# #12117 Store at -20C

# ITCH (D8Q6D) Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: WB, IP	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 105	Source/Isotype: Rabbit IgG	UniProt ID: #Q96J02	Entrez-Gene Id: 83737
Product Usage Information	Application			Dilution		
	We	stern Blotting		1:1000		
	Imr	nunoprecipitation		1:200		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at $-20^{\circ}$ C. Do not aliquot the antibody.				
Specificity / Sensitiv	ity ITCI	ITCH (D8Q6D) Rabbit mAb recognizes endogenous levels of total ITCH protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp125 of human ITCH protein.				

### **Background**

ITCH is a HECT domain-containing E3 ubiquitin ligase, first identified in genetic studies of the mouse *agouti* locus, in which mutations result in characteristic coat color changes. One particular *agouti* mutation (non-agouti-lethal 18H) is notable for the development of immunological defects not observed in other *agouti* mutant mice; these include lymphoid hyperplasia and chronic stomach, lung and skin inflammation (manifest as constant itching). The 18H *agouti* mutation was traced to a chromosomal inversion that disrupted expression of an adjacent gene in the *agouti* locus, subsequently termed *ltch* to reflect the chronic itching phenotype (1-3).

Further characterizations revealed that *Itch* encoded a NEDD4-like E3-ubiquitin ligase capable of catalyzing Lys29, Lys48, and/or Lys63-linked ubiquitination of target proteins, leading to their degradation by the proteosome pathway (4-6). The distinct phenotypes of *Itch* mutant mice led to the identification of an important regulatory role for ITCH-mediated ubiquitination in inflammatory signaling pathways. For example, ITCH-mediated ubiquitination of the transcription factor JunB was shown to play a direct inhibitory role in regulating expression of the proinflammatory cytokine IL-4. ITCH-null T lymphocytes consequently exhibit increased production of IL-4, leading to biased differentiation of naive CD4+ cells towards the proinflammatory Th2 lineage (7). In accordance with the findings from mutant *Itch* mouse models, a genetic linkage study in humans identified loss-of-function mutations in *ITCH* as a direct cause of syndromic multisystem autoimmune disease (SMAD) (8).

Notably, targets of ITCH-mediated ubiquitination are not restricted to immune signaling pathways. For example, key mediators of the Hedgehog (9,10), Wnt/ $\beta$ -catenin (11), Hippo (12), and Notch signaling pathways (13,14) have been identified as important targets of ITCH-mediated ubiquitination (2).

### **Background References**

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- 7. Fang, D. et al. (2002) Nat Immunol 3, 281-7.
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### **Species Reactivity**

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

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**Western Blot Buffer** 

ITCH (D8Q6D) Rabbit mAb (#12117) Datasheet Without Images Cell Signaling Technology IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry

milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

WB: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

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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