## NDUFA12 Antibody

PACO17787



Product Information	
Size:	Protein Background:
50ul	This gene encodes a protein which is part of mitochondrial complex 1, part of the
Reactivity:	oxidative phosphorylation system in mitochondria. Complex 1 transfers electrons to ubiquinone from NADH which establishes a proton gradient for the generation of ATP. Mutations in this gene are associated with Leigh syndrome due to mitochondrial complex 1 deficiency. Pseudogenes of this gene are located on chromosomes 5 and 13.
Human, Mouse	
Source:	Alternative splicing results in multiple transcript variants.
Rabbit	Gene ID:
lsotype:	NDUFA12
lgG	Uniprot
Applications:	Q9UI09
ELISA, IHC	Synonyms:
Recommended dilutions:	NADH dehydrogenase (ubiquinone) 1 alpha subcomplex, 12
ELISA:1:1000-1:10000, IHC:1:50-1:200	Immunogen:
	Synthetic peptide of human NDUFA12.
	Storage:

-20° C, pH7.4 PBS, 0.05% NaN3, 40% Glycerol



The image on the left is immunohistochemistry of paraffin-embedded Human brain tissue using PACO17787(NDUFA12 Antibody) at dilution 1/50, on the right is treated with synthetic peptide. (Original magnification: x—200).