

Novo Nordisk is pleased to provide this overview of non-profit organizations that can provide information, resources, and support to endocrinologists, their patients, and their families.

Please note, this is not intended to be an exhaustive list of resources, and Novo Nordisk is not endorsing or recommending any particular organization listed below.

Foundation for Prader-Willi Research

The mission of FPWR is to eliminate the challenges of Prader-Willi syndrome through the advancement of research and therapeutic development.

fpwr.org

888-322-5487 | info@fpwr.org



Partnership for Prescription Assistance

PhRMA's Medicine Assistance Tool (MAT) is a search engine designed to help patients, caregivers and health care providers learn more about the resources available through the various biopharmaceutical industry programs.

medicineassistancetool.org



Human Growth Foundation

The HGF is a global leader in endocrine research, education, patient advocacy, and support.

hgfound.org

800-451-6434 | hgf1@hgfound.org



Patient Access Network

PAN helps federally and commercially insured people living with life-threatening, chronic, and rare diseases with the out-of-pocket costs for their prescribed medications.

panfoundation.org

866-316-7263 | info@panfoundation.org



MAGIC Foundation

The MAGIC Foundation provides support services for the families of children afflicted with a wide variety of chronic and/or critical disorders, syndromes and diseases that affect a child's growth.

magicfoundation.org

800-362-4423 | contactus@magicfoundation.org



Patient Services, Inc.

PSI provides financial support and guidance for qualified patients with specific, rare chronic diseases. Through PSI assistance programs patients and their families rediscover hope and health.

patientservicesinc.org

800-366-7741



National Organization for Rare Disorders

NORD is a patient advocacy organization dedicated to individuals with rare diseases and the organizations that serve them.

rarediseases.org

800-634-7207



Pituitary Network Association

The PNA disseminates information on matters regarding early detection, symptoms, treatments and resources available to patients with pituitary disease.

pituitary.org

805-499-9973 | info@pituitary.org



Needy Meds

NeedyMeds connects people to programs that will help them afford their medications and other healthcare costs.

needymeds.org

800-503-6897



The Pituitary Society

The Pituitary Society provides scientists, clinicians and trainees with the most up-to-date, reliable research and education on the biology of the pituitary and the causes, diagnosis, and treatment of pituitary disease, in order to improve patient care, and to provide a uniquely international forum for professional growth and scientific exchange.

pituitarysociety.org

310-988-9486



Noonan Syndrome Foundation

The mission of the NSF is to support, educate, and advocate for those with Noonan Syndrome.

teamnoonan.org

info@teamnoonan.org



Prader-Willi Syndrome Association

The PWSA provides support for individuals, families, professionals and organizations to be a resource for education and information about Prader-Willi syndrome.

pwsausa.org

800-926-4797



RASopathiesNet



The mission of RASopathiesNet is to advance research of the RASopathies by bringing together families, clinicians and scientists.

rasopathiesnet.org
info@rasopathiesnet.org

Ray Tye Medical Aid Foundation

The Ray Tye Medical Aid Foundation funds in-hospital life saving medical treatment and surgeries for those who do not have medical insurance, and for which no other financial resources are available.



rtmaf.org
781-356-4300 | rtmaf.info@gmail.com

Turner Syndrome Foundation



TSF supports research initiatives and facilitates education programs that increase professional awareness and enhance medical care of those affected by Turner syndrome.

turnersyndromefoundation.org
800-594-4585 | info@tsfusa.org

Turner Syndrome Society of the United States



The TSSUS advances knowledge, facilitates research, and provides support for all persons touched by Turner syndrome. Staff members answer questions via phone and e-mail, work with local groups, create resources, plan events, and keep their website up to date and relevant.

turnersyndrome.org
800-365-9944 / 832-912-6006 | info@turnersyndrome.org