# abcam

### Product datasheet

## Anti-Von Willebrand Factor antibody [EPR12011] - BSA and Azide free ab240177



#### 1 References 3 Images

#### Overview

**Product name** Anti-Von Willebrand Factor antibody [EPR12011] - BSA and Azide free

**Description** Rabbit monoclonal [EPR12011] to Von Willebrand Factor - BSA and Azide free

**Host species** Rabbit

Suitable for: WB **Tested applications** 

Unsuitable for: ICC/IF,IHC-P or IP

Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Human serum, Human plasma, Rat plasma, and Mouse platelet lysates.

**General notes** ab240177 is the carrier-free version of ab174290.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® is a trademark of Fluidigm Canada Inc.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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#### **Properties**

Form Liquid

**Storage instructions** Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EPR12011

**Isotype** IgG

#### **Applications**

The Abpromise quarantee Our Abpror

Our **Abpromise guarantee** covers the use of ab240177 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Predicted molecular weight: 309 kDa.

**Application notes** Is unsuitable for ICC/IF,IHC-P or IP.

#### **Target**

Function Important in the maintenance of hemostasis, it promotes adhesion of platelets to the sites of

vascular injury by forming a molecular bridge between sub-endothelial collagen matrix and platelet-surface receptor complex GPlb-IX-V. Also acts as a chaperone for coagulation factor VIII, delivering it to the site of injury, stabilizing its heterodimeric structure and protecting it from

premature clearance from plasma.

Tissue specificity Plasma.

Involvement in disease Defects in VWF are the cause of von Willebrand disease (VWD) [MIM:277480]. VWD defines a

group of hemorrhagic disorders in which the von Willebrand factor is either quantitatively or qualitatively abnormal resulting in altered platelet function. Symptoms vary depending on severity and disease type but may include prolonged bleeding time, deficiency of factor VIII and impaired platelet adhesion. Type I von Willebrand disease is the most common form and is characterized by partial quantitative plasmatic deficiency of an otherwise structurally and functionally normal Willebrand factor; type II is associated with a qualitative deficiency and functional anomalies of the

Willebrand factor; type Ill is the most severe form and is characterized by total or near-total absence of Willebrand factor in the plasma and cellular compartments, also leading to a profound

deficiency of plasmatic factor VIII.

Sequence similarities Contains 1 CTCK (C-terminal cystine knot-like) domain.

Contains 4 TIL (trypsin inhibitory-like) domains.

Contains 3 VWFA domains. Contains 3 VWFC domains. Contains 4 VWFD domains. **Domain**The von Willebrand antigen 2 is required for multimerization of vWF and for its targeting to

storage granules.

Post-translational

All cysteine residues are involved in intrachain or interchain disulfide bonds.

modifications

N- and O-glycosylated.

**Cellular localization** 

Secreted. Secreted > extracellular space > extracellular matrix. Localized to storage granules.

#### **Images**



Western blot - Anti-Von Willebrand Factor antibody [EPR12011] - BSA and Azide free (ab240177) **All lanes :** Anti-Von Willebrand Factor antibody [EPR12011] (ab174290) at 1/1000 dilution (Purified)

Lane 1 : Human serum lysates
Lane 2 : Human plasma lysates
Lane 3 : Rat plasma lysates

Lane 4: Mouse platelet lysates

Lysates/proteins at 20 µg per lane.

#### **Secondary**

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

**Predicted band size:** 309 kDa **Observed band size:** 309 kDa

This data was developed using <u>ab174290</u>, the same antibody clone in a different buffer formulation.

We are unsure how to define the extra bands.

150— 150— 75—

Western blot - Anti-Von Willebrand Factor antibody [EPR12011] - BSA and Azide free (ab240177) **All lanes :** Anti-Von Willebrand Factor antibody [EPR12011] (ab174290) at 1/1000 dilution

Lane 1 : Human serum lysate
Lane 2 : Human plasma lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 309 kDa

This data was developed using <u>ab174290</u>, the same antibody clone in a different buffer formulation.



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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