abcam

ab307194

Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free

Recombinant | RabMAb

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	pH: 7.2 - 7.4 Constituents: PBS
Form	Liquid
Clonality	Monoclonal
Immunogen	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	EPR27094-16
Purification technique	Affinity purification Protein A
Concentration	1 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

Flow Cyt (Intra)

Tested

Species	Mouse, Human
Dilution info	-

Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

ICC/IF

Tested

Species	Human
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Mouse
Dilution info	-
Notes	-

IHC-P

Tested

Species	Mouse
Dilution info	-
Notes	This product is not suitable for detection in IHC-P using a fluorescent secondary antibody.

Species	Rat
Dilution info	-
Notes	This product is not suitable for detection in IHC-P using a fluorescent secondary antibody.

Species	Human
Dilution info	-
Notes	This product is not suitable for detection in IHC-P using a fluorescent secondary antibody.

WB

Tested

Species	Rat, Mouse, Human
Dilution info	-
Notes	-

Target data

See full target information TBX21

Function

Lineage-defining transcription factor which initiates Th1 lineage development from naive Th precursor cells both by activating Th1 genetic programs and by repressing the opposing Th2 and Th17 genetic programs (PubMed:10761931). Activates transcription of a set of genes important for Th1 cell function, including those encoding IFN-gamma and the chemokine receptor CXCR3. Induces permissive chromatin accessibilty and CpG methylation in IFNG (PubMed:33296702). Activates IFNG and CXCR3 genes in part by recruiting chromatin remodeling complexes including KDM6B, a SMARCA4-containing SWI/SNF-complex, and an H3K4me2-methyltransferase complex to their promoters and all of these complexes serve to establish a more permissive chromatin state conducive with transcriptional activation (By similarity). Can activate Th1 genes also via recruitment of Mediator complex and P-TEFb (composed of CDK9 and CCNT1/cyclin-T1) in the form of the super elongation complex (SEC) to super-enhancers and associated genes in activated Th1 cells (PubMed:27292648). Inhibits the Th17 cell lineage commitment by blocking RUNX1-mediated transactivation of Th17 cell-specific transcriptinal regulator RORC. Inhibits the Th2 cell lineage commitment by suppressing the production of Th2 cytokines, such as IL-4, IL-5, and IL- 13, via repression of transcriptional regulators GATA3 and NFATC2. Protects Th1 cells from amplifying aberrant type-I IFN response in an IFN-gamma abundant microenvironment by acting as a repressor of type-I IFN transcription factors and type-I IFN-stimulated genes. Acts as a regulator of antiviral B-cell responses; controls chronic viral infection by promoting the antiviral

antibody IgG2a isotype switching and via regulation of a broad antiviral gene expression program (By similarity). Required for the correct development of natural killer (NK) and mucosal-associated invariant T (MAIT) cells (PubMed:33296702).

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C
Storage information	Do Not Freeze

Notes

Patented technology
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

What are the advantages of a recombinant monoclonal antibody?
This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free batch production

For more information, read more on recombinant antibodies.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with 1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

Compatibility
This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® is a trademark of Fluidigm Canada Inc.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	T-bet also known as Tbx21 is a T-box transcription factor that plays an important role in regulating immune responses. T-bet has a molecular mass of approximately 58 kDa. It is extensively expressed in T-helper 1 (Th1) cells and natural killer (NK) cells facilitating these cells' development and function. T-bet guides the transcriptional activities necessary for cellular differentiation and immune response by binding to specific DNA sequences. Its expression is tightly controlled reflecting its central role in immune regulation. "9d bet" is also referenced in scientific research although its specific context requires precise understanding within a given study.
Biological function summary	This transcription factor is pivotal for manufacturing interferon-gamma (IFN-γ) initiating and sustaining Th1 cell immunity. T-bet doesn't act alone; it functions as part of larger protein complexes interacting with other transcription factors and cofactors to exert its effects. The capacity to influence IFN-γ production makes T-bet a significant player in immune responses helping coordinate responses against intracellular pathogens. Moreover the expression of T-bet is not limited solely to T cells but also impacts other immune cells like CD8+ T cells and B cells. Anti-bet or anti-T antibodies frequently target T-bet in research to explore immune system intricacies better.
Pathways	T-bet integrates into the immune signaling network through pathways such as the JAK-STAT pathway and the Th1 differentiation pathway. It activates transcription of the IFNG gene working closely with related proteins like STAT4. T-bet's influence extends as it cooperates with STAT1 enabling a feed-forward loop that amplifies and stabilizes Th1 responses. "Bet t products" could refer to those derived from such transcriptional activation. Additionally T-bet involvement goes beyond transcriptional regulation affecting cytokine transport and secretion processes vital for an effective immune response.
Associated diseases and disorders	Researchers have linked T-bet to autoimmune diseases and infectious diseases. Aberrant expression of T-bet can lead to enhanced Th1 responses intensifying autoimmune conditions such as multiple sclerosis (MS). Its role in diseases connects it with NF-κB pathways and other transcription factors influencing inflammatory processes. Furthermore "anti bet" studies often involve understanding T-bet's role in infectious diseases seeking to manipulate its function to adjust immune responses in chronic infections. Studies targeting its pathways or interactions with disease-specific proteins aim to develop therapeutic strategies turning an element of dysregulation into a novel treatment approach.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

