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TBK1/NAK Rabbit mAb

Catalog No.: A3458 Recombinant 21 Publications

Basic Information

Observed MW
84kDa

Calculated MW
84kDa

Category
Primary antibody

Applications
WB,IF/ICC,IHC-P,IP,ELISA

Cross-Reactivity
Human, Mouse, Rat

CloneNo number
ARC0778

Background

The NF-kappa-B (NFKB) complex of proteins is inhibited by I-kappa-B (IKB) proteins, which inactivate NFKB by trapping it in the cytoplasm. Phosphorylation of serine residues on the IKB proteins by IKB kinases marks them for destruction via the ubiquitination pathway, thereby allowing activation and nuclear translocation of the NFKB complex. The protein encoded by this gene is similar to IKB kinases and can mediate NFKB activation in response to certain growth factors. The protein is also an important kinase for antiviral innate immunity response.

Recommended Dilutions

WB	1:1000 - 1:4000
IF/ICC	1:100 - 1:800
IHC-P	1:500 - 1:1000
IP	0.5μg-4μg antibody for 200μg-400μg extracts of whole cells
ELISA	Recommended starting concentration is 1 μg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID
29110

Swiss Prot
Q9UHD2

Immunogen
Synthetic peptide. This information is considered to be commercially sensitive.

Synonyms
NAK; T2K; IIAE8; FTDALS4; TBK1/NAK

Product Information

Source
Rabbit

Isotype
IgG

Purification
Affinity purification

Storage
Store at -20℃. Avoid freeze / thaw cycles.
Buffer: PBS with 0.09% Sodium azide,0.05% BSA,50% glycerol,pH7.3.

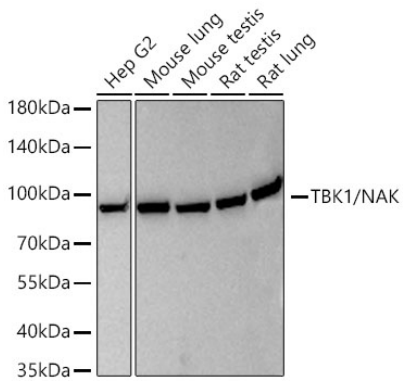
Contact

400-999-6126

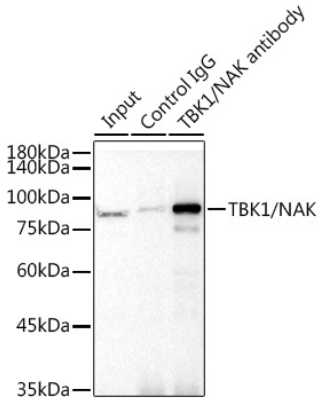
cn.market@abclonal.com.cn

www.abclonal.com.cn

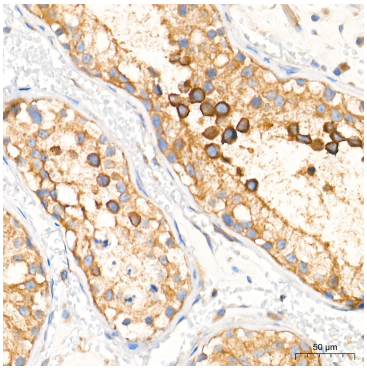
Validation Data



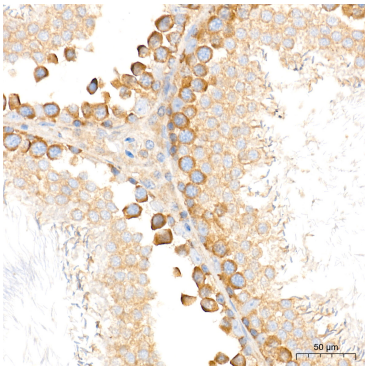
Western blot analysis of various lysates using TBK1/NAK Rabbit mAb (A3458) at 1:4000 dilution incubated at room temperature for 1.5 hours.
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
Lysates/proteins: 25 µg per lane.
Blocking buffer: 3% nonfat dry milk in TBST.
Detection: ECL Basic Kit (RM00020).
Exposure time: 90s.



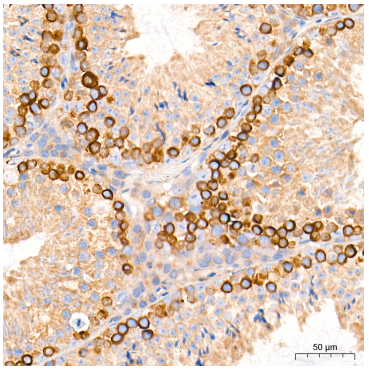
Immunoprecipitation analysis of 300 µg extracts from 293T cells using 3 µg TBK1/NAK Rabbit mAb (A3458). Western blot was performed from the immunoprecipitate using TBK1/NAK Rabbit mAb (A3458) at a dilution of 1:1000.



Immunohistochemistry analysis of paraffin-embedded Human testis tissue using TBK1/NAK Rabbit mAb (A3458) at a dilution of 1:750 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

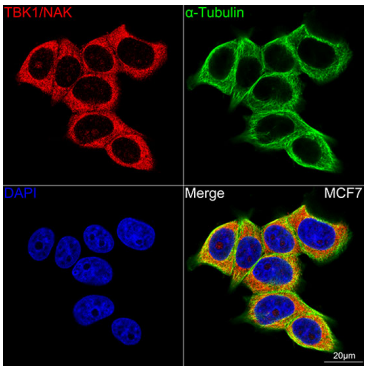


Immunohistochemistry analysis of paraffin-embedded Rat testis tissue using TBK1/NAK Rabbit mAb (A3458) at a dilution of 1:750 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

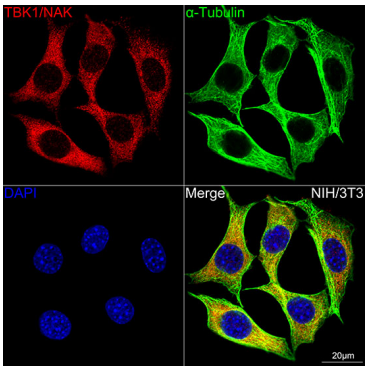


Immunohistochemistry analysis of paraffin-embedded Mouse testis tissue using TBK1/NAK Rabbit mAb (A3458) at a dilution of 1:750 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Validation Data



Confocal imaging of MCF7 cells using TBK1/NAK Rabbit mAb (A3458,at dilution of 1:100) (Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.



Confocal imaging of NIH/3T3 cells using TBK1/NAK Rabbit mAb (A3458,at dilution of 1:100) (Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.