

**ITPRIPL1 Antibody (N-term) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP16909a****Specification**

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**ITPRIPL1 Antibody (N-term) Blocking Peptide - Product Information**Primary Accession [Q6GPH6](#)**ITPRIPL1 Antibody (N-term) Blocking Peptide - Additional Information****Gene ID** 150771**Other Names**

Inositol 1, 5-trisphosphate receptor-interacting protein-like 1, ITPRIPL1, KIAA1754L

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**ITPRIPL1 Antibody (N-term) Blocking Peptide - Protein Information****Name** ITPRIPL1**Synonyms** KIAA1754L**Cellular Location**

Membrane; Single-pass type I membrane protein

**ITPRIPL1 Antibody (N-term) Blocking Peptide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

**ITPRIPL1 Antibody (N-term) Blocking Peptide - Images****ITPRIPL1 Antibody (N-term) Blocking Peptide - Background**

ITPRIPL1 (inositol 1,4,5-trisphosphate receptor-interacting protein-like 1), also known as KIAA1754L, is a 555 amino acid protein belonging to the ITPRIP family. ITPRIPL1 is a single-pass type I membrane protein expressed as two isoforms produced by alternative splicing events. The gene

that encodes ITPRIPL1 maps to human chromosome 2, the second largest human chromosome, consisting of 237 million bases encoding over 1,400 genes and making up approximately 8% of the human genome. A number of genetic diseases are linked to genes on chromosome 2. Harlequin ichthyosis, a rare and morbid skin deformity, is associated with mutations in the ABCA12 gene. The lipid metabolic disorder sitosterolemia is associated with ABCG5 and ABCG8. An extremely rare recessive genetic disorder, Alström syndrome is due to mutations in the ALMS1 gene. It has been hypothesized that human chromosome 2 is the result of an ancient fusion of two ancestral chromosome due to its composition of a vestigial second centromere and vestigial telomeres.

#### **ITPRIPL1 Antibody (N-term) Blocking Peptide - References**

Lim, J., et al. Cell 125(4):801-814(2006)