

BMP-4 rabbit pAb

Catalog No :	YT7841
Reactivity :	Human;Rat;Mouse;
Applications :	WB;ELISA
Target :	BMP-4
Fields :	>>Cytokine-cytokine receptor interaction;>>TGF-beta signaling pathway;>>Hippo signaling pathway;>>Signaling pathways regulating pluripotency of stem cells;>>Thyroid hormone signaling pathway;>>Pathways in cancer;>>Basal cell carcinoma;>>Fluid shear stress and atherosclerosis
Gene Name :	BMP4 BMP2B DVR4
Protein Name :	BMP-4
Human Gene Id :	652
Human Swiss Prot No :	P12644
Mouse Gene Id :	12159
Mouse Swiss Prot No :	P21275
Rat Swiss Prot No :	Q06826
Immunogen :	Synthesized peptide derived from human BMP-4 AA range: 261-310
Specificity :	This antibody detects endogenous levels of Human BMP-4
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Polyclonal, Rabbit,IgG
Dilution :	WB 1:1000-2000 ELISA 1:5000-20000
Purification :	The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration : 1 mg/ml

Storage Stability : -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight : 45kD

Background : disease:Defects in BMP4 are the cause of microphthalmia syndromic type 6 (MCOPS6) [MIM:607932]; also known as microphthalmia and pituitary anomalies or microphthalmia with brain and digit developmental anomalies. Microphthalmia is a clinically heterogeneous disorder of eye formation, ranging from small size of a single eye to complete bilateral absence of ocular tissues (anophthalmia). In many cases, microphthalmia/anophthalmia occurs in association with syndromes that include non-ocular abnormalities. MCOPS6 is characterized by microphthalmia/anophthalmia associated with facial, genital, skeletal, neurologic and endocrine anomalies.,function:Induces cartilage and bone formation. Also act in mesoderm induction, tooth development, limb formation and fracture repair.,online information:Bone morphogenetic protein 4 entry,similarity:Belongs to the TGF-beta family.,subunit:Homodimer; disulfide-linked (By similarity). Interacts with GREM2 (By similarity) and SOSTDC1. Part of a complex consisting of TWSG1 and CHRDL1.,tissue specificity:Expressed in the lung and lower levels seen in the kidney. Present also in normal and neoplastic prostate tissues, and prostate cancer cell lines.

Function : skeletal system development, ossification, angiogenesis, ovarian follicle development, blood vessel development,osteoblast differentiation, eye development, urogenital system development, metanephros development, ureteric bud development, branching involved in ureteric bud morphogenesis, formation of primary germ layer, mesoderm formation, cell fate specification, cell fate determination, mesodermal cell fate commitment, induction of an organ,morphogenesis of a branching structure, kidney development, regulation of protein amino acid phosphorylation,positive regulation of protein amino acid phosphorylation, vasculature development, morphogenesis of an epithelium,lens development in camera-type eye, lens morphogenesis in camera-type eye, immune system development,regionalization, reproductive developmental process, regulation of transcription, DNA-dependent, regulation of transcription fr

Subcellular Location : Secreted, extracellular space, extracellular matrix.

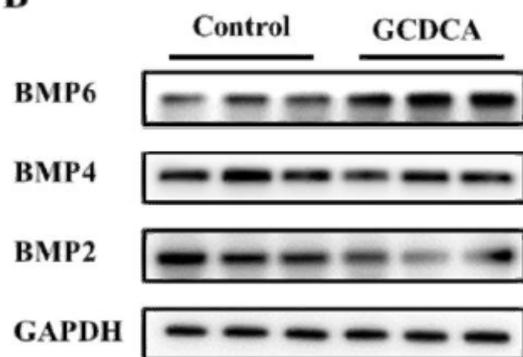
Expression : Expressed in the lung and lower levels seen in the kidney. Present also in normal and neoplastic prostate tissues, and prostate cancer cell lines.

Tag : orthogonal

Sort : 2800

No4 :	<u>1</u>
Host :	<u>Rabbit</u>
Modifications :	<u>Unmodified</u>

Products Images

B

Glycochenodeoxycholate Affects Iron Homeostasis via Up-Regulating Heparin Expression Nutrients Long-jiao Wang WB Mouse