

Recombinant

DGRmAb®

DDX39B (DGR36011) Rabbit mAb (PBS Only)

db16119-PBS

Package : 10µg 100µg

Product Name : DDX39B (DGR36011) Rabbit mAb (PBS Only)**Cat.No.:** db16119-PBS**Synonyms** : BAT1; UAP56; D6S81E**Application** : WB, IHC-P, ICC/IF, FC**Reactivity** : Human,Mouse,Rat**Host species** : Rabbit**Background**

This gene encodes a member of the DEAD box family of RNA-dependent ATPases that mediate ATP hydrolysis during pre-mRNA splicing. The encoded protein is an essential splicing factor required for association of U2 small nuclear ribonucleoprotein with pre-mRNA, and it also plays an important role in mRNA export from the nucleus to the cytoplasm. This gene belongs to a cluster of genes localized in the vicinity of the genes encoding tumor necrosis factor alpha and tumor necrosis factor beta. These genes are all within the human major histocompatibility complex class III region. Mutations in this gene may be associated with rheumatoid arthritis. Alternative splicing results in multiple transcript variants. Related pseudogenes have been identified on both chromosomes 6 and 11. Read-through transcription also occurs between this gene and the upstream ATP6V1G2 (ATPase, H⁺ transporting, lysosomal 13kDa, V1 subunit G2) gene. [provided by RefSeq, Feb 2011]

Immunogen

Recombinant protein of human DDX39B

Gene ID

7919

Swiss Prot

Q13838

Synonyms

BAT1; UAP56; D6S81E

Reactivity

Human,Mouse,Rat

Application

WB, IHC-P, ICC/IF, FC

Calculated MW

49 kDa

Observed MW

49 kDa

Host species

Rabbit

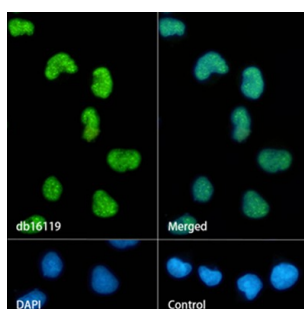
Clonality

Monoclonal

Clonality No.

DGR36011

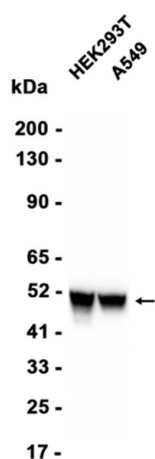
Isotype	IgG
Purity	Affinity Purification
Conjugation	Un-conjugated
Concentration	1 mg/mL
Formulation	PBS Only
Storage Stability	Store at -20°C. Recommended to aliquot into single-use vials. Supplied in 1X PBS (pH 7.4). BSA and Azide Free. Stable for 12 months from date of receipt.



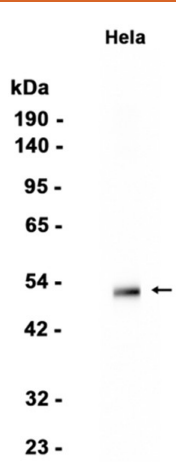
Immunofluorescence analysis of HeLa cells labelling DDX39B with [db16119](#).

The cells were fixed with cold 100% methanol (10min, 4°C) and blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween 20 for 1h. The cells were then incubate with [db16119](#) (1:100) at room temprature for 1h, followed by a further incubation at room temperature for 45min with Goat Anti Rabbit IgG (H+L)-AF488 [db10005](#), shown in green). Nuclear DNA was labeled in blue with DAPI.

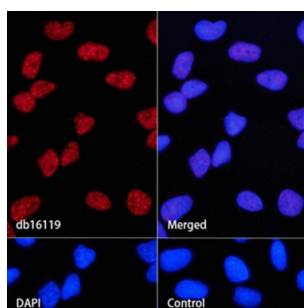
Control: Secondary antibody only.



Western blot analysis of extracts from HEK293T, A549 cells using [db16119](#) at 1:500.



Western blot analysis of extracts from HeLa cells using [db16119](#) at 1:1000.



Immunofluorescence analysis of HeLa cells labelling DDX39B with [db16119](#).

The cells were fixed with 4% PFA (10min, RT) followed by treatment with 0.1% Triton X-100 (10min, RT), and blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween 20 for 1h. The cells were then incubate with [db16119](#) (1:100) at room temprature for 1h, followed by a further incubation at room temperature for 45min with Goat Anti Rabbit IgG (H+L)-AF647[db10006](#), shown in red). Nuclear DNA was labeled in blue with DAPI.

Control: Secondary antibody only.