

Chicken polyclonal antibody to human GFRa-1

Catalogue #	303-100
Immunogen:	human GFRa-1
Immunogen Description:	Recombinant His-tagged human GFRa-1 protein produced using CHO-based Icosagen Cell factory Ltd. proprietary suspension cell line. For production of hGFRa-1, glycosylphosphatidylinositol GPI-anchor was removed and protein was secreted to the cell culture supernatant. Protein was purified by Ni-affinity chromatography following gel-filtration from cell culture supernatant
Alternative Names:	RET ligand 1, TGF-beta related neurotrophic factor receptor 1
Uniprot ID:	P56159
Clonality:	Polyclonal
Class:	IgY
Reactivity:	Human GFRa-1 Doesn't cross-react with hGFRa-2, hGFRa-3, hGFRa-4
Application:	ELISA, WB, IF
Protocol:	Polyclonal antibody working titer has to be established practically for each particular antigen and assay format
ELISA:	0,02-0,05 µg/ml
IF:	0,5-1 µg/ml
Purification:	GFRa-1 affinity purified
Buffer:	Ammonium sulphate, saturated (PBS pH 7.4)
Shipping:	This product is shipped in non-frozen liquid form in ambient conditions
Storage:	Store at +4 °C upon receipt. As product is (NH ₄) ₂ SO ₄ precipitate, mix well by pipetting or vortexing prior use

Background: Glial cell line-derived neurotrophic factor (GDNF) and neurturin (NTN) are two structurally related, potent neurotrophic factors that play key roles in the control of neuron survival and differentiation. The GFR α -1 is a member of the GDNF receptor family. It is a glycosylphosphatidylinositol (GPI)-linked cell surface receptor for both GDNF and NTN, and mediates activation of the RET tyrosine kinase receptor (www.ncbi.nlm.nih.gov/gene/2674)

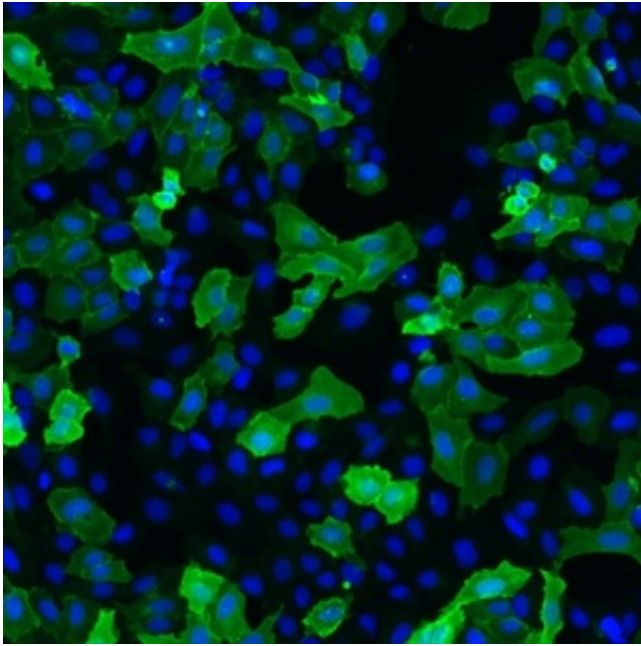


Figure 1. Immunofluorescence detection of human GFR α -1 expressed in U2OS cells. GFR α -1 was visualized using Chicken polyclonal antibody to human GFR α -1 (dilution 1:2000) and Goat anti-chicken IgY – H&L DyLight 550 (Abcam, ab96952) (dilution 1:200) as secondary antibody. For nuclear staining DAPI was used. ArrayScan VTI platform (Thermo Scientific) was used for image acquisition (10x objective). Composite picture was generated using pseudocolors green for GFR α -1 specific signal and blue for nuclei.

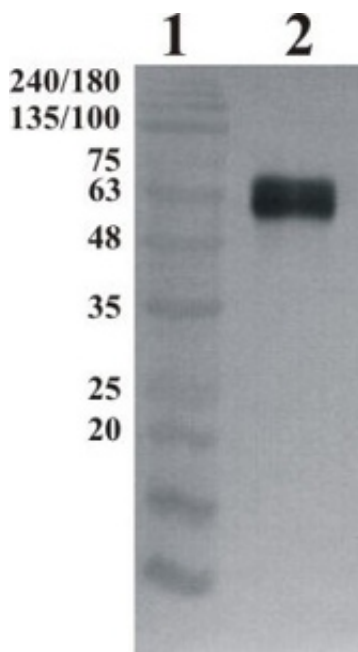


Figure 2. Western-Blot detection of human GFR?1, using Chicken polyclonal antibody to human GFR?1. GFR?1 was C-terminally His tagged. Protein was expressed by CHO cell culture. 100 ng of purified protein was loaded per lane. Line 1. Prestained protein ladder (Naxo 8003). Line 2. hGFR?1. Primary antibody dilution 1:5000 was used. Rabbit anti-chicken antibody (Icosagen) was used as secondary antibody.