

Product datasheet

anti-Vimentin mouse monoclonal, VIM 3B4, liquid, purified

Short overview

Cat. No. 690013 **Quantity** 1 ml

Concentration 50 μg/ml (50 μg)

Product description

Host Mouse
Antibody Type Monoclonal
Isotype IgG2a kappa
Clone VIM 3B4

Immunogen Vimentin purified from bovine lens

Formulation PBS pH 7.4 with 0.09% sodium azide and 0.5% BSA **UniprotID** P48616 (Bovine), P09654 (Chicken), P08670 (Human)

Synomym Vimentin, VIM Conjugate Unconjugated

Purification Affinity chromatography

Storage Short term at 2-8°C; long term storage in aliquots at -20°C; avoid freeze/thaw cycles

Intended useResearch use onlyApplicationICC/IF, IHC, WB

Reactivity Amphibia, Bovine, Chicken, Human, Monkey, Mouse

Applications

Immunocytochemistry (ICC) Assay dependent

Immunohistochemistry (IHC) - frozen 1:100-1:200 (250-500 ng/ml)

Immunohistochemistry (IHC) - paraffin 1:100-1:200 (250-500 ng/ml, protease treatment and/or microwave

treatment recommended)

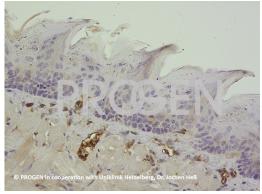
Western Blot (WB) 1:500-1:5,000 (10-100 ng/ml)

Background

The antibody is highly specific for the intermediate filament protein vimentin which is present in all cells of mesenchymal origin. VIM 3B4 has turned out to be the most avid mab to vimentin. Polypeptide reacting: 57 kDa intermediate filament protein (vimentin) of mesenchymal cells. Tumors specifically detected: sarcoma (including myosarcoma), lymphoma, melanoma. The binding region of monoclonal antibody VIM3B4 has been characterized by Bohn et al. (1992). According to these authors, the epitope has been localized on the alpha-helical part of vimentin (rod domain coil 2). Due to an aa substitution at position of aa 353 in murine vimentin (that could explain for the weak cross-reaction of the antibody with murine vimentin) they were able to narrow down the binding region around position 353. These findings were confirmed by truncation mutagenesis experiments using human vimentin (Rogers et al., 1995).

Bohn W, Wiegers W, Beuttenmüller M, Traub P: Species-specific recognition patterns of monoclonal antibodies directed against vimentin. Exp Cell Res 201: 1-7 (1992). Rogers KR, Eckelt A, Nimmrich V, Janssen K-P, Schliwa M, Herrmann H, Franke WW: Truncation mutagenesis of the non-alpha-helical carboxyterminal tail domain of vimentin reveals contributions to cellular localization but not to filament assembly. Eur J Cell Biol 66: 136-150 (1995).

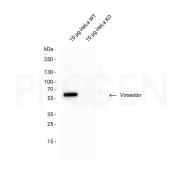
Product images



Mouse tongue (courtesy of J.Heß, University Hospital Heidelberg)



Human skin (courtesy of J.Heß, University Hospital Heidelberg)



Western blot analysis of HeLa lysate with anti-Vimentin antibody. Western blot analysis was performed on 10 μ g wild type (WT) and 10 μ g Vimentin knockout (KO) HeLa lysate. The PVDF membrane was blocked with 5% milk in PBST (PBS + 0.1% Tween 20) for 1 h at RT. The primary antibody anti-Vimentin mouse monoclonal, VIM 3B4 (Cat. No. 690013) was diluted in blocking buffer (antibody concentration 33 ng/ml) and incubated for 1 h at RT. The secondary antibody anti-mouse IgG, HRP conjugate was also diluted in blocking buffer (antibody concentration 200 ng/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using PierceTM ECL Western Blotting Substrate.

References

Publication	Species	Application
Soglia, F. et al. The evolution of vimentin and desmin in Pectoralis major muscles of broiler chickens supports their	chicken	WB
essential role in muscle regeneration. Front. Physiol. 13,		
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caveolin-1-dependent invasiveness, and Mol. Oncol. 12,		
1735–1752 (2018).		
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Zayas-Santiago, A. et al. Unidirectional	caiman	IHC
photoreceptor-to-Mýller glia coupling and unique K+ channel		
expression in Caiman retina. PLoS One 9, (2014).		