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Welcome to the US Hereditary Angioedema Association, a non-profit advocacy organization serving people with Hereditary Angioedema (HAE) and their caregivers. HAE is a very rare and potentially life-threatening genetic condition involving swelling of various parts of the body, hands, feet, or face. Our association is a community of people affected by HAE and their loved ones who are leading the fight in HAE research, advocacy and finding a cure. Through a passionate commitment to the HAE community, we offer a wide variety of services and resources that further HAE education, clinical research, community engagement, access to medications, personalized support networks, and a wide range of services to help people living with HAE lead a normal life.



**WE DELIVER THE LATEST HAE NEWS
AND RESEARCH ADVANCEMENTS.**



**WE ARE YOUR DESTINATION FOR THE
LATEST HAE INFORMATION AND
RESOURCES.**

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OUR MISSION

To lead a nationwide advocacy movement that focuses on increasing HAE awareness and education, empowering access to suitable treatment, and fostering ground-breaking research that includes searching for a cure.



OUR VISION

Unrestricted access to therapy so people affected by HAE are unburdened by symptoms and able to experience life to the fullest.



OUR VALUES

To fulfill our community's highest priority needs with innovative programs, services, and activities that are delivered with an extraordinary level of empathy, kindness, and compassion.

TAKE ACTION

Your contribution helps fund our many programs and resources for people living with HAE and their loved ones

[DONATE](#)

Join our efforts to maintain a strong political advocacy presence on Capitol Hill

[ADVOCATE](#)

Get involved and help serve our community

[VOLUNTEER](#)

Participate in one of our many ongoing initiatives, or create your own!

[FUNDRAISE](#)

TAKE ACTION

Educate medical staff
about HAE

[GET AN ER TOOLKIT](#)

Create and host an event as a fun way to get involved and help educate your community about HAE

[CREATE AN EVENT](#)

Register as a patient or caregiver member of the HAEA

[JOIN US](#)

Participate in camps, activities and other youth events!

[JOIN THE YOUTH PROGRAM](#)

IN OUR OWN WORDS



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Living with HAE



[RICHARD](#)
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HAEA.org

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The US Hereditary Angioedema Association (HAEA) is a 501(c)(3) non-profit organization.

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The HAEA is an advocacy and research organization committed to actively engaging our community in a wide variety of grassroots activities that promote HAE education and awareness. We provide personalized services to address the unique needs of people with HAE and their families, which includes helping them secure access to and reimbursement for modern HAE medicines. Our great success in supporting clinical research has resulted in a variety of FDA-approved therapeutic options. We work closely with expert physicians to continuously upgrade our community's quality of life through improving diagnosis and knowledge of the condition, and encourage an individualized approach to selecting an optimal treatment. The HAEA is product and company neutral, and continues to enthusiastically support drug discovery research aimed at the next generation of HAE therapies.