

**PRDM11 Antibody (C-term) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP1211b****Specification**

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**PRDM11 Antibody (C-term) Blocking Peptide - Product Information**

Primary Accession [O9NOV5](#)  
Other Accession [NP\\_064614](#)

**PRDM11 Antibody (C-term) Blocking Peptide - Additional Information**

**Gene ID** 56981

**Other Names**

PR domain-containing protein 11, 211-, PRDM11, PFM8

**Target/Specificity**

The synthetic peptide sequence used to generate the antibody [AP1211b](/product/products/AP1211b) was selected from the C-term region of human PRDM11. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

**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.

**PRDM11 Antibody (C-term) Blocking Peptide - Protein Information**

**Name** PRDM11

**Synonyms** PFM8

**Function**

May be involved in transcription regulation.

**Cellular Location**

Nucleus. Cytoplasm

**Tissue Location**

Highly expressed in lung, including bronchial epithelial cells and airway smooth muscle cells, as well as in peripheral blood mononuclear cells. In tonsils, expressed in B-cell types, including naive B-cells, centroblasts, centrocytes and memory B- cells (at protein level). In benign hyperplastic

lymph nodes, expressed in germinal center cells in both the dark and light zones, as well as in scattered cells in the mantle zone and the interfollicular area (at protein level).

### **PRDM11 Antibody (C-term) Blocking Peptide - Protocols**

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

- [Blocking Peptides](#)

### **PRDM11 Antibody (C-term) Blocking Peptide - Images**

### **PRDM11 Antibody (C-term) Blocking Peptide - Background**

Similar to acetylation and phosphorylation, histone methylation at the N-terminal tail has emerged as an important role in regulating chromatin dynamics and gene activity. Histone methylation occurs on arginine and lysine residues and is catalyzed by two families of proteins, the protein arginine methyltransferase family and the SET-domain-containing methyltransferase family. Five members have been identified in the arginine methyltransferase family. About 27 are grouped into the SET-domain family, and another 17 make up the PR domain family that is related to the SET domain family. The retinoblastoma protein-interacting zinc finger gene RIZ1 is a tumor suppressor gene and a FOUNDRING member of the PR domain family. RIZ1 inactivation is commonly found in many types of human cancers and occurs through loss of mRNA expression, frame shift mutation, chromosomal deletion, and missense mutation. RIZ1 is also a tumor susceptibility gene in mice. The loss of RIZ1 mRNA in human cancers was shown to associate with DNA methylation of its promoter CpG island. Methylation of the RIZ1 promoter strongly correlated with lost or decreased RIZ1 mRNA expression in breast, liver, colon, and lung cancer cell lines as well as in liver cancer tissues.

### **PRDM11 Antibody (C-term) Blocking Peptide - References**

Xiao, B., et al., Curr. Opin. Struct. Biol. 13(6):699-705 (2003). Jiang, G.L., et al., Histol. Histopathol. 15(1):109-117 (2000).