# abcam

### **Product datasheet**

## Anti-TTC5 antibody ab36855

#### <u>1 References</u> 4 图像

概述		
产品名称	Anti-TTC5抗体	
描述	兔多克隆抗体to TTC5	
宿主	Rabbit	
经测试应 <b>用</b>	适用于: WB, IHC-P, ICC/IF	
种属反应性	<b>与反</b> 应: Mouse, Human	
免疫原	A 14 amino acid peptide from near the carboxy terminus of human TTC5.	
<b>阳性</b> 对照	RAW264.7 cell lysate.	
<b>常</b> 规说 <b>明</b>	The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.	
	If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As	

# FxdLiquid**fxbüng**Shipped at 4°C. Store at +4°C.**fki?a**PH: 7.2<br/>Preservative: 0.02% Sodium azide<br/>Constituent: PBS**fxb**Imunogen affinity purified**fxb5bjb**<

#### 应用

性能

The Abpromise guarantee Abpromise™承

Abpromise™承诺保证使用ab36855于以下的经测试应用

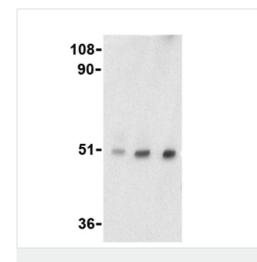
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use a concentration of 0.5 - 2 µg/ml. Detects a band of approximately 49 kDa (predicted molecular weight: 49 kDa).
IHC-P		Use a concentration of 10 µg/ml.
ICC/IF		Use at an assay dependent concentration.

<b>靶</b> 标	
相关性	Tetratricopeptide repeat protein 5 (TTC5) is a member of a diverse group of functionally distinct proteins that are characterized by containing one or more tetratricopeptide repeats. Each motif consists of two anti-parallel a-helices such that tandem arrays of TPR motifs generate a right-handed helical structure with an amphipathic channel that may serve to accommodate the complementary region of a target protein. While the exact function of TTC5 remains unclear, it is thought that the TPR motifs serve to mediate protein-protein interactions such as those seen with protein chaperones HSP70 and HSP90 and some proteins involved in cell stress response signaling pathways such as protein phosphatase 5, suggesting that TTC5 may also function via protein-protein interactions mediated by its TPR motifs.
细 <b>胞定位</b>	Nucleus. Cytoplasm. Note: Phosphorylation at Ser-203 results in nuclear localization, while unphosphorylated protein localizes to the cytoplasm

#### 图片

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Western blot - Anti-TTC5 antibody (ab36855)

Lane 1 : Anti-TTC5 antibody (ab36855) at 0.5 μg/ml Lane 2 : Anti-TTC5 antibody (ab36855) at 1 μg/ml Lane 3 : Anti-TTC5 antibody (ab36855) at 2 μg/ml

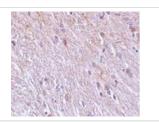
Lane 1 : RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) whole cell lysate Lanes 2-3 : RAW264.7 cell lysate

Lysates/proteins at 15 µg/ml per lane.

#### Secondary

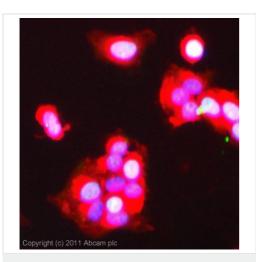
All lanes : anti-rabbit IgG HRP at 1/10000 dilution

Predicted band size: 49 kDa Observed band size: 49 kDa



ab36855 at 10 $\mu$ g/ml staining TCC5 in mouse brain tissue by IHC

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TTC5 antibody (ab36855)



Immunocytochemistry/ Immunofluorescence - Anti-TTC5 antibody (ab36855) ICC/IF image of ab36855 stained MCF7 cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab36855, 5µg/ml) overnight at +4°C. The secondary antibody (green) was **ab96899** Dylight 488 goat anti-rabbit IgG (H+L) used at a 1/250 dilution for 1h. Alexa Fluor®594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

Immunocytochemistry/ Immunofluorescence - Anti-TTC5 antibody (ab36855) Immunofluorescence of TTC5 in Mouse Brain cells using ab36855 at 20 ug/ml.

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

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