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HES1 (D6P2U) Rabbit mAb

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Entrez-Gene ID #3280 UniProt ID #Q14469

100 μl (10 western blots)

For Research Use Only. Not For Use In Diagnostic Procedures.

Applications	
W, IP, IHC-P	
Endogenous	

Species Cross-Reactivity*

Molecular Wt. 30 kDa Isotype Rabbit IgG**

rev. 06/17/14

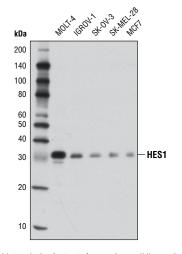
Background: HES1 (Hairy and Enhancer of Split 1) is one of seven members of the HES family of basic helix-loop-helix (bHLH) transcription factors which function primarily to repress transcription of bHLH-dependent genes (1). HES1 is understood to play an important conserved role in maintaining pluripotency of embryonic and adult stem/progenitor cells via the transcriptional repression of genes that promote differentiation (1,2). HES1 is particularly well known as a repressive mediator of the canonical Notch signaling pathway (3). HES1 plays a key role in mediating Notch-dependent T cell lineage commitment (4), and has been reported to be an essential mediator of Notch-induced T cell acute lymphoblastic leukemia (T-ALL) (4,5). HES1 is also reported to mediate Notch-induced repression of differentiation in a number of cancer cell types. A conditional deletion of HES1 from intestinal tumor cells in APC-mutant mice reduced tumor cell proliferation, while promoting differentiation toward epithelial lineages (6). Overexpression of HES1 in a human osteosarcoma (OS) cell line was shown to repress expression of the Notch antagonist Dtx1, leading to increased OS cell invasiveness (7). Other genes subject to transcriptional repression by HES1 include Neurogenin-2, Math1/Atoh1 and the NOTCH ligands DLL1 and Jagged1 (6,8,9).

Specificity/Sensitivity: HES1 (D6P2U) Rabbit mAb recognizes endogenous levels of total HES1 protein.

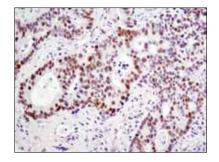
Source/Purification: Monoclonal antibody is produced by immunizing animals with recombinant protein specific to human HES1 protein.

Background References:

- (1) Kageyama, R. et al. (2007) Development 134, 1243-51.
- (2) Hatakeyama, J. et al. (2004) Development 131, 5539-50.
- (3) Kobayashi, T. and Kageyama, R. (2010) *Genes Cells* 15, 689-98.
- (4) Wendorff, A.A. et al. (2010) Immunity 33, 671-84.
- (5) Espinosa, L. et al. (2010) *Cancer Cell* 18, 268-81.
- (6) Ueo, T. et al. (2012) *Development* 139, 1071-82.
- (7) Zhang, P. et al. (2010) *Oncogene* 29, 2916-26.
- (8) Kageyama, R. et al. (2008) *Dev Growth Differ* 50 Suppl 1, S97-103.
- (9) Kobayashi, T. et al. (2009) Genes Dev 23, 1870-5.



Western blot analysis of extracts from various cell lines using HES1 (D6P2U) Rabbit mAb.



Immunohistochemical analysis of paraffin-embedded human lung carcinoma using HES1 (D6P2U) Rabbit mAb.

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

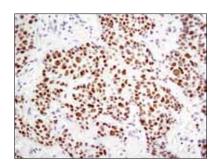
- *Species cross-reactivity is determined by western blot.
- **Anti-rabbit secondary antibodies must be used to detect this antibody.

Recommended Antibody Dilutions:

Western blotting 1:1000
Immunoprecipitation 1:200
Immunohistochemistry (Paraffin) 1:6400†
Unmasking buffer: Citrate
Antibody diluent: SignalStain® Antibody Diluent #8112
Detection reagent: SignalStain® Boost (HRP, Rabbit) #8114
† Optimal IHC dilutions determined using SignalStain® Boost

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IHC Detection Reagent.



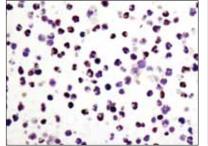
Immunohistochemical analysis of paraffin-embedded human breast carcinoma using HES1 (D6P2U) Rabbit mAb.

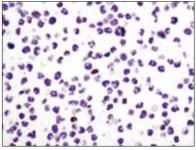
IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v BSA, 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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Tween®20 at 4°C with gentle shaking, overnight.





Immunohistochemical analysis of paraffin-embedded CUTTL1 cell pellets, control (left) or Compound E-treated (right), using HES1 (D6P2U) Rabbit mAb.