



THE SWEDISH INFORMATION CENTRE FOR RARE DISEASES

# The Swedish Rare Disease Information Database

[www.socialstyrelsen.se/rarediseases](http://www.socialstyrelsen.se/rarediseases)

**THE SWEDISH INFORMATION CENTRE FOR RARE DISEASES** produces a database that currently includes detailed information on nearly 300 diagnoses. The Information Centre also serves as a helpline and a source of guidance to those directly or indirectly affected by rare diseases.

**THE DATABASE** is produced in cooperation with leading medical experts in Sweden. Patient organizations are also important partners. New diagnoses are continuously added and the information is updated on a regular basis. The database is published in Swedish, but many of the diagnoses are also available in English. The site is easily accessible to everyone, and there are currently more than 50,000 Swedish and international visitors every month.

**TARGET GROUPS** include people with rare diseases and their families, parent and patient organisations, professionals, researchers and public authorities.

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