



# Mouse anti-Human BEST1 monoclonal antibody, clone F77 (CABT-B9850)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Immunogen</b>	A synthetic peptide (conjugated with KLH) corresponding to human BEST1.
<b>Isotype</b>	IgG1
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Clone</b>	F77
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB,IF
<b>Sequence Similarities</b>	KDHMDPYWALENRDEAHS
<b>Format</b>	Liquid
<b>Size</b>	100 µl
<b>Buffer</b>	In buffer containing 0.1% sodium azide
<b>Storage</b>	Store at -20°C or -80°C.

## BACKGROUND

<b>Introduction</b>	This gene encodes a member of the bestrophin gene family. This small gene family is characterized by proteins with a highly conserved N-terminus with four to six transmembrane domains. Bestrophins may form chloride ion channels or may regulate voltage-gated L-type calcium-ion channels. Bestrophins are generally believed to form calcium-activated chloride-ion
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channels in epithelial cells but they have also been shown to be highly permeable to bicarbonate ion transport in retinal tissue. Mutations in this gene are responsible for juvenile-onset vitelliform macular dystrophy (VMD2), also known as Best macular dystrophy, in addition to adult-onset vitelliform macular dystrophy (AVMD) and other retinopathies. Alternative splicing results in multiple variants encoding distinct isoforms.[provided by RefSeq, Nov 2008]

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<b>Keywords</b>	BEST1; bestrophin 1; ARB; BMD; BEST; RP50; VMD2; TU15B; bestrophin-1; Best disease; Best1V1Delta2; vitelliform macular dystrophy protein 2;
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## GENE INFORMATION

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<b>Entrez Gene ID</b>	<a href="#">7439</a>
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<b>UniProt ID</b>	<a href="#">O76090</a>
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<b>Function</b>	contributes_to chloride channel activity; ion channel activity
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