

## SDHB Polyclonal Antibody

<b>Catalog No :</b>	YT5450
<b>Reactivity :</b>	Human;Mouse;Rat;Fish
<b>Applications :</b>	WB;IHC;IF;ELISA
<b>Target :</b>	SDHB
<b>Fields :</b>	>>Citrate cycle (TCA cycle);>>Oxidative phosphorylation;>>Metabolic pathways;>>Carbon metabolism;>>Thermogenesis;>>Non-alcoholic fatty liver disease;>>Alzheimer disease;>>Parkinson disease;>>Amyotrophic lateral sclerosis;>>Huntington disease;>>Prion disease;>>Pathways of neurodegeneration - multiple diseases;>>Chemical carcinogenesis - reactive oxygen species;>>Diabetic cardiomyopathy
<b>Gene Name :</b>	SDHB
<b>Protein Name :</b>	Succinate dehydrogenase [ubiquinone] iron-sulfur subunit mitochondrial
<b>Human Gene Id :</b>	6390
<b>Human Swiss Prot No :</b>	P21912
<b>Mouse Gene Id :</b>	67680
<b>Mouse Swiss Prot No :</b>	Q9CQA3
<b>Rat Gene Id :</b>	298596
<b>Rat Swiss Prot No :</b>	P21913
<b>Immunogen :</b>	The antiserum was produced against synthesized peptide derived from the Internal region of human SDHB. AA range:131-180
<b>Specificity :</b>	SDHB Polyclonal Antibody detects endogenous levels of SDHB protein.
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1:500 - 1:2000. IHC: 1:100-1:300. ELISA: 1:20000.. IF 1:50-200
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)
<b>Observed Band :</b>	31kD
<b>Cell Pathway :</b>	Citrate cycle (TCA cycle);Oxidative phosphorylation;Alzheimer's disease;Parkinson's disease;Huntington's disease;
<b>Background :</b>	Complex II of the respiratory chain, which is specifically involved in the oxidation of succinate, carries electrons from FADH to CoQ. The complex is composed of four nuclear-encoded subunits and is localized in the mitochondrial inner membrane. The iron-sulfur subunit is highly conserved and contains three cysteine-rich clusters which may comprise the iron-sulfur centers of the enzyme. Sporadic and familial mutations in this gene result in paragangliomas and pheochromocytoma, and support a link between mitochondrial dysfunction and tumorigenesis. [provided by RefSeq, Jul 2008],
<b>Function :</b>	catalytic activity:Succinate + ubiquinone = fumarate + ubiquinol.,cofactor: Binds 1 2Fe-2S cluster.,cofactor: Binds 1 3Fe-4S cluster.,cofactor: Binds 1 4Fe-4S cluster.,disease: Defects in SDHB are a cause of Cowden-like syndrome [MIM:612359]. Cowden-like syndrome is a cancer predisposition syndrome associated with elevated risk for tumors of the breast, thyroid, kidney and uterus.,disease: Defects in SDHB are a cause of paraganglioma and gastric stromal sarcoma [MIM:606864]; also called Carney-Stratakis syndrome. Gastrointestinal stromal tumors may be sporadic or inherited in an autosomal dominant manner, alone or as a component of a syndrome associated with other tumors, such as in the context of neurofibromatosis type 1 (NF1). Patients have both gastrointestinal stromal tumors and paragangliomas. Susceptibility to the tumors was inherited in an apparently autosomal dominant manner, with inc
<b>Subcellular Location :</b>	Mitochondrion inner membrane; Peripheral membrane protein; Matrix side.
<b>Expression :</b>	Brain,Fibroblast,Liver,

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