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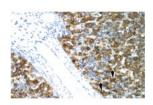
## HIGH PERFORMANCE ANTIBODIES ... AND MORE

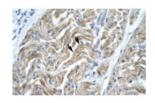
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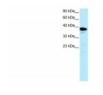
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## **NFYA Antibody**

CATALOG NUMBER: 27-329







Antibody used in IHC on Human Liver cell lysates.

Antibody used in IHC on Human Muscle.

Antibody used in WB on Human Jurkat cells at 0.2-1 ug/ml.

Specifications	
SPECIES REACTIVITY:	Dog, Human, Mouse, Rat
TESTED APPLICATIONS:	ELISA, IHC, WB
APPLICATIONS:	NFYA antibody can be used for detection of NFYA by ELISA at 1:62500. NFYA antibody can be used for detection of NFYA by western blot at 0.2-2.0 ug/mL, and HRP conjugated secondary antibody should be diluted 1:50,000 - 100,000.
USER NOTE:	Optimal dilutions for each application to be determined by the researcher.
POSITIVE CONTROL:	1) Cat. No. 1205 - Jurkat Cell Lysate
PREDICTED MOLECULAR WEIGHT:	37 kDa, 34 kDa
IMMUNOGEN:	Antibody produced in rabbits immunized with a synthetic peptide corresponding a region of human NFYA.
HOST SPECIES:	Rabbit
Dyanastica	
Properties	
PURIFICATION:	Antibody is purified by peptide affinity chromatography method.
PHYSICAL STATE:	Lyophilized
BUFFER:	Antibody is lyophilized in PBS buffer with 2% sucrose. Add 50 uL of distilled water. Final antibody concentration is 1 mg/mL.
CONCENTRATION:	1 mg/ml
STORAGE CONDITIONS:	For short periods of storage (days) store at 4°C. For longer periods of storage, store NFYA antibody at -20°C. As with any antibody avoid repeat freeze-thaw cycles.
CLONALITY:	Polyclonal
CONJUGATE:	Unconjugated
Additional Info	
ALTERNATE NAMES:	NFYA, HAP2, CBF-A, CBF-B, NF-YA
ACCESSION NO.:	NP_002496
PROTEIN GI NO.:	4505389

OFFICIAL SYMBOL:	NFYA
GENE ID:	4800
Background	
BACKGROUND:	NFYA is one subunit of a trimeric complex, forming a highly conserved transcription factor that binds to CCAAT motifs in the promoter regions in a variety of genes. Subunit A associates with a tight dimer composed of the B and C subunits, resulting in a trimer that binds to DNA with high specificity and affinity. The sequence specific interactions of the complex are made by the A subunit, suggesting a role as the regulatory subunit. In addition, there is evidence of post-transcriptional regulation in this gene product, either by protein degradation or control of translation. Further regulation is represented by alternative splicing in the glutamine-rich activation domain, with clear tissue-specific preferences for the two isoforms.
REFERENCES:	1) Zhu, Q.S., et al., (2004) J.Biol.Chem.279(29), 29902-29910.

## FOR RESEARCH USE ONLY

December 12, 2016