

NF-E2 (G-2): sc-514429

BACKGROUND

The nuclear DNA binding protein NF-E2 regulates expression of globulin genes in developing erythroid cells through interaction with upstream AP-1-like recognition sites. More specifically, NF-E2 recognizes a site containing an intact AP-1 binding motif, preceded by a G residue two base pairs upstream. NF-E2 is an obligate heterodimer composed of NF-E2 p45 and NF-E2 p18. NF-E2 p18, also known as NF-E2U or MAFK, is a ubiquitously expressed component that is related to the v-Maf oncogene. It contains a basic-leucine zipper domain that functions in DNA binding and dimerization. In addition, NF-E2 p18 may play a role in erythroid differentiation. The major component of NF-E2 is a polypeptide, designated NF-E2 p45, that belongs to the basic region-leucine zipper family of transcription factors. This subunit of NF-E2 is specifically expressed at low level in hematopoietic progenitor cells and differentiated cells of the erythroid, megakaryocyte and mast cell lineages.

REFERENCES

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2. Philipsen, S., et al. 1990. The β -globin dominant control region: hyper-sensitive site 2. *EMBO J.* 9: 2159-2167.
3. Ney, P.A., et al. 1990. Tandem AP-1-binding sites within the human β -globin dominant control region function as an inducible enhancer in erythroid cells. *Genes Dev.* 4: 993-1006.
4. Jarman, A.P., et al. 1991. Characterization of the major regulatory element upstream of the human α -globin gene cluster. *Mol. Cell. Biol.* 11: 4679-4689.
5. Andrews, N.C., et al. 1993. Erythroid transcription factor NF-E2 is a haematopoietic-specific basic-leucine zipper protein. *Nature* 362: 722-728.
6. Peters, L.L., et al. 1993. Mouse microcytic anaemia caused by a defect in the gene encoding the globin enhancer-binding protein NF-E2. *Nature* 362: 768-770.
7. Igarashi, K., et al. 1994. Regulation of transcription by dimerization of erythroid factor NF-E2 p45 with small Maf proteins. *Nature* 367: 568-572.
8. Kataoka, K., et al. 1995. Small Maf proteins heterodimerize with Fos and may act as competitive repressors of the NF-E2 transcription factor. *Mol. Cell. Biol.* 15: 2180-2190.

CHROMOSOMAL LOCATION

Genetic locus: NFE2 (human) mapping to 12q13.13.

SOURCE

NF-E2 (G-2) is a mouse monoclonal antibody raised against amino acids 1-230 of NF-E2 of human origin.

PRODUCT

Each vial contains 200 μ g IgG κ kappa light chain 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-514429 X, 200 μ g/0.1 ml.

APPLICATIONS

NF-E2 (G-2) is recommended for detection of NF-E2 of 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) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for NF-E2 siRNA (h): sc-36046, NF-E2 shRNA Plasmid (h): sc-36046-SH and NF-E2 shRNA (h) Lentiviral Particles: sc-36046-V.

NF-E2 (G-2) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

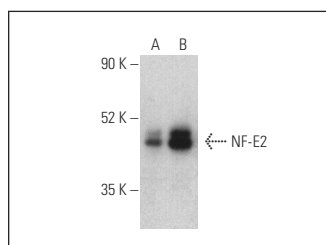
Molecular Weight of NF-E2: 45 kDa.

Positive Controls: K-562 nuclear extract: sc-2130 or MEG-01 nuclear extract: sc-2150.

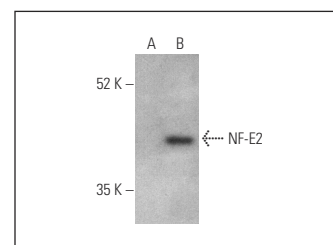
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 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 m-IgG κ BP-FITC: sc-516140 or m-IgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA



NF-E2 (G-2): sc-514429. Western blot analysis of NF-E2 expression in K-562 (A) and MEG-01 (B) nuclear extracts. Detection reagent used: m-IgG κ BP-HRP: sc-525408.



NF-E2 (G-2): sc-514429. Western blot analysis of NF-E2 expression in non-transfected: sc-117752 (A) and mouse NF-E2 transfected: sc-122031 (B) 293T whole cell lysates.

STORAGE

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

RESEARCH USE

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