CEBPA Rabbit pAb

Catalog No.: A28512



Basic Information

Observed MW

42 kDa

Calculated MW

37 kDa/36 kDa/26 kDa/41 kDa

Category

Primary antibody

Applications

WB,IF-P,IHC-P,ELISA

Cross-Reactivity

Human, Mouse, Rat

Background

This intronless gene encodes a transcription factor that contains a basic leucine zipper (bZIP) domain and recognizes the CCAAT motif in the promoters of target genes. The encoded protein functions in homodimers and also heterodimers with CCAAT/enhancer-binding proteins beta and gamma. Activity of this protein can modulate the expression of genes involved in cell cycle regulation as well as in body weight homeostasis. The use of alternative in-frame non-AUG (CUG) and AUG start codons results in several protein isoforms with different lengths. Differential translation initiation is mediated by an out-of-frame, upstream open reading frame which is located between the CUG and the first AUG start codons.

Recommended Dilutions

WB 1:1000 - 1:3000

IF-P 1:100 - 1:200

IHC-P 1:100 - 1:400

ELISA Recommended starting concentration is 1 μg/mL.

Please optimize the concentration based on your specific assay requirements. For highratio antibody dilutions (≥1:10000)a sequential dilution method is strongly recommended to ensure measurement

accuracy.

Immunogen Information

Gene ID Swiss Prot 12606 P53566

Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

Synonyms

Cebp; CBF-A; C/ebpalpha

Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

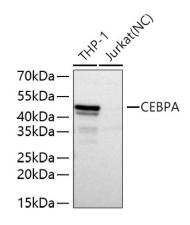
Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide,50% glycerol,pH7.3.

Contact

2	400-999-6126
\bowtie	cn.market@abclonal.com.cr
•	www.abclonal.com.cr



Western blot analysis of various lysates using CEBPA Rabbit pAb (A28512) at 1:3000 dilution incubated overnight at 4° C.

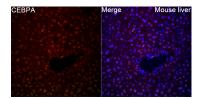
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

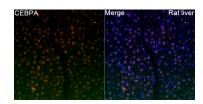
Lysates/proteins: 25 µg per lane.

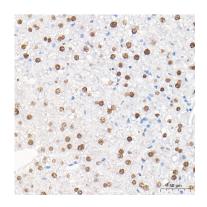
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020). Negative control (NC): Jurkat.

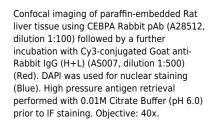
Exposure time: 30 s.



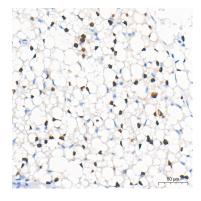




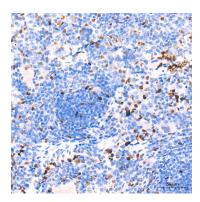
Confocal imaging of paraffin-embedded Mouse liver tissue using CEBPA Rabbit pAb (A28512, dilution 1:100) followed by a further incubation with Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IF staining. Objective: 40x.



Immunohistochemistry analysis of paraffinembedded Mouse liver tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse fat tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

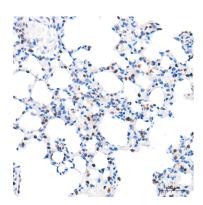


Immunohistochemistry analysis of paraffinembedded Mouse spleen tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

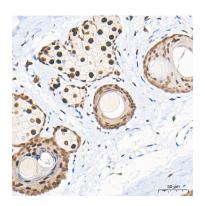


Immunohistochemistry analysis of paraffinembedded Rat liver tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Validation Data



Immunohistochemistry analysis of paraffinembedded Rat lung tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat skin tissue using CEBPA Rabbit pAb (A28512) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.