



## **Product Datasheet**

PDGFB Antibody (orb1272340)



## www.biorbyt.com

**Description**nts. PDGFB Antibody

Species/Host Rabbit

Reactivity Human

**Conjugation** Unconjugated

Tested ELISA, NeA, WB

**Applications** 

**Immunogen** Produced from sera of rabbits pre-immunized with highly

pure (>98%) recombinant hPDGF-BB.

Target PDGFB

Form/Appearance Lyophilized

Concentration batch dependent

**Storage** PDGF-BB antibody is stable for at least 2 years from date

of receipt at -20°C. The reconstituted antibody is stable for at least two weeks at 2-8°C. Frozen aliquots are stable for at least 6 months when stored at -20°C. Avoid

repeated freeze-thaw cycles.

**Note** For research use only

**Clonality** Polyclonal

Uniprot ID P01127

NCBI P01127

**Dilution Range** Neutralization: To yield one-half maximal inhibition

[ND50] of the biological activity of hPDGF-BB (10.0 ng/mL), a concentration of 0.05 - 0.07  $\mu$ g/mL of this antibody is required. ELISA:To detect hPDGF-BB by direct

ELISA (using 100 μL/well antibody solution) a

concentration of at least 0.5  $\mu$ g/mL of this antibody is required. This antigen affinity purified antibody, in conjunction with compatible secondary reagents, allows the detection of 0.2 - 0.4 ng/well of recombinant hPDGF-BB. Sandwich:To detect hPDGF-BB by sandwich ELISA (using 100  $\mu$ L/well antibody solution) a concentration of 0.5 - 2.0  $\mu$ g/mL of this antibody is required. This antigen

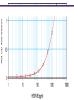
affinity purified antibody, in conjunction with our Biotinylated Anti-Human PDGF-BBas a detection antibody,

allows the detection of at least 0.2 - 0.4 ng/well of

recombinant hPDGF-BB.Western Blot:To detect hPDGF-BB by Western Blot analysis this antibody can be used at a concentration of 0.1 - 0.2  $\mu$ g/mL. Used in conjunction with compatible secondary reagents the detection limit for recombinant hPDGF-BB is 1.5 - 3.0 ng/lane, under either

reducing or non-reducing conditions.

Backing 1998



To detect hPDGF-BB by sandwich ELISA (us...



To detect hPDGF-BB by Western Blot analy...



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