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The aHUS Diagnosis Survey 2024 is now live.

In a period of some very significant news for the aHUS community, we are adding to it by launching a major Global Poll of aHUS patients and their families. We are seeking to find out what it means to have a very rare disease in 2024.

We last did this in 2014 so we will be able to compare the results to provide valuable insights.

aHUS patients, parents and other family members can participate by completing a survey questionnaire which has been put on our website and can be accessed at [this link](#).

There are 10 language versions of the questionnaire to choose from, including Arabic, English, French, German, Hindi, Italian, Portuguese, Spanish, Dutch and Turkish.

The aim is to publish the results of the Poll in a health/medical journal to spread awareness of aHUS as a rare disease through our the world.

Our community's participation is important, follow [the link](#) to find out more.

aHUS alliance Global Action



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