Stronger Together

The Tennessee Council on Developmental Disabilities brings people together to grow opportunities and communities

A Special Advertising Supplement
For the People, 
By the People

The Council educates and connects citizens and government for change that matters.

The Tennessee Council on Developmental Disabilities exists to improve the lives of people with disabilities and their families. With its mix of experts and citizens, it has successfully filled that mission for nearly 50 years.

The Developmental Disabilities Act of 1970 paved the way for state councils; the Tennessee council was established by executive order from the governor’s office in 1975.

“The interesting thing about the Council is it’s a state agency, but it includes both policymakers and private citizens--it brings together educators, policymakers, legislators and private citizens to address disability issues,” said Wanda Willis, the Council’s executive director for 28 years. “The membership of the Council is just electrifying — and that’s intentional.”

By law, the diverse group must include representatives from state departments of education, health, labor, vocational rehabilitation, mental health and Medicaid, as well as from both of Tennessee’s Centers of Excellence (the Kennedy Center at Vanderbilt and the Boling Center at The University of Tennessee). “We also have a representative from our state protection and advocacy system, Disability Rights Tennessee,” Willis added.

But the most important contribution comes from the 21 citizen members who have direct experience with intellectual and physical disabilities. “Council members reflect our state’s diversity, and they come from every area of the state, both urban and rural,” Willis said.

The Council’s chair is one of those citizen members. Tecia Puckett Pryor is an attorney and mediator whose 16-year-old son Harrison is on the autism spectrum. “He is a very friendly and loving kid and he really brightens people’s days by being with him,” Pryor said. “I take delight in knowing that others find joy in him.” She knows she has come a long way since she first learned of his disability. “I feel like the main transformation for me has been becoming aware of what’s really important for him.”

Education and employment quickly topped that list, which led to her work with the Council. “Our mission is to give people tools so they can be active and knowledgeable and promote change,” Pryor said. “We are educating people so they can be leaders in their communities and across the state.”

The Council’s direct connection to both citizens’ needs and state government allows it to bring the right people together to work on big issues. One example is the Employment Roundtable, which the Council hosts to share information and build partnerships across state programs working on employment for people with disabilities. Businesses are seeing the benefits of hiring employees who have a disability.

Willis remembers a recent Roundtable guest speaker who employs people with disabilities. “He said, ‘I wish I’d known all this years ago. These are the best employees I’ve ever had, and it completely changed the culture of the workplace.’ That’s the kind of change we’re working for.”
Partners in Policymaking

Free training gives individuals and families the tools to live their best lives

Faith Henshaw and David Frizzell are more than just siblings; they’re friends. They like to sing in the car, exercise at the YMCA and go to movies and concerts together. Like most big sisters, Faith always has been very involved in her brother’s life. Unlike most siblings, she will need to continue some level of care-giving into the future: David was born with Down syndrome.

The Tennessee Council on Developmental Disabilities’ Partners in Policymaking® program is helping her prepare for that future. The free training teaches adults with disabilities and their families advocacy strategies like how to talk to lawmakers, how to exercise educational rights and how to find employment opportunities. It also provides ongoing support through friendships and networking.

“A sibling, that’s your longest-lasting relationship; you’re going to have it for life,” Faith said. “I know one day when my parents are gone, he will be my sole responsibility. And while that’s something I willingly choose, it’s terrifying. … I’m not alone, there are other individuals who have the same fears as me.”

Their mother, Virginia Frizzell, is also a graduate of the Partners in Policymaking program. She said the training has made her more aware of what she can do to advocate on her son’s behalf and how to find resources that enable him to live as independently as possible.

“I feel confident that if something comes up in the future, I know where to go to find help.”

VIRGINIA FRIZZELL Partners in Policymaking graduate

“Being a teacher of 23 years, I felt I knew it all,” Virginia said. “But I was so surprised when I got there and went through the class at how much I didn’t know and how much more there was out there to learn.”

While finding resources and programs is important, Faith says the friendships and networking she’s gained through Partners training is just as invaluable.

