

Store at  
-20°C  
#47061**CELSR2 (D2M9H) XP® Rabbit mAb**

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

| Applications: | Reactivity: | Sensitivity: | MW (kDa): | Source/Isotype: | UniProt ID: | Entrez-Gene Id: |
|---------------|-------------|--------------|-----------|-----------------|-------------|-----------------|
| W, IHC-P      | H           | Endogenous   | 320       | Rabbit IgG      | #Q9HCU4     | 1952            |

**Product Usage Information****Application**

Western Blotting  
Immunohistochemistry (Paraffin)

**Dilution**

1:1000  
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 #19983.

**Specificity/Sensitivity**

CELSR2 (D2M9H) XP® Rabbit mAb recognizes endogenous levels of total CELSR2 protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding His1781 of human CELSR2 protein.

**Background**

CELSR2 (cadherin EGF LAG seven-pass G-type receptor, also known as flamingo homolog 3 or epidermal growth factor-like protein 2) is a member of the flamingo subfamily of non-classical cadherins, part of the cadherin superfamily. CELSR2 is a 7-transmembrane helix receptor that contains nine cadherin-like domains, seven EGF-like repeats, and 2 laminin A G-type repeats (1). It shares structural characteristics of both an adhesion molecule and a G protein-coupled receptor, suggesting putative roles in both cell-cell adhesion and juxtacrine signaling. Its function has been associated with dendrite morphogenesis (2), neural plate anterior-posterior pattern formation (3), and regulation of transcription via the Wnt signaling pathway (4). In a loss-of-function mouse model, Celsr2 deletion resulted in defects in the planar organization of ependymal cilia, leading to defective cerebrospinal fluid dynamics and hydrocephalus (5). In humans, SNPs in the CELSR2 gene cluster on chromosome 1 have been associated with enhanced risk of coronary artery disease (6).

**Background References**

1. Vincent, J.B. et al. (2000) *DNA Res* 7, 233-5.
2. Shima, Y. et al. (2004) *Dev Cell* 7, 205-16.
3. Devenport, D. and Fuchs, E. (2008) *Nat Cell Biol* 10, 1257-68.
4. Qu, Y. et al. (2014) *Proc Natl Acad Sci U S A* 111, E2996-3004.
5. Tissir, F. et al. (2010) *Nat Neurosci* 13, 700-7.
6. Zhou, Y.J. et al. (2015) *Int J Clin Exp Pathol* 8, 9543-51.

**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 BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key**

**W:** Western Blotting **IHC-P:** Immunohistochemistry (Paraffin)

**Cross-Reactivity Key**

**H:** Human

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