports

Maine Developmental Disabilities Council monitors available data about services and supports available to Mainers with developmental disabilities and gathers input about gaps in the system.

Adult Developmental Services

Upon reaching adulthood, the availability of individual and family supports changes from a model of entitlement to eligibility. To be eligible for Developmental Services, an individual must demonstrate substantial functional limitations in three (3) or more of the areas of major life activity (such as self-care, receptive and expressive language, learning, mobility, self-direction, capacity for independent living, and economic self-sufficiency).

MDDC has noted with appreciation the attention to the state's addressing the waitlists for waiver services. However, there continues to be a significant, growing, need for services. MDDC also notes the challenges encountered by eligible persons in actually accessing those services, particularly people with challenging behaviors. Waitlist data was obtained by the Office of Aging and Disability Services and its predecessors.



Children's Services and Special Education

Children with developmental disabilities are entitled to a fairly robust array of services and supports. However, Maine families experience very significant challenges in accessing those services, especially the areas of:

Early intervention: Young children who are at risk of developmental disabilities need early intervention services. They are not receiving timely and effective services due to (de facto) waitlists, high rates of staff turnover, insufficient funding, lack of inter-department coordination and erosion of service delivery capacity due to systematic regulatory changes.

Obtaining community supports: At times, families do not receive needed services because qualified providers are not available. This is noteworthy in long waits for Section 28 services. One father spoke to MDDC of having to decide whether an unreliable support person was preferable to no support person and of changing holiday plans based upon the availability of support. Another spoke of having no choice but to pursue residential placement (due to the lack of availability of support). Another parent spoke of a child receiving speech therapy once every other week for a 90-minute session rather than the 20 minutes three times a week that the child needed.

Inadequate transition planning: One mother's comments reflect many other families' concerns. She said:

"...Transition is one thing, but if there is little of quality or even adequate to transition to, the point of transitioning is completely lost. My experience and observations are that a sufficient and qualified pool of support people doesn't exist, aren't paid or supported well enough, etc. Right now, my husband and I feel like the most realistic plan for our daughter's future is that we must never become ill and must never die OR we should clone ourselves. I know that seems absurd, but it is not said necessarily in jest. I'm sure many other families relate to those kinds of sentiments, nor is it new to you."

Other Services

While the Office of Aging and Disability Services (OADS) provides developmental services to approximately 6000 individuals, there are thousands more Mainers who meet the federal criteria of having a developmental disability. Some receive appropriate services through another section of Maine Care such as Section 19, 20, or 50. Some have multiple diagnoses and receive services such as Section 65 or 96 which may create emphasis on alternative diagnoses, leading to less than effective/efficient care. Some do well with no services at all. Some do not, Some are in the criminal justice system. MDDC hears from/of many Mainers with significant intellectual and developmental impairments and their family members who simply do not qualify for services. They struggle to obtain and retain housing, transportation, employment, and healthcare. These people are disproportionately poor and have been impacted by such issues as Maine's response to Medicaid expansion, affordable housing policy, and income-based supports such as SNAP, TANF, general assistance, and heating subsidies.

Other Issues

Maine Developmental Disabilities Council monitors other issues that significantly impact Maine families affected by developmental disabilities.

Poverty³ **32.4% of working-age Mainers with disabilities live at or below the federal poverty level (compared to 7.7% for people without disabilities).**

Unemployment

Historically, the unemployment rate⁴ of people with disabilities is double that of the general unemployment rate. While numbers specific to people with developmental disabilities are not available, relevant data such as the National Core Indicators⁵ suggest that the numbers for people with developmental disabilities are considerably worse, with only 30% of people employed. For individuals with developmental disabilities who do not receive waiver-level services, access to ongoing employment supports is extremely limited. This significantly limits people's ability to work.

Aging Caregivers

The aging of Maine's population stands to pose serious challenges in meeting the support needs of persons with developmental disabilities. Families continue to be the primary providers of care. According to national statistics, an estimated 76% of individuals with developmental disabilities live at home. In 25% of these households, the family caregiver was age 60 or older and the average age of the family member with a disability was 38 years. There are an estimated 641,000 adults over age 60 who have developmental and related disabilities, a number that is projected to double by 2030.⁶ **In Maine it is estimated that approximately 4,230 persons with developmental disabilities are living at home with caregivers over the age of 60.**⁷

Dental Care

Limited or lack of access to both preventative and restorative dental care is a barrier to the health of people with developmental disabilities who are not institutionalized. Lack of access is even more significant for those who do not have access to waiver-level services.

