

Catalog: OM105579



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## Rabbit anti-IRF6 polyclonal antibody - middle region

Catalog: OM105579	
	100ug

## **Product profile**

Product name Rabbit anti-IRF6 polyclonal antibody - middle region

Antibody Type Primary Antibodies

Immunogen The immunogen for anti-IRF6 antibody: synthetic peptide directed towards the middle region of human IR

F6

## **Key Feature**

**Clonality** Polyclonal

**Isotype** lgG

Host Species Rabbit

Tested Applications WB ,IHC

Species Reactivity Bovine Dog Horse Human Mouse Pig Rabbit Rat Sheep Zebra Fish

Concentration 1 mg/ml

Purification Affinity purified

### **Target Information**

Gene Symbol IRF6

Gene Synonyms LPS; OFC6; PIT; PPS; VWS; VWS1

Gene Full Name Interferon regulatory factor 6

Gene Summary IRF6 is a member of the interferon regulatory transcription factor (IRF) family. Family members share a hig

hly-conserved N-terminal helix-turn-helix DNA-binding domain and a less conserved C-terminal protein-bin ding domain. Mutations in its gene can cause van der Woude syndrome and popliteal pterygium syndrom

e. This protein is involved in palate formation. The protein encoded by this gene shares strong similarity with Saccharomyces cerevisiae Cdc23, a protein essential for cell cycle progression through the G2/M transi

 $tion. This \ protein \ is \ a \ component \ of \ an aphase-promoting \ complex \ (APC), which \ is \ composed \ of \ eight \ pro$ 

tein subunits and highly conserved in eucaryotic cells. APC catalyzes the formation of cyclin B-ubiquitin c

onjugate that is responsible for the ubiquitin-mediated proteolysis of B-type cyclins. This protein and 3 ot

her members of the APC complex contain the TPR (tetratricopeptide repeat), a protein domain important for protein-protein interaction. This gene encodes a member of the interferon regulatory transcription factor (IRF) family. Family members share a highly-conserved N-terminal helix-turn-helix DNA-binding domain and a less conserved C-terminal protein-binding domain. Mutations in this gene can cause van der Woude syndrome and popliteal pterygium syndrome. This protein is involved in palate formation. Publication Note: This Ref Seq record includes a subset of the publications that are available for this gene. Please see the Entrez Gene record to access additional publications.

Alternative Names LPS, OFC6, PIT, PPS, VWS, VWS1

Molecular Weight (MW) 53kDa

Sequence 467 amino acids

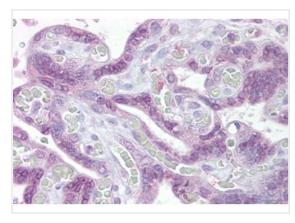
#### **Database Links**

Entrez Gene 3664

SwissProt ID 014896

Protein Accession NP\_006138

### **Application**



#### Immunohistochemistry

Application Notes WB:1:500~1:2000

IHC:1:50~1:200

**Notes**:Optimal dilutions/concentrations should be determined by the researcher.

### **Additional Information**

Form Liquid

Storage Instructions Aliquot and store at -20°C. Avoid repeated freeze / thaw cycles

**Storage Buffer** phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.

**Note** The product is for research use only,not for use in diagnostic or therapeutic procedures.

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