

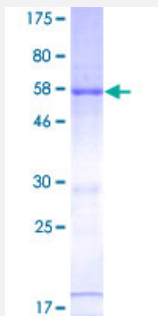
Full-Length

RNASEH2A (Human) Recombinant Protein (P01)

Catalog # H00010535-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human RNASEH2A full-length ORF (AAH11748, 1 a.a. - 299 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MDLSELERDNTGRCRLSSPVPVAVCRKEPCVLGVDEAGRGPVLGPMVYAICYCPLPRLADLEALK
VADSKTLLESERERLFAKMEDTDFVGWALDVLSPNLISTSMLGRVKYNLSLSDTATGLIQYALD
QGVNVTQVFVDTVGMPETYQARLQSFPGIEVTVKAKADALYPVVSAAISAKVARDQAVKKWQ
FVEKLQDLTDYSGGYNDPKTKAWLKEHVEPVFGFPQFVRFSWRTAQ TILEKEAEDVIWEDSA
SENQEGLRKITSYFLNEGSQARPRSSHRYFLERGLSATSL

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

58.63

Interspecies Antigen Sequence

Rat (84)

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 — RNASEH2A

Entrez GeneID[10535](#)**GeneBank Accession#**[BC011748](#)**Protein Accession#**[AAH11748](#)**Gene Name**

RNASEH2A

Gene Alias

AGS4, JUNB, RNASEHI, RNHIA, RNHL

Gene Description

ribonuclease H2, subunit A

Omim ID[606034 610333](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

The protein encoded by this gene is a component of the heterotrimeric type II ribonuclease H enzyme (RNaseH2). RNaseH2 is the major source of ribonuclease H activity in mammalian cells and endonucleolytically cleaves ribonucleotides. It is predicted to remove Okazaki fragment RNA primers during lagging strand DNA synthesis and to excise single ribonucleotides from DNA-DNA duplexes. Mutations in this gene cause Aicardi-Goutieres Syndrome (AGS), a an autosomal recessive neurological disorder characterized by progressive microcephaly and psychomotor retardation, intracranial calcifications, elevated levels of interferon-alpha and white blood cells in the cerebrospinal fluid

Other Designations

ribonuclease H2, large subunit|ribonuclease HI, large subunit

Pathway

- [DNA replication](#)