

Chicken pAb to human RET

Catalogue # 306-100

Immunogen: human RET

Immunogen Recombinant His-tagged extracellular fragment

Description: of human RET protein produced using CHO-

based Icosagen Cell factory Ltd. proprietary

suspension cell line. The extracellular fragment of hRET was expressed and secreted to the cell culture supernatant. Protein was purified by Niaffinity chromatography following gel-filtration

from cell culture supernatant

Alternative Names: Proto-oncogene tyrosine-protein kinase receptor

Ret, Cadherin family member 12

Proto-oncogene c-Ret

Uniprot ID: P07949

Clonality: Polyclonal

Class: IgY

Reactivity: Human RET

Application: ELISA, WB, IF

Protocol: Polyclonal antibody working titer has to be

established practically for each particular antigen

and assay format

ELISA: 8-12,5 ng/ml

IF: $2-4 \mu g/ml$

Purification: RET 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

(NH4)2SO4 precipitate, mix well by pipetting or

vortexing prior use

Background: The RET proto-oncogene is a receptor tyrosine

kinase for members of the glial cell line-derived neurotrophic factor family of extracellular signalling molecules

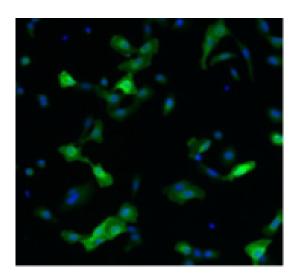


Figure 1. Immunofluorescence detection of human RET expressed in U2OS cells. RET was visualized using Chicken polyclonal antibody to human RET (dilution 1:500) and Goat anti-chicken IgY – H&L DyLight 550 (Abcam, ab96952) (dilution 1:200) as secondary antibody. For nuclear staining DAPI was used. ArrayScan VTI platform (Thermo Scientific) was used for image acquisition (10x objective). Composite picture was generated using pseudocolors green for RET specific signal and blue for nuclei

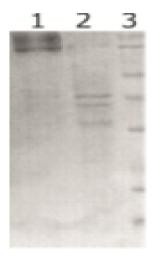


Figure 2. Western-Blot detection of human RET expressed in CHO cells. **Line 1.** hRET-containing CHOEBNALT85 cell culture supernatant. **Line 2.** Negative control – CHOEBNALT85 cell culture supernatant. 5 ?I of supernatant were loaded per line. Primary antibody dilution 1:5000 was used. Rb pAb to Chk IgY ab6753, was used as secondary antibody.