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## Datasheet for ABIN6244147 anti-SKOR2 antibody (C-Term)

2 Images



## Overview

Quantity:	400 µL
Target:	SKOR2
Binding Specificity:	AA 741-775, C-Term
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SKOR2 antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF)

## Product Details

Immunogen:	This SKOR2 antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 741-775 amino acids from the C-terminal region of human SKOR2.
Clone:	RB51023
Isotype:	Ig Fraction
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.

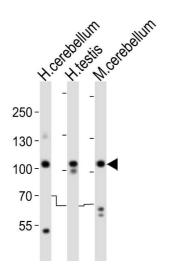
## Target Details

Target:	SKOR2
Alternative Name:	SKOR2 (SKOR2 Products)
Background:	Exhibits transcriptional repressor activity (By similarity). Acts as a TGF-beta antagonist in the

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Target Details	
	nervous system.
Molecular Weight:	104235
UniProt:	Q2VWA4
Application Details	
Application Notes:	IF: 1:2000. WB: 1:1000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Purified polyclonal antibody supplied in PBS with 0.09 $\%$ (W/V) sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Expiry Date:	6 months

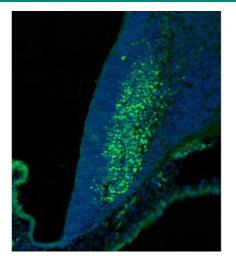
### Images



#### Western Blotting

**Image 1.** Western blot analysis of lysates from human cerebellum, human testis, mouse cerebellum tissue lysate (from left to right), using SKOR2 Antibody (C-term) (ABIN6244147 and ABIN6577773). (ABIN6244147 and ABIN6577773) was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20 µg per lane.

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#### Immunofluorescence

**Image 2.** (ABIN6244147 and ABIN6577773) staining SKOR2 in mouse cerebellum at embryonic stage 12.5 sections by Immunohistochemistry (IHC-P - 4 % paraformaldehydefixed, cryopreserved in 30 % sucrose/1X PBS for 30 hours, and then cut at a cryotome). Tissue was fixed with formaldehyde and blocked with 3 % BSA for 0.5 hour at room temperature. Samples were incubated with primary antibody (1/2000) for 1 hours at 37 °C.Immunofluorescence image showing purkinje cell neurons staining on mouse cerebellum tissue.The nuclear counter stain is DI (blue).

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