

Anti-C/EBP delta/epsilon Antibody

Catalog # AP53659

Product Information

ApplicationWB, IFPrimary AccessionP49716Other AccessionQ15744

Reactivity Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 28467

Additional Information

Gene ID 1052

Other Names CEBPD; CCAAT/enhancer-binding protein delta; C/EBP delta; Nuclear factor

NF-IL6-beta; NF-IL6-beta; CEBPE; CCAAT/enhancer-binding protein epsilon;

C/EBP epsilon

Target/Specificity Recognizes endogenous levels of C/EBP delta/epsilon protein.

Dilution WB~~1/500 - 1/1000 IF~~1/50 - 1/200

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name CEBPD

Function Transcription activator that recognizes two different DNA motifs: the CCAAT

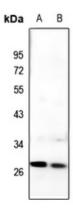
homology common to many promoters and the enhanced core homology common to many enhancers (PubMed:<u>16397300</u>). Important transcription factor regulating the expression of genes involved in immune and

inflammatory responses (PubMed:<u>16397300</u>, PubMed:<u>1741402</u>). Transcriptional activator that enhances IL6 transcription alone and as

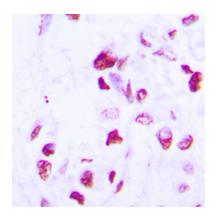
heterodimer with CEBPB (PubMed: 1741402).

Cellular Location Nucleus.

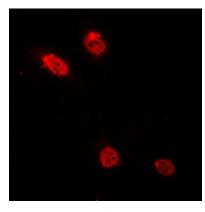
Background



Western blot analysis of C/EBP delta/epsilon expression in CT26 (A), Hela (B) whole cell lysates.



Immunohistochemical analysis of C/EBP delta/epsilon staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of C/EBP delta/epsilon staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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