Rare diseases are often chronically debilitating and sometimes life threatening. They each affect fewer than 200,000 people nationwide. Rare diseases may seem like a small corner of health care, but together, they affect nearly 10 percent of the US population. That’s about equal to the number of Americans diagnosed with diabetes. Yet most rare diseases are not studied enough. As a result, patients with rare diseases often face years of testing or ineffective treatments.

Rare diseases affect about 25 MILLION Americans.

Source: Centers for Disease Control and Prevention

Research Addressing Questions That Matter

PCORI funds comparative clinical effectiveness research (CER) to determine which healthcare options work best for which patients, based on their needs and preferences. CER produces evidence that helps people make better-informed healthcare choices.

PATIENT
I have lupus, which causes my immune system to attack healthy organs. My lupus led me to develop kidney disease. To prevent kidney failure, I need to choose between several medications. But several factors affect how each drug will work with my body. How can I know which of the available treatments will work best for me?

CAREGIVER
My daughter has cerebral palsy and attends physical therapy once a week for two hours to help develop her motor skills. As she’s gotten older, I’ve seen less progress. Which would better help her to continue to progress or even speed up her development: attending longer weekly sessions or attending more sessions per week?

STUDY SPOTLIGHTS

Comparing Pediatric Crohn’s Disease Treatments

Crohn’s disease affects about 38,000 children in the United States. This study is comparing the experience of patients who receive a treatment called anti-TNF with those who get a combination of anti-TNF and an anti-inflammatory drug called methotrexate. It assesses how well each treatment reduces or eliminates disease symptoms and how many drug-related side effects occur. Details about this project appear at www.pcori.org/Kappelman026.

Comparing Treatments for Adults with Sickle Cell Disease

Around 100,000 Americans have sickle cell disease, a blood cell disorder that can cause episodes of severe pain and sometimes even lead to death. People with sickle cell disease often face negative attitudes and long delays when seeking treatment for pain crises in emergency departments. This study is comparing the timeliness and quality of care experienced by patients who go to emergency departments versus those who visit infusion centers. Details about this project appear at www.pcori.org/Lanzkron027.
PCORI’s Commitment to Rare Disease Research

PCORI is committed to changing the culture of research by bringing together a variety of stakeholders to help drive useful, relevant research about what works best for individual patients. Part of this commitment specifically focuses on funding research that addresses outstanding questions about the prevention, diagnosis, and treatment of rare diseases. Learn more about our rare disease resources at www.pcori.org/RD-resources.

PCORI’s Advisory Panel on Rare Disease seeks to bring voices from across health care and the rare disease community into our work on rare conditions. The panel provides a wide range of stakeholder guidance on everything from refining and prioritizing research questions for potential funding to engaging patients as partners in research. More details are available at www.pcori.org/RDAP.

As of August 2018, PCORI has awarded $87 MILLION TO FUND 29 comparative clinical effectiveness research studies related to rare diseases.

BY THE NUMBER OF PROJECTS

These studies focus on 19 diseases and conditions. Learn more at www.pcori.org/RD-projects.

- Acute myeloid leukemia
- Cerebral palsy
- Chiari type I malformation and syringomyelia
- Disorders of sex development
- Duarte galactosemia
- Eosinophilic esophagitis
- Hydrocephalus
- Idiopathic subglottic stenosis
- Kawasaki disease
- Lupus nephritis
- Myasthenia gravis
- Non-CF bronchiectasis
- Pediatric Crohn’s disease
- Pediatric transverse myelitis
- Polyarticular juvenile idiopathic arthritis
- Sickle cell disease
- Spinal cord injury and spina bifida
- Systemic scleroderma
- Urea cycle disorders

These conditions affect the following body systems:

- Cardiovascular/Circulatory
- Digestive/Excretory
- Endocrine
- Immune
- Muscular/Skeletal
- Nervous
- Reproductive
- Respiratory

PCORnet’s Promise in Improving Rare Disease Care

PCORnet is investing in people and infrastructure to support high-quality patient-centered research. PCORnet, the National Patient-Centered Clinical Research Network, enables us to learn from the data and experiences of 100 million Americans. Rare diseases are well represented in PCORnet’s 33 partner networks. Several of the Patient-Powered Research Networks, which are operated and governed by groups of patients and their partners, focus on one or more such illnesses. And each of the 13 health-system-based Clinical Data Research Networks has a rare disease as one of its conditions of primary focus.

Bringing Patients Together to Drive Research

Patient-Powered Research Networks (PPRNs) that focus on rare diseases include:

- Community-Engaged Network for All (multiple rare conditions)
- DuchenneConnect (Duchenne and Becker muscular dystrophy)
- NephCure (nephrotic syndrome)
- PARTNERS (childhood rheumatic diseases)
- Phelan-McDermid Syndrome Data Network
- PI-CONNECT (primary immunodeficiencies)
- Rare Epilepsy Network
- Vasculitis PPRN