# 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

HUMAN GROWTH FOUNDATION

### **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

# **Human Growth Foundation**

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

hafound.ora 800-451-6434 | haf1@hafound.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 arowth.



magicfoundation.org 800-362-4423 | contactus@magicfoundation.org

### **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

## **Needy Meds**

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

needymeds.org 800-503-6897

### **Noonan Syndrome** Foundation

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

teamnoonan.org info@teamnoonan.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

## 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

## Patient Services, Inc.

PSI provides financial support



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

patientservicesinc.org 800-366-7741

## **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

# 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

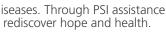
310-988-9486

## **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





Pituitary Network

Association

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National Organization

for Rare Disorders

# **RASopathiesNet**

# **RAS**opathiesNet

THE

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WE WILL NEVER STOP CARING

RAY TYE

MEDICAL AID

Turner

Society

Svndrome

of the United States

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.



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

#### **Turner Syndrome** Foundation

TSF supports research initiatives and facilitates TURNER SYNDROME FOUNDATION

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

