

# Mouse/Rat ALCAM/CD166 Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF1172

DESCRIPTION			
Species Reactivity	Mouse/Rat		
Specificity	Detects mouse ALCAM in direct ELISAs and Western blots. In direct ELISAs, approximately 40% cross-reactivity with recombinant human (rh) ALCAM is observed, and less than 10% cross-reactivity with rhBCAM, recombinant mouse (rm) OCAM, and rmMAdCAM-1 is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse ALCAM/CD166 Trp28-Lys527 Accession # AAC06342		
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.		

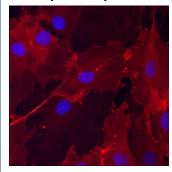
#### **APPLICATIONS**

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	0.1 μg/mL	Recombinant Mouse ALCAM/CD166 Fc Chimera (Catalog # 1172-AL)
Flow Cytometry	2.5 μg/10 <sup>6</sup> cells	Mouse splenocytes treated with PHA
Immunocytochemistry	5-15 μg/mL	See Below
Immunohistochemistry	5-15 μg/mL	See Below

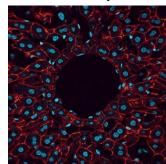
#### DATA

#### Immunocytochemistry



ALCAM/CD166 in Rat Mesenchymal Stem Cells. ALCAM/CD166 was detected in immersion fixed undifferentiated rat mesenchymal stem cells using Goat Anti-Mouse/Rat ALCAM/CD166 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1172) at 10 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm. View our protocol for Fluorescent ICC Staining of Cells on Coverslips.

#### Immunohistochemistry



ALCAM/CD166 in Mouse Liver.
ALCAM/CD166 was detected in perfusion fixed frozen sections of mouse liver using Goat Anti-Mouse/Rat ALCAM/CD166
Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1172) at 1.7 µg/mL overnight at 4 °C. Tissue was stained using the NorthemLights ™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). View our protocol for Fluorescent IHC Staining of Frozen Tissue Sections.

### PREPARATION AND STORAGE

**Reconstitution** Reconstitute at 0.2 mg/mL in sterile PBS.

Shipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

## Stability & Storage

Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.

#### BACKGROUND

ALCAM, activated leukocyte cell adhesion molecule, is a type I membrane glycoprotein and a member of the immunoglobulin supergene family. It is also known as CD166, MEMD, SC-1/DM-GRASP/BEN in the chicken, and KG-CAM in the rat. ALCAM is expressed on thymic epithelial cells, activated B and T cells, and monocytes. ALCAM can bind itself homotypically and is also capable of binding CD6, NgCAM, and other, as of yet, unidentified brain proteins. ALCAM/CD6 interaction may be involved in T cell development and T cell regulation. Additionally, ALCAM/CD6 and ALCAM/NgCAM interactions may play roles in the nervous system. ALCAM has also been observed to be upregulated on highly metastasizing melanoma cell lines and may play a role in tumor migration. ALCAM is a 583 amino acid (aa) protein consisting of a 27 aa signal peptide, a 500 aa extracellular domain, a 24 aa transmembrane domain, and a 32 aa cytoplasmic domain. The extracellular domain of ALCAM contains 5 lg-like domains of which the amino-terminal V1 domain is essential for ligand binding and ALCAM-mediated cell aggregation (1-4). The ECD of mouse ALCAM shares 97.5% aa sequence identity with rat ALCAM ECD.

## References:

- 1. Bowen, M.A. et al. (1995) J. Exp. Med. 181:2213.
- 2. Aruffo, A. et al. (1997) Immunol. Today 18:498.
- 3. Degen, W.G. et al. (1998) Am. J. Pathol. 152:805.
- 4. Van Kempen, L. et al. (2001) J. Biol. Chem. 276:25783.



Rev. 11/8/2012 Page 1 of 1