

CD98 (4F2): sc-59145

BACKGROUND

CD98 (4F2, CD98, MDU1, 4F2HC, 4T2HC, NACAE, SLC3A2) is a disulfide-linked heterodimer composed of a glycosylated heavy chain and a non-glycosylated light chain. CD98 is a scaffolding protein that interacts with basolaterally expressed amino acid transporters and $\beta 1$ integrins and can alter amino acid transport and cell adhesion, migration and branching morphogenesis. The heavy chain is a type II integral membrane protein. CD98 is expressed on T cells and is upregulated upon T cell activation. CD98 is also present on monocytes and at lower levels on granulocytes, platelets and lymphocytes. Evidence suggests that CD98 may play a role in the regulation of T cell activation and proliferation. Alternate transcriptional splice variants, encoding different isoforms exist for the human CD98 gene.

CHROMOSOMAL LOCATION

Genetic locus: SLC3A2 (human) mapping to 11q12.3.

SOURCE

CD98 (4F2) is a mouse monoclonal antibody raised against the H0ON pre-B leukemia cell line of human origin.

PRODUCT

Each vial contains 200 μ g IgG_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

CD98 (4F2) is available conjugated to agarose (sc-59145 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-59145 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-59145 PE), fluorescein (sc-59145 FITC), Alexa Fluor® 488 (sc-59145 AF488), Alexa Fluor® 546 (sc-59145 AF546), Alexa Fluor® 594 (sc-59145 AF594) or Alexa Fluor® 647 (sc-59145 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-59145 AF680) or Alexa Fluor® 790 (sc-59145 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

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APPLICATIONS

CD98 (4F2) is recommended for detection of CD98 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and flow cytometry (1 μ g per 1 x 10⁶ cells).

Suitable for use as control antibody for CD98 siRNA (h): sc-35033, CD98 shRNA Plasmid (h): sc-35033-SH and CD98 shRNA (h) Lentiviral Particles: sc-35033-V.

Molecular Weight of CD98: 125 kDa.

Positive Controls: Jurkat whole cell lysate: sc-2204, Hep G2 cell lysate: sc-2227 or HeLa whole cell lysate: sc-2200.

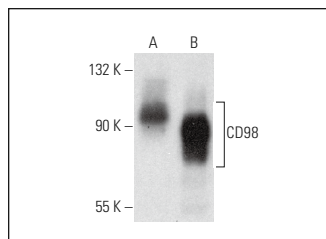
STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

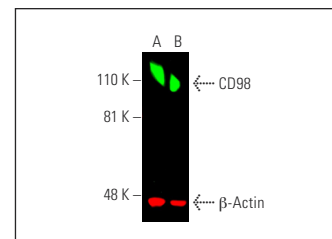
RESEARCH USE

For research use only, not for use in diagnostic procedures.

DATA



Western blot analysis of CD98 expression in Jurkat whole cell lysate (A) and Jurkat whole cell lysate immunoprecipitated with CD98 (4F2): sc-59145 (B) and detected with CD98 (C-20): sc-7095. Immunoprecipitation reagent used: Protein A-Agarose: sc-2001.



Simultaneous direct near-infrared western blot analysis of CD98 expression, detected with CD98 (4F2) Alexa Fluor® 680: sc-59145 AF680 and β -Actin expression, detected with β -Actin (C4) Alexa Fluor® 790: sc-47778 AF790 in HeLa (A) and Hep G2 (B) whole cell lysates. Blocked with UltraCruz® Blocking Reagent: sc-516214.

SELECT PRODUCT CITATIONS

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3. Janpipatkul, K., et al. 2014. Downregulation of LAT1 expression suppresses cholangiocarcinoma cell invasion and migration. *Cell. Signal.* 26: 1668-1679.
4. Kim, M.Y., et al. 2016. Molecular association of CD98, CD29, and CD147 critically mediates monocytic U937 cell adhesion. *Korean J. Physiol. Pharmacol.* 20: 515-523.
5. Gong, S., et al. 2018. Quantitative algorithm-based paired imaging measurement for antibody-triggered endocytosis in cultured cells. *SLAS Discov.* 23: 832-841.
6. Saito, Y., et al. 2022. Polarity protein SCRIB interacts with SLC3A2 to regulate proliferation and tamoxifen resistance in ER⁺ breast cancer. *Commun. Biol.* 5: 403.
7. Liu, D., et al. 2022. Both *in situ* and circulating SLC3A2 could be used as prognostic markers for human lung squamous cell carcinoma and lung adenocarcinoma. *Cancers* 14: 5191.
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9. Lee, M.J., et al. 2024. Senescence of endothelial cells increases susceptibility to Kaposi's sarcoma-associated herpesvirus infection via CD109-mediated viral entry. *J. Clin. Invest.* 135: e183561.

PROTOCOLS

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