



Oifig an Stiúirthóir Cúnta Náisiúnta,
Foireann Míchumais Náisiúnta,
An Chéad Urlár - Oifigí 13, 14, 15,
Áras Phlásóg na Rós, Coimpléasc Gnó na hOllscoile,
Páirc Náisiúnta Teicneolaíochta,
Caladh an Treoigh,
Luimneach.

Office of the Assistant National Director,
National Disability Team,
First Floor- Offices 13, 14, 15,
Roselawn House, University Business Complex,
National Technology Park,
Castletroy,
Limerick.

23rd July 2024

Deputy Catherine Murphy,
Dail Eireann,
Leinster House,
Kildare Street,
Dublin 2.
E-mail: catherine.murphy@oireachtas.ie

Dear Deputy Murphy,

The Health Service Executive has been requested to reply directly to you in the context of the following parliamentary question, which was submitted to this department for response.

PQ: 31989/24

To ask the Minister for Children; Equality; Disability; Integration and Youth the number of new ASD diagnostic teams that have been developed within the State; the details of those teams; and the referral path

HSE Response

The implementation of the Progressing Disability Services for Children & Young People (PDS) programme is agreed Government and HSE policy. This policy supports the reconfiguration of children's disability services to provide equitable, child and family centred services based on need rather than diagnosis. This aligns with the UN Convention on the Rights of People with Disabilities.

The Progressing Disability Services for Children and Young People (PDS) model addresses the previous inequity in service provision whereby there may have been an excellent service for some children and little or no service for others. This variance may have been linked to diagnosis, age group or geography. Under the PDS programme children's disability services are changing from diagnosis based to needs based, so that all children with a disability or developmental delay have access to the right service based on their needs no matter where they live.

PDS is a significant change programme for the provision of services and supports for children from birth to 18 years of age, in line with Sláintecare and the Programme for Government, in order to:

- Provide a clear pathway and fairer access to services for all children with a disability
- Make the best use of available resources for the benefit of all children and their families
- Ensure effective teams are working in partnership with families and with education staff to support children with a disability to reach their full potential.



PDS aligns with two clear objectives of The Sláintecare Report to:

- Provide the majority of care at or as close to home as possible
- Create an integrated system of care with healthcare professionals working closely together.

The National Policy on Access to Services for Children & Young People with Disability & Developmental Delay ensures that children are directed to the appropriate service based on the complexity of their presenting needs rather than based on diagnosis. Many children with a disability who have support needs can be effectively supported within mainstream health services. This policy provides a single point of entry, signposting parents and referrers to the most appropriate service (Primary Care for non-complex functional difficulties and Children's Disability Network Teams for complex functional difficulties).

Children's Disability Network Teams (CDNTs)

93 Children's Disability Network Teams (CDNTs) are aligned to 96 Community Healthcare Networks (CHNs) across the country and are providing services and supports for children aged from birth to 18 years of age.

The establishment of CDNTs is intended to facilitate the provision of equitable, child and family centred services for all children with complex needs associated with their disability or developmental delay.

Regardless of the nature of their disability, where they live, or the school they attend, every child with complex needs and their families have access to a range of family centred services and supports of their CDNT according to their individual needs. This includes universal, targeted and specialist supports, such as individual therapeutic intervention and access to specialist consultation and assessment when needed. Supports are provided as is feasible in the child's natural environments - their home, school and community.

The model of service for all CDNTs is family-centred and based on the needs of the child. This includes universal, targeted and specialised supports and interventions, as appropriate to the individual child and family. It is based on the objectives of empowering and supporting parents and others who are with the child on a daily basis to facilitate the child's developmental needs.

The CDNTs are currently providing services and supports for over 46,000 children and strategies and supports for urgent cases on the waitlist where staffing resources allow. However, there are significant challenges for CDNTs including:

- Significant staffing vacancies
- Growth in numbers of children with complex need
- Growth in demand for Assessment of Need, diverting further resources away from interventions.

Roadmap for Service Improvement 2023 – 2026, Disability Services for Children and Young People

The HSE's Roadmap for Service Improvement 2023 – 2026, Disability Services for Children and Young People, approved by the HSE Board and launched by the Government and the HSE in October 2023 is a targeted Service Improvement Programme to achieve a quality, accessible, equitable and timely service for all children with complex needs as a result of a disability and their families.

It sets out the overall aim for Children's Disability Services, provided by the HSE and its partner agencies, for every child to have a childhood of inclusive experiences where they can have fun, learn, develop interests and skills, and form positive relationships with others in a range of different settings.

The Roadmap, which is now in its implementation phase, has established four Working Groups which report into a Service Improvement Programme Board every month which in turn reports to the Roadmap Oversight Group chaired by the Minister of State, Ann Rabbitte.

The WGs have wide membership including HSE, Section 38 and 39 front line disabilities staff and management, parent voices, staff reps, Primary Care, CAMHS and Department of Children, Equality, Disability Integration and Youth.

The Roadmap contains 60 actions of which 12 have been completed and the majority of remaining actions are in train. These include a robust suite of 21 staff retention and recruitment actions.



The HSE is entirely committed to the delivery of the Roadmap actions over the lifetime of the plan.

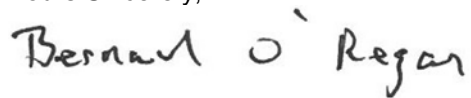
With regard to the specific question asked, as outlined above, children's disability services provide equitable, child and family centred services based on need rather than diagnosis.

With regard to an Assessment of Need under the Disability Act 2005, during 2023, the HSE developed AON Administrative Hubs in Community Healthcare Areas (CHOs) 1, 4, 5, 7, 8 and established AON Assessment Hubs in CHOs 3, 6 and 9.

Under the *Roadmap for Service Improvement 2023-2026, Disability Services for Children and Young People*, these AON Hubs will be amalgamated into 6 AON Assessment Hubs aligned to the 6 Regional Health Areas.

It is important to note that children do not require an Assessment of Need as defined by the Disability Act (2005) in order to access a CDNT or Primary Care service. They can be referred by a healthcare professional or parent/carer to the CDNT for children with complex needs as a result of their disability, or to Primary Care for children with non-complex needs.

Yours Sincerely,



Bernard O'Regan
Assistant National Director
National Disability Team

