

Who can refer to the Lanarkshire Service?

The service has an open referral system, which means direct referrals can be made by those impacted, a family member or a third party with consent. Please email sha-admin@hdscotland.org, call our Lanarkshire office on **01698 753 955** or get in touch using the numbers below:



Services for Lanarkshire

Senior HD Specialist
07714 245 740

HD Specialist
07985 252 004

Admin/Resource worker
01698 753 955

Youth Service
07983 724 201

Financial Wellbeing Service
07710 391 621

Lanarkshire office

Scottish Huntington's Association
Hunter Community Health Centre
Andrew Street
East Kilbride
G74 1AD
Telephone: 01698 753955

National office

Scottish Huntington's Association
Business First
Burnbrae Road
Paisley
PA1 2FB
Telephone: 0141 848 0308
Email: sha-admin@hdscotland.org

Further information can also be found on our website at hdscotland.org



Specialist support for Huntington's disease families

hdscotland.org

Scottish Charity No: SC010985
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Specialist support for Huntington's disease families

Lanarkshire Service



Huntington's disease: How we can support you and your family



What is Huntington's disease?

Huntington's disease is caused by an inherited faulty gene that damages the brain over time.

People with the disease can eventually lose the ability to walk, talk, eat, drink and care for themselves, requiring specialist support from those who understand the condition.

The disease is genetic, meaning it is passed down from one generation to the next. It therefore impacts entire families over generations rather than individuals alone. Anyone with a parent who has Huntington's has a 50% risk of inheriting the condition from them.

As Huntington's progresses it can affect:

Movement: People can experience repetitive involuntary movements resulting in a decline in mobility, balance and coordination as well as difficulties with speech and swallowing.

Thinking processes: As the disease advances, people can lose their ability to process information, make decisions, solve problems, plan, organise and care for themselves.

Mental health: People can also experience a decline in mental health. Depression, anxiety, irritability, obsessive pre-occupations and apathy are amongst the most common mental health problems experienced. Psychosis may also occur.

About Scottish Huntington's Association

Scottish Huntington's Association is the only charity in the country exclusively dedicated to providing expert and personalised support for those impacted by Huntington's disease.

Our personalised support reduces unnecessary hospital admissions, supports carers and other family members; lowers household poverty; and alleviates wellbeing risks to children and young people living in Huntington's families.

We are commissioned by NHS Boards and Health and Social Care Partnerships throughout the country to share our expertise with front line staff and build support for improved services and higher standards of care for every family impacted by this devastating disease.

What services do we offer?

HD Specialists

Our HD Specialists provide vital physical health, mental health and wellbeing support, and care co-ordination for all adults within a Huntington's family – whether they be symptomatic, asymptomatic, at risk or carers. They liaise with, amongst others, occupational therapists, speech therapists, physiotherapists, care homes, dieticians and community mental health teams. Additional HD Specialist support includes specialist assessment, symptom management, emotional support, help to plan for future care needs and enabling carers to build skills and resilience. Our expertise is invaluable to health and social care professionals who access training and advice from our HD Specialist teams. Most health and social care staff have no experience of Huntington's disease at all, and so depend upon the support of our specialist teams.

Youth Service

Our Specialist Youth Advisors provide personalised support, age-appropriate information and resources, opportunities to build friendships and enjoy respite experiences that would otherwise be impossible. They also provide specialist education, support and guidance to other professionals who work with children and young people who are impacted by Huntington's disease.

Young people growing up in Huntington's families see the disease changing their affected parent, with many knowing they have a 50% chance of inheriting the same condition. Many become young carers for siblings or their parents, while others are unable to live with their mum or dad. Some are grieving for loved ones, others are struggling with genetic testing. Huntington's disease impacts their home lives, education and social lives.

Financial Wellbeing Service

Our Financial Wellbeing advisors help families to reduce debt and maximise income through access to benefits and financial products. The service's work includes support with household budgeting, planning for the future by organising power of attorney, and providing specialist energy advice to help cut domestic fuel costs. Since 2015, the service has secured more than £6 million in financial gain for families across Scotland. Huntington's disease cuts working lives short, leaving many households we support living below the poverty line.

Where do we see people?

Wherever is most comfortable, whether that be at home or in another suitable venue.

HD Clinical Leads

HD Clinical Leads are senior NHS employed doctors (such as psychiatrists, neurologists or geneticists) designated to providing specialist medical advice, care and treatment to people with Huntington's. They work with Scottish Huntington's Association staff and other health and social care colleagues to coordinate care across their local NHS Board area. They run specialist clinics supported and facilitated by our HD Specialist staff, provide home visits to those unable to attend clinic and provide access to HD research programmes. In light of our expertise Scottish Huntington's Association coordinates and chairs the national network of HD Clinical Leads.

HD Management Clinics

Clinics are held every Tuesday. Please contact the Greater Glasgow and Clyde service for more details.

Support Groups

These provide opportunities to meet informally with others whose lives are impacted by Huntington's disease. Again, please get in touch with the Greater Glasgow and Clyde service for more details.

National Care Framework for Huntington's Disease

We have published a Scottish Government backed National Care Framework for Huntington's Disease which outlines the care and support families should be entitled to receive throughout the country. It can be viewed at

<http://care.hdscotland.org>

