

Basic Information

Product Name	Anti-ZEB1 Antibody	
Gene Name	ZEB1	
Source	Rabbit	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, IHC, ICC/IF, FCM	
Contents	500 ug/ml antibody with PBS , 0.02% NaN3 , 1 mg BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence of human ZEB1 (LLKAYYALNAQPSAEELSKIADSVNLPLDVVKKWFEKMQ).	
concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	200KD	
Dilution Ratios	Western blot(WB): 1:500-2000 Immunohistochemistry in paraffin section (IHC): 1:50-400 Immunocytochemistry in fixed cells: 1:50-400 Flow cytometry (FCM): 1-3 μ g/1x10 ⁶ cells (Boiling the paraffin sections in 10mM citrate buffer,pH6.0,or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

ZEB1 (Zinc Finger E Box-Binding Homeobox 1), also called TCF8, NIL2A or DELTA-EF1, is a protein that in humans is encoded by the ZEB1 gene. Fluorescence in situ hybridization localized the ZEB1 gene to chromosome 10p11.2. Krafchak et al. (2005) demonstrated a complex (core plus secondary) binding site for TCF8 in the promoter of the COL4A3 gene, mutant in Alport syndrome and which encodes collagen type IV alpha-3. They detected expression of TCF8 in cornea. Nishimura et al. (2006) found that delta-Ef1 was upregulated during differentiation in a mouse smooth muscle cell (SMC) line.

Reference

Anti-ZEB1 Antibody被引用在1文献中。

Selected Validation Data

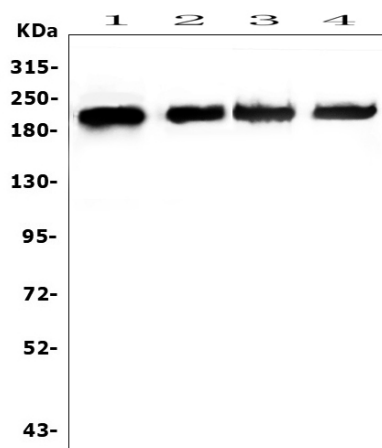


Figure 1. Western blot analysis of anti-ZEB1 antibody (A00548-2).. The sample well of each lane was loaded with 50ug of sample under reducing conditions. Lane 1: human SGC-7901 whole cell lysates, Lane 2: human U-87MG whole cell lysates, Lane 3: human HEK293 whole cell lysates, Lane 4: human PC-3 whole cell lysates. Use rabbit anti- ZEB1 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002). A specific band was detected for ZEB1 at approximately 200KD. The expected band size for ZEB1 is at 124KD.

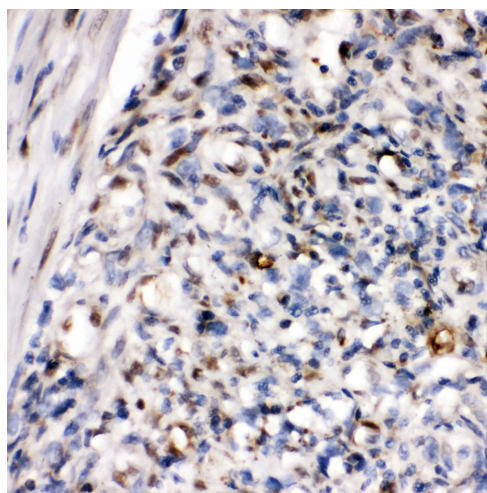


Figure 2. IHC analysis using anti- ZEB1 antibody (A00548-2). detected in paraffin-embedded section of human melanoma tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

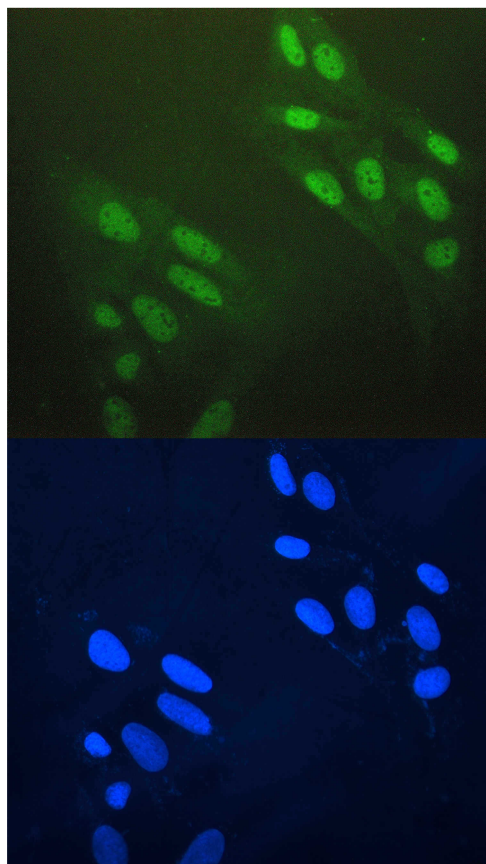


Figure 4. ICC analysis using anti-ZEB1 antibody (A00548-2) was detected in immersion fixed U2OS cell line. Cells were stained using the Dylight488-conjugated Anti-rabbit IgG Secondary Antibody (green)(Catalog#BA1127) and counterstained with DAPI (blue).

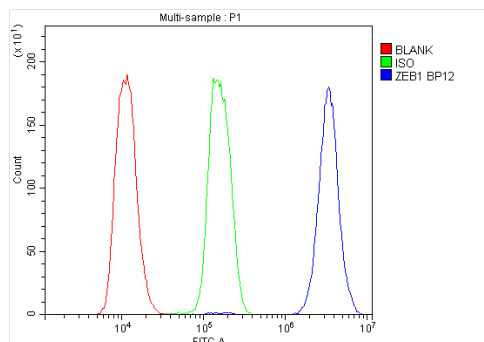


Figure 5. Flow cytometry analysis of U2OS cell (1×10^6). DyLight 488 conjugated goat anti-rabbit IgG (blue) was used as secondary antibody. Isotype control antibody (Green line) was rabbit IgG DyLight 488. Unlabelled sample (Red line).