



IκB-α (phospho Tyr305) Polyclonal Antibody

Catalog No	YP-Ab-01360
Isotype	IgG
Reactivity	Human;Mouse;Rat;Monkey
Applications	WB;IHC;IF;ELISA
Gene Name	NFKBIA IKBA MAD3 NFKBI
Immunogen	The antiserum was produced against synthesized peptide derived from human IκappaB-α around the phosphorylation site of Tyr305. AA range:268-317
Specificity	Phospho-IκB-α (Y305) Polyclonal Antibody detects endogenous levels of IκB-α protein only when phosphorylated at Y305.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB: 1/500 - 1/2000. IHC: 1/100 - 1/300. ELISA: 1/5000.. IF 1:50-200
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	IκB-α;NFKBIA IKBA MAD3 NFKBI;NF-kappa-B inhibitor alpha;NFKBIA;IKBA;MAD3;NFKBI;l-kappa-B-alpha;IκB-alpha;IκappaBalpha;Major histocompatibility complex enhancer-binding protein MAD3
Observed Band	about 40kd
Calculated Molecular Weight	about 40kd
Cell Pathway	Cytoplasm. Nucleus. Shuttles between the nucleus and the cytoplasm by a nuclear localization signal (NLS) and a CRM1-dependent nuclear export. .
Tissue Specificity	Brain,Kidney,Lymph node,Monocyte,
Function	disease:Defects in NFKBIA are the cause of ectodermal dysplasia anhidrotic with T-cell immunodeficiency autosomal dominant (AEDDAID) [MIM:612132]. Ectodermal dysplasia defines a heterogeneous group of disorders due to abnormal development of two or more ectodermal structures. AEDDAID is an ectodermal dysplasia associated with decreased production of pro-inflammatory cytokines and certain interferons, rendering patients susceptible to infection., function:Inhibits the activity of dimeric NF-kappa-B/REL complexes by trapping REL dimers in the cytoplasm through masking of their nuclear localization signals. On cellular stimulation by immune and proinflammatory responses,



becomes phosphorylated promoting ubiquitination and degradation, enabling the dimeric RELA to translocate to the nucleus and activate transcription., induction: Induced in adherent monocytes., online information: NFKBIA mutation

Background

This gene encodes a member of the NF-kappa-B inhibitor family, which contain multiple ankyrin repeat domains. The encoded protein interacts with REL dimers to inhibit NF-kappa-B/REL complexes which are involved in inflammatory responses. The encoded protein moves between the cytoplasm and the nucleus via a nuclear localization signal and CRM1-mediated nuclear export. Mutations in this gene have been found in ectodermal dysplasia anhidrotic with T-cell immunodeficiency autosomal dominant disease. [provided by RefSeq, Aug 2011]

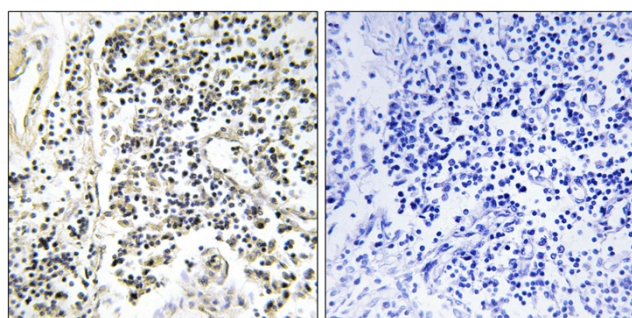
matters needing attention

Avoid repeated freezing and thawing!

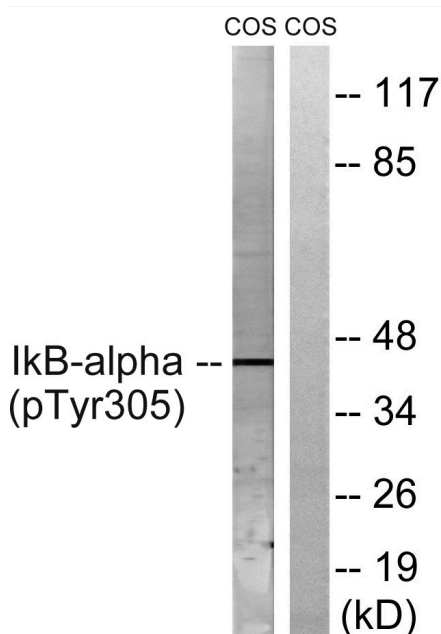
Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Immunohistochemistry analysis of paraffin-embedded human lymph node, using IkappaB-alpha (Phospho-Tyr305) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from COS7 cells treated with nocodazole 1ug/ml 16h, using IkappaB-alpha (Phospho-Tyr305) Antibody. The lane on the right is blocked with the phospho peptide.