



## Perforin 1 rabbit pAb

Cat#: orb771385 (Manual)

For research use only. Not intended for diagnostic use.

**Product Name** Perforin 1 rabbit pAb

**Host species** Rabbit

**Applications** WB;IHC;ELISA

**Species Cross-Reactivity** Human; Rat; Mouse;

**Recommended dilutions** WB 1:500-2000, IHC 1:50-200, ELISA 1:10000-20000

**Immunogen** The antiserum was produced against synthesized peptide derived from the C-

terminal region of human PRF1. AA range:451-500

Perforin 1 Polyclonal Antibody detects endogenous levels of Perforin 1 **Specificity** 

**Formulation** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium

azide..

Store at -20°C. Avoid repeated freeze-thaw cycles. **Storage** 

**Protein Name** Perforin 1

Gene Name PRF1

Cellular localization Cytolytic granule . Secreted. Cell membrane ; Multi-pass membrane protein .

Endosome lumen . Stored in cytolytic granules of cytolytic T-lymphocytes and secreted into the cleft between T-lymphocyte and target cell (PubMed:20038786). Inserts into the cell membrane of target cells and forms

pores (PubMed:20889983). Membrane insertion and pore formation requires a major conformation change (PubMed:20889983). May be taken up via endocytosis involving clathrin-coated vesicles and accumulate in a first time

in large early endosomes (PubMed:20038786). .



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Purification The antibody was affinity-purified from rabbit antiserum by affinity-

epitope-specific immunogen. chromatography using

**Clonality** Polyclonal

Concentration 1 mg/ml

**Observed band** 61kD

**Human Gene ID** 5551

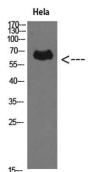
**Human Swiss-Prot Number** P14222

Perforin-1 (P1) (Cytolysin) (Lymphocyte pore-forming protein) (PFP) **Alternative Names** 

**Background** 

The protein encoded by this gene has structural and functional similarities to complement component 9 (C9). Like C9, this protein creates transmembrane tubules and is capable of lysing non-specifically a variety of target cells. This protein is one of the main cytolytic proteins of cytolytic granules, and it is known to be a key effector molecule for T-cell- and natural killer-cell-mediated cytolysis. Defects in this gene cause familial hemophagocytic lymphohistiocytosis type 2 (HPLH2), a rare and lethal autosomal recessive disorder of early childhood. Alternative splicing results in multiple transcript variants encoding the same protein [provided by RefSeq. Jul 2008]

variants encoding the same protein. [provided by RefSeq, Jul 2008],

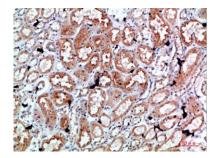


Western Blot analysis of Hela cells using Perforin 1 Polyclonal Antibody diluted at 1:500. Secondary antibody(catalog#:RS0002) was diluted at 1:20000

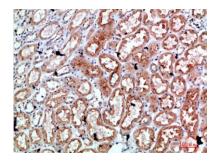




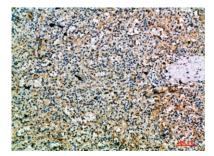
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Immunohistochemical analysis of paraffin-embedded human-kidney, antibody was diluted at 1:200



Immunohistochemical analysis of paraffin-embedded human-kidney, antibody was diluted at 1:200



 $Immunohistochemical \ analysis \ of \ paraffin-embedded \ human-spleen, \ antibody \ was \ diluted \ at 1:200$