

OUR MISSION




HCU Network America strives to inform and provide resources for patients and families, create connections, influence state and federal policy, and support advancement of diagnosis and treatment for HCU and related disorders.



CONNECT WITH US



**HCU
NETWORK
AMERICA**

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-  @HCU America

HCU Network America is a registered 501c3 non profit organization dedicated to helping patients and their families affected by homocystinuria (HCU), methylenetetrahydrofolate reductase (MTHFR) and selected cobalamin deficiencies.

**EMPOWERING
PEOPLE AFFECTED BY**

**Classical
Homocystinuria**

**Cobalamin
Disorders**

**Severe
MTHFR**

**THROUGH
EDUCATION &
AWARENESS
& FIND A
CURE**



OUR GOALS

SUPPORT RESEARCH

that improves diagnosis and treatment including a cure for the disease.

CREATE CONNECTIONS

across the community and facilitate sharing of information and best practices through in-person and virtual events and discussions complications

EARLY & EFFICIENT DIAGNOSIS

to enable access to care and avoid complications

PROVIDE EDUCATIONAL RESOURCES

to patients and caregivers along with physicians and the clinics they work with to better manage the disease

- > HCU treatment and dietary approaches
- > General disease management
- > Increased access for treatment and supplements
- > Research findings and opportunities for clinical trials

CLINIC RESOURCES

Clinic Education and Resource Packets

Clinical Trial Information

PATIENT RESOURCES

New Patient Toolkits

Website & Social Media

Infographics, Checklists and Guides

Podcast, Webinars and Educational Recordings

