

USHER SYNDROME COALITION

You are the **EXPERT**

TOGETHER WE CAN MAKE USHER SYNDROME HISTORY

JOIN US:

- *Register* with the **USH Trust** - be informed and involved in the latest research.
- *Access* genetic testing through **Unraveling USH**.
- *Find support* from the global community through the **USH Blue Book**.
- *Locate experts* in the **USH Yellow Book** - a centralized directory of researchers and resources worldwide.
- *Network* at **USH Connections Conferences**, the largest annual gathering of the Usher syndrome community.
- *Stay informed* through **USH Talks**, providing the latest in research developments.

THE USHER SYNDROME COALITION'S MISSION

To raise awareness and accelerate research for the most common genetic cause of combined deafness and blindness, while providing information and support to individuals and families affected by Usher syndrome.

LEARN MORE: [USHER-SYNDROME.ORG](https://usher-syndrome.org)

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USHER SYNDROME COALITION

Join the **USH TRUST**

UNRAVELING USHER SYNDROME **ONE PERSON AT A TIME**

Usher syndrome is the most common genetic cause of combined deafness and blindness. Researchers are working to find treatments and a cure. Their biggest challenge is finding enough people to participate in studies.

Of the estimated 400,000 people worldwide living with Usher syndrome, researchers are in touch with less than 1%.

The Usher Syndrome Coalition is working to change that with these valuable resources:

THE USH TRUST

The **largest international registry** of people with Usher syndrome.

- Simple - only requires your name, email and type of Usher syndrome, if known.
- Confidential - You log in. You control your information.
- HIPAA compliant and secure.
- The place to be to learn about clinical trials.

THE USH BLUE BOOK

A **global network of support** for the Usher syndrome community.

THE USH YELLOW BOOK

A **centralized directory** of researchers and resources worldwide.

JOIN NOW: [USHER-REGISTRY.ORG](https://usher-registry.org)

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