

PARENTAL QUALITY OF LIFE AND SOCIAL SUPPORT IN EDS/HSD



Study for parents/guardians of individuals
with Ehlers-Danlos Syndrome
(EDS)/Hypermobility Spectrum Disorder
(HSD) looking at their quality of life and
perceptions of the social support that they
receive.

For more information about the study or to participate:

-Scan the QR code

-Or follow this link

https://surreyfahs.eu.qualtrics.com/jfe/form/SV_87

[wCvDud9NzwsKk](https://surreyfahs.eu.qualtrics.com/jfe/form/SV_87)

-Or email ec00864@surrey.ac.uk



Requirements to take part:

- Must be a parent/guardian of an individual (of any age) with Ehlers-Danlos Syndrome/Hypermobility Spectrum Disorder
- Must live with or be in close contact with the individual who has Ehlers-Danlos Syndrome/Hypermobility Spectrum Disorder
- Must **NOT** have a diagnosis of Ehlers-Danlos Syndrome/Hypermobility Spectrum Disorder themselves or suspect that they may have it.
- Must have a good ability to read and understand English.



What the study will involve:

- Filling out an online questionnaire, which should take approximately 15-20 minutes. You will be asked questions surrounding your quality of life and the social support that you receive.



This study was reviewed and
given a favourable ethical opinion
by the University of Surrey Ethics
Committee. Ref: FHMS 22-23 274

EGA