Are you a patient?

Patients don't just benefit from our work - they shape it!
Read more about how to contribute.

Newsletter

Do you want to stay up-to-date?
Sign up for our newsletter!

Contact

eurreb.eu

registries@lumc.nl

regular online drop-in sessions

European Registries for Rare
Endocrine and Bone Conditions

EuRREB

European Registries for Rare
Endocrine and Bone conditions

more data, better outcomes

Endo-ERN
European Reference Network on Rare Endocrine Conditions

ERN BOND
EUROPEAN REFERENCE NETWORK ON RARE BONE DISEASES

Leiden University Medical Center

Funded by the European Union

EuRREB

Europe's Resource for Rare Endocrine and Bone Conditions

EuRREB supports healthcare professionals, researchers, and patients in collecting data through patient-centered registries and enabling collaborative research across Europe to improve care for those affected by rare endocrine and bone diseases.

Our Registries

e-REC is an electronic reporting programme that captures new clinical encounters.

In the **Core Registry** we collect more detailed and longitudinal data.

Do you want to contribute? Register here:

See which centres are already participating!

Research

Research drives EuRREB

Through patient-centered registries we enable collaborative research across Europe.

Condition-specific modules

These cover aspects of specific rare conditions such as diagnosis, therapy, and monitoring, and are built into the Core Registry for dynamic and longitudinal collection of data.

Active studies

A study is conducted by experts, including patient representatives, with data from the different condition-specific modules.

Publications

Read the publications of our studies.

Do you want to use our data? Send us an email: **registries@lumc.nl**