“The world is hard and scary as is. When you add a special need or disability, it makes it harder and more scary,” she said. “This just gives you the aid and comfort you need. It makes it manageable.”

For more information and to apply for the Partners in Policymaking program, visit www.tn.gov/cdd/training-and-news or call 615-532-6556.
Raising an Inclusive Generation

Kids who grow and learn together build a society where everyone is valued

Alison and Brad Bynum believe inclusive education is essential for their 11-year-old daughter, Charlotte. But they also believe it is the way the world will become a better place.

“I feel strongly societal change starts with kids, so they can grow into a generation of inclusivity,” said Alison, who is also newly appointed to the Council.

“If they grow up together, when a person is in a position of offering employment, they will have grown up with people with disabilities and view them as employable. We have to start young with the people who will be hiring someday, living in the community, providing health care, if we want to change the culture.”

This commitment to inclusion started almost immediately after the Bynums began receiving multiple diagnoses for Charlotte, whom they adopted as an infant. They enrolled Charlotte – who has complex medical needs, uses a wheelchair and doesn’t speak – in an early intervention program when she was 6 months old. From ages 15 months to 6 years, she attended an early learning center and preschool that included both children with and without disabilities. Students without disabilities served as “peer models” in the same classroom.

But things changed in elementary school. Although Charlotte was on the rolls of a kindergarten general education classroom, she spent most of her time at school in a comprehensive development classroom with only students with disabilities.

“Because of our daughter’s disabilities and medical needs, it is challenging to assess what she is learning. She is still working to master an eye gaze communication device, though she does communicate using facial expressions, her body and her head. We cannot prove that she doesn’t understand, so we presume competence,” Alison said.

The Bynums worked together with school administrators and almost a dozen professionals, including teachers, therapists and nurses to develop an Individualized Education Plan. The team agreed that Charlotte could benefit from more time in her general education classroom, giving her the opportunity to hear grade level content alongside her peers. The social and emotional growth that has followed has been as important to Charlotte as the academic goals.

Charlotte is now a fifth grader. Her mother describes her as loving horses – she rides every week – as well as reading and spending time with her friends at church. “She’s a great listener,” Alison said with a laugh. “She knows all our secrets – if she learns to talk, we’ll all be toast!”

Charlotte is also loved by her classmates. Alison spends a few minutes with them at the beginning of each school year, explaining Charlotte’s disabilities. She tells them, “Asking respectful questions to learn is always okay.” Their understanding and relationships with Charlotte have flourished.

In fact, a classmate wrote a letter to the principal requesting playground equipment Charlotte could access with her wheelchair. Said Alison, “This type of outcome isn’t measured, but it’s priceless.”

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ALISON BYNUM
Mother and TCDD member

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INCLUSIVE EDUCATION RESOURCES

Advocacy Agencies
Disability Rights Tennessee: A statewide organization that advocates for the rights of people with disabilities; 800-342-1660; disabilityrightstn.org.

The ARC of Tennessee: A statewide organization founded by parents of children with intellectual and developmental disabilities; 800-835-7077; thearctn.org.

Support and Training for Exceptional Parents (STEP): A statewide family-to-family program that provides information, advocacy training and support to parents; 423-639-0125; tnstep.org.

Tennessee Voices for Children: Advocacy for students with challenging behaviors; 800-670-9882; tnvvoices.org.

Governing Organizations
The Tennessee Department of Education: A state agency that oversees K-12 public education; 615-741-5158; tn.gov/education.

Informational Websites
Wrightslaw: A user-friendly website for parents, teachers, advocates and attorneys that provides information about special education law and advocacy for children with disabilities; wrightslaw.com.
Danny Dee Simmons not only landed a good job for himself – at a tractor company he has revered since he was a boy – but he has paved the way for others to follow in his footsteps, earning real money for real work.

Simmons’ journey began in 2012, when St. John’s Community Services began converting from sheltered to fully integrated employment. “Danny had worked in a workshop environment for many years,” recalled Dwayne Webb, the program director for day and employment services. “But he started seeing some of his friends go out in the community and get jobs...so he stopped me in the breezeway one day and said, ‘I want a job – I want to work at John Deere.’ It was all his own initiative.”

