At Horizon, our support of the urea cycle disorder (UCD) community is part of our overall commitment to people living with rare diseases.

We actively partner with leading professional societies and patient advocacy organizations to support people living with UCDs and their families. If you or your loved one lives with a UCD, we want you to be aware of all of the resources available to you from Horizon and other organizations that Horizon supports.

Support and Assistance

Patient Support
and assistance in place so that eligible patients are able to receive our medicines regardless of their ability to pay*

* Assumes utilization of Horizon financial support programs

Connection
to help with travel to see a specialist, support through life’s transitions (changing schools, transition to adult care, changing employment), and more

UCD Genetic Testing
to speed diagnosis and improve access to genetic testing at no charge to patients through our partnership with Invitae

Community Engagement

Website and Facebook page co-created with people living with UCDs and their families provides education and resources

- Educational fact sheets and animations
- Expert videos
- Low-protein recipes
- School resources

10+
low-protein cooking classes specifically designed for people living with UCDs held each year along with tools to manage UCD including online recipes and a cookbook

50+
rare disease events sponsored and attended each year by Horizon employees

patient advocacy group partnerships globally, including leading UCD advocacy organizations
Clinical trials are research studies performed in people that are aimed at evaluating whether a medicine is safe and effective. Less than 12 percent of medicines that enter the clinical trial phase of study result in an approved medicine.
Connection to Urea Cycle Disorder and Rare Disease Support and Services

Horizon provides support to many organizations that provide a wide range of crucial programs and services for people living with urea cycle disorders and other rare diseases.

**National Organization for Rare Disorders (RareDiseases.org)**
Provides many resources for people living with rare diseases, their families and other advocates, including but not limited to:

- **Co-pay Program**: assists eligible individuals who have health insurance to cover health insurance deductibles, copayments and coinsurance costs associated with the care of urea cycle disorders.
- **Medical Assistance Program**: assists eligible individuals who are uninsured or for whom coverage has been denied, with out-of-pocket costs for medical expenses such as medical visits, laboratory and diagnostic testing, physician prescribed medical foods & supplements, travel to and from a disease-related medical appointment and other expenses for the care of this disease.
- **Emergency Relief Program**: May assist eligible individuals and/or families who need financial support when faced with limited resources to pay for unexpected or emergency non-medical expenses.

For information about each of the above programs, send an e-mail to UCD@rarediseases.org.

**Global Genes (GlobalGenes.org)**
Develops educational resources, programs, and events that unite patients, advocates, and industry experts.

- **Global Genes RARE Patient Advocacy Summit**: The largest rare patient advocate event worldwide.
- **Scholarship to the RARE Patient Advocacy Summit**: With the help of generous sponsorships, advocates from all over the globe are afforded travel and lodging scholarships to attend the Summit. E-mail events@globalgenes.org for scholarship information.

**Miracle Flights (MiracleFlights.org)**
Provides financial assistance to low-income children for commercial air travel to obtain special medical care.

**The Sing Me a Story Foundation (SingMeAStory.org)**
Combines the imaginations of children with the talents of songwriters to create stories and songs that bring joy to all those involved.

**Good Days (GoodDays.org)**
Works to improve the health and quality of life of patients with chronic disease, cancer, or other life-altering conditions. The organization may help provide families with financial and travel assistance.

**The HealthWell Foundation (HealthWellFoundation.org)**
Independent non-profit dedicated to reducing financial barriers to care for underinsured Americans with chronic and life-altering medical conditions. HealthWell offers a financial lifeline to adults and children who desperately need critical medical treatments but can’t afford them by assisting with their cost-sharing obligations.

*costs for medications are not eligible for NORD programs.

If you would like to speak with Horizon about our work – or have ideas you would like to share – please send an e-mail to connect@HorizonTherapeutics.com.

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**About Horizon**
Horizon is focused on researching, developing and commercializing medicines that address critical needs for people impacted by rare and rheumatic diseases. Our pipeline is purposeful: we apply scientific expertise and courage to bring clinically meaningful therapies to patients. We believe science and compassion must work together to transform lives. For more information on how we go to incredible lengths to impact lives, please visit [www.horizontherapeutics.com](http://www.horizontherapeutics.com), follow us @HorizonNews on Twitter, like us on Facebook or explore career opportunities on LinkedIn.