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# Datasheet for ABIN6257219 anti-GPR153 antibody (Internal Region)

3 Images



Overview

Quantity:	100 µL
Target:	GPR153
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GPR153 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunohistochemistry (IHC), Immunocytochemistry (ICC)

## Product Details

Immunogen:	A synthesized peptide derived from human GPR153, corresponding to a region within the internal amino acids.
Isotype:	lgG
Specificity:	GPR153 Antibody detects endogenous levels of total GPR153.
Predicted Reactivity:	Pig,Zebrafish,Bovine,Horse,Sheep,Dog,Chicken,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

## Target Details

GPR153

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## Target Details

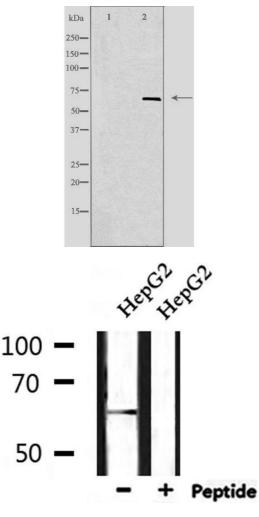
Alternative Name:	GPR153 (GPR153 Products)
Background:	Description: Orphan receptor. Gene: GPR153
Molecular Weight:	65 kDa
Molecular Weight: Gene ID:	65 kDa 387509

### **Application Details**

Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

# Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
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:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

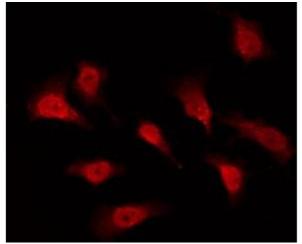


### Western Blotting

**Image 1.** Western blot analysis of extracts from K562 cells, using GPR153 antibody. The lane on the left is treated with the antigen-specific peptide.

#### Western Blotting

**Image 2.** Western blot analysis of extracts from HepG2, using GPR153 Antibody.



#### Immunofluorescence (fixed cells)

**Image 3.** ABIN6276016 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25<sub>j</sub>ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37<sub>j</sub>ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod

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