Webb quickly reached out to Mitch Hobby, the finance and incentive manager for Tennessee Tractor LLC, which is the John Deere dealer in town. “We met with Mitch, and I introduced them...but then I pushed back from the table and let them have their interview,” said Webb. Simmons was immediately hired to do janitorial work – cleaning, sweeping, mopping, dusting the showroom – and it has been nothing short of a win-win. “I work there three days a week,” Simmons said proudly. “I like John Deere tractors, I like the customers, I like having friends and I like making money.”

But his seven-year presence at Tennessee Tractor also has been important for everyone. “We love having him here,” Hobby said. “He’s a morale booster – he’s always coming in with a good attitude. And he comes from this rural area, so he knows everyone – staff and customers.”

The town, which is home for Simmons, was actually overjoyed to see him. “People didn’t know what happened to Danny when his parents passed,” said Wendy Bratton, the residential program coordinator. “So he’s been able to reunite with a lot of people he grew up with.”

Simmons’ work ethic impressed Hobby so much, he now has St. John’s participants employed at five of the 10 Tennessee Tractor locations. “I firmly believe this: Everyone can do something; you can contribute,” Hobby said. “It makes people see us in a different light, but also ‘helping’ in a different light... These employees obviously provide a service that we need.”

Webb hopes other businesses will see similar advantages to inclusive hiring practices. “When customers come in, they might have someone in their family with developmental disabilities – or know someone who has a family member – and it helps them all think outside the box,” he said. “Danny set the bar for his peers but is also setting the path for people coming up.”
At New Vision Baptist Church in Murfreesboro, Tenn., 24-year old Terrell Brown frequently stands at the front door, greeting congregants as they enter the building.

“They like to say, ‘Good morning,’” he said. “I like to help.”

Brown’s mother, Linda Brown, said Terrell, who has autism, took it upon himself recently to welcome his fellow church members as they arrived for small group meetings.

“He likes to be the door greeter,” said Linda. “He loves that.”

Linda said that even before joining New Vision, she and her husband, Jeff, sought out congregations that would not only minister to Terrell’s special needs, but also allow him, as he grew older, to become an active church volunteer.

“It was hard for Terrell to sit in the sanctuary,” she said. “It was a sensory thing. But he always liked children and preschoolers.”

So at their previous church, Linda brainstormed with a church leader about possible activities Terrell could do during church services. At first, a church leader suggested a special needs group that taught Bible stories.

“But Terrell was past that,” she said. “He was slowly becoming more independent.”

So the church arranged for Terrell to volunteer in the preschool children’s church department, she says. There, Terrell helped with Bible lessons, passed out snacks and took youngsters outside to play.

“And if they were crying,” Linda said, “Terrell would sit and hold them. He just had that spirit of comfort.

“He got the opportunity to learn and become more independent and responsible by working with the children,” she added.

Meanwhile, to become a better advocate for Terrell, Linda in 2010 enrolled in the Partners in Policymaking® Leadership Institute, a national program hosted by the Tennessee Council on Developmental Disabilities that teaches people with disabilities and their families how to advocate for themselves to affect changes in local, state and federal policies.

“It made all the difference in the world,” she said. “The biggest thing that happened was that I got over the fear of speaking up for Terrell. I used to be so concerned about hurting someone else’s feelings or offending them. But then, I thought, that’s not good. We have to do what’s best for Terrell.”

At New Vision Baptist where the Browns joined in 2018, Terrell participates in a small group for adults with special needs. But, Linda said she plans to inquire about Terrell officially joining the church greeters and serving with fellow congregants at a local food bank or charity.

She adds churches can take steps to help those with disabilities find ways to serve.

“Talk with the person with the disability,” she said. “Talk to the parents, and be willing to think outside the box.”
Inclusive higher education programs open doors for students with developmental disabilities

Like any freshman, Morgan Thompson was excited about starting college, moving into the dorms and being on her own for the first time. It was a dream that was far from certain for Morgan, who has language delays. In the past, higher education was out of reach for many people with intellectual and developmental disabilities. But in Tennessee, several universities now offer inclusive programs designed for such students.

