



Anti-DNAJC30 monoclonal antibody (DCABH-11303)

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

PRODUCT INFORMATION

Antigen Description	This intronless gene encodes a member of the DNAJ molecular chaperone homology domain-containing protein family. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23.
Immunogen	A synthetic peptide of human DNAJC30 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

Gene Name	DNAJC30 DnaJ (Hsp40) homolog, subfamily C, member 30 [Homo sapiens]
Official Symbol	DNAJC30

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Synonyms	DNAJC30; DnaJ (Hsp40) homolog, subfamily C, member 30; WBSCR18, Williams Beuren syndrome chromosome region 18; dnaJ homolog subfamily C member 30; Williams Beuren syndrome chromosome region 18; williams-Beuren syndrome chromosomal region 18 protein; WBSCR18; MGC12943;
Entrez Gene ID	84277
Protein Refseq	NP 115693
UniProt ID	B3KSU4
Chromosome Location	7q11.23
Function	heat shock protein binding; unfolded protein binding;