



Anti-human Endocan/ESM-1 monoclonal antibody

Clone MEP08 (C-Terminal)

Essential Notes

Cat. Number : LIA-0901

Clone : MEP08

Concentration : 1 mg/mL

Size : 100 µg

Formulation : PBS pH 7.4

Storage : 4°C / -20°C

Immunogen : *E. coli* derived C-Ter peptide (60-165)

Specificity : human and monkey endocan

Source : mouse

Ig isotype : IgG2a, K

Applications : IHC

■ Preparation/Source

Endocan/ESM-1 is a 165 amino acid peptide that carries a dermatan sulfate chain. Anti-endocan/ESM-1 antibodies clone MEP08 were produced from a hybridoma resulting from the fusion of mouse myeloma Sp2/0 cells with B cells obtained from mouse immunized with a *E. coli* derived C-terminal peptide (60-165) from recombinant human endocan. They were purified by protein A affinity chromatography.

■ Formulation

Solution in phosphate buffer saline 1x, pH 7.4

■ Concentration

The concentration of MEP08 was 1 mg/mL as determined by measurement of protein.

■ Purity

Purity > 90%, as determined by SDS-PAGE and as visualized by silver staining.

■ Specificity

Specificity is determined by ability to recognize **human and monkey endocan**.

■ Storage

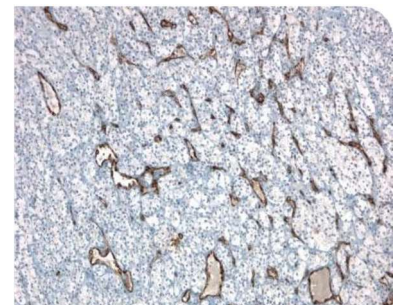
Antibody can be stored at 2°C - 8°C for 6 months without loss of activity. They can be easily aliquoted and stored frozen from -20°C to -80°C for long term storage. Avoid repeated freeze-thaw cycles.

■ Applications

Immunohistochemistry (IHC) :

Anti-human endocan antibody clone MEP08 is recommended to detect human endocan in paraffin-embedded tissues. Recommended working dilutions were determined to be 5 µg/ mL. Optimal dilutions should be determined according to tissue origins.

Other : to be determined.



Tumor vessels expressing endocan (brown) in kidney cancers as detected by IHC using the anti-endocan/ESM-1 antibody clone MEP08.

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■ Bibliography related to MEP08 Antibody Applications

- Calderaro J, et al. **ESM1** as a Marker of Macrotrabecular-Massive Hepatocellular Carcinoma. Clin Cancer Res. 2019 Oct 1;25(19):5859-5865
- Rudini N, et al. Phenotypic and molecular changes in nodule-in-nodule hepatocellular carcinoma with pathogenetic implications. Histopathology. 2018 Oct;73(4):601-611.
- Lin LY, et al. **Endocan** expression is correlated with poor progression-free survival in patients with pancreatic neuroendocrine tumors. Medicine (Baltimore). 2017 Oct;96(41):e8262.
- Suzuki H, et al. Increased **endocan** expression in lesional skin and decreased **endocan** expression in sera in atopic dermatitis. J Dermatol. 2017 Dec;44(12):1392-1395.
- Aung PP, et al Microvessel density, lymphovascular density, and lymphovascular invasion in primary cutaneous melanoma-correlation with histopathologic prognosticators and BRAF status. Hum Pathol. 2015 Feb;46(2):304-12.
- Priya MK, et al. Tipping off endothelial tubes: nitric oxide drives tip cells. Angiogenesis 2015 Apr;18(2):175-89.
- Ziol M, et al. **ESM-1** expression in stromal cells is predictive of recurrence after radiofrequency ablation in early hepatocellular carcinoma. J Hepatol. 2013 Dec;59(6):1264-70.
- Frahm KA, et al **Endocan** immunoreactivity in the mouse brain: method for identifying nonfunctional blood vessels. J Immunol Methods. 2013 Dec 15;398-399:27-32.
- Roudnicky F, et al. **Endocan** is upregulated on tumor vessels in invasive bladder cancer where it mediates VEGF-A-induced angiogenesis. Cancer Res. 2013 Feb 1;73(3):1097-106.
- Li S, et al. Detection on dynamic changes of endothelial cell specific molecule-1 in acute rejection after renal transplantation. Urology. 2012 Sep;80(3):738.e1-8.
- Cornelius A, et al. Endothelial expression of **endocan** is strongly associated with tumor progression in pituitary adenoma. Brain Pathol. 2012 Nov;22(6):757-64.
- Carrillo LM, et al. Immunolocalization of **endocan** during the endothelial-mesenchymal transition process. Eur J Histochem. 2011;55(2):e13.
- Kang YH, et al. **ESM-1** silencing decreased cell survival, migration, and invasion and modulated cell cycle progression in hepatocellular carcinoma. Amino Acids. 2011 Mar;40(3):1003-13. doi: 10.1007/s00726-010-0729-6. Epub 2010 Sep 7.
- Strasser GA, et al. Microarray analysis of retinal endothelial tip cells identifies CXCR4 as a mediator of tip cell morphology and branching. Blood. 2010 Jun 17;115(24):5102-10.
- Leroy X, et al. Vascular **endocan (ESM-1)** is markedly overexpressed in clear cell renal cell carcinoma. Histopathology. 2010 Jan;56(2):180-7.
- Maurage CA, et al **Endocan** expression and localization in human glioblastomas. J Neuropathol Exp Neurol. 2009 Jun;68(6):633-41.
- Rennel E, et al. **Endocan** is a VEGF-A and PI3K regulated gene with increased expression in human renal cancer. Exp Cell Res. 2007 Apr 15;313(7):1285-94. Epub 2007 Feb 6.
- Grigoriu BD, et al **Endocan** expression and relationship with survival in human non-small cell lung cancer. Clin Cancer Res. 2006 Aug 1;12(15):4575-82.

■ Background

Endocan, also known as endothelial cell-specific molecule (ESM-1), was originally discovered by Lassalle and collaborators in endothelial cells. Structurally, endocan is a dermatan sulfate proteoglycan of 50 kDa that is freely circulating in blood. Endocan binds CD11a/CD18 integrin (also called LFA-1 for Leukocyte Function-associated Antigen-1) on human leukocytes inhibiting consequently its binding to ICAM-1 and transendothelial migration. Moreover, endocan has been recently described as a biomarker of tip cells and neoangiogenesis. The expression of endocan is upregulated by pro-inflammatory molecules such as tumor necrosis factor alpha, and pro-angiogenic molecules such as vascular endothelial growth factor and fibroblast growth factor 2. Endocan binds via its dermatan sulfate chain to hepatocyte growth factor/ scatter factor. Endocan appears as a pertinent biomarker of endothelial dysfunction.

■ Companion products

- Anti-human endocan/ESM-1 mAb (C-ter) ; clone MEP14 : **LIA-1001**
- Anti-murine endocan/ESM-1 mAb (N-ter) ; clone GGR222 : **LIA-0905**
- Anti-human endocan/ESM-1 mAb (N-ter) ; clone MEP21 : **LIA-0902**
- Human recombinant endocan/ESM-1 (50 kDa) : **LIP-1001**
- DIYEK H1 (Do It Yourself Elisa Kit for Human Endocan quantification) : **LIK-1101**
- JDIEK H1 (Just Do It Elisa Kit for Human Endocan quantification) : **LIK-1201**

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