abcam

Product datasheet

Anti-Tuberin antibody [Y320] - BSA and Azide free ab178441

敲除 验证 重组 RabMAb

6 References 7 图**像**

概述	
产 品名称	Anti-Tuberin 抗体 [Y320] - BSA and Azide free
描述	兔 单 克隆抗体 [Y320] to Tuberin - BSA and Azide free
宿主	Rabbit
特异性	This antibody recognises Tuberous sclerosis complex-2 (TCS2 also known as Tuberin. It is predicted to detect isoform 2-4 based on sequence homology.
经 测 试应 用	适用于: Flow Cyt (Intra), IHC-P, WB
种属反应性	与反应: Mouse, Rat, Human
免疫原	Synthetic peptide within Human Tuberin aa 1750-1850 (C terminal). The exact sequence is proprietary.
阳性 对照	WB: HAP1 and HeLa whole cell lysates. Rat and mouse brain tissue lysates; Flow Cyt (intra): Jurkat cells; IHC: Rat and mouse liver tissue. Human lung carcinoma tissue.
常 规说 明	ab178441 is the carrier-free version of ab32554 .
	Our <u>carrier-free</u> 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.
	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 <u>conjugation kits</u> 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.
	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.
	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 production For more information <u>see here</u> . Our RabMAb [®] technology is a patented hybridoma-based technology for making rabbit

性能	
形式	Liquid
存 放 说明	Shipped at 4°C. Store at +4°C. Do Not Freeze.
存储溶液	Constituent: PBS
无载体	是
纯 度	Protein A purified
克隆	单 克隆
克 隆 编号	Y320
同种型	lgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab178441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
Flow Cyt (Intra)		Use at an assay dependent concentration. <u>ab199376</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Detects a band of approximately 220 kDa (predicted molecular weight: 200 kDa).

靶标

功能	In complex with TSC1, inhibits the nutrient-mediated or growth factor-stimulated phosphorylation of S6K1 and EIF4EBP1 by negatively regulating mTORC1 signaling. Acts as a GTPase-activating protein (GAP) for the small GTPase RHEB, a direct activator of the protein kinase activity of mTORC1. Implicated as a tumor suppressor. Involved in microtubule-mediated protein transport, but this seems to be due to unregulated mTOR signaling. Stimulates weakly the intrinsic GTPase activity of the Ras-related proteins RAP1A and RAB5 in vitro. Mutations in TSC2 lead to constitutive activation of RAP1A in tumors.
组织 特异性	Liver, brain, heart, lymphocytes, fibroblasts, biliary epithelium, pancreas, skeletal muscle, kidney, lung and placenta.
疾病相关	Defects in TSC2 are the cause of tuberous sclerosis type 2 (TSC2) [MIM:613254]. TSC2 is an autosomal dominant multi-system disorder that affects especially the brain, kidneys, heart, and skin. It is characterized by hamartomas (benign overgrowths predominantly of a cell or tissue type

that occurs normally in the organ) and hamartias (developmental abnormalities of tissue combination). Clinical symptoms can range from benign hypopigmented macules of the skin to profound mental retardation with intractable seizures to premature death from a variety of disease-associated causes.
 Defects in TSC2 are a cause of lymphangioleiomyomatosis (LAM) [MIM:606690]. LAM is a progressive and often fatal lung disease characterized by a diffuse proliferation of abnormal smooth muscle cells in the lungs. It affects almost exclusively young women and can occur as an isolated disorder or in association with tuberous sclerosis complex.
 序列相似性 Contains 1 Rap-GAP domain.
 翻译后修饰 Phosphorylation at Ser-1387, Ser-1418 or Ser-1420 does not affect interaction with TSC1. Phosphorylation at Ser-939 and Thr-1462 by PKB/AKT1 is induced by growth factor stimulation.

图片

	1	2	3	4	5	6	
			=				
250 kDa 🗕	-	-	-	-	100	and a	 → Tuberin
150 kDa 🗕	-	_	-	_	_	_	
100 kDa 🗕	-	-	-				
75 kDa 🗕		-		-			
50 kDa 🗕				-			
37 kDa 🗕							Copyright (c) 2020 Abeam plo
25 kDa 🗕							0 Ab
20 kDa 🗕							202
15 kDa 🗕							ht (c)
10 kDa 🗕							yrig
							Cop

Western blot - Anti-Tuberin antibody [Y320] - BSA and Azide free (ab178441) All lanes : Anti-Tuberin antibody [Y320] (<u>ab32554</u>) at 1/5000 dilution (purified)

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate boiled

Lane 2 : HeLa (Human cervix adenocarcinoma epithelial cell)

whole cell lysate unboiled

Lane 3 : Mouse brain lysate boiled

Lane 4 : Mouse brain lysate unboiled

Lane 5 : Rat brain lysate boiled

Lane 6 : Rat brain lysate unboiled

Lysates/proteins at 20 µg per lane.

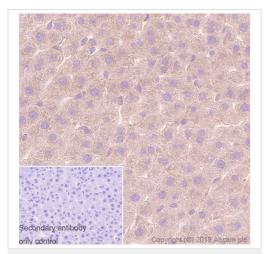
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 200 kDa Observed band size: 200 kDa

The unboiled lysate are recommended to be used in SDS-PAGE, in order to prevent membrane protein aggregation.

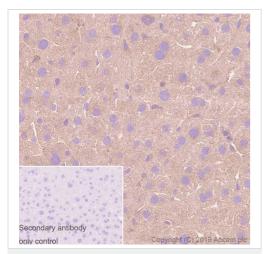
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab32554</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Tuberin antibody [Y320] -BSA and Azide free (ab178441)

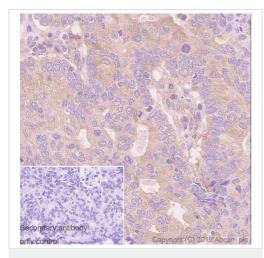
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat liver tissue sections labeling Tuberin with purified <u>ab32554</u> at 1/100 dilution (2.97 µg/mL). Heat mediated antigen retrieval using Bond[™] Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab32554**).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Tuberin antibody [Y320] -BSA and Azide free (ab178441) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse liver tissue sections labeling Tuberin with purified <u>ab32554</u> at 1/100 dilution (2.97 µg/mL). Heat mediated antigen retrieval using Bond[™] Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