

# Anti-TACC3 antibody [EPR7756] - BSA and Azide free ab224525

KO VALIDATED Recombinant RabMAb<sup>®</sup>

7 Images

### Overview

|                            |  |
|----------------------------|--|
| <b>Product name</b>        | Anti-TACC3 antibody [EPR7756] - BSA and Azide free   |
| <b>Description</b>         | Rabbit monoclonal [EPR7756] to TACC3 - BSA and Azide free  |
| <b>Host species</b>        | Rabbit   |
| <b>Tested applications</b> | <b>Suitable for:</b> IHC-P, ICC/IF, WB   |
| <b>Species reactivity</b>  | <b>Reacts with:</b> Mouse, Rat, Human  |
| <b>Immunogen</b>           | Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.  |
| <b>Positive control</b>    | WB: HEK293T and HCT116 cell lysates. ICC/IF: HeLa cells. IHC-P: Human tonsil and testis tissues.   |
| <b>General notes</b>       | <p>ab224525 is the carrier-free version of <a href="#">ab134154</a>.</p> <p>Our <b>carrier-free</b> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.</p> <p>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.</p> <p>Use our <b>conjugation kits</b> for antibody conjugates that are ready-to-use in as little as 20 minutes with &lt;1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"><li>- High batch-to-batch consistency and reproducibility</li><li>- Improved sensitivity and specificity</li><li>- Long-term security of supply</li><li>- Animal-free production</li></ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p> |

## Properties

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|                             |   |
|-----------------------------|---|
| <b>Form</b>                 | Liquid  |
| <b>Storage instructions</b> | Shipped at 4°C. Store at +4°C. Do Not Freeze. |
| <b>Storage buffer</b>       | pH: 7.2<br>Constituent: PBS                   |
| <b>Carrier free</b>         | Yes   |
| <b>Purity</b>               | Protein A purified                            |
| <b>Clonality</b>            | Monoclonal                                    |
| <b>Clone number</b>         | EPR7756                                       |
| <b>Isotype</b>              | IgG   |

## Applications

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**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab224525 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application   | Abreviews | Notes  |
|---------------|-----------|--|
| <b>IHC-P</b>  |           | Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.<br>See <b><u>IHC antigen retrieval protocols</u></b> . |
| <b>ICC/IF</b> |           | Use at an assay dependent concentration.   |
| <b>WB</b>     |           | Use at an assay dependent concentration. Predicted molecular weight: 90 kDa.   |

## Target

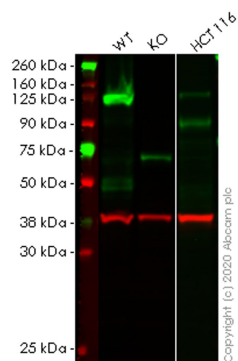
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|                              |  |
|------------------------------|--|
| <b>Function</b>              | Plays a role in the microtubule-dependent coupling of the nucleus and the centrosome. Involved in the processes that regulate centrosome-mediated interkinetic nuclear migration (INM) of neural progenitors (By similarity). May be involved in the control of cell growth and differentiation. May contribute to cancer. |
| <b>Sequence similarities</b> | Belongs to the TACC family.  |
| <b>Cellular localization</b> | Cytoplasm.   |

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## Images

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Western blot - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

**All lanes** : Anti-TACC3 antibody [EPR7756] ([ab134154](#)) at 1/1000 dilution

**Lane 1** : Wild-type HEK293T cell lysate

**Lane 2** : TACC3 knockout HEK293T cell lysate

**Lane 3** : HCT116 cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

**All lanes** : Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) at 1/10000 dilution

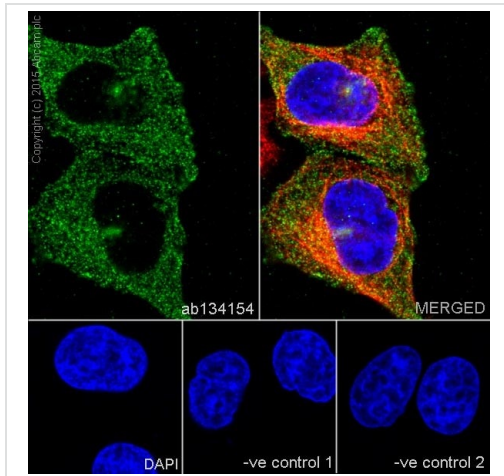
**Predicted band size:** 90 kDa

**Observed band size:** 125 kDa

This data was developed using the same antibody clone in a different buffer formulation ([ab134154](#)).

**Lanes 1-3:** Merged signal (red and green). Green - [ab134154](#) observed at 125 kDa. Red - loading control [ab8245](#) observed at 36 kDa.

[ab134154](#) Anti-TACC3 antibody [EPR7756] was shown to specifically react with TACC3 in wild-type HEK293T cells. Loss of signal was observed when knockout cell line [ab266316](#) (knockout cell lysate [ab257723](#)) was used. Wild-type and TACC3 knockout samples were subjected to SDS-PAGE. [ab134154](#) and Anti-GAPDH antibody [6C5] - Loading Control ([ab8245](#)) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed ([ab216776](#)) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

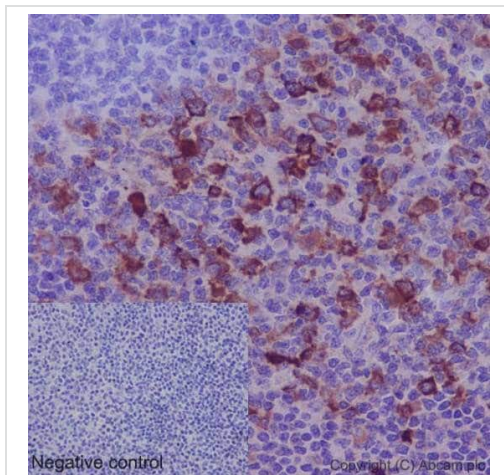


Immunocytochemistry/ Immunofluorescence - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling TACC3 with purified **ab134154** at 1/200. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. An Alexa Fluor® 555-conjugated goat anti-rabbit IgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

Control: primary antibody (1/200) and secondary antibody, **ab150113**, an Alexa Fluor® 488-conjugated goat anti-mouse IgG (1/500).

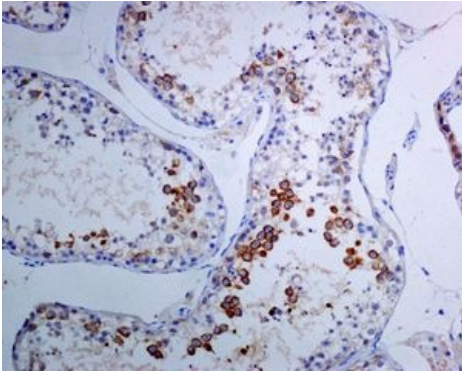
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab134154**).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling TACC3 with purified **ab134154** at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody. Negative control using PBS instead of primary antibody. Counterstained with Hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab134154**).

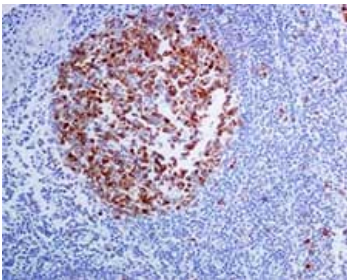


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human testis tissue labelling TACC3 with unpurified **ab134154** at 1/50 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab134154**).

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

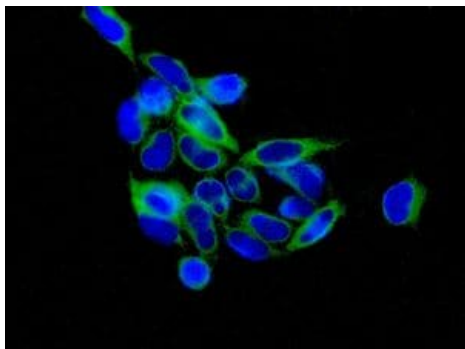


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling TACC3 with unpurified **ab134154** at 1/50 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab134154**).

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling TACC3 with unpurified **ab134154** at 1/50 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab134154**).

Immunocytochemistry/ Immunofluorescence - Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

### Why choose a recombinant antibody?



**Research with confidence**  
Consistent and reproducible results



**Long-term and scalable supply**  
Recombinant technology



**Success from the first experiment**  
Confirmed specificity



**Ethical standards compliant**  
Animal-free production

Anti-TACC3 antibody [EPR7756] - BSA and Azide free (ab224525)

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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