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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

CSH1 (Human) Recombinant Protein

Catalog Number: P9029

Regulation Status: For research use only (RUO)

Product Description: Human CSH1 (P0DML2) recombinant protein with Ala at N-Terminus expressed in *Escherichia coli*.

Host: *Escherichia coli*

Theoretical MW (kDa): 22.4

Protocols: See our web site at <http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Form: Lyophilized

Preparation Method: *Escherichia coli* expression system

Purity: > 99% by gel filtration analysis and SDS PAGE

Activity: Placental Lactogen Human is biologically active as evidenced by inducing proliferation of Nb2 cells.

Storage Buffer: Lyophilized from 0.02-0.03% NaHCO₃.

Storage Instruction: Store at -20°C. Aliquot the product after reconstitution to avoid repeated freezing/thawing cycles.

Entrez GeneID: 1442

Gene Symbol: CSH1

Gene Alias: CSA, CSMT, FLJ75407, PL

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]