

HALO Survey

The **HA**ematology Lived Experience and **O**utcomes (**HALO**) study aims to improve care by learning more about the long-term health, wellbeing, and quality of life of people diagnosed with beta thalassaemia, sickle cell disease, or leukaemia

Eligible participants in the North of England have been invited by post to complete the HALO study survey

By taking part, participants will help researchers to understand what life is like in adulthood, after being diagnosed with these conditions in childhood

This research will guide recommendations to help healthcare providers understand what matters to patients

We encourage anyone who is invited to share their lived experience in the HALO survey to help shape better healthcare, education, and support services

With participants' permission we will access routinely collected information using NHS England sources, national registries, educational and social care data sources

To find out more about the HALO study



www.halohaemstudy.org.uk

