

# **Anti-PU.1 Antibody**

Rabbit polyclonal antibody to PU.1 Catalog # AP60516

## **Specification**

## **Anti-PU.1 Antibody - Product Information**

Application WB, IF/IC, IHC

Primary Accession P17947
Other Accession P17433

Reactivity Human, Mouse, Rat, Zebrafish, Pig,

Chicken Rabbit Polyclonal 31083

Host Clonality Calculated MW

## **Anti-PU.1 Antibody - Additional Information**

### **Gene ID 6688**

## **Other Names**

Transcription factor PU.1; 31 kDa-transforming protein

# **Target/Specificity**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human PU.1. The exact sequence is proprietary.

#### **Dilution**

WB~~WB (1/500 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500) IF/IC~~N/A IHC~~1:100~500

### **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

## **Anti-PU.1 Antibody - Protein Information**

## Name SPI1

#### **Function**

Pioneer transcription factor, which controls hematopoietic cell fate by decompacting stem cell heterochromatin and allowing other transcription factors to enter otherwise inaccessible genomic sites. Once in open chromatin, can directly control gene expression by binding genetic regulatory elements and can also more broadly influence transcription by recruiting transcription factors, such as interferon regulatory factors (IRFs), to otherwise inaccessible genomic regions



(PubMed:<a href="http://www.uniprot.org/citations/23658224" target="\_blank">23658224</a>, PubMed:<a href="http://www.uniprot.org/citations/33951726" target="\_blank">33951726</a>). Transcriptionally activates genes important for myeloid and lymphoid lineages, such as CSF1R (By similarity). Transcriptional activation from certain promoters, possibly containing low affinity binding sites, is achieved cooperatively with other transcription factors. FCER1A transactivation is achieved in cooperation with GATA1 (By similarity). May be particularly important for the pro- to pre-B cell transition (PubMed:<a href="http://www.uniprot.org/citations/33951726" target="\_blank">33951726</a>). Binds (via the ETS domain) onto the purine-rich DNA core sequence 5'-GAGGAA-3', also known as the PU-box (PubMed:<a

href="http://www.uniprot.org/citations/33951726" target="\_blank">33951726</a>). In vitro can bind RNA and interfere with pre-mRNA splicing (By similarity).

## **Cellular Location**

Nucleus {ECO:0000255|PROSITE-ProRule:PRU00237, ECO:0000269|PubMed:33951726}

#### **Tissue Location**

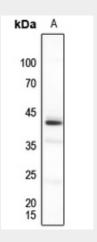
In the bone marrow, concentrated in hematopoietic stem cell, lymphoid progenitor, myeloid lineage (granulocyte macrophage progenitors, classical dendritic cells, monocytes) and B-cell clusters Among B-cells, predominantly expressed in pre-B1 cells (PubMed:33951726). Expressed in germinal center B-cells (PubMed:23166356).

## **Anti-PU.1 Antibody - Protocols**

Provided below are standard protocols that you may find useful for product applications.

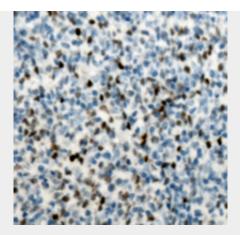
- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

## **Anti-PU.1 Antibody - Images**

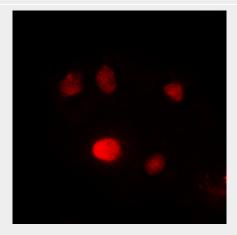


Western blot analysis of PU.1 expression in zebrafish (A) whole cell lysates.





Immunohistochemical analysis of PU.1 staining in human lymph node 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 PU.1 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  $^{\circ}$ 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.

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