alpha-Actinin 1



VENDOR: R&D Systems

CAT #: MAB8279

SIMPLE WESTERN CERTIFIED ANTIBODY DATASHEET

<u>View</u> Antibody Link

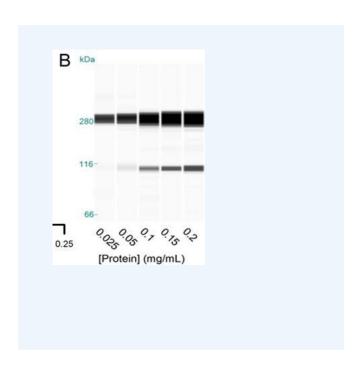


Figure: Detection and quantification of SI protein. (Panels A) and (Panels B) show the concentration-dependence of SI and α actinin signals (Panel C) using automated capillary Western blotting of Caco-2/TC7 lysates. (Panel D) shows Caco-2/TC7 cell surface protein fraction purified after biotinylation from cells cultured on 6-well Transwell filters in glucose or sucrose for 21 days and treated on the apical side for the final 3 days with 1.5 mg/mL OLE. (Panel E) shows the total cellular SI, relative to αactinin with data normalised to the glucose control and presented as mean ± SEM with n = 12, analysed in duplicate runs. The size was determined relative to a protein standard ladder for SI in the total lysate relative to α-actinin (F), normalised and presented as mean ± SEM with n = 24. * p < 0.05, ** p < 0.01, *** p < 0.001 from ANOVA followed by Tukey's post-hoc test. There was no change in the α-actinin loading control between the groups.

PROTEIN TARGET/ANTIBODY	
Protein Target	alpha-Actinin 1
Protein Isoform	Unmodified
Antibody Type	Primary
Host Species/Clonality	Mouse Monoclonal
ASSAY	
Sample Type	Caco-2/TC-7
Sample Concentration	Not_Stated
Antibody Concentration/Dilution	Not_Stated
Antibody Diluent	
Detection Mode	Chemiluminescence
Separation Type	Size
Matrix	66-440kDa
Observed kDa	Not_Stated

^{*}Image collected and cropped by CiteAb from the following publication https://www.mdpi.com/2072-6643/11/7/1505 licensed under a CC-BY license

<u>PUBLICATIONS</u>	
1. Pyner, A., Chan, S. Y., et al. Indirect Chronic Effects of an Oleuropein-Rich Olive Leaf Extract on Sucrase-Isomaltase In Vitro and In Vivo.	
Nutrients. 2019 Jul 1;11(7):NULL. 10.3390/NU11071505. PMID:31266155.	
2.	
3.	
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6.	

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