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## CEND1 (D6A6) Rabbit mAb

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

<b>Applications:</b> W, IP, IF-F	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 22	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q8N111	<b>Entrez-Gene Id:</b> 51286
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### Product Usage Information

#### Application

Western Blotting  
Immunoprecipitation  
Immunofluorescence (Frozen)

#### Dilution

1:1000  
1:100  
1:400

### 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.

### Specificity/Sensitivity

CEND1 (D6A6) Rabbit mAb recognizes endogenous levels of total CEND1 protein.

### Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human CEND1 protein.

### Background

The progression of progenitor cells towards neuronal differentiation is regulated by cell cycle control and the transition from proliferative to neurogenic cell divisions. Cell cycle exit and neuronal differentiation 1 (CEND1) is a neuronal protein widely expressed in the adult nervous system (1). It is implicated in the synchronization of cell cycle exit and differentiation of neuronal precursors in the developing nervous system, and its expression marks the exit of proliferative cells from the cell cycle (2,3). Levels of CEND1 expression in the subventricular zone of the adult nervous system are critical for cell cycle control and neuronal differentiation mechanisms during neonatal SVZ neurogenesis (4). It has recently been shown that neural progenitor cells (NPCs) that overexpress CEND1 display increased neuronal differentiation in a mouse model of brain injury, suggesting its potential use as a therapeutic intervention for neurodegenerative diseases and brain injury (5).

### Background References

1. Patsavoudi, E. et al. (1995) *J Neurosci Res* 40, 506-18.
2. Politis, P.K. et al. (2007) *Proc Natl Acad Sci USA* 104, 17861-6.
3. Koutmani, Y. et al. (2004) *Eur J Neurosci* 20, 2509-23.
4. Katsimpardi, L. et al. (2008) *Stem Cells* 26, 1796-807.
5. Makri, G. et al. (2010) *Stem Cells* 28, 127-39.

### 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 **IP:** Immunoprecipitation **IF-F:** Immunofluorescence (Frozen)

### Cross-Reactivity Key

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

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