

## **Product datasheet for AP53317PU-N**

## PLEKHH2 (C-term) Rabbit Polyclonal Antibody

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** FC, IHC, WB

Recommended Dilution: ELISA: 1/1000.

**Western Blot:** 1/100-1/500. **Flow Cytometry:** 1/10-50.

**Immunohistochemistry on Paraffin Sections:** 1/50-1/100.

Reactivity: Human
Host: Rabbit

**Isotype:** lg

Clonality: Polyclonal

Immunogen: KLH conjugated synthetic peptide between 1057~1086 amino acids from the C-terminal

region of human PLEKHH2

**Specificity:** This antibody recognizes Human PLEKHH (C-term).

Formulation: PBS containing 0.09% (W/V) Sodium Azide as preservative

State: Aff - Purified

State: Liquid purified Ig fraction

**Concentration:** lot specific

**Purification:** Protein A column, followed by peptide affinity purification

Conjugation: Unconjugated

Storage: Store 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: pleckstrin homology, MyTH4 and FERM domain containing H2

Database Link: Entrez Gene 130271 Human

Q8IVE3

Synonyms: KIAA2028

Note: Molecular Weight: 168229 Da



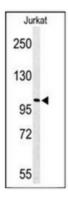
**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

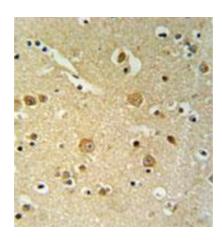
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## **Product images:**

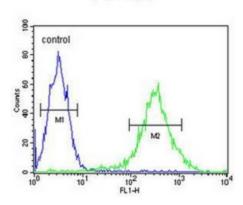


Western blot analysis of PLEKHH2 Antibody (Cterm) in Jurkat cell line lysates (35ug/lane). PKHH2 (arrow) was detected using the purified Pab.



Immunohistochemistry analysis in formalin fixed and paraffin embedded brain tissue reacted with PLEKHH2 Antibody (C-term), which was peroxidase conjugated to the secondary antibody and followed by DAB staining. This data demonstrates the use of the APKHH2 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.

## Jurkat



Flow cytometric analysis of Jurkat cells using PLEKHH2 Antibody (C-term) (right histogram) compared to a negative control cell (left histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.