

GeneReviews®



Free, current, expert-authored and peer-reviewed descriptions of heritable disorders

ABOUT GENEREVIEW®

- The busy clinician's #1 source for diagnosis, management, and genetic counseling information
- 725+ chapters with links to disease-specific and general consumer resources
- Includes chapters with a single-gene or -phenotype focus and overviews covering the genetic causes of common conditions
- Educational materials to clarify genetics concepts for patients and general clinicians
- Resources for genetics professionals on genetic testing, including founder variants and direct-to-consumer testing
- New and/or updated chapters published weekly
- Content linked to ClinVar, MedGen, GTR®, PheGenI, Medical Genetics Summaries, and dbGaP

Search GeneReviews® By:



Disorder



Text



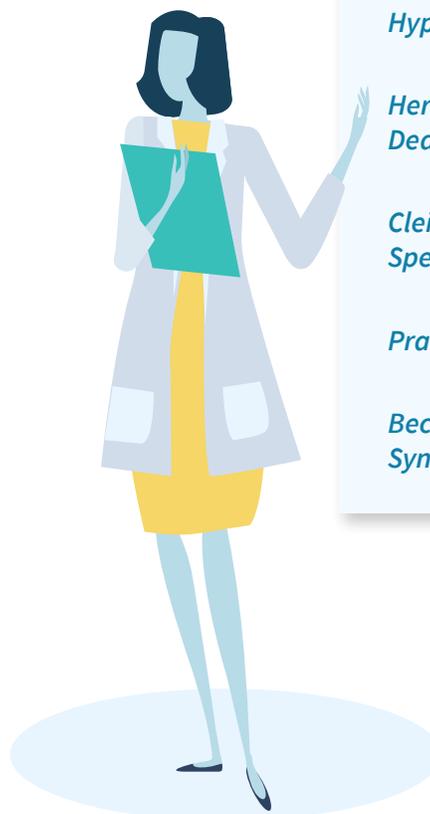
Author



Gene Symbol



Protein Name



GeneReviews® Chapters

*Ehlers-Danlos Syndrome,
Hypermobility Type*

*Hereditary Hearing Loss and
Deafness Overview*

*Cleidocranial Dysplasia
Spectrum Disorder*

Prader-Willi Syndrome

*Beckwith-Wiedemann
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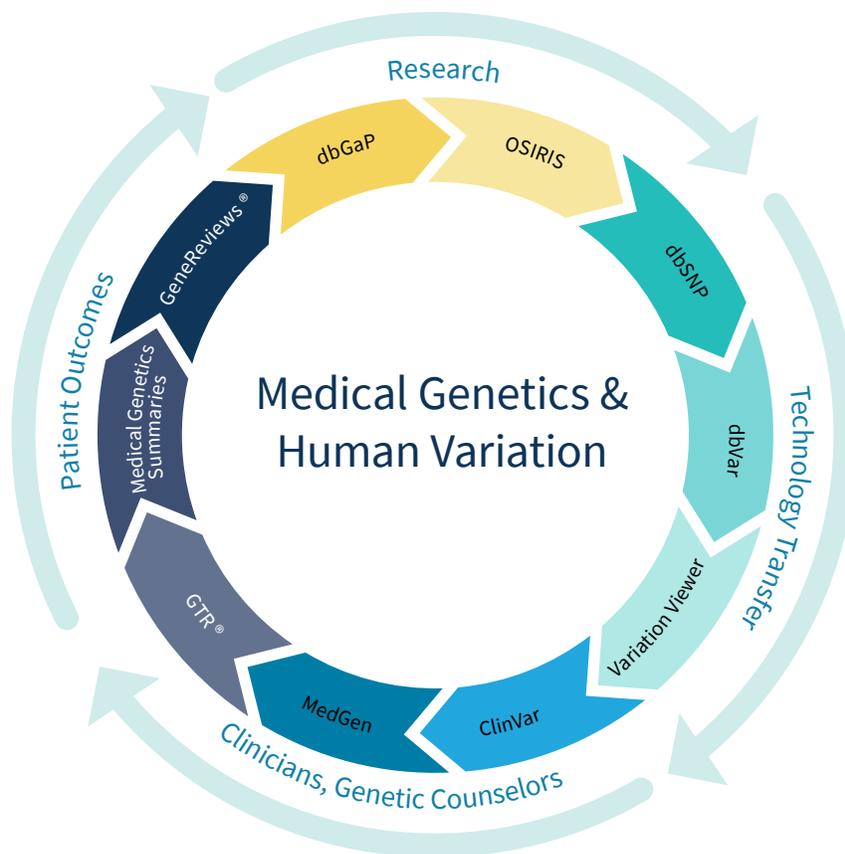
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ClinVar

user-submitted database for information about genomic variation and its relationship to human health.



dbGaP

database for genotype and phenotype research studies.



dbSNP and dbVar

databases of small and large genomic variants including both common variations and clinical mutations.



GTR®

provider-submitted database of clinical and research molecular, cytogenetic and biochemical genetic tests and supporting information.



MedGen

aggregates information from and provides access to authoritative medical genetics resources.



Medical Genetics Summaries and GeneReviews®

up-to-date, peer-reviewed, medically actionable summaries for heritable diseases and pharmacogenetics.



OSIRIS

Open source short tandem repeat (STR) analysis tool for forensic, clinical and research use.



Variation Viewer

interactive browser for examination of nucleotide variants in a genomic context.