

ER α (C-3): sc-514857

BACKGROUND

Estrogen receptors (ER) are members of the steroid/thyroid hormone receptor superfamily of ligand-activated transcription factors. Estrogen receptors, including ER α and ER β , contain DNA binding and ligand binding domains and are critically involved in regulating the normal function of reproductive tissues. They are located in the nucleus, though some estrogen receptors associate with the cell surface membrane and can be rapidly activated by exposure of cells to estrogen. ER α and ER β have been shown to be differentially activated by various ligands. Receptor-ligand interactions trigger a cascade of events, including dissociation from heat shock proteins, receptor dimerization, phosphorylation and the association of the hormone activated receptor with specific regulatory elements in target genes. Evidence suggests that ER α and ER β may be regulated by distinct mechanisms even though they share many functional characteristics.

REFERENCES

1. Mason, B.H., et al. 1983. Progesterone and estrogen receptors as prognostic variables in breast cancer. *Cancer Res.* 43: 2985-2990.
2. Evans, R.M. 1988. The steroid and thyroid hormone receptor superfamily. *Science* 240: 889-895.
3. Danielian, P.S., et al. 1992. Identification of a conserved region required for hormone dependent transcriptional activation by steroid hormone receptors. *EMBO J.* 11: 1025-1033.

CHROMOSOMAL LOCATION

Genetic locus: ESR1 (human) mapping to 6q25.1; Esr1 (mouse) mapping to 10 A1.

SOURCE

ER α (C-3) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 575-599 at the C-terminus of ER α of mouse origin.

PRODUCT

Each vial contains 200 μ g IgG₃ in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-514857 X, 200 μ g/0.1 ml.

ER α (C-3) is available conjugated to agarose (sc-514857 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-514857 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; and to either phycoerythrin (sc-514857 PE), fluorescein (sc-514857 FITC), Alexa Fluor[®] 488 (sc-514857 AF488) or Alexa Fluor[®] 647 (sc-514857 AF647), 200 μ g/ml, for IF, IHC(P) and FCM.

Blocking peptide available for competition studies, sc-514857 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

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STORAGE

Store at 4 $^{\circ}$ C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

ER α (C-3) is recommended for detection of ER α of mouse, rat and human origin by Western Blotting (starting dilution 1:100, 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), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for ER α siRNA (h): sc-29305, ER α siRNA (m): sc-29306, ER α siRNA (r): sc-45949, ER α shRNA Plasmid (h): sc-29305-SH, ER α shRNA Plasmid (m): sc-29306-SH, ER α shRNA Plasmid (r): sc-45949-SH, ER α shRNA (h) Lentiviral Particles: sc-29305-V, ER α shRNA (m) Lentiviral Particles: sc-29306-V and ER α shRNA (r) Lentiviral Particles: sc-45949-V.

ER α (C-3) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of ER α long isoform: 66 kDa.

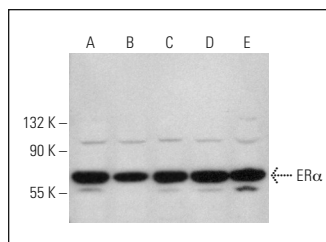
Molecular Weight of ER α short isoform: 54 kDa.

Molecular Weight of ER46: 48 kDa.

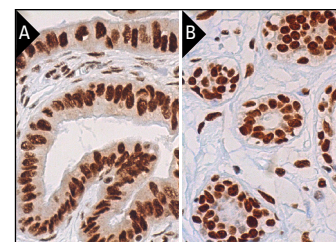
Molecular Weight of ER36: 36 kDa.

Positive Controls: BT-20 cell lysate: sc-2223, MOLT-4 cell lysate: sc-2233 or Raji whole cell lysate: sc-364236.

DATA



ER α (C-3): sc-514857. Western blot analysis of ER α expression in AN3CA (A), BT-20 (B), MOLT-4 (C), Raji (D) and T-47D (E) whole cell lysates.



ER α (C-3): sc-514857. Immunoperoxidase detection of ER α in formalin fixed, paraffin-embedded human fallopian tube tissue, showing nuclear staining of glandular cells (A) and human breast tissue, showing nuclear staining of glandular cells and myoepithelial cells. Detection reagent used: m-IgG κ BP-HRP: sc-516102 (B).

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.