

### **Product Data Sheet**

# Anti-HEN1/2 Antibody

Catalog # Source Reactivity Applications

CPA3994 Rabbit H, M, R, C WB, IH, IF/IC

**Description** Rabbit polyclonal antibody to HEN1/2

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human HEN1/2. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of HEN1/2 protein.

**Clonality** Polyclonal

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500)

Gene Symbol NHLH1; NHLH2

Alternative Names NHLH1; BHLHA35; HEN1; Helix-loop-helix protein 1; HEN-1; Class A basic

helix-loop-helix protein 35; bHLHa35; Nescient helix loop helix 1; NSCL-1; NHLH2;

BHLHA34; HEN2; KIAA0490; Helix-loop-helix protein 2; HEN-2; Class A basic

helix-loop-helix protein 34; bHLHa34; Nescient helix loop helix 2; NSCL-2

Entrez Gene 4807, 4808 (Human); 18071, 18072 (Mouse)

SwissProt Q02575, Q02577 (Human); Q02576, Q64221 (Mouse)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

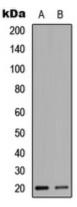
freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

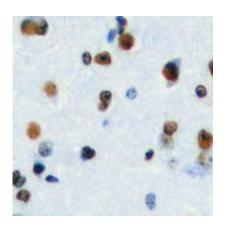
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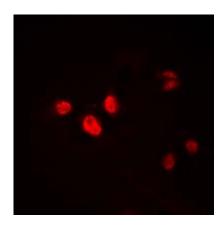
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Western blot analysis of HEN1/2 expression in HEK293T (A), NIH3T3 (B) whole cell lysates.



Immunohistochemical analysis of HEN1/2 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. w



Immunofluorescent analysis of HEN1/2 staining in HEK293T cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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