

XL α s (M-14): sc-18993

BACKGROUND

The GTP-binding proteins are well-known regulators of cellular functions, including vesicular transport. XL α s, for extra large α s, is a paternally expressed, plasma membrane-associated protein consisting of a novel XL domain followed by a α s domain. XL α s is specifically associated with the *trans*-Golgi network and occurs selectively in cells containing both the regulated and the constitutive pathway of protein secretion. Like G α_s , XL α s undergoes a conformational change upon binding of GTP γ S. In neuroendocrine cells, the two related G proteins, G α_s and XL α s, exhibit distinct properties with regard to receptor-mediated activation, but converge onto the same effector system, adenylyl cyclase. XL α s is found in adult neuroendocrine tissue, with a particularly high level of expression in the pituitary. The human GNAS gene maps to chromosome 20q13.32.

REFERENCES

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CHROMOSOMAL LOCATION

Genetic locus: GNAS (human) mapping to 20q13.32; *Gnas* (mouse) mapping to 2 H4.

SOURCE

XL α s (M-14) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of XL α s of human origin.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-18993 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

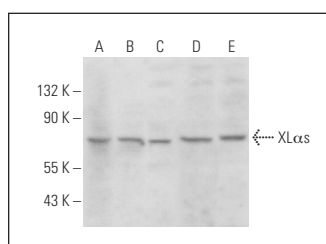
XL α s (M-14) is recommended for detection of XL α s of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); non cross-reactive with G α_s .

Positive Controls: A549 cell lysate: sc-2413, Jurkat whole cell lysate: sc-2204 or HEK293 whole cell lysate: sc-45136.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



XL α s (M-14): sc-18993. Western blot analysis of XL α s expression in A549 (A), Jurkat (B), HEK293 (C), NCI-H460 (D) and MCF7 (E) whole cell lysates.

SELECT PRODUCT CITATIONS

1. Krechowec, S.O., Burton, K.L., Newlaczyl, A.U., Nunn, N., Vlatkovic, N. and Plagge, A. 2012. Postnatal changes in the expression pattern of the imprinted signalling protein XL α s underlie the changing phenotype of deficient mice. *PLoS ONE* 7: e29753.

RESEARCH USE

For research use only, not for use in diagnostic procedures.