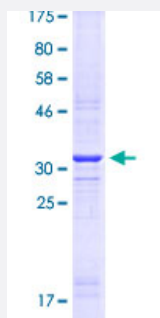


MSX1 (Human) Recombinant Protein (Q01)

Catalog # H00004487-Q01

Size 25 ug, 10 ug

Applications



Specification

Product Description	Human MSX1 partial ORF (NP_002439.1, 216 a.a. - 297 a.a.) recombinant protein with GST tag at N-terminal.
Sequence	NRRAKAKRLQEAELEKLKMAAKPMLPPAAFGLSFPLGGPAAVAAAAGASLYGASGPFQRAALP VAPVGLYTAHVGYSMYHLT
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	34.65
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — MSX1

Entrez GeneID [4487](#)

GeneBank Accession# [NM_002448.1](#)

Protein Accession# [NP_002439.1](#)

Gene Name MSX1

Gene Alias HOX7, HYD1

Gene Description msh homeobox 1

Omim ID [106600](#) [142983](#) [189500](#) [608874](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes a member of the muscle segment homeobox gene family. The encoded protein functions as a transcriptional repressor during embryogenesis through interactions with components of the core transcription complex and other homeoproteins. It may also have roles in limb-pattern formation, craniofacial development, particularly odontogenesis, and tumor growth inhibition. Mutations in this gene, which was once known as homeobox 7, have been associated with nonsyndromic cleft lip with or without cleft palate 5, Witkop syndrome, Wolf-Hirschorn syndrome, and autosomal dominant hypodontia. [provided by RefSeq]

Other Designations OTTHUMP00000115387|homeobox 7|msh homeobox 1|msh homeobox homolog 1

Disease

- [Breast Neoplasms](#)
- [Cleft Lip](#)
- [Cleft Palate](#)

- [Disease Models](#)
- [Genetic Predisposition to Disease](#)
- [Multiple System Atrophy](#)
- [Parkinson disease](#)
- [Sleep Apnea](#)