What are the Ehlers-Danlos syndromes?

The Ehlers-Danlos syndromes are a group of genetic connective tissue disorders with symptoms affecting the whole body.

Those affected face challenges to their physical and mental health. Symptoms are widespread and can be disabling. One of the rarer types significantly reduces life expectancy. Hypermobility spectrum disorders (HSD) have similar symptoms to the most common type of Ehlers-Danlos syndrome, the hypermobile type. People now diagnosed with HSD may have been classified in the past as having Ehlers-Danlos syndrome.

The Ehlers-Danlos syndromes affect at least 13,000 people in the UK (1 in 5,000) and could affect up to 1 in 500 people.

They affect women, men and children of all races.



is the average time to diagnosis in the UK.

The Ehlers-Danlos syndromes can be difficult to recognise because some symptoms can be mistaken for other conditions.

Family history and genetics

Ehlers-Danlos syndromes are genetic conditions, meaning they can be passed on to children.

Depending on the type, there is a 25% to 50% chance the condition will be passed on.

A person can also have one of the Ehlers-Danlos syndromes when there is no family history.



Symptoms

The symptoms of the most common types include:

Fatigue

Chronic pain

regulating blood pressure

Gut, bowel problems

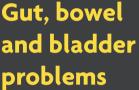
Prolapsing

Fragile, stretchy skin which damages easily

Loose joints which dislocate easily

The rarer types have additional distinctive signs and symptoms.

Difficulty







Support

The Ehlers-Danlos Support UK supports people with Ehlers-Danlos syndromes and hypermobility spectrum disorders. We provide information and support through physical and virtual support groups, a helpline, website and events. We educate medical professionals to recognise the conditions and we promote and fund research to advance knowledge about them.

Call our helpline today: **0800 907 8518**



Visit: ehlers-danlos.org