

## Product datasheet

### anti-EP-CAM mouse monoclonal, VU-1D9, ascites fluid

#### Short overview

<b>Cat. No.</b>	16114
<b>Quantity</b>	1 ml

#### Product description

<b>Host</b>	Mouse
<b>Antibody Type</b>	Monoclonal
<b>Isotype</b>	IgG1
<b>Clone</b>	VU-1D9
<b>Immunogen</b>	Isolated from small cell lung carcinoma-derived cell line (NC1-H69)
<b>Formulation</b>	Contains 0.09% sodium azide
<b>Note</b>	Centrifuge prior to opening
<b>Conjugate</b>	Unconjugated
<b>Purification</b>	Ascites
<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>	ICC/IF, IHC, WB
<b>Reactivity</b>	Human

#### Applications

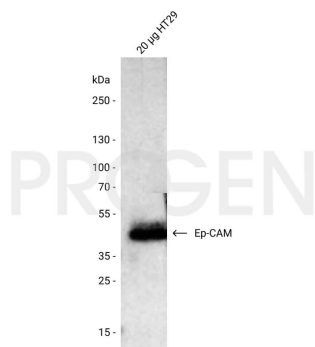
<b>Immunocytochemistry (ICC)</b>	Assay dependent
<b>Immunohistochemistry (IHC) - frozen</b>	1:10-1:20
<b>Immunohistochemistry (IHC) - paraffin</b>	1:10-1:20 (microwave treatment recommended)
<b>Western Blot (WB)</b>	1:200-1:5.000

#### Background

Ep-CAM (homophilic cell-cell adhesion molecule, also named ESA, EGP40, 17-1A antigen, KSA, GA7333-2) is a 40 kD epithelial protein expressed on the baso-lateral cell surface of many epithelial tissues. It is a transmembrane glycoprotein with 3 potential glycosylation sites. The extracellular domain has a cysteine-rich repeat and a small domain with homology to nidogen. Antibody stains most epithelial cells and carcinoma derived therefrom.

Positive control: breast carcinoma.

#### Product images



Western blot analysis of human HT29 cell lysate with anti-Ep-CAM antibody. Western blot analysis was performed on 20 µg of HT29 lysate. Cells were lysed with RIPA buffer. The PVDF membrane was blocked with 5% dry milk in PBST (PBS + 0.1% Tween 20) for 1 h at RT. The primary antibody anti-EP-CAM mouse monoclonal, VU-1D9 (Cat. No. 16114) was diluted in blocking buffer (1:5.000) and incubated for 1 h at RT. The secondary antibody anti-mouse IgG goat polyclonal, HRP conjugate was also diluted in blocking buffer (antibody concentration 0.2 µg/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using Pierce™ ECL Western Blotting Substrate.