

Product Data Sheet

Flag-Tag Monoclonal Antibody

Catalog Number: ORF.FLAGMAB-50

Product Details		
Product Name	Flag-Tag Monoclonal Antibody	
Catalog Number	ORF.FLAGMAB-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 DYKDDDDK tag, also known as the FLAG® tag, is an eight-amino acid epitope sequence (Asp-Tyr-Lys-Asp-Asp-Asp-Asp-Lys) used to facilitate protein purification, detection, and localization. It can be fused to either the N-or C-terminus of a target protein and enables efficient isolation and identification using anti-FLAG antibodies. Optimized for compatibility, the FLAG tag is highly hydrophilic, minimizing the risk of protein denaturation or loss of activity upon tagging. Its versatility makes it widely applicable in protein-protein interaction studies, protein expression analysis, and cellular localization assays.		kDa 200 140 100 80 60 50 40 30 20	
Applications	WB: 1:2000 – 1:5000 IF/IC: 1:200 – 1:500		
Reactivity	IP: 1:100 – 1:200 N/A		
Specificity	Recognizes C-terminal, internal, and N-terminal FLAG-tag fusion proteins		

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

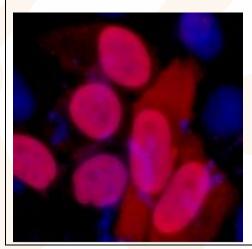
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Immunogen	KLH-conjugated synthetic peptide encompassing a sequence of FLAG-tag. The exact sequence is proprietary.
Description	Mouse monoclonal antibody to FLAG-tag
Buffer	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, 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 FLAG-tag staining in 293T cells transfected with a Flag-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).

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