

## **Product datasheet**

# anti-Keratin Type II mouse monoclonal, Ks pan1-8, liquid, purified, sample

#### Short overview

 Cat. No.
 690006S

 Quantity
 200 µl

Concentration 50 μg/ml (10 μg)

### **Product description**

HostMouseAntibody TypeMonoclonalIsotypeIgG2aCloneKs pan1-8

**Immunogen** Cytoskeletal proteins from cultured human MCF-7 cells **Formulation** PBS pH 7.4 with 0.09% sodium azide and 0.5% BSA

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 use Research use only
Application ICC/IF, IHC, WB

Reactivity Amphibia, Bovine, Human, Mouse, Rat

## **Applications**

Immunocytochemistry (ICC)Assay dependentImmunohistochemistry (IHC) - frozen1:10-1:100 (0.5-5 μg/ml)

Immunohistochemistry (IHC) - paraffin1:10-1:100 (0.5-5 μg/ml, microwave treatment recommended)

Western Blot (WB) 1:500 (0.1 μg/ml)

### Background

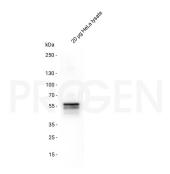
Ks pan1-8 represents an excellent marker for distinguishing carcinomas from non-epithelial tumors. Polypeptides reacting: Mr 52,500-Mr 68,000 keratins (type II keratins K1-K8; formerly also designated cytokeratins 1-8) of human epithelial cells. Tumors specifically detected: all epithelium-derived neoplasms.

Reactivity on cultured cell lines MCF-7, RT 112, HT-29, Detroit 562, RPMI 2650, SSC-12.

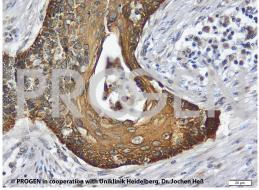
#### **Product images**



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



Western blot analysis of HeLa lysate with anti-Keratin Type II antibody. Western blot analysis was performed on 20  $\mu$ g HeLa 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-Keratin Type II mouse monoclonal, Ks pan1-8 (Cat. No. 690006) was diluted in blocking buffer (antibody concentration 0.1  $\mu$ g/ml) 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  $\mu$ g/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using PierceTM ECL Western Blotting Substrate.



IHC of human HNSCC tissue (courtesy of J.Heß, University Hospital Heidelberg)

# References

Publication	Species	Application
Kasai, Y. et al. A stable protocol for the fabrication of transplantable human oral mucosal epithelial cell sheets for	human	FACS
clinical application. Regen Ther. 14, 87-94(2020).		
Kimelman, D. et al. Regulation of posterior body and epidermal morphogenesis in zebrafish by localized Yap1 and Wwtr1. Elife. 6, (2017).	zebrafish	whole mount
Hatzold, J. et al. Tumor suppression in basal keratinocytes via dual non-cell-autonomous functions of a Na,K-ATPase beta subunit.eLife, 5 (2016).	zebrafish	whole mount
Fischer, B. et al. p53 and TAp63 Promote Keratinocyte Proliferation and Differentiation in Breeding Tubercles of the Zebrafish 6 Cologne Excellence Cluster on Cellular Stress Responses in Aging-Associated Diseases. PLoS Genet 10, (2014).	zebrafish	IHC
Montpetit, A. et al. Disruption of AP1S1, Causing a Novel Neurocutaneous Syndrome, Perturbs Development of the Skin and Spinal Cord. PLoS Genet 4, (2008).	zebrafish	whole mount