α-actinin (H-2): sc-17829



The Power to Question

BACKGROUND

The spectrin gene family encodes a diverse group of cytoskeletal proteins that include spectrins, dystrophins and α -actinins. There are four tissue-specific α -actinins, namely α -actinin-1, α -actinin-2, α -actinin-3 and α -actinin-4, which are localized to muscle and non-muscle cells, including skeletal, cardiac and smooth muscle cells, as well as within the cytoskeleton. Each α -actinin protein contains one Actin-binding domain, two calponin-homology domains, two EF-hand domains and four spectrin repeats, through which they function as bundling proteins that can cross-link F-Actin, thus anchoring Actin to a variety of intracellular structures. Defects in the gene encoding α -actinin-4 are the cause of focal segmental glomerulosclerosis 1 (FSGS1), a common renal lesion characterized by decreasing kidney function and, ultimately, renal failure.

REFERENCES

- 1. Youssoufian, H., et al. 1990. Cloning and chromosomal localization of the human cytoskeletal α -actinin gene reveals linkage to the β -spectrin gene. Am. J. Hum. Genet. 47: 62-71.
- 2. Nishiyama, M., et al. 1990. Expression of human α -actinin in human hepatocellular carcinoma. Cancer Res. 50: 6291-6294.

SOURCE

 α -actinin (H-2) is a mouse monoclonal antibody raised against amino acids 593-892 mapping at the C-terminus of α -actinin-1 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.

 α -actinin (H-2) is available conjugated to agarose (sc-17829 AC), 500 $\mu g/0.25$ ml agarose in 1 ml, for IP; to HRP (sc-17829 HRP), 200 $\mu g/ml$, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-17829 PE), fluorescein (sc-17829 FITC), Alexa Fluor* 488 (sc-17829 AF488), Alexa Fluor* 546 (sc-17829 AF546), Alexa Fluor* 594 (sc-17829 AF594) or Alexa Fluor* 647 (sc-17829 AF647), 200 $\mu g/ml$, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor* 680 (sc-17829 AF680) or Alexa Fluor* 790 (sc-17829 AF790), 200 $\mu g/ml$, for Near-Infrared (NIR) WB, IF and FCM.

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APPLICATIONS

 $\alpha\text{-}actinin$ (H-2) is recommended for detection of $\alpha\text{-}actinin$ isoforms of mouse, rat and human origin by Western Blotting (starting dilution 1:500, dilution range 1:500-1:5,000), immunoprecipitation [1-2 μg per 100-500 μg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffinembedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

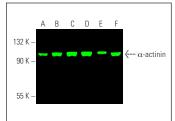
Molecular Weight of α -actinin: 100 kDa.

Positive Controls: Jurkat whole cell lysate: sc-2204, HeLa whole cell lysate: sc-2200 or K-562 whole cell lysate: sc-2203.

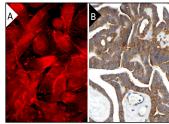
STORAGE

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

DATA



 α -actinin (H-2): sc-17829. Near-Infrared western blot analysis of α -actinin expression in Hela (Al), Jurkat (B), Hela (C), RT-4 (D), SuRH30 (E) and K-562 (F) whole cell lysates. Blocked with UltraCruz® Blocking Reagent: sc-516214. Detection reagent used: m-lgG $_1$ BP-CFL 680: ss-533665



 $\alpha\text{-actinin}$ (H-2) Alexa Fluor \$94: sc-17829 AF594. Direct immunofluorescence staining of formalin-fixed SW480 cells showing membrane and focal adhesions localization. Blocked with UltraCruz* Blocking Reagent: sc-516214 (A). $\alpha\text{-actinin}$ (H-2): sc-17829. Immunoperoxidase staining of formalin fixed, paraffin-embedded human ovarian cancer showing cytoplasmic staining of tumor cells. Kindly provided by The Swedish Human Protein Atlas (HPA) program (B).

SELECT PRODUCT CITATIONS

- 1. Lan, S., et al. 2003. Direct interaction between α -actinin and hepatitis C virus NS5B. FEBS Lett. 554: 289-294.
- 2. Matte, I., et al. 2015. Ovarian cancer ascites enhance the migration of patient-derived peritoneal mesothelial cells via cMet pathway through HGF-dependent and -independent mechanisms. Int. J. Cancer 137: 289-298.
- 3. Black, J.C., et al. 2016. Regulation of transient site-specific copy gain by microRNA. J. Biol. Chem. 291: 4862-4871.
- 4. Bernadzki, K.M., et al. 2017. Liprin-α-1 is a novel component of the murine neuromuscular junction and is involved in the organization of the postsynaptic machinery. Sci. Rep. 7: 9116.
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- Mournetas, V., et al. 2021. Myogenesis modelled by human pluripotent stem cells: a multi-omic study of Duchenne myopathy early onset.
 J. Cachexia Sarcopenia Muscle 12: 209-232.

RESEARCH USE

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