

**DMP4 rabbit pAb**

<b>Catalog No :</b>	YT6873
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB
<b>Target :</b>	DMP4
<b>Gene Name :</b>	FAM20C DMP4
<b>Protein Name :</b>	DMP4
<b>Human Gene Id :</b>	56975
<b>Human Swiss Prot No :</b>	Q8IXL6
<b>Mouse Gene Id :</b>	80752
<b>Mouse Swiss Prot No :</b>	Q5MJS3
<b>Immunogen :</b>	Synthesized peptide derived from human DMP4 AA range: 328-378
<b>Specificity :</b>	This antibody detects endogenous levels of DMP4 at Human/Mouse
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1[?]500-2000
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)

**Molecularweight :** 64kD

**Background :** This gene encodes a member of the family of secreted protein kinases. The encoded protein binds calcium and phosphorylates proteins involved in bone mineralization. Mutations in this gene are associated with the autosomal recessive disorder Raine syndrome. [provided by RefSeq, Apr 2014],

**Function :** disease:Defects in FAM20C are the cause of Raine syndrome (RNS) [MIM:259775]. RNS is an autosomal recessive osteosclerotic bone dysplasia with neonatal lethal outcome. Clinical features include generalized osteosclerosis, craniofacial dysplasia and microcephaly.,function:Calcium-binding protein which may play a role in dentin mineralization.,similarity:Belongs to the FAM20 family.,tissue specificity:Widely expressed.,

**Subcellular Location :** Secreted . Golgi apparatus .

**Expression :** Widely expressed.

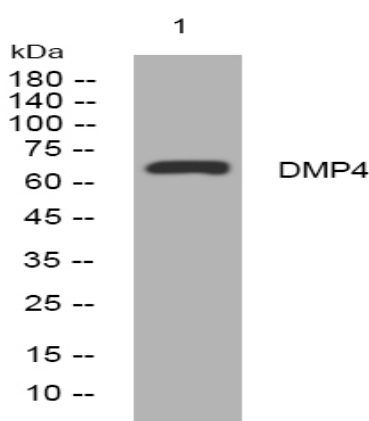
**Sort :** 5165

**No4 :** 1

**Host :** Rabbit

**Modifications :** Unmodified

## Products Images



Western blot analysis of lysates from 293T cells, primary antibody was diluted at 1:1000, 4 ° over night