



## Funding research

to conquer Alport syndrome by providing more than \$2 million in research funds with strategic partners



## Encouraging investment

into new therapies to treat Alport syndrome, resulting in Phase 2 and Phase 3 clinical trials



## Facilitating collaboration

in Alport syndrome research through international partnerships with all stakeholders



## Supporting the patient registry (ASTOR)

to collect data to better understand how Alport syndrome impacts lives



## Connecting patients and families

through an online support group, the Peer Support Network, and regional Alport Family Meetings



## Ensuring patients receive top-level care

by educating medical professionals with recommendations for treatment based on the latest research, and developing an accredited continuing medical education (CME) webinar for physicians



## Empowering patients

with the resources to advocate for themselves through webinars, online resources, guides and brochures



## Ensuring the patient perspective

is understood by researchers, industry, and the FDA



## Advocating for the needs of patients

with Alport syndrome on Capitol Hill



## Reaching teens and young adults,

the hardest hit age group, by connecting young patients and including special programs to instill hope and encouragement

Together we are making a difference for the future of those living with Alport syndrome. Please help us continue to advance research and empower patients and families.