

Basic Information

Product Name	Anti-MT-CO1 Antibody	
Gene Name	MT-CO1	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, ICC/IF	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence at the C-terminus of human MTCO1.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	37 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 Immunocytochemistry/Immunofluorescence (ICC/IF): 1:50-400 (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied.

Background Information

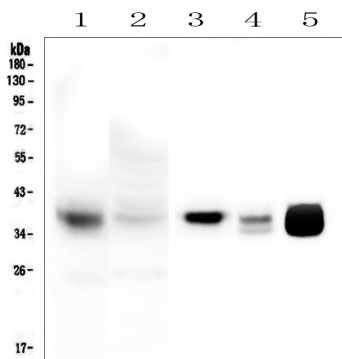
Cytochrome c oxidase subunit I (CO1 or MTCO1) is 1 of 3 mitochondrial DNA (mtDNA) encoded subunits (MTCO1, MTCO2, MTCO3) of respiratory Complex IV. Complex IV is located within the mitochondrial inner membrane and is the third and final enzyme of the electron transport chain of mitochondrial oxidative phosphorylation. It is composed of 13 polypeptides. Subunits I, II, and III (MTCO1, MTCO2, MTCO3) are encoded by mtDNA while subunits IV, Va, Vb, VIa, VIb, VIc, VIIa, VIIb, VIIc, and VIII are nuclear encoded. The cytochrome c oxidase family of enzymes have 4 redox centers, 2 hemes and 2 copper centers. In mitochondrial Complex IV, the 2 hemes are a and a₃ and the 2 coppers are CuA and CuB. The 2 hemes and CuB are bound to subunit I. Acin-Perez et al. (2003) identified a cell line containing single and double missense mutations in the cytochrome c oxidase (COX) subunit I gene of mouse mitochondrial DNA. And they hypothesized that deleterious mutations can arise and become predominant; cultured cells can maintain several mtDNA haplotypes at stable frequencies; the respiratory chain has little spare COX capacity; and that the size of a cavity in the vicinity of val421 in MTCO1 of animal COX may affect the function of the

enzyme.

Reference

Anti-MT-CO1 Antibody被引用在2文献中。

Selected Validation Data



Western blot analysis of MT-CO1 using anti-MT-CO1 antibody (BA2149).

The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human HeLa mitochondria lysates at 20ug,

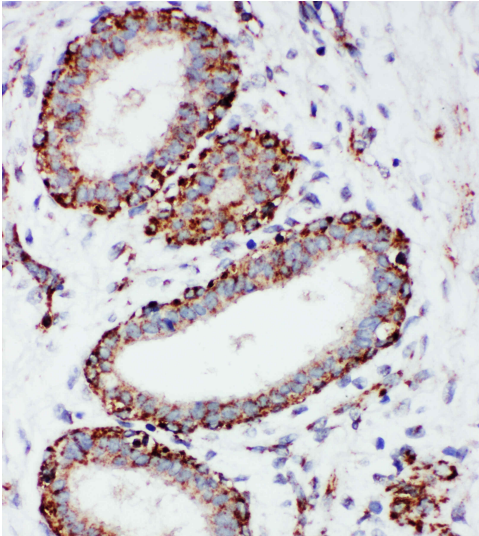
Lane 2: human HeLa whole cell lysates at 20ug,

Lane 3: human Caco-2 whole cell lysates at 50ug,

Lane 4: rat heart tissue lysates at 50ug,

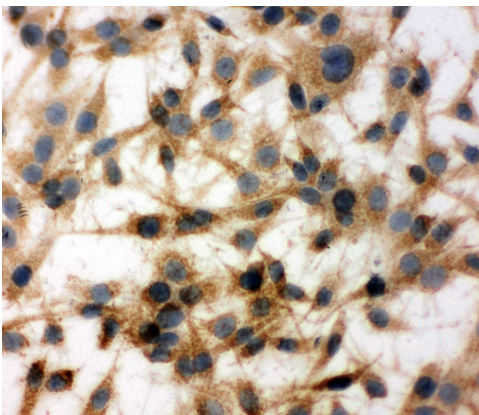
Lane 5: mouse heart tissue lysates at 50ug.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-MT-CO1 antigen affinity purified polyclonal antibody (BA2149) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for MT-CO1 at approximately 37 kDa. The expected band size for MT-CO1 is at 57 kDa.



IHC analysis of MT-CO1 using anti-MT-CO1 antibody (BA2149).

MT-CO1 was detected in a paraffin-embedded section of human mammary cancer tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-MT-CO1 Antibody (BA2149) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



ICC analysis of MT-CO1 using anti-MT-CO1 antibody (BA2149).

MT-CO1 was detected in an immunocytochemical section of C6 cells. The section was incubated with rabbit anti-MT-CO1 Antibody (BA2149) at a dilution of 1:100. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The section was developed using Streptavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.