

COMP rabbit pAb

Catalog_no :	<u>AT6588</u>
Applications :	<u>WB</u>
Reactivity :	<u>Human, Mouse,Rat</u>
Category :	<u>抗原抗体</u>
Size :	<u>100µg/50µg/20µg</u>
Gene_name :	<u>COMP</u>
Protein_name :	<u>COMP</u>
Humangene_id	<u>1311</u>
:	
Humanswissprot	<u>P49747</u>
_no :	
Mousegene_id :	<u>12845</u>
Mouseswissprot	<u>Q9R0G6</u>
_no :	
Ratgene_id :	<u>25304</u>
Ratswissprot_no	<u>P35444</u>
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Immunogen :	<u>Synthesized peptide derived from human COMP</u>
Specificity :	<u>This antibody detects endogenous levels of COMP at Human/Mouse/Rat</u>
Formulation :	<u>Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.</u>
Source :	<u>Rabbit</u>
Dilution :	<u>WB 1 : 500-2000</u>
Purification :	<u>The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.</u>
Concentration :	<u>1 mg/ml</u>
Storage_stability	<u>-20°C/1 year</u>
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Background :	<u>The protein encoded by this gene is a noncollagenous extracellular matrix (ECM) protein. It consists of five identical glycoprotein subunits, each with EGF-like and calcium-binding (thrombospondin-like) domains. Oligomerization results from</u>

formation of a five-stranded coiled coil and disulfides. Binding to other ECM proteins such as collagen appears to depend on divalent cations. Contraction or expansion of a 5 aa aspartate repeat and other mutations can cause pseudoachondroplasia (PSACH) and multiple epiphyseal dysplasia (MED). [provided by RefSeq, Jul 2016],
