

DDX20 Antibody (C-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP14870b**Specification**

DDX20 Antibody (C-term) - Product Information

Application	WB,E
Primary Accession	O9UHI6
Other Accession	NP_009135.4
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	92241
Antigen Region	713-741

DDX20 Antibody (C-term) - Additional Information**Gene ID** 11218**Other Names**

Probable ATP-dependent RNA helicase DDX20, Component of gems 3, DEAD box protein 20, DEAD box protein DP 103, Gemin-3, DDX20, DP103, GEMIN3

Target/Specificity

This DDX20 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 713-741 amino acids from the C-terminal region of human DDX20.

Dilution

WB~~1:1000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

DDX20 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

DDX20 Antibody (C-term) - Protein Information**Name** DDX20**Synonyms** DP103, GEMIN3

Function The SMN complex catalyzes the assembly of small nuclear ribonucleoproteins (snRNPs), the building blocks of the spliceosome, and thereby plays an important role in the splicing of cellular pre- mRNAs. Most spliceosomal snRNPs contain a common set of Sm proteins SNRNPB, SNRPD1, SNRPD2, SNRPD3, SNRPE, SNRPF and SNRPG that assemble in a heptameric protein ring on the Sm site of the small nuclear RNA to form the core snRNP (Sm core). In the cytosol, the Sm proteins SNRPD1, SNRPD2, SNRPE, SNRPF and SNRPG are trapped in an inactive 6S pICln-Sm complex by the chaperone CLNS1A that controls the assembly of the core snRNP. To assemble core snRNPs, the SMN complex accepts the trapped 5Sm proteins from CLNS1A forming an intermediate. Binding of snRNA inside 5Sm triggers eviction of the SMN complex, thereby allowing binding of SNRPD3 and SNRNPB to complete assembly of the core snRNP. May also play a role in the metabolism of small nucleolar ribonucleoprotein (snoRNPs).

Cellular Location

Cytoplasm. Nucleus, gem Note=Localized in subnuclear structures next to coiled bodies, called Gemini of Cajal bodies (Gems).

Tissue Location

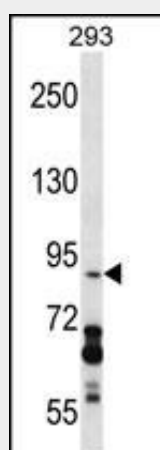
Ubiquitous.

DDX20 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

DDX20 Antibody (C-term) - Images



DDX20 Antibody (C-term) (Cat. #AP14870b) western blot analysis in 293 cell line lysates (35ug/lane). This demonstrates the DDX20 antibody detected the DDX20 protein (arrow).

DDX20 Antibody (C-term) - Background

DEAD box proteins, characterized by the conserved motif

Asp-Glu-Ala-Asp (DEAD), are putative RNA helicases. They are implicated in a number of cellular processes involving alteration of RNA secondary structure such as translation initiation, nuclear and mitochondrial splicing, and ribosome and spliceosome assembly. Based on their distribution patterns, some members of this family are believed to be involved in embryogenesis, spermatogenesis, and cellular growth and division. This gene encodes a DEAD box protein, which has an ATPase activity and is a component of the survival of motor neurons (SMN) complex. This protein interacts directly with SMN, the spinal muscular atrophy gene product, and may play a catalytic role in the function of the SMN complex on RNPs.

DDX20 Antibody (C-term) - References

Todd, A.G., et al. J. Mol. Biol. 401(5):681-689(2010)
Sun, X., et al. Cell Stress Chaperones 15(5):567-582(2010)
Wilker, E.H., et al. Environ. Health Perspect. 118(7):943-948(2010)
Boni, V., et al. Pharmacogenomics J. (2010) In press :
Ye, Y., et al. Cancer Prev Res (Phila) 1(6):460-469(2008)