



EHE Global Patient Registry

EHRegistry.iamrare.org

JOIN TODAY

EHE Research - Powered by YOU

You can participate in vital Epithelioid Hemangioendothelioma (EHE) research by joining the EHE Global Patient Registry and detailing your unique experience in a natural history study of EHE.

Because EHE is so incredibly rare, this is a powerful opportunity for you to **directly impact EHE research** to improve understanding of this ultra-rare cancer and accelerate research toward improved treatments and ultimately a cure.

Who can join the Registry?

Anyone diagnosed with EHE anywhere in the world, including:

- ✓ Adults
- ✓ Children or minors (represented by a parent or guardian)
- ✓ Deceased persons (included by a legal adult representative)

How to Join

1.

Visit
EHRegistry.iamrare.org
to create your profile.

2.

Consent to provide
your information to
the registry.

3.

Answer questions about
your EHE experience, such
as symptoms, diagnosis
and treatment.

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The EHE Global Patient Registry is hosted by the National Organization for Rare Disorders (NORD®) on the IAMRARE® secure platform, and is sponsored by The EHE Foundation, a 501(c)(3) organization dedicated to pursuing effective treatments for EHE and supporting patients and their families. To learn more visit fightEHE.org.