



Product Data Sheet

VSV-G-Tag Monoclonal Antibody

Catalog Number: ORF.VSVGMAb-50

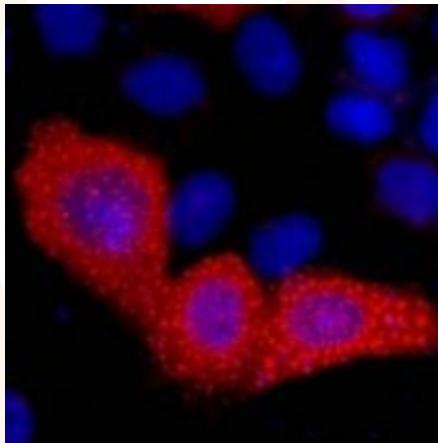
Product Details	
Product Name	VSV-G-Tag Monoclonal Antibody
Catalog Number	ORF.VSVGMAb-50
Size	50 µL
Concentration	1 mg/mL
Clonality	Monoclonal
Source	Mouse
Isotype	IgG
Purification	The antibody was purified by immunogen affinity chromatography.

Product Description	Product Image
The Vesicular Stomatitis Virus (VSV) G tag is derived from the glycoprotein of VSV, an enveloped RNA virus belonging to the Rhabdoviridae family. VSV G plays a critical role in viral assembly and budding from the plasma membrane of host cells. The commonly used VSV-G epitope tag corresponds to the sequence YTDIEMNRLGK, located in the extracellular membrane-proximal stem region of the glycoprotein, which is essential for efficient budding. As a fusion tag, VSV-G enables convenient detection, localization, and characterization of recombinant proteins, especially when protein-specific antibodies are not available. This monoclonal antibody specifically recognizes the VSV-G epitope and is optimized for Western blotting, providing sensitive and reliable detection of VSV-G-tagged proteins in research applications.	
Product Specifications and Product Specific Information	
Applications	WB: 1:2000 – 1:5000 IF/IC: 1:200 – 1:500 IP: 1:100 – 1:200
Reactivity	N/A

For research applications only. Not for diagnostic or therapeutic use.

Specificity	Recognizes C-terminal, internal and N-terminal VSV-G-tag fusion proteins.
Immunogen	KLH-conjugated synthetic peptide encompassing a sequence of VSV-G-tag. The exact sequence is proprietary.
Description	Mouse monoclonal antibody to VSV-G-tag
Buffer	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Storage and Stability		
	Temperature	Storage Time
Short Term	4°C	1 month
Long Term	-20°C	12 months
Avoid repeated freeze-thaw cycles.		

Product Data	
	Immunofluorescent analysis of VSV-G-tag staining in 293T cells transfected with a VSV-G-tag protein. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

For research applications only. Not for diagnostic or therapeutic use.