# C/EBP $\delta$ (C-22): sc-151



The Power to Question

## **BACKGROUND**

The transcription factor C/EBP  $\alpha$  (CCAAT-enhancer binding protein) is a heat-stable, sequence-specific DNA-binding protein first purified from rat liver nuclei that binds avidly to several different  $\emph{cis}$ -regulatory DNA sequences commonly associated with viral and cellular genes transcribed by RNA polymerase II. C/EBP  $\alpha$  regulates gene expression in a variety of tissues including liver, adipose, lung and intestine. C/EBP  $\alpha$  uses a bipartite structural motif to bind DNA. Two protein chains dimerize through a set of amphipathic  $\alpha$  helices termed the leucine zipper. Highly basic polypeptide regions emerge from the zipper to form a linked set of DNA contact surfaces. C/EBP  $\alpha$  appears to function exclusively in terminally differentiated, growth-arrested cells. Additional family members include C/EBP  $\beta$ , C/EBP  $\gamma$ , C/EBP  $\delta$  and C/EBP  $\epsilon$ , all of which exhibit similar DNA-binding specificities and affinities to C/EBP  $\alpha$ . Furthermore, C/EBP  $\beta$  and C/EBP  $\delta$  readily form heterodimers both with each other as well as with C/EBP  $\alpha$ .

## CHROMOSOMAL LOCATION

Genetic locus: CEBPD (human) mapping to 8q11.21; Cebpd (mouse) mapping to 16 A2.

#### SOURCE

C/EBP  $\delta$  (C-22) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the C-terminus of C/EBP  $\delta$  of mouse origin.

# **PRODUCT**

Each vial contains 200  $\mu$ g lgG 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-151 X, 200  $\mu$ g/0.1 ml.

Blocking peptide available for competition studies, sc-151 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## **APPLICATIONS**

C/EBP  $\delta$  (C-22) is recommended for detection of C/EBP  $\delta$  of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu g$  per 100-500  $\mu g$  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 C/EBP  $\delta$  siRNA (h): sc-37722, C/EBP  $\delta$  siRNA (m): sc-37723, C/EBP  $\delta$  shRNA Plasmid (h): sc-37722-SH, C/EBP  $\delta$  shRNA Plasmid (m): sc-37723-SH, C/EBP  $\delta$  shRNA (h) Lentiviral Particles: sc-37722-V and C/EBP  $\delta$  shRNA (m) Lentiviral Particles: sc-37723-V.

C/EBP  $\delta$  (C-22) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

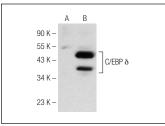
Molecular Weight of C/EBP δ: 28 kDa.

Positive Controls: C/EBP (h): 293T Lysate: sc-176938 or Sol8 nuclear extract: sc-2157.

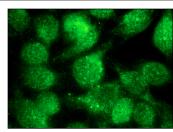
#### **STORAGE**

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

## **DATA**



C/EBP  $\delta$  (C-22): sc-151. Western blot analysis of C/EBP  $\delta$  expression in non-transfected: sc-117752 (**A**) and human C/EBP  $\delta$  transfected: sc-176938 (**B**) 293T whole cell beautic



C/EBP  $\delta$  (C-22): sc-151. Immunofluorescence staining of methanol-fixed HeLa cells showing nuclear and cytoplasmic localization.

# **SELECT PRODUCT CITATIONS**

- 1. Ford, A.M., et al. 1996. Regulation of the myeloperoxidase enhancer binding proteins PU.1, C/EBP  $\alpha$ ,  $\beta$ , and  $\delta$  during granulocyte-lineage specification. Proc. Natl. Acad. Sci. USA 93: 10838-10843.
- 2. Steger, D.J., et al. 2010. Propagation of adipogenic signals through an epigenomic transition state. Genes Dev. 24: 1035-1044.
- 3. Ceccarelli, V., et al. 2011. Eicosapentaenoic acid demethylates a single CpG that mediates expression of tumor suppressor CCAAT/enhancer-binding protein δ in U937 leukemia cells. J. Biol. Chem. 286: 27092-27102.
- 4. Watanabe, M., et al. 2015. The E3 ubiquitin ligase TRIM23 regulates adipocyte differentiation via stabilization of the adipogenic activator PPARy. Elife 4: e05615.
- 5. Kim, Y.M., et al. 2015. The anti-obesity effects of a tuna peptide on 3T3-L1 adipocytes are mediated by the inhibition of the expression of lipogenic and adipogenic genes and by the activation of the Wnt/β-catenin signaling pathway. Int. J. Mol. Med. 36: 327-334.
- 6. Hu, Y.J., et al. 2015. Transcriptional and post-transcriptional control of adipocyte differentiation by Jumonji domain-containing protein 6. Nucleic Acids Res. 43: 7790-7804.

#### **RESEARCH USE**

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



Try **C/EBP**  $\delta$  **(C-6):** sc-365546 or **C/EBP**  $\delta$  **(D-1):** sc-515028, our highly recommended monoclonal alternatives to C/EBP  $\delta$  (C-22). Also, for AC, HRP, FITC, PE, Alexa Fluor<sup>®</sup> 488 and Alexa Fluor<sup>®</sup> 647 conjugates, see **C/EBP**  $\delta$  **(C-6):** sc-365546.