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## anti-GHRHR antibody (C-Term)

**Images** 



Overview	
Quantity:	100 μL
Target:	GHRHR
Binding Specificity:	C-Term
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GHRHR antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human GHRHR, corresponding to a region within C-terminal amino acids.
Isotype:	IgG
Specificity:	GHRHR Antibody detects endogenous levels of total GHRHR.
Predicted Reactivity:	Pig,Horse,Rabbit,Dog,Chicken
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	GHRHR

## **Target Details**

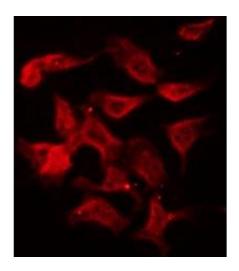
Alternative Name:	GHRHR (GHRHR Products)
Background:	Description: Receptor for GRF, coupled to G proteins which activate adenylyl cyclase.  Stimulates somatotroph cell growth, growth hormone gene transcription and growth hormone secretion.  Gene: GHRHR
Molecular Weight:	47 kDa
Gene ID:	2692
UniProt:	Q02643
Pathways:	Intracellular Steroid Hormone Receptor Signaling Pathway, Hormone Transport, Regulation of Intracellular Steroid Hormone Receptor Signaling, cAMP Metabolic Process

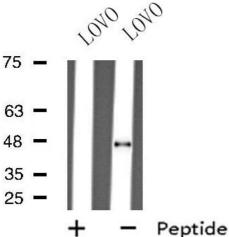
## Application Details

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

### Handling

Hariding	
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





### Immunofluorescence (fixed cells)

**Image 1.** ABIN6275998 staining LOVO 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¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod

#### **Western Blotting**

**Image 2.** Western blot analysis of extracts from LOVO cells, using GHRHR antibody.