

## **IDH1 Polyclonal Antibody**

Catalog No: YT5416

**Reactivity:** Human; Mouse; Rat

**Applications:** WB;IHC;IF;ELISA

Target: IDH1

**Fields:** >>Citrate cycle (TCA cycle);>>Glutathione metabolism;>>Metabolic

pathways;>>Carbon metabolism;>>2-Oxocarboxylic acid

metabolism;>>Biosynthesis of amino acids;>>Peroxisome;>>Central carbon

metabolism in cancer

Gene Name: IDH1

**Protein Name:** Isocitrate dehydrogenase [NADP] cytoplasmic

075874

O88844

Human Gene Id: 3417

**Human Swiss Prot** 

No:

Mouse Gene Id: 15926

**Mouse Swiss Prot** 

No:

Rat Gene ld: 24479

Rat Swiss Prot No: P41562

**Immunogen:** Synthesized peptide derived from the N-terminal region of human IDH1.

**Specificity:** IDH1 Polyclonal Antibody detects endogenous levels of IDH1 protein.

**Formulation:** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution :** WB 1:500 - 1:2000. IHC: 1:100-1:300. ELISA: 1:20000.. IF 1:50-200

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**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 46kD

**Cell Pathway :** Citrate cycle (TCA cycle);Glutathione metabolism;

**Background:** Isocitrate dehydrogenases catalyze the oxidative decarboxylation of isocitrate to

2-oxoglutarate. These enzymes belong to two distinct subclasses, one of which utilizes NAD(+) as the electron acceptor and the other NADP(+). Five isocitrate dehydrogenases have been reported: three NAD(+)-dependent isocitrate

dehydrogenases, which localize to the mitochondrial matrix, and two

NADP(+)-dependent isocitrate dehydrogenases, one of which is mitochondrial and the other predominantly cytosolic. Each NADP(+)-dependent isozyme is a homodimer. The protein encoded by this gene is the NADP(+)-dependent

isocitrate dehydrogenase found in the cytoplasm and peroxisomes. It contains the PTS-1 peroxisomal targeting signal sequence. The presence of this enzyme in peroxisomes suggests roles in the regeneration of NADPH for intraperoxisomal

reductions, such as the conversion of 2, 4-dienoyl-CoAs to

Function: catalytic activity: Isocitrate + NADP(+) = 2-oxoglutarate + CO(2) +

NADPH.,catalytic activity:Oxalosuccinate + NADP(+) = 2-oxoglutarate + CO(2) +

NADPH., cofactor: Binds 1 magnesium or manganese ion per

subunit., disease: Defects in IDH1 are a cause of glioblastoma multiforme (GBM) [MIM:137800]; also called familial glioma of brain. Gliomas are central nervous

system neoplasms derived from glial cells and comprise astrocytomas,

glioblastoma multiforme, oligodendrogliomas, and

ependymomas.,miscellaneous:Cancer mutations affecting Arg-132 are tissuespecific, and suggest that this residue plays a unique role in the development of

high-grade gliomas., online information: Isocitrate dehydrogenase

entry, similarity: Belongs to the isocitrate and isopropylmalate dehydrogenases

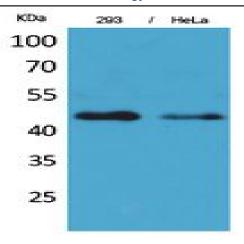
family., subunit: Homodimer.,

Subcellular Location:

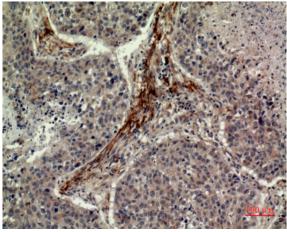
Cytoplasm, cytosol . Peroxisome .

**Expression:** Brain, Cajal-Retzius cell, Fetal brain cortex, Human endometri

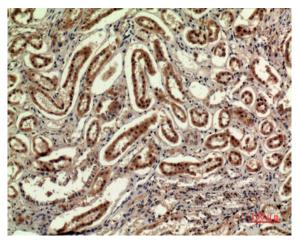
## **Products Images**



Western Blot analysis of 293, HeLa cells using IDH1 Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Immunohistochemical analysis of paraffin-embedded humanlung, antibody was diluted at 1:100



Immunohistochemical analysis of paraffin-embedded humankidney, antibody was diluted at 1:100