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## anti-NDUFA7 antibody (AA 27-38)



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Quantity:	100 μg
Target:	NDUFA7
Binding Specificity:	AA 27-38
Reactivity:	Human
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This NDUFA7 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA

#### **Product Details**

Purpose:	NDUFA7 (aa27-38)	
Immunogen:	Peptide with sequence C-RYQEISKRTQPP, from the internal region of the protein sequence according to NP_004992.2.	
Sequence:	RYQEISKRTQ PP	
Isotype:	IgG	
Cross-Reactivity:	Cow, Human, Pig	
Purification:	Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.	
Grade:	Verified	

### **Target Details**

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Target:	NDUFA7	
Alternative Name:	NDUFA7 (NDUFA7 Products)	
Background:	NDUFA7, NADH dehydrogenase (ubiquinone) 1 alpha subcomplex, 7, 14.5 kDa, B14.5a, Cl-B14.5a, NADH dehydrogenase [ubiquinone] 1 alpha subcomplex subunit 7, NADH-ubiquinone oxidoreductase subunit B14.5a, complex I B14.5a subunit	
Gene ID:	4701	
NCBI Accession:	NP_004992	
Application Details		
Application Notes:	Western Blot: Approx 14 kDa band observed in Human Heart lysates (calculated MW of 12.6 kDa according to NP_004992.2). Recommended concentration: 0.1-0.3 µg/mL. Peptide ELISA: antibody detection limit dilution 1:128000.	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	0.5 mg/mL	
Buffer:	Supplied at 0.5 mg/mL in Tris saline, 0.02 % sodium azide, pH 7.3 with 0.5 % bovine serum albumin.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Handling Advice:	Minimize freezing and thawing.	
Storage:	-20 °C	
Storage Comment:	Aliquot and store at -20°C, with minimal freeze/thawing. A working aliquot may be refrigerate at 4°C for a few weeks and still remain viable.	

250kDa 150kDa 100kDa 75kDa 50kDa 37kDa 25kDa 20kDa 15kDa

#### **Western Blotting**

**Image 1.** ABIN1049507 (0.1 $\mu$ g/ml) staining of Human Heart lysate (35 $\mu$ g protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.