With funding from the Tennessee Council on Developmental Disabilities, Lipscomb University in Nashville developed its Igniting the Dream of Education and Access at Lipscomb (IDEAL) certificate program. The two-year program supports students through peer mentorship, career training, classwork accommodations, independent living skills and professional internships.

“They help you be independent, learn new skills and it gives us job opportunities and college opportunities,” Morgan said. “[They had] small classes and … if I didn’t understand something in the classroom, I could get help with it.”

Morgan graduated from the IDEAL program in 2019 with a certificate in career exploration. She took college classes in biology and communications, and completed internships with the university’s Office of Alumni Relations and College of Education. She also worked at off-campus internships with the Department of Intellectual and Developmental Disabilities and the state’s Department of Education. Through that hands-on experience, she discovered a love for working in an office.

“I like doing clerical work,” she said. “For the Department of Intellectual and Developmental Disabilities, I did lots of video editing, taking pictures with a camera and going to conferences. I enjoyed doing that as well.”

Social opportunities are also a big part of the college experience, something Morgan enjoyed during her time at Lipscomb.

“I was one of the first IDEAL female students to live in the dorms. [I liked] going to events on campus like sporting events,” she said. “I was in a sorority called Phi Sigma and I enjoyed that and making lots of friends.”

Morgan says she learned a lot through the IDEAL program, most importantly how to be more independent.

“So I can live on my own one day — make decisions, budget money and do things on my own,” she said. “Someday I plan on going into an office and doing clerical work. I would love that.”

For more information on the IDEAL program, visit www.lipscomb.edu/academics/undergraduate-studies/ideal-program.
30 Year Anniversary

Our nation recently marked 30 years since the signing of the Americans with Disabilities Act. Members of the Council on Developmental Disabilities shared their thoughts on the ADA as part of a #DisabilityMeans social media campaign.

For more #DisabilityMeans stories, follow the Council on Facebook and Twitter.

#DisabilityMeans life, liberty, and the pursuit of happiness.

“I am proud to be part of the ADA generation. That means I grew up with the privileges and the rights that the ADA has afforded people with disabilities. I do not take them for granted.

So much of this country still wants to focus on awareness of people with disabilities. They’ve had 30 years to be aware that I exist – and more than that, if you count the fact that we’ve existed since people have existed. I’m not asking for awareness any more.

I am asking for acceptance and respect. I don’t think we’ve gotten there overall as a country. The Declaration of Independence still does not apply to people with disabilities in its entirety. But I think we are getting there. And I look forward to every day that I get to spend to help push us farther.”

Jean-Marie Lawrence, member of the Council on Developmental Disabilities, Southeast Development District

#DisabilityMeans freedom to be a part of my community.

“I was 1 year old when the ADA came into place. I was never a part of that other world. To me, the ADA means freedom. It’s almost like a memorandum of understanding between our communities and people with disabilities. Before the ADA, we were confined to segregation. Now, as Americans with disabilities, we are able to be part of our community.

In the next 30 years, I hope that we can have a more inclusive world to live in. I hope that people will take the time to get to know people with disabilities before they make preconceived notions. In today’s world, there is a lot of prejudgment. I want people to know that we have a place at the table. Our voice needs to be heard. Our disability is a part of who we are, but it’s not who we are.”

Clancy Hopper, member of the Council on Developmental Disabilities, Mid Cumberland District

#DisabilityMeans working hard to get from here to there.

“I really appreciate the ADA. It made us think about basic accessibility. But the truth is, we still have a long way to go. I can get into a restaurant, but the table legs keep me from pulling my wheelchair up to the table. There is an accessible stall in the bathroom, but I have to be able to pull doors open to get there. It’s hard to find accessible housing and transportation. If you want to really be accessible, hire people with disabilities, so they can help you think about what’s needed. Disability is a part of being human. We need to plan for it so everyone can join in.”

Martez Williams, member-at-large of the Council on Developmental Disabilities