

Store at  
-20C  
#64326**CD133 (D2V8Q) XP<sup>®</sup> Rabbit mAb**

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

<b>Applications:</b> W, IHC-Bond, IHC-P, IF-IC	<b>Reactivity:</b> H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 133	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #O43490	<b>Entrez-Gene Id:</b> 8842
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**Product Usage Information****Application**

Western Blotting  
IHC Leica Bond  
Immunohistochemistry (Paraffin)  
Immunofluorescence (Immunocytochemistry)

**Dilution**

1:1000  
1:200 - 1:800  
1:200 - 1:800  
1:200 - 1:800

**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 #51917.

**Specificity/Sensitivity**

CD133 (D2V8Q) XP<sup>®</sup> Rabbit mAb recognizes endogenous levels of total CD133 protein. This antibody is not sensitive to the glycosylation status of CD133.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the 1st extracellular loop of human CD133 protein. The epitope has been mapped to a region spanning amino acid residues 303-312.

**Background**

CD133, also known as Prominin, was first described as a cell surface marker recognized by monoclonal antibody AC133 on putative hematopoietic stem cells (1). Subsequent cDNA cloning indicated that CD133 is a five-transmembrane protein with a predicted molecular weight of 97 kDa. Due to heavy glycosylation, its apparent molecular weight is 130 kDa as determined by SDS-PAGE analysis (2). Besides blood stem cells, CD133 is expressed on and used to isolate other stem cells, including cancer stem cells (3-7). A deletion mutation in CD133 produces aberrant protein localization and may result in retinal degeneration in humans (8).

**Background References**

1. Yin, A.H. et al. (1997) *Blood* 90, 5002-12.
2. Miraglia, S. et al. (1997) *Blood* 90, 5013-21.
3. Handgretinger, R. et al. (2003) *Ann N Y Acad Sci* 996, 141-51.
4. Monzani, E. et al. (2007) *Eur J Cancer* 43, 935-46.
5. O'Brien, C.A. et al. (2007) *Nature* 445, 106-10.
6. Ricci-Vitiani, L. et al. (2007) *Nature* 445, 111-5.
7. Singh, S.K. et al. (2004) *Nature* 432, 396-401.
8. Maw, M.A. et al. (2000) *Hum. Mol. Genet.* 9, 27-34.

**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 **IHC-Bond:** IHC Leica Bond **IHC-P:** Immunohistochemistry (Paraffin) **IF-IC:** Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key**

**H:** Human

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