

DNAXPAb

Hard-to-Find
Antibody

CSH1 DNAXPab

Catalog # H00001442-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human CSH1 DNA using DNAX™ Immune technology.
Technology	DNAX™ Immune
Immunogen	Full-length human DNA
Sequence	MAPGSRTSLLAFALLCLPWLQEAGAVQTVPLSRLFDHAMLQAHRAHQLAIDTYQEFEETYIPKD QKYSFLHDSQTSFCFSDSIPTPSNMEETQQKSNLELLRISLLIESWLEPVRFRLRSMFANNLVYDTS DSDDYHLLKDLEEGIQTLMGRLEDGSRRTGQILKQTYSKFDTNSHNHDALLKNYGLLYCFRKDMD KVETFLRMVQCRSVEGSCGF
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — CSH1

Entrez GeneID [1442](#)

GeneBank Accession# [NM_001317.3](#)

Protein Accession# [NP_001308.1](#)

Gene Name CSH1

Gene Alias CSA, CSMT, FLJ75407, PL

Gene Description chorionic somatomammotropin hormone 1 (placental lactogen)

Omim ID [150200](#)

Gene Ontology [Hyperlink](#)

Gene Summary

The protein encoded by this gene is a member of the somatotropin/prolactin family of hormones and plays an important role in growth control. The gene is located at the growth hormone locus on chromosome 17 along with four other related genes in the same transcriptional orientation; an arrangement which is thought to have evolved by a series of gene duplications. Although the five genes share a remarkably high degree of sequence identity, they are expressed selectively in different tissues. Alternative splicing generates additional isoforms of each of the five growth hormones, leading to further diversity and potential for specialization. This particular family member is expressed mainly in the placenta and utilizes multiple transcription initiation sites. Expression of the identical mature proteins for chorionic somatomammotropin hormones 1 and 2 is upregulated during development, although the ratio of 1 to 2 increases by term. Mutations in this gene result in placental lactogen deficiency and Silver-Russell syndrome. [provided by RefSeq]

Other Designations CS-1|choriomammotropin|chorionic somatomammotropin A|chorionic somatomammotropin hormone 1|hCS-A|placental lactogen

Pathway

- [Cytokine-cytokine receptor interaction](#)
- [Jak-STAT signaling pathway](#)
- [Neuroactive ligand-receptor interaction](#)

Disease

- [Birth Weight](#)

- [Body Weight](#)
- [Breast cancer](#)
- [Breast Neoplasms](#)
- [Genetic Predisposition to Disease](#)
- [Metabolic Syndrome X](#)