CBFA1 Antibody

Catalog No: #31165

Package Size: #31165-1 50ul #31165-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

Description

Product Name	CBFA1 Antibody	
Host Species	Rabbit	
Clonality	Polyclonal	
Applications	ELISA WB IHC	
Species Reactivity	Hu Ms	
Specificity	The antibody detects endogenous level of total CBFA1 protein.	
Immunogen Type	Peptide	
Immunogen Description	Synthetic peptide corresponding to a region derived from 320-336 amino acids of Human Core-binding factor	
	subunit alpha-1	
Target Name	CBFA1	
Other Names	Core-binding factor subunit alpha-1 , CCD, AML3, CCD1, OSF2, OSF-2, PEA2aA, PEBP2A1, PEBP2A2,	
	PEBP2aA, PEBP2aA1	
Accession No.	Swiss-Prot:Q13950Gene ID:860;	
Uniprot	Q13950	
GeneID	860;	
Concentration	1.0mg/ml	
Formulation	Rabbit IgG in pH7.4 PBS, 0.05% NaN3, 40% Glycerol.	
Storage	Store at -20°C/1 year	

Application Details

Predicted MW: 54kd	
ELISA: 1:500-1:5000	
Western blotting: 1:200-1:1000	
Immunohistochemistry: 1:10-1:50	

Images

$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Gel: 10%SDS-PAGE Lane1: Human renal cancer tissue lysate Lane2: Human leg malignant fibrous histiotoma tissue lysate Lysates: 10 ug per lane Primary antibody: 1/400 dilution Secondary antibody: Donkey anti Rabbit IgG - H&L (HRP) at 1/5000 dilution Exposure time: 1 minute
40 — 35 —	Exposure time: 1 minute



The image on the left is immunohistochemistry of paraffin-embedded Human brain tissue using 31165(CBFA1 Antibody) at dilution 1/10, on the right is treated with the synthetic peptide.

Background

This gene is a member of the RUNX family of transcription factors and encodes a nuclear protein with an Runt DNA-binding domain. This protein is essential for osteoblastic differentiation and skeletal morphogenesis and acts as a scaffold for nucleic acids and regulatory factors involved in skeletal gene expression. The protein can bind DNA both as a monomer or, with more affinity, as a subunit of a heterodimeric complex. Mutations in this gene have been associated with the bone development disorder cleidocranial dysplasia (CCD). Transcript variants that encode different protein isoforms result from the use of alternate promoters as well as alternate splicing.

Note: This product is for in vitro research use only