Dysadherin (D-2): sc-166782



The Power to Question

BACKGROUND

Dysadherin (FXYD domain-containing ion transport regulator 5) is a cancer-associated cell membrane glycoprotein. Dysadherin downregulates the expression of E-cadherin, the prime mediator of cell-cell adhesion in epithelial cells, by a posttranscriptional mechanism. Decreasing intercellular adhesiveness facilitates the metastasis of cancer cells. Dysadherin is present in spleen, lung, skeletal muscle and testis tissue, and maps to human chromosome 19q13.12.

REFERENCES

- Ino, Y., et al. 2002. Dysadherin, a cancer-associated cell membrane glycoprotein, downregulates E-cadherin and promotes metastasis. Proc. Natl. Acad. Sci. USA 99: 365-370.
- Shimamura, T., et al. 2003. Dysadherin overexpression in pancreatic ductal adenocarcinoma reflects tumor aggressiveness: relationship to E-cadherin expression. J. Clin. Oncol. 21: 659-667.
- Hirohashi, S. and Kanai, Y. 2003. Cell adhesion system and human cancer morphogenesis. Cancer Sci. 94: 575-581.
- 4. Sato, H., et al. 2003. Dysadherin: expression and clinical significance in thyroid carcinoma. J. Clin. Endocrinol. Metab. 88: 4407-4412.
- Wu, D., et al. 2004. Prognostic significance of dysadherin expression in cervical squamous cell carcinoma. Pathol. Oncol. Res. 10: 212-218.

CHROMOSOMAL LOCATION

Genetic locus: FXYD5 (human) mapping to 19q13.12.

SOURCE

Dysadherin (D-2) is a mouse monoclonal antibody raised against amino acids 1-178 representing full length Dysadherin of human origin.

PRODUCT

Each vial contains 200 μg lgG_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

Dysadherin (D-2) is recommended for detection of Dysadherin isoform 1 and 2 of human origin by Western Blotting (starting dilution 1:100, 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 solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for Dysadherin siRNA (h): sc-45745, Dysadherin shRNA Plasmid (h): sc-45745-SH and Dysadherin shRNA (h) Lentiviral Particles: sc-45745-V.

Molecular Weight (predicted) of Dysadherin: 19 kDa.

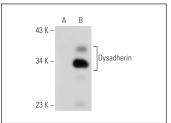
Molecular Weight (observed) of Dysadherin: 35/47 kDa.

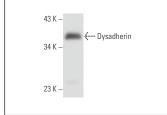
Positive Controls: Dysadherin (h): 293 Lysate: sc-111349, CCRF-CEM cell lysate: sc-2225 or JAR cell lysate: sc-2276.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker Molecular Weight Standards: sc-2035, UltraCruz* Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz* Mounting Medium: sc-24941 or UltraCruz* Hard-set Mounting Medium: sc-359850.

DATA





Dysadherin (D-2): sc-166782. Western blot analysis of Dysadherin expression in non-transfected: sc-110760 (A) and human Dysadherin transfected: sc-111349 (B) 293 whole cell Ivsates.

Dysadherin (D-2): sc-166782. Western blot analysis of Dysadherin expression in CCRF-CEM whole cell lysate.

SELECT PRODUCT CITATIONS

- 1. Tokhtaeva, E., et al. 2016. The O-glycosylated ectodomain of FXYD5 impairs adhesion by disrupting cell-cell trans-dimerization of Na,K-ATPase β 1 subunits. J. Cell Sci. 129: 2394-2406.
- 2. Lubarski-Gotliv, I., et al. 2016. FXYD5 has a pro-inflammatory role in epithelial cells. J. Biol. Chem. 291: 11072-11082.
- 3. Besso, M.J., et al. 2019. FXYD5/dysadherin, a biomarker of endometrial cancer myometrial invasion and aggressiveness: its relationship with TGF- β 1 and NF κ B pathways. Front. Oncol. 9: 1306.
- Hostrup, M., et al. 2023. High-intensity training represses FXYD5 and glycosylates Na,K-ATPase in type II muscle fibres, which are linked with improved muscle K+ handling and performance. Int. J. Mol. Sci. 24: 5587.

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.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.