Transportation

Most people with developmental disabilities do not drive. In a state that is largely rural with little public transit infrastructure, this severely limits the ability to travel. People with developmental disabilities rely on a patchwork of service providers, family and friends, public transit, taxis, and volunteer drivers. Individuals with developmental disabilities report that obtaining transportation to social and recreational opportunities, particularly individual activities, is especially difficult. Community transportation (public/regional and volunteer) is limited geographically, making short trips complicated and limiting vocational and social opportunities, and is essentially non-existent in the evening or on weekends. People with physical disabilities have additional difficulties with transportation in that taxis and private vehicles of family, friends, and volunteer drivers may not be accessible, public transit has limited accessible seating, and in the winter, people may have difficulty negotiating icy conditions.

MDDC has noted with concern the access and safety challenges of persons with developmental disabilities utilizing Non-Emergency Medical Transportation.

³ <https://www.disabilitystatistics.org/reports/acs.cfm?statistic=7>

⁴ www.maine.gov/labor/cwri/disabilities/index.html

⁵ https://www.nationalcoreindicators.org/upload/core-indicators/ME_IPS_state_508.pdf

⁶ Heller, Tamar Ph.D. (2011) Rehabilitation Research and Training Center on Aging with Developmental Disabilities, Clearinghouse on Aging and Developmental Disabilities, Chicago, IL

⁷ Braddock, et al. (2015) The State of the States in Developmental Disabilities Tenth Edition, The University of Colorado

IV. Maine Developmental Disabilities Council Projects and Initiatives 2022 Highlights

Supporting Self Advocacy

Speaking Up for Us (SUFU) is Maine's self-advocacy organization run by and for people with I/DD. Created in 1993 to help ensure that individuals living with I/DD in Maine are equal, thriving members of their communities. SUFU's mission is to create new opportunities for adults living with I/DD to discover their abilities, exercise freedom and contribute to their communities. Supporting self-advocacy has been MDDC's most important initiative for many years in terms of fiscal effort and organizational support. In FY22 MDDC provided funding and technical assistance for self-advocates to develop leadership and advocacy skills, to increase understanding of and participation in policymaking and expanding awareness of and membership in self-advocacy, including youth and underserved populations. SUFU, like many organizations, began the year operating on a virtual platform and transitioned back into some in-person activities by the year's end. After overcoming very significant barriers to the digital shift of recent years, this hybrid format presented operational challenges. Self-advocate feedback was positive, although some members had challenges with participation throughout the year due to continued public health concerns.

Advocacy

In accordance with its responsibilities under the federal DD Act, Maine Developmental Disabilities Council has provided ongoing information and support to legislators and other public policymakers in the development and review of proposed legislation, rules, and other policies affecting individuals with developmental disabilities and their families. MDDC has provided information and participated in the revision of statutes and rules affecting civil rights, education, healthcare, employment, housing, and other matters of concern to individuals with disabilities, parents, and other family members. The legislative and rulemaking process continued to take place virtually. This seems to have decreased travel related barriers, increasing geographic diversity of stakeholders in all types of policymaking

Addressing Sexual Abuse

Early in 2022, MDDC received a request for evidence-supported methods to treat people with DD who have been sexually assaulted. We conducted a literature review and community canvas and learned that trauma therapists and other practitioners express concern that they do not have the necessary skills to work with persons who have DD and may turn them away because they feel as though they won't be effective. Therapists fluent in working with persons with DD feel that they don't have the skill to treat the trauma. This leaves the individual with DD without many places to find help. MDDC realized that there wasn't an easy answer and therefore conceived the Community of Praxis as a way to increase clinical capacity and professional ability to support people with I/DD who have experienced sexual abuse. MDDC gathered 7 clinicians with expertise treating diverse individuals with I/DD, including persons with limited communication skills, and expertise in trauma informed for sexual abuse and an Advisory Committee composed of people with lived experience and representatives from the self-advocacy and sexual assault response organizations. This Community of Praxis created a resource compendium of existing best or emerging practice and recommendations to improve the care of individuals with DD who have experienced or witnessed sexual assault.

Documenting our History/Telling Our Stories

The history of the treatment of people with developmental disabilities, both in Maine and elsewhere, is a story of stigmatization, isolation, and abuse – but within that tragic history live stories of

perseverance, joy, and progress towards something better. In 1908, the Maine School for the Feeble-Minded opened in New Gloucester. Over the years, its name changed several times, and eventually became known as Pineland Center, but the idea behind its creation remained: that the proper place for people with developmental disabilities was in an institution, shut away from the rest of society. When the rampant abuses and neglect in Pineland and elsewhere were exposed in the middle of the 20th century, a movement for disability rights was born. Self-advocates and their allies fought to close Pineland (which finally happened in 1996) and create a system of services that would serve people with developmental disabilities in their homes and communities. To mark the 25th anniversary of the closure of Pineland that took place in 2021, MDDC has created this online exhibit to document the history of the system's response to people with developmental disabilities in Maine. Please visit <https://shadowsofpineland.org/>

Early Intervention

The National Center for Disease Control and Prevention's Act Early program aims to improve early identification of children with autism and other developmental disabilities so children and families can get needed services and support to improve functional outcomes. MDDC's Executive Director chaired the Independent Advisory Review Committee on Maine's Early Childhood Special Education Services and oversaw the implantation of Help Me Grow, system so that young children can be screened for disabilities and families can access what they need. MDDC will continue to serve as a long-term resource to this program.

