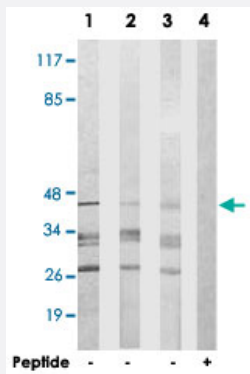


GSC2 polyclonal antibody

Catalog # PAB17336 Size 100 ug

Applications



Western Blot (Cell lysate)

Western blot analysis of extracts from 293 cells (Lane 1), HepG2 cells (Lane 2) and COLO 205 cells (Lane 3 and 4), using GSC2 polyclonal antibody (Cat # PAB17336).

Peptide "+" means "with peptide blocking".

Specification

Product Description	Rabbit polyclonal antibody raised against synthetic peptide of GSC2.
Immunogen	A synthetic peptide corresponding to internal of human GSC2.
Host	Rabbit
Reactivity	Human, Mouse
Specificity	This antibody detects endogenous levels of total GSC2 protein.
Form	Liquid
Recommend Usage	Western Blot (1:500-1:1000) ELISA (1:20000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.4 (150mM NaCl, 0.02% sodium azide, 50% glycerol)
Storage Instruction	Store at -20°C. Aliquot to avoid repeated freezing and thawing.

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot (Cell lysate)

Western blot analysis of extracts from 293 cells (Lane 1), HepG2 cells (Lane 2) and COLO 205 cells (Lane 3 and 4), using GSC2 polyclonal antibody (Cat # PAB17336).

Peptide "+" means "with peptide blocking".

Gene Info — GSC2

Entrez GeneID	2928
---------------	----------------------

Protein Accession#	O15499
--------------------	------------------------

Gene Name	GSC2
-----------	------

Gene Alias	GSCL
------------	------

Gene Description	goosecoid homeobox 2
------------------	----------------------

Omim ID	601845
---------	------------------------

Gene Ontology	Hyperlink
---------------	---------------------------

Gene Summary	Goosecoidlike (GSCL), a homeodomain-containing gene, resides in the critical region for VCFS/DGS on 22q11. Velocardiofacial syndrome (VCFS) is a developmental disorder characterized by conotruncal heart defects, craniofacial anomalies, and learning disabilities. VCFS is phenotypically related to DiGeorge syndrome (DGS) and both syndromes are associated with hemizygous 22q11 deletions. Because many of the tissues and structures affected in VCFS/DGS derive from the pharyngeal arches of the developing embryo, it is believed that haploinsufficiency of a gene involved in embryonic development may be responsible for its etiology. The gene is expressed in a limited number of adult tissues, as well as in early human development. [provided by RefSeq]
--------------	---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

Other Designations	goosecoid-like
--------------------	----------------

Publication Reference

- [Characterization and mutation analysis of goosecoid-like \(GSCL\), a homeodomain-containing gene that maps to the critical region for VCFS/DGS on 22q11.](#)

Funke B, Saint-Jore B, Puech A, Sirotkin H, Edelmann L, Carlson C, Raft S, Pandita RK, Kucherlapati R, Skoultschi A, Morrow BE.

Genomics 1997 Dec; 46(3):364.

- [The DiGeorge syndrome minimal critical region contains a goosecoid-like \(GSCL\) homeobox gene that is expressed early in human development.](#)

Gottlieb S, Emanuel BS, Driscoll DA, Sellinger B, Wang Z, Roe B, Budarf ML.

American Journal of Human Genetics 1997 May; 60(5):1194.