

# 0644

# ITM2B/Bri2 (E6O3Y) Rabbit mAb



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

<b>Applications:</b> W, IP, IF-IC	Reactivity:	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 42, 84	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q9Y287	Entrez-Gene Id: 9445
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence (Immunocytochemistry)			<b>Dilution</b> 1:1000 1:50 1:800 - 1:3200	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		ITM2B/Bri2 (E6O3Y) Rabbit mAb recognizes endogenous levels of total human ITM2B/Bri2 protein. Bands at 84 kDa and 21 kDa are occasionally detected, likely representing oligomers of ITM2B/Bri2 and cleavage products, respectively. This antibody does not cross-react with rodent orthologs of ITM2B/Bri2 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala105 of human ITM2B/Bri2 protein.				
Background		Integral membrane protein 2b (ITM2B), also known as Bri2, is a type II membrane protein. ITM2B is expressed as a precursor immature form and is processed by furin proteases to produce the mature ITM2B/Bri2 protein and a soluble C-terminal fragment (1,2). The membrane-bound ITM2B/Bri2 protein can be further processed by ADAM10 and intramembrane proteases (1). Several studies have implicated the <i>ITM2B</i> gene with familial forms of dementia and neurodegenerative diseases, including Alzheimer's disease (AD). Mutations in the human <i>ITM2B</i> gene are linked to several familial British and Danish dementia disorders (3,4). <i>ITM2B</i> gene mutations cause abnormal processing of the ITM2B/Bri2 protein, suggesting that the products of ITM2B/Bri2 protein cleavage might contribute directly to disease etiology (4). Interestingly, cleavage of disease-linked mutant <i>ITM2B</i> -encoded proteins generates peptides (ABri and ADan) that are more prone to deposit as amyloid fibrils, a pathological hallmark of many neurodegenerative diseases (2,4). Additionally, ITM2B/Bri2 interacts with Aβ-precursor protein (APP), a gene/protein linked to AD, and may alter APP processing and fibril formation (5-7). ITM2B/Bri2 is one of a family of proteins containing a BRICHOS domain, identified by their ability to inhibit Aβ fibril formation (8). The specific function of ITM2B/Bri2 is unclear, but it may contribute to normal synaptic function via an unknown mechanism (9).				
Background Refe	rences	<ol> <li>Tsachaki, M. et al. (2011) Glycobiology 21, 1382-8.</li> <li>Kim, S.H. et al. (1999) Nat Neurosci 2, 984-8.</li> <li>Vidal, R. et al. (1999) Nature 399, 776-81.</li> <li>Vidal, R. et al. (2000) Proc Natl Acad Sci U S A 97, 4920-5.</li> <li>Fotinopoulou, A. et al. (2005) J Biol Chem 280, 30768-72.</li> <li>Matsuda, S. et al. (2005) J Biol Chem 280, 28912-6.</li> <li>Matsuda, S. et al. (2008) J Neurosci 28, 8668-76.</li> <li>Matsuda, S. et al. (2011) Neurobiol Aging 32, 1400-8.</li> <li>Yao, W. et al. (2019) Sci Rep 9, 4862.</li> </ol>				

**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** 

 $\textbf{W:} \ \textbf{Western Blotting IP:} \ \textbf{Immunoprecipitation IF-IC:} \ \textbf{Immunofluorescence (Immunocytochemistry)}$ 

**Cross-Reactivity Key** 

H: Human

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