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Keratin 17 (D73C7) Rabbit mAb



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Applications: WB, IF-IC, FC-FP	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 48	Source/Isotype: Rabbit IgG	UniProt ID: #Q04695	Entrez-Gene Id 3872
Product Usage Information	Application					Dilution
	We	Western Blotting				1:1000
	Imr	Immunofluorescence (Immunocytochemistry)				1:200
	Flo	Flow Cytometry (Fixed/Permeabilized)				1:200
Storage		plied in 10 mM sodi 1% sodium azide. St	cerol and less than			
Specificity / Sensitiv	/ity Kera	Keratin 17 (D73C7) Rabbit mAb detects endogenous levels of keratin 17 protein.				

Species predicted to react based on 100% sequence homology: Monkey, Dog

Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to amino acids near the carboxy terminus of human keratin 17.

Background

Keratins (cytokeratins) are intermediate filament proteins that are mainly expressed in epithelial cells. Keratin heterodimers composed of an acidic keratin (or type I keratin, keratins 9 to 23) and a basic keratin (or type II keratin, keratins 1 to 8) assemble to form filaments (1,2). Keratin isoforms demonstrate tissueand differentiation-specific profiles that make them useful as research biomarkers (1). Research studies have shown that mutations in keratin genes are associated with skin disorders, liver and pancreatic diseases, and inflammatory intestinal diseases (3-6).

Keratin 17 is involved in wound healing and cell growth, two processes that require rapid cytoskeletal remodeling (7). Keratinocytes deficient in keratin 17 exhibit abnormal Akt/mTOR signaling and fail to produce an increase in translation, cell size, or growth; these cells also exhibit abnormal 14-3-3σ localization. As 14-3-3σ typically associates with keratin 17, these results imply that Akt/mTOR signaling results in sequestration of 14-3-3 σ with keratin 17 in the cytosol, which is required for translation and cell growth. Phosphorylation of keratin 17 on Ser44 may provide a docking site for 14-3-3σ binding (8).

Background References

- 1. Moll, R. et al. (1982) Cell 31, 11-24.
- 2. Chang, L. and Goldman, R.D. (2004) Nat Rev Mol Cell Biol 5, 601-13.
- 3. Ramaekers, F.C. and Bosman, F.T. (2004) J Pathol 204, 351-4.
- 4. Lane, E.B. and McLean, W.H. (2004) J Pathol 204, 355-66.
- 5. Zatloukal, K. et al. (2004) J Pathol 204, 367-76.
- 6. Owens, D.W. and Lane, E.B. (2004) J Pathol 204, 377-85.
- 7. Paladini, R.D. et al. (1996) J. Cell Biol. 132, 381-397.
- 8. Kim, S. et al. (2006) Nature 441, 362-365.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS,

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

FC-FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key

Keratin 17 (D73C7) Rabbit mAb (#4543) Datasheet Without Images Cell Signaling Technology

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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