

MaxPab®

# GP1BB purified MaxPab mouse polyclonal antibody (B01P)

Catalog # H00002812-B01P Size 500 ug

Specification	
Product Description	Mouse polyclonal antibody raised against a full-length human GP1BB protein.
Immunogen	GP1BB (AAl60146.1, 1 a.a. ~ 206 a.a) full-length human protein.
Sequence	MGSGPRGALSLLLLLAPPSRPAAGCPAPCSCAGTLVDCGRRGLTWASLPTAFPVDTTELVLTG NNLTALPPGLLDALPALRTAHLGANPWRCDCRLVPLRAWLAGRPERAPYRDLRCVAPPALRGRL LPYLAEDELRAACAPGPLCWGALAAQLALLGLGLLHALLLVLLLCRLRRLRARARARAAARLSLT DPLVAERAGTDES
Host	Mouse
Reactivity	Human
Interspecies Antigen Sequence	Mouse (90); Rat (84)
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## **Applications**

Western Blot (Transfected lysate)

**Protocol Download** 

### Gene Info — GP1BB

Entrez GenelD 2812



### **Product Information**

GeneBank Accession#	BC160146.1
Protein Accession#	<u>AAI60146.1</u>
Gene Name	GP1BB
Gene Alias	CD42c
Gene Description	glycoprotein lb (platelet), beta polypeptide
Omim ID	<u>138720</u> <u>231200</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Platelet glycoprotein lb (GPlb) is a heterodimeric transmembrane protein consisting of a disulfide-linked 140 kD alpha chain and 22 kD beta chain. It is part of the GPlb-V-IX system that constitutes the receptor for von Willebrand factor (VWF), and mediates platelet adhesion in the arterial circul ation. GPlb alpha chain provides the VWF binding site, and GPlb beta contributes to surface expression of the receptor and participates in transmembrane signaling through phosphorylation of its intracellular domain. Mutations in the GPlb beta subunit have been associated with Bernard-Souli er syndrome, velocardiofacial syndrome and giant platelet disorder. The 206 amino acid precurso r of GPlb beta is synthesized from a 1.0 kb mRNA expressed in plateletes and megakaryocytes. A 411 amino acid protein arising from a longer, unspliced transcript in endothelial cells has been described; however, the authenticity of this product has been questioned. Yet another less abundant GPlb beta mRNA species of 3.5 kb, expressed in nonhematopoietic tissues such as endothelium, brain and heart, was shown to result from inefficient usage of a non-consensus polyA signal within a separate gene (septin 5) located upstream of this gene. In the absence of polyadenylation from its own imperfect site, the septin 5 gene uses the consensus polyA signal of this gene. [provided by RefSeq
Other Designations	glycoprotein lb, beta polypeptide nuclear localization signal deleted in velocardiofacial syndrome

## Pathway

- ECM-receptor interaction
- Hematopoietic cell lineage

#### Disease

- Bernard-Soulier Syndrome
- Blood Platelet Disorders
- Thalassemia
- Thrombocytopenia



• von Willebrand Disease