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# anti-HBG2 antibody (C-Term)





Overview

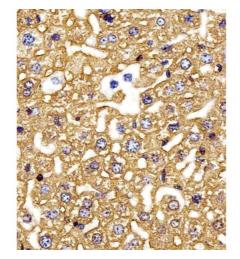
Quantity:	400 μL
Target:	HBG2
Binding Specificity:	AA 101-133, C-Term
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This HBG2 antibody is un-conjugated
Application:	Western Blotting (WB), Flow Cytometry (FACS), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))
Product Details	
Immunogen:	This HBG2 antibody is generated from a rabbit immunized with a KLH conjugated synthetic
Immunogen:	This HBG2 antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 101-133 amino acids from the C-terminal region of human HBG2.
Immunogen: Isotype:	
	peptide between 101-133 amino acids from the C-terminal region of human HBG2.
Isotype:	peptide between 101-133 amino acids from the C-terminal region of human HBG2.  Ig Fraction
Isotype: Purification:	peptide between 101-133 amino acids from the C-terminal region of human HBG2.  Ig Fraction
Isotype: Purification: Target Details	peptide between 101-133 amino acids from the C-terminal region of human HBG2.  Ig Fraction  This antibody is purified through a protein A column, followed by peptide affinity purification.

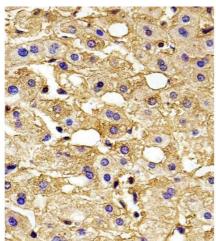
# **Target Details**

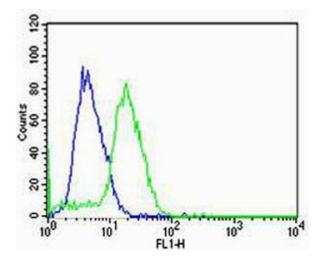
Molecular Weight:	16 kDa
Gene ID:	3048
UniProt:	P69892

Application Details	
Application Notes:	For IHC-P starting dilution is: 1:25
	For FACS starting dilution is: 1:25
	For WB starting dilution is: 1:1000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	0.35 mg/mL

Format:	Liquid
Concentration:	0.35 mg/mL
Buffer:	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
Storage Comment:	Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high temperatures.







## **Immunohistochemistry**

**Image 1.** Immunohistochemical analysis of paraffinembedded M. liver section using HBG2 Antibody . Antibody was diluted at 1:100 dilution. A peroxidase-conjugated goat anti-rabbit IgG at 1:400 dilution was used as the secondary antibody, followed by DAB staining.

### **Immunohistochemistry**

**Image 2.** Immunohistochemical analysis of paraffinembedded H. liver section using HBG2 Antibody . Antibody was diluted at 1:100 dilution. A peroxidase-conjugated goat anti-rabbit IgG at 1:400 dilution was used as the secondary antibody, followed by DAB staining.

## **Flow Cytometry**

**Image 3.** Flow cytometric analysis of U-87 MG cells using HBG2 Antibody (green) compared to an isotype control of rabbit IgG(blue). Antibody was diluted at 1:25 dilution. An Alexa Fluor 488 goat anti-rabbit IgG at 1:400 dilution was used as the secondary antibody.