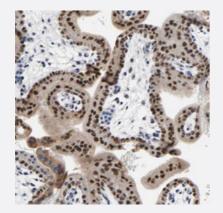


GRHL2 polyclonal antibody

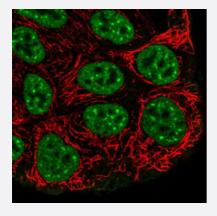
Catalog # PAB30713 Size 100 uL

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human placenta with GRHL2 polyclonal antibody (Cat # PAB30713) shows strong nuclear positivity in trophoblastic cells.



Immunofluorescence

Immunofluorescent staining of RT-4 cells with GRHL2 polyclonal antibody (Cat # PAB30713) (Green) shows positivity in plasma membrane and nucleus but excluded from the nucleoli.

Specification	
Product Description	Rabbit polyclonal antibody raised against partial recombinant human GRHL2.
Immunogen	Recombinant protein corresponding to human GRHL2.
Sequence	NRVQVLKTVPVNLSLNQDHLENSKREQYSISFPESSAIIPVSGITVVKAEDFTPVFMAPPVHYPRG DGEEQRVVIFEQTQYDVPSLATHSAYLKDDQRSTPDSTYSESFKDAATEKFRSASVGAEEYMYD QTSSGTFQYTLEATKSLRQK
Host	Rabbit



Product Information

Reactivity	Human
Form	Liquid
Purification	Antigen affinity purification
Isotype	lgG
Recommend Usage	Immunofluorescence (1-4 ug/mL) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (1:500-1:1000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide).
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Gene Info — GRHL2	
Entrez GenelD	<u>79977</u>
Protein Accession#	Q6ISB3
Gene Name	GRHL2
Gene Alias	BOM, DFNA28, FLJ11172, FLJ13782, MGC149294, MGC149295, TFCP2L3
Gene Description	grainyhead-like 2 (Drosophila)
Omim ID	<u>608576</u> <u>608641</u>
Gene Ontology	<u>Hyperlink</u>



Product Information

Gene Summary	The protein encoded by this gene is a transcription factor that can act as a homodimer or as a het erodimer with either GRHL1 or GRHL3. Defects in this gene are a cause of non-syndromic senso rineural deafness autosomal dominant type 28 (DFNA28)
Other Designations	transcription factor CP2-like 3

Disease

- Genetic Predisposition to Disease
- Presbycusis
- Tobacco Use Disorder