



**Orders:** 877-616-CELL (2355)  
orders@cellsignal.com

**Support:** 877-678-TECH (8324)

**Web:** info@cellsignal.com  
cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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## IDH1 (D2H1) Rabbit mAb

**For Research Use Only. Not for Use in Diagnostic Procedures.**

<b>Applications:</b> W, IP, IF-IC, FC-FP	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 46	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #O75874	<b>Entrez-Gene Id:</b> 3417
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### Product Usage Information

#### Application

Western Blotting  
Immunoprecipitation  
Immunofluorescence (Immunocytochemistry)  
Flow Cytometry (Fixed/Permeabilized)

#### Dilution

1:1000  
1:50  
1:400  
1:200

### Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

For a carrier free (BSA and azide free) version of this product see product #55269.

### Specificity/Sensitivity

IDH1 (D2H1) Rabbit mAb recognizes endogenous levels of total IDH1 protein. This antibody does not recognize endogenous IDH2 protein, but does recognize IDH2 when recombinantly overexpressed.

### Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Arg222 of human IDH1 protein.

### Background

IDH1 is one of three isocitrate dehydrogenases that catalyze the oxidative decarboxylation of isocitrate to α-ketoglutarate (α-KG). These enzymes exist in two distinct subclasses that utilize either NAD or NADP<sup>+</sup> respectively, as an electron acceptor (1). IDH1 is the NADP<sup>+</sup>-dependent isocitrate dehydrogenase found in the cytoplasm and peroxisomes. IDH2 and 3 are mitochondrial enzymes that also function in the Krebs cycle. IDH1 is inactivated by phosphorylation at Ser113 and contains a clasp-like domain wherein both polypeptide chains in the dimer interlock (2,3). IDH1 is expressed in a wide range of species and also in organisms that lack a complete citric acid cycle. Mutations in IDH1 have been reported in glioblastoma (4), acute myeloid leukemia (5,6), and other malignancies (7). IDH1 appears to function as a tumor suppressor that, when mutationally inactivated, contributes to tumorigenesis in part through induction of the HIF-1 pathway (8).

### Background References

1. Ramachandran, N. and Colman, R.F. (1980) *J Biol Chem* 255, 8859-64.
2. Bennett, P.M. and Holms, W.H. (1975) *J Gen Microbiol* 87, 37-51.
3. Hurley, J.H. et al. (1990) *Science* 249, 1012-6.
4. Bleeker, F.E. et al. (2009) *Hum Mutat* 30, 7-11.
5. Abbas, S. et al. (2010) *Blood* 116, 2122-6.
6. Paschka, P. et al. (2010) *J Clin Oncol* 28, 3636-43.
7. Watanabe, T. et al. (2009) *Am J Pathol* 174, 1149-53.
8. Zhao, S. et al. (2009) *Science* 324, 261-5.

### Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

### Western Blot Buffer

**IMPORTANT:** For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

### Applications Key

**W:** Western Blotting **IP:** Immunoprecipitation **IF-IC:** Immunofluorescence (Immunocytochemistry) **FC-FP:** Flow Cytometry (Fixed/Permeabilized)

### Cross-Reactivity Key

**H:** Human **M:** Mouse **R:** Rat **Mk:** Monkey

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