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## 53BP1 Rabbit pAb

Catalog No.: A5757 5 Publications

## **Basic Information**

#### **Observed MW**

450kDa

## **Calculated MW**

214kDa

#### Category

Primary antibody

## **Applications**

ELISA, WB, IHC-P

#### **Cross-Reactivity**

Human, Mouse, Rat

## **Background**

This gene encodes a protein that functions in the DNA double-strand break repair pathway choice, promoting non-homologous end joining (NHEJ) pathways, and limiting homologous recombination. This protein plays multiple roles in the DNA damage response, including promoting checkpoint signaling following DNA damage, acting as a scaffold for recruitment of DNA damage response proteins to damaged chromatin, and promoting NHEJ pathways by limiting end resection following a double-strand break. These roles are also important during V(D)J recombination, class switch recombination and at unprotected telomeres. Alternative splicing results in multiple transcript variants encoding different isoforms.

## **Recommended Dilutions**

**WB** 1:500 - 1:1000

**IHC-P** 1:50 - 1:200

## **Immunogen Information**

 Gene ID
 Swiss Prot

 7158
 Q12888

## **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 895-1031 of human TP53BP1 (NP\_005648.1).

### **Synonyms**

p202; 53BP1; TDRD30; p53BP1

#### **Contact**

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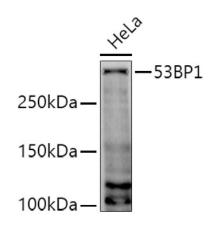
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

## Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.



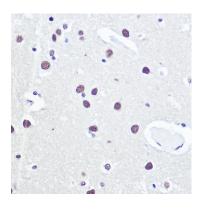
Western blot analysis of extracts of HeLa cells, using 53BP1 antibody (A5757) at 1:1000 dilution.

Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Enhanced Kit (RM00021).

Exposure time: 90s.



Immunohistochemistry analysis of paraffin-embedded human brain using 53BP1 Rabbit pAb (A5757) at dilution of 1:100 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.