



DDX3Y Polyclonal Antibody

catalog number: E-AB-63587

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human; Mouse; Rat

Immunogen Recombinant fusion protein of human DDX3Y (NP 001116137.1).

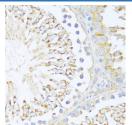
Host Rabbit Isotype IgG

Purification Affinity purification

Buffer Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

Applications	Recommended Dilution
IHC	1:50-1:100
IF	1:50-1:100

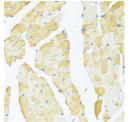
Data



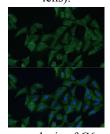


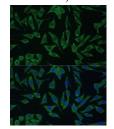
Immunohistochemistry of paraffin-embedded Rat testis using Immunohistochemistry of paraffin-embedded Rat brain using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens). DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens).





Immunohistochemistry of paraffin-embedded Mouse heart Immunohistochemistry of paraffin-embedded Mouse brain using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens) lens).





Immunofluorescence analysis of C6 cells using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

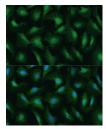
Immunofluorescence analysis of L929 cells using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

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Immunofluorescence analysis of U-2 OS cells using DDX3Y Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

Preparation & Storage

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Shipping The product is shipped with ice pack, upon receipt, store it immediately at the

temperature recommended.

Background

The protein encoded by this gene is a member of the DEAD-box RNA helicase family, characterized by nine conserved motifs, included the conserved Asp-Glu-Ala-Asp (DEAD) motif. These motifs are thought to be involved in ATP binding, hydrolysis, RNA binding, and in the formation of intramolecular interactions. This protein shares high similarity to DDX3X, on the X chromosome, but a deletion of this gene is not complemented by DDX3X. Mutations in this gene result in male infertility, a reduction in germ cell numbers, and can result in Sertoli-cell only sydrome. Pseudogenes sharing similarity to both this gene and the DDX3X paralog are found on chromosome 4 and the X chromosome. Alternative splicing results in multiple transcript variants encoding different isoforms.

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