

Product Data Sheet

Anti-NDUFV2 Antibody

Catalog # Source Reactivity Applications

CPA1792 Rabbit H, M, R, P WB, IH

Description Rabbit polyclonal antibody to NDUFV2

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human NDUFV2. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of NDUFV2 protein.

Clonality Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/50 - 1/100)

Gene Symbol NDUFV2

Alternative Names NADH dehydrogenase [ubiquinone] flavoprotein 2 mitochondrial; NADH-ubiquinone

oxidoreductase 24 kDa subunit

Entrez Gene 4729 (Human); 72900 (Mouse); 81728 (Rat)

SwissProt P19404 (Human); Q9D6J6 (Mouse); P19234 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

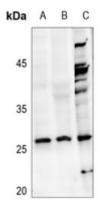
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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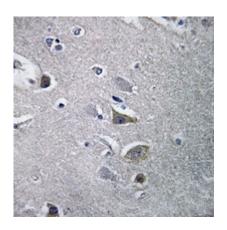
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Western blot analysis of NDUFV2 expression in mouse brain (A), rat stomach (B), THP1 (C) whole cell lysates. (Predicted band size: 27 kD; Observed band size: 27 kD)



Immunohistochemical analysis of NDUFV2 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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