

## Product datasheet

### anti-Vimentin mouse monoclonal, XL-VIM-14.13, supernatant

#### Short overview

<b>Cat. No.</b>	65189
<b>Quantity</b>	5 ml

#### Product description

<b>Host</b>	Mouse
<b>Antibody Type</b>	Monoclonal
<b>Isotype</b>	IgG1
<b>Clone</b>	XL-VIM-14.13
<b>Immunogen</b>	Vimentin from cytoskeletal fraction of XLKE cells (cultured <i>Xenopus laevis</i> kidney epithelial cells)
<b>Formulation</b>	Contains 0.09% sodium azide
<b>Conjugate</b>	Unconjugated
<b>Purification</b>	Hybridoma cell culture supernatant
<b>Storage</b>	Short term at 2-8°C; long term storage in aliquots at -20°C; avoid freeze/thaw cycles
<b>Intended use</b>	Research use only
<b>Application</b>	IEM, IHC, WB
<b>Reactivity</b>	Carp, Human, Trout
<b>No reactivity</b>	Bovine

#### Applications

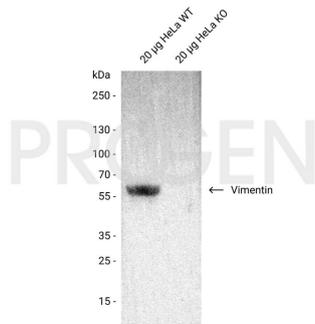
<b>Cell-based Assay</b>	Assay dependent
<b>Immunohistochemistry (IHC) - frozen</b>	Ready-to-use
<b>Western Blot (WB)</b>	1:10-1:20

#### Background

The anti-Vimentin antibody detects an epitope within the rod domain of xenopus and trout vimentin. Vimentin of amphibia and fish, predominantly found in glial and white blood cells.

Polypeptide reacting: MW 53,325 (pI 4.95) intermediate filament protein (vimentin) of *Xenopus laevis* (epitope presumably located between amino acids 79 and 88 within rod domain).

#### Product images



Western blot analysis of HeLa lysate with anti-Vimentin antibody. Western blot analysis was performed on 20 µg wild type (WT) and 20 µg 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, XL-VIM-14.13 (Cat. No. 65189) was diluted in blocking buffer (1:10) 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 Pierce™ ECL Western Blotting Substrate.

## References

Publication	Species	Application
<a href="#">Zirwes, R. F., Kouzmenko, A. P., Peters, J. M., Franke, W. W. &amp; Schmidt-Zachmann, M. S. Topogenesis of a nucleolar protein: determination of molecular segments directing nucleolar association. Mol. Biol. Cell 8, 231-48 (1997).</a>	xenopus	WB,ICC-IF,IEM
<a href="#">Herrmann, H., Munick, M. D., Brettel, M., Fouquet, B. &amp; Markl, J. Vimentin in a cold-water fish, the rainbow trout: highly conserved primary structure but unique assembly properties. J. Cell Sci. 109 ( Pt 3, 569-78 (1996).</a>	trout	WB,IHC (frozen)
<a href="#">Herrmann, H., Hofmann, I. &amp; Franke, W. W. Identification of a nonapeptide motif in the vimentin head domain involved in intermediate filament assembly. J. Mol. Biol. 223, 637-50 (1992).</a>	xenopus	ICC-IF