

## **Product datasheet for AP50446PU-N**

## OriGene Technologies, Inc.

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## C12orf48 (PARPBP) (C-term) Rabbit Polyclonal Antibody

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** FC, IHC, WB

Recommended Dilution: ELISA: 1/1000.

Western blotting: 1/100 - 1/500.

Immunohistochemistry on paraffin sections: 1/50 - 1/100.

Flow Cytometry: 1/10 - 1/50.

Reactivity: Human Rabbit

**Isotype:** lg

Clonality: Polyclonal

**Immunogen:** KLH conjugated synthetic peptide between 529-558 amino acids from the C-terminal region

of human C12orf48

**Specificity:** This antibody reacts to C12orf48.

**Formulation:** PBS containing 0.09% (W/V) sodium azide as preservative

State: Aff - Purified

State: Liquid purified Ig fraction

**Concentration:** lot specific

**Purification:** Affinity chromatography on Protein A

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

**Gene Name:** PARP1 binding protein

Database Link: Entrez Gene 55010 Human

Q9NWS1

Synonyms: C12orf48, PARI, PARP-1 binding protein, PARP1-binding protein, PCNA-interacting partner

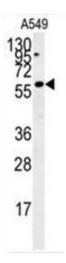
Note: Molecular Weight: 65054 Da



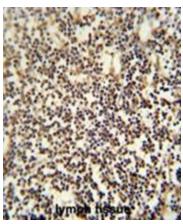


**Protein Families:** Druggable Genome

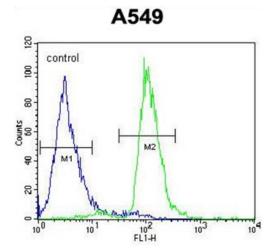
## **Product images:**



C12orf48 Antibody (C-term) western blot analysis in A549 cell line lysates (35ug/lane). This demonstrates the C12orf48 antibody detected the FBXO34 protein (arrow).



C12orf48 antibody (C-term) immunohistochemistry analysis in formalin fixed and paraffin embedded human lymph tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of the C12orf48 antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.



C12orf48 Antibody (C-term) flow cytometric analysis of A549 cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