Health

People with developmental disabilities experience a variety of health disparities including but not limited to being more likely to report being in poor health, having shorter average life expectancy than the general population, being less likely to receive preventative care, having higher rates of undiagnosed hearing and vision impairments, higher rates of/poorly managed chronic health conditions, and being prescribed higher rates of psychotropic medication. There are a variety of contributing factors, including societal/attitudinal barriers (stigma), barriers due to disability related conditions and/or functional limitations related to his/her disability, and other barriers. MDDC is working with Maine Primary Care Association to address these issues through a multipronged approach, including increasing understanding of adverse occurrences experienced by individuals with DD, planning and implementing a response to address adverse occurrences, increasing clinical capacity to address the needs of individuals with DD through evidence-based practices (project ECHO), and modifying typical clinical processes/practices to address specific needs of individuals with DD through evidence-based practices (project ECHO). Part of that work included a "practice readiness document", outlining how to create a safe and supportive environment for these patients. ECRI, a nationally renowned institute for safe medication practices, became a partner in testing that document. The Administration for Research and Health Quality (AHRQ) invited MPCA, ECRI, and MDDC to present on the partnerships and readiness brief at the national conference. The presentation was so well received, and interest in the readiness brief so positive, AHRQ subsequently honored the work by completing an impact case study on the project which can be found at their website. As a result of this work MDDC has been recognized by HRSA for its role in innovation in utilizing a patient safety organization to partner with a local agency on a shared need (in this case, how to best provide care for patients with developmental disabilities).

Post-Secondary Education

Despite many years of effort with the state colleges and universities, Maine does not have any residential postsecondary education options for youth with ID. Thus, MDDC was excited to support a small liberal arts college that had identified this need to conduct program planning, in-person and virtual outreach, education, and technical assistance to increase understanding of and interest in campus based post-secondary education to youth with intellectual disabilities and their family members.

Reducing Restraint and Seclusion

MDDC supported the Coalition Against Restraint & Seclusion (CARS), a workgroup consisting of family members, self-advocates, & other advocates, and funded a pilot project, in which a service provider who runs both a special purpose program and residential programs engages in a quality improvement and fidelity assessment process study to determine whether restraint, seclusion, and isolation can be reduced when staff receives increased training and technical assistance in the program. MDDC worked the AUCD to complete a formal program evaluation. Despite significant programmatic challenges due to the pandemic, results showed that fidelity was, indeed, associated with a reduction of incident reports by twofold in residential settings & threefold in educational settings. MDDC will use these promising outcomes to inform its continued advocacy aimed at eliminating the use of restraint and seclusion.

Self-Direction

CMS says: "Self-directed Medicaid services means that participants, or their representatives if applicable, have decision-making authority over certain services and take direct responsibility to manage their services with the assistance of a system of available supports. The self-directed service delivery model is an alternative to traditionally delivered and managed services, such as an agency delivery model. Self-direction of services allows participants to have the responsibility for managing all aspects of service delivery in a person-centered planning process."

MDDC contracted with Applied Self Direction to provide technical assistance and with the state's community service provider organization to support this effort and has worked with a coalition of advocates, service providers, and state agency representatives to provide input towards and information about a system that allows people with developmental disabilities more say and flexibility in their services. After many years of advocacy, MDDC is thrilled to report that a self-directed option is available to recipients of one of the waivers.

Appendix A –MDDC Five-Year State Plan ✦ 2022-2026 ✦ Goals and Objectives

GOAL # 1 People with developmental disabilities, families, and communities will have increased opportunities for choice, self-determination, and community membership.

- *Objective 1.1:* MDDC will promote the civil rights of individuals with developmental disabilities.
- *Objective 1.2:* Support Maine's self-advocacy organization and promote development of diverse leaders among people with developmental disabilities and families.

GOAL # 2 Improve systems of support for people with developmental disabilities and families.

- *Objective 2.1:* Decrease health and educational disparities experienced by people with developmental disabilities.
- *Objective 2.2:* Improve access, quality, and integration of services and supports for individuals with developmental disabilities and families.
- *Objective 2.3:* Support efforts to expand access to information and services for underserved individuals with developmental disabilities and families.