We do not recommend this combination. It is not covered by our product promise.

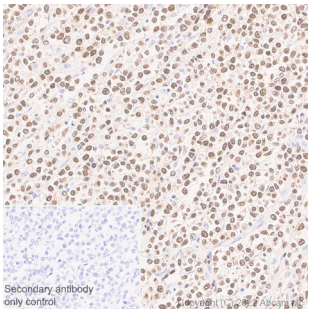
We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

Full details and terms and conditions can be found here:
Terms & Conditions.

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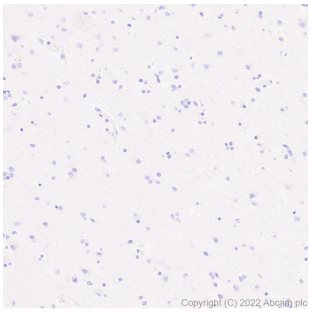
15 product images



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

This data was developed using ab307193, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human non-Hodgkin's tissue labeling T-bet / Tbx21 with ab307193 at 1/500 (1.044 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection) was used. Nuclear staining on human non-Hodgkin's lymphoma (T cell). The section was incubated with ab307193 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution 2) for 20 mins

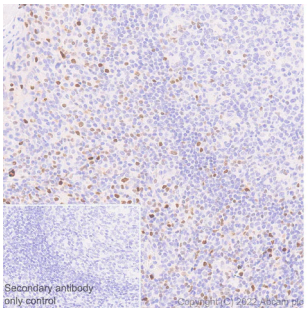
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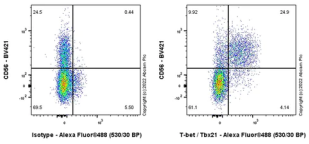
This data was developed using ab307193, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human cerebrum tissue labeling T-bet / Tbx21 with ab307193 at 1/500 (1.044 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection) was used. Negative control: No staining on human cerebrum. The section was incubated with ab307193 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution 2) for 20 mins

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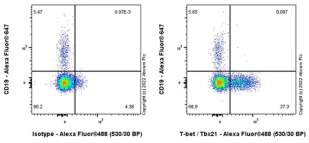
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This data was developed using ab307193, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human tonsil tissue labeling T-bet / Tbx21 with ab307193 at 1/500 (1.044 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection) was used. Nuclear staining on human tonsil. The section was incubated with ab307193 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution 2) for 20 mins



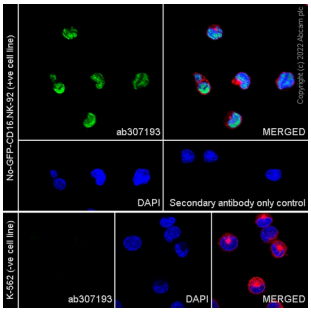
Flow Cytometry (Intracellular) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

This data was developed using ab307193, the same antibody clone in a different buffer formulation. Flow cytometric analysis of 2% paraformaldehyde fixed 0.1% Tween-20 permeabilized Human peripheral blood mononuclear cell (PBMC) cells labelling T-bet / Tbx21 with ab307193 at 1/500 dilution (0.1µg)/ Right (Red) compared with a Rabbit monoclonal IgG (ab172730) / Left isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Live cells were stained with anti-CD56 conjugated to BV421. Fixed with 2% PFA for 10 min followed by intracellularly staining with rabbit IgG or ab307193.



Flow Cytometry (Intracellular) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

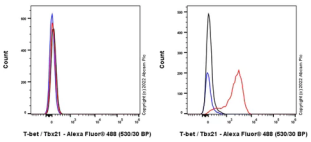
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Immunocytochemistry/ Immunofluorescence - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

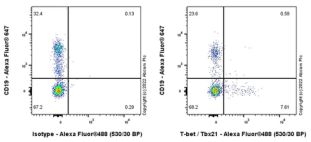
This data was developed using ab307193, the same antibody clone in a different buffer formulation. Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized No-GFP-CD16.NK-92 (CD16 ectopically expressed natural killer (NK-92®) cell with no GFP tag) cells labeling T-bet / Tbx21 with ab307193 at 1/500 (1.044 µg/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 2µg/ml dilution (Green). Confocal image showing nuclear staining in No-GFP-CD16.NK-92 cell line. Negative control: K562. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 2.5µg/ml dilution (Red). The Nuclear

counterstain was DAPI (Blue). Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 2µg/ml dilution.



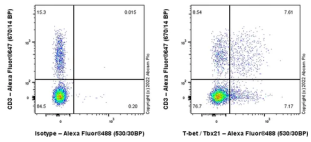
Flow Cytometry (Intracellular) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

This data was developed using ab307193, the same antibody clone in a different buffer formulation. Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized K562 (human chronic myelogenous leukemia lymphoblast, Left) / No-GFP-CD16.NK-92 (CD16 ectopically expressed natural killer(NK-92®) cell with no GFP tag, Right) cells labelling T-bet / Tbx21 with ab307193 at 1/500 dilution (0.1µg) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Negative control: K562.



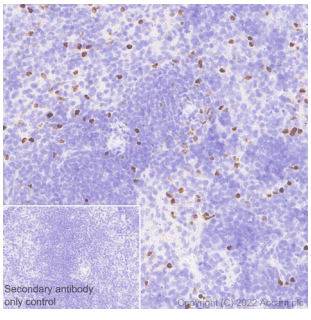
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Flow Cytometry (Intracellular) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

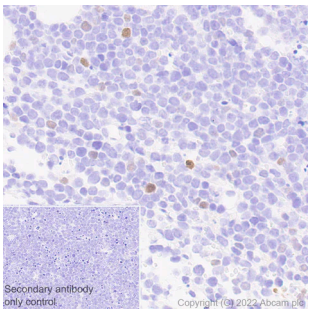
This data was developed using ab307193, the same antibody clone in a different buffer formulation. Flow cytometric analysis of 2% paraformaldehyde fixed 0.1% Tween-20 permeabilized Mouse splenocytes cells labeling T-bet / Tbx21 with ab307193 at 1/500 dilution (0.1µg)/ Right (Red) compared with a Rabbit monoclonal IgG (ab172730) / Left isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Live cells were stained with anti-CD3 conjugated to Alexa Fluor® 647. Fixed with 2% PFA for 10 min followed by intracellularly staining with rabbit IgG or ab307193.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

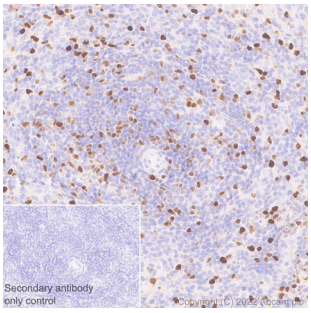
This data was developed using ab307193, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Mouse spleen tissue labeling T-bet / Tbx21 with ab307193 at 1/500 (1.044 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection) was used. Nuclear staining on mouse spleen. The section was incubated with ab307193 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution 2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

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Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

This data was developed using 307193, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST Negative control: K-562 In lane 1, the lysate was freshly made and used for Western Blotting immediately to minimize protein degradation. The identity of the bands below 50 kDa are unknown. Exposure time: 15 seconds
All lanes:
Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] (ab307193) at 1/1000 dilution

Lane 1:
No-GFP-CD16.NK-92 (human malignant non-Hodgkins lymphoma natural killer cell) whole cell lysate 20 µg

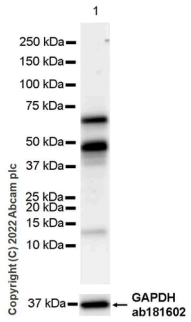
Lane 2:
K-562 (human chronic myelogenous leukemia lymphoblast) whole cell lysate 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 60 kDa

Exposure time: 15s



Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

This data was developed using 307193, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: The lysate was freshly made and used for Western Blotting immediately to minimize protein degradation. The identity of the bands below 50 kDa are unknown. 103 seconds Exposure time:
All lanes:
Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] (ab307193) at 1/1000 dilution

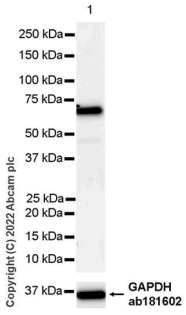
All lanes:
Rat spleen tissue lysate 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 60 kDa

Exposure time: 103s



Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] - BSA and Azide free (ab307194)

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All lanes:
Western blot - Anti-T-bet / Tbx21 antibody [EPR27094-16] (ab307193) at 1/1000 dilution

All lanes:
Mouse spleen tissue lysate 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 60 kDa

Exposure time: 180s

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.