THE GRIN TEAM LEIPZIG

We are a team of geneticists interested in the causes of GRIN-associated diseases and have contributed significantly to the current state of knowledge of GRIN-associated disease research since 2013.
In addition to research, we also have a focus on the diagnosis of GRIN-associated diseases. This is used as part of a broad screening of patients with developmental disorders and can also be performed specifically, for example, as part of prenatal diagnostics.
In the context of diagnostics or also in the case of externally diagnosed GRIN-associated disease, we have established a special consultation hour/video consultation hour for affected persons and relatives (if necessary also online).

CONSULTATION REGISTRATION:

phone: (+49) 341 9723840

Genetic Consultation Leipzig

https://www.uniklinikum-leipzig.de/einrichtungen/humangenetik/genetische-sprechstunde

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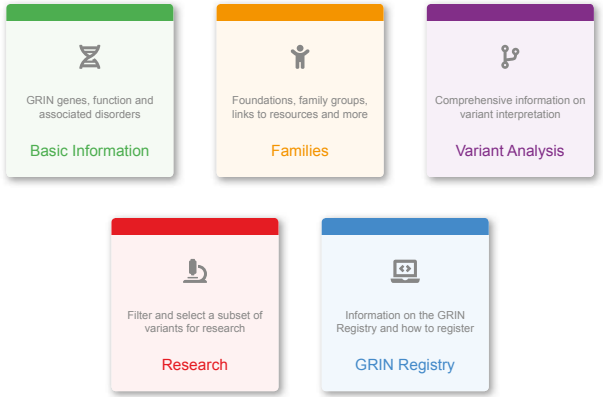
Universitätsklinikum Leipzig
Medizin ist unsere Berufung.

Information on GRIN-associated diseases

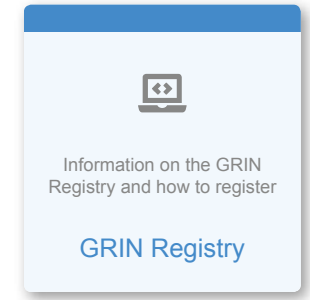
UNIVERSITÄT LEIPZIG
Medizinische Fakultät

THE GRIN PORTAL

The GRIN portal is a new, central information platform for parents, clinicians and scientists.



http://grin-portal.broadinstitute.org/



WHY A REGISTRY?

In the GRIN Registry, clinical and genetic data from affected individuals with GRIN-associated diseases are collected and analyzed in a centralized manner. This should lead to a better understanding and new insights into GRIN-associated diseases.

WHO CAN ENTER THE DATA?

The registration can be made by parents, but also by attending physicians with the consent of the parents.

WHERE CAN I FIND THE REGISTRY?

REGISTER NOW!

GRIN Registry

https://redcap.medizin.uni-leipzig.de/redcap/surveys/?s=PRAEF9N7J7

WHO IS „GEMEINSAM GRIN"?

"Gemeinsam GRIN" is a family association of people affected by GRIN-associated diseases in German-speaking countries..

WHAT DOES „GEMEINSAM GRIN"?

For affected parents and relatives who have a child diagnosed with a GRIN-associated disease, this group wants to be a contact point.

The goal is to support other affected families with questions that may arise due to the special needs of the children and to share experiences with each other.

To provide the public with knowledge about GRIN-associated diseases due to a genetic alteration in one of the GRIN genes (e.g., GRIN1, GRIN2A, GRIN2B, or GRIN2D).

You are welcome to contact us via:
Mail: Gemeinsam-grin@gmx.de
Facebook: Gemeinsam GRIN