Goat anti-human von Willebrand Factor
Peroxidase Conjugated IgG
0.15 mg

**Product Specifications**

**Description:**
Vial containing XXXX ml of IgG conjugated to horseradish peroxidase (HRP) through carbohydrate groups. Total protein is 0.15 mg.

**Format:**
IgG-HRP conjugate as a clear, slightly red-brown liquid.

**Host Animal:**
Goat

**Immunogen:**
Human vWF purified from plasma.

**Concentration:**
IgG-HRP concentration is XXXX mg/ml, determined by absorbance using an extinction coefficient (E1%280) of 14.

**Buffer:**
A buffered stabilizer solution containing 50% (v/v) glycerol.

**Storage:**
Store between –10 and –20°C. Product will become viscous but will not freeze. Avoid storage in frost-free freezers. Keep vial tightly capped. Allow product to warm to room temperature and gently mix before use. Avoid exposure to sodium azide as this is an inhibitor of peroxidase activity.

**Specificity:**
Prior to conjugation, this antibody was specific for vWF as demonstrated by immunoelectrophoresis and ELISA.

**Applications:**
Suitable as a source of peroxidase-labeled antibodies to vWF.

**Rz Ratio (Reinheitszahl, A403/A280):**
XXX as determined spectrophotometrically.

**Related Products:**
- Cat #: GAVWF-IG Goat anti-human vWF, whole IgG from antiserum
- Cat #: GAVWF-AP Goat anti-human vWF, affinity purified IgG
- Cat #: VWF-EIA Paired antibody set for ELISA of vWF, 5 x 96 wells
- Cat #: VWF-DE Plasma deficient in vWF/F.VIII, immune depleted

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**Limited Warranty:** This product is warranted to perform in accordance with its labeling and literature. Affinity Biologicals Inc. disclaims any implied warranty of merchantibility or fitness for any other purposes, and in no event will Affinity Biologicals Inc. be liable for any consequential damages arising out of aforesaid express warranty.

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**Description of von Willebrand Factor**

von Willebrand Factor (vWF, also previously referred to as Factor VIII related antigen) is a large adhesive protein produced in endothelial cells and megakaryocytes. There are two critical functions of vWF, the first being its involvement in the process of platelet adhesion and aggregation through interaction with platelet receptor glycoprotein Ib, the second being the binding and stabilization of Factor VIII (antihemophilic factor) for secretion and transport in plasma. The vWF precursor protein is synthesized with a 95,000 dalton propeptide (also known as vWF antigen-II), believed to be involved in the intracellular multimerization of the vWF subunits. The mature vWF multimers are then packed into storage organelles within the cell (Weibel-Palade bodies) after which the propeptide is cleaved and released. vWF circulates as multimers of disulfide linked 220,000 dalton subunits and the molecular weight of these multimers ranges from 0.5-20 million daltons.

The plasma concentration of vWF is typically 10 µg/ml, but increased levels are often observed in pregnancy and other conditions of physiological stress. von Willebrand’s disease (vWD) is perhaps the most common inherited bleeding disorder in humans and is the result of either quantitative deficiencies of vWF (vWD Types I & III), or one of a number of qualitative disorders of vWF structure and function (vWD Type II).

**Review Articles**

2. Sadler JE; von Willebrand Factor (Minireview); JBC 266:34, pp 22777-22780, 1991.