#### **Anti-NFIL3 Rabbit Polyclonal Antibody**



## **Catalog #: 51459**

#### **Aliases**

E4BP4; IL3BP1; Nuclear factor interleukin-3-regulated protein; E4 promoter-binding protein 4; Interleukin-3 promoter transcriptional activator; Interleukin-3-binding protein 1; Transcriptional activator NF-IL3A

## **Background**

Gene Name: NFIL3 NCBI Gene Entry: 4783 UniProt Entry: Q16649

# **Application Information**

Molecular Weight: Predicted, 51 kDa; observed, 51 kDa

Clonality: Rabbit polyclonal antibody

Species Reactivity: Human, mouse, rat, bovine, chicken, monkey

Applications Tested: Western blotting (WB), immunohistochemistry (IHC), immunocytochemistry

(IC)

## **Immunogen**

A synthesized peptide derived from human NFIL3

## **Isotype**

Rabbit IgG

# **Storage Buffer**

Supplied in PBS (pH 7.3) containing 30% glycerol, and 0.01% sodium azide.

## **Storage**

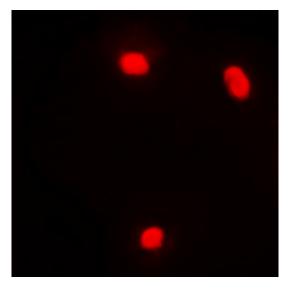
Store at -20 °C for one year.

#### **Recommended Dilutions**

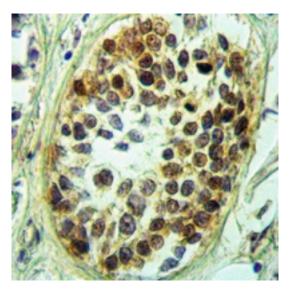
Western Blotting (WB): 1:500-1:1,000 Immunohistochemistry (IHC): 1:50-1:100 Immunocytochemistry (IC): 1:50-1:200

**Note:** This product is for research use only.

#### Validation Data

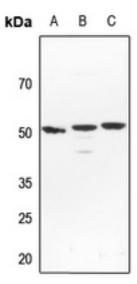


Immunocytochemical analysis of NFIL3 staining in K562 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).



Immunohistochemical analysis of NFIL3 staining in human kidney 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.

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Western blotting analysis of NFIL3 expression in Hela (A), mouse lung (B), rat lung (C) whole cell lysates.