

🔇 Tel: 400-999-8863 🛎 Email:3631691544@qq.com 🛛 🙆 Website: www.upingBio.com

NDUFS6 Mouse mAb

YP-mAb-18404
lgG
Human,Mouse,Rat
WB
Recombinant fusion protein containing a sequence corresponding to amino acids 29-124 of human NDUFS6 (NP_004544.1)
Affinity purification
WB 1:500 - 1:2000
1 mg/ml
≥90%
-20°C/1 year
14kDa
This gene encodes a subunit of the NADH:ubiquinone oxidoreductase (complex I), which is the first enzyme complex in the electron transport chain of mitochondria. This complex functions in the transfer of electrons from NADH to the respiratory chain. The subunit encoded by this gene is one of seven subunits in the iron-sulfur protein fraction. Mutations in this gene cause mitochondrial complex I deficiency, a disease that causes a wide variety of clinical disorders, including neonatal disease and adult-onset neurodegenerative disorders.
Avoid repeated freezing and thawing!

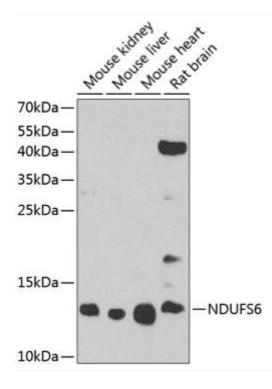


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Western blot analysis of extracts of various cell lines, using NDUFS6 antibody (A3985) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µ g per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 90s.