



## Anti-C10ORF2 monoclonal antibody (DCABH-10779)

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

## **PRODUCT INFORMATION**

Antigen Description	This gene encodes a hexameric DNA helicase which unwinds short stretches of double-stranded DNA in the 5 to 3 direction and, along with mitochondrial single-stranded DNA binding protein and mtDNA polymerase gamma, is thought to play a key role in mtDNA replication. The protein localizes to the mitochondrial matrix and mitochondrial nucleoids. Mutations in this gene cause infantile onset spinocerebellar ataxia (IOSCA) and progressive external ophthalmoplegia (PEO) and are also associated with several mitochondrial depletion syndromes. Alternative splicing results in multiple transcript variants encoding distinct isoforms
Immunogen	A synthetic peptide of human C10orf2 is used for rabbit immunization.
Isotype	IgG
Source/Host	Rabbit
Species Reactivity	Human
Purification	Protein A
Conjugate	Unconjugated
Applications	Western Blot (Transfected lysate); ELISA
Buffer	In 1x PBS, pH 7.4
Preservative	None
Storage	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## **GENE INFORMATION**

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Gene Name	C10orf2 chromosome 10 open reading frame 2 [ Homo sapiens ]
Official Symbol	C10orf2
Synonyms	C10ORF2; chromosome 10 open reading frame 2; infantile onset spinocerebellar ataxia (autosomal recessive), IOSCA; twinkle protein, mitochondrial; FLJ21832; PEO; PEO1; twinkle; TWINKLE; TWINL; ataxin 8; mitochondrial twinkle protein; T7-like mitochondrial DNA helicase; progressive external ophthalmoplegia 1 protein; T7 gp4-like protein with intramitochondrial nucleoid localization; SCA8; ATXN8; IOSCA; PEOA3; SANDO; MTDPS7;
Entrez Gene ID	<u>56652</u>
Protein Refseq	NP_001157284
UniProt ID	Q96RR1
Chromosome Location	10q24
Function	5-3 DNA helicase activity; ATP binding; hydrolase activity; nucleotide binding; protease binding; single-stranded DNA binding;