

## DESCRIPTION

<b>Species Reactivity</b>	Human
<b>Specificity</b>	Detects human vWF-A2 in direct ELISAs and Western blots. Detects an epitope N-terminal to the ADAMTS13 cleavage site between aa 1498-1605. Another antibody (clone 210909, Catalog # <a href="#">MAB2764</a> ) recognizes an epitope on the C-terminal side (aa 1606-1665) of the ADAMTS13 cleavage site.
<b>Source</b>	Monoclonal Mouse IgG <sub>2B</sub> Clone # 210905
<b>Purification</b>	Protein A or G purified from hybridoma culture supernatant
<b>Immunogen</b>	<i>E. coli</i> -derived recombinant human vWF-A2 Asp1498-Val1665 Accession # P04275
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose and Mannitol. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.

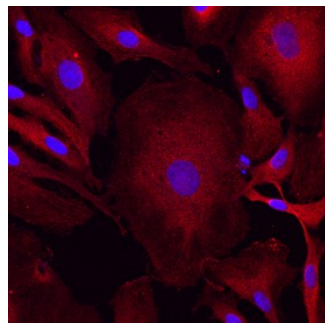
## APPLICATIONS

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	<b>Recommended Concentration</b>	<b>Sample</b>
<b>Western Blot</b>	1 µg/mL	Recombinant Human vWF-A2 (Catalog # <a href="#">2764-WF</a> )
<b>Immunocytochemistry</b>	8-25 µg/mL	See Below
<b>Immunoprecipitation</b>	25 µg/mL	Cell lysates spiked with Recombinant Human vWF-A2 (Catalog # <a href="#">2764-WF</a> ), see our <a href="#">available Western blot detection antibodies</a>

## DATA

### Immunocytochemistry



**vWF-A2 in HUVEC Human Cells.**  
vWF-A2 was detected in immersion fixed HUVEC human umbilical vein endothelial cells using Mouse Anti-Human vWF-A2 Monoclonal Antibody (Catalog # MAB27641) at 10 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Mouse IgG Secondary Antibody (red; Catalog # [NL007](#)) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm. View our protocol for [Fluorescent ICC Staining of Cells on Coverslips](#).

## PREPARATION AND STORAGE

<b>Reconstitution</b>	Reconstitute at 0.5 mg/mL in sterile PBS.
<b>Shipping</b>	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
<b>Stability &amp; Storage</b>	<b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b> <ul style="list-style-type: none"> <li>12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>6 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>

**BACKGROUND**

von Willebrand Factor (vWF) is a large, multimeric glycoprotein made by endothelial cells and megakaryocytes. The pre-pro-vWF protein contains 2813 amino acids (aa), which consists of a 22 aa signal peptide, a 741 aa propeptide and a mature vWF monomer of 2050 aa (1-4). The pro-vWF undergoes dimerization in the endoplasmic reticulum (ER) through a C-terminal "cysteine-knot" (CK) domain. The pro-vWF dimers are transported to the Golgi and associate into multimers by forming disulfide bonds in the amino-terminal region of the mature form. Proteolytic processing of the pro-region also occurs in the Golgi. Mature vWF is stored in Weibel-Pallade bodies in endothelial cells and granules in megakaryocytes and platelets. The unusually-large vWF (ulvWF) multimers released from cells are very efficient in binding to platelets to form thrombus. The population of these highly active ulvWF multimers is controlled by a specific protease, ADAMTS13, which cleaves between residues Tyr1605 and Met1606 in the A2 domain of vWF. In the plasma, vWF appears as a series of large and intermediate size multimers with molecular masses from several thousand to 500 kDa. vWF also performs hemostatic functions (3-5). In a high shear-stressed environment, vWF undergoes conformational changes to expose a binding site for glycoprotein Iba. As a result, vWF facilitates the aggregation of platelets. In addition to platelet binding, vWF binds coagulation factor VIII to increase the lifetime of FVIII in plasma. The purified rhvWF-A2 contains the A2 domain of vWF.

**References:**

1. Sadler, J. E. (1998) *Annu. Rev. Biochem.* **67**:395.
2. Ruggeri, Z. M. (2003) *Cur. Opin. Hemat.* **10**:142.
3. Michiels, J. J. *et al.* (2006) *Clin. Appl. Thromb. Hemost.* **12**:397.
4. Groot, E. *et al.* (2007) *Cur. Opin. Hemat.* **14**:284.
5. Lenting, P. J. *et al.* (2007) *J. Thromb. Haemos.* **5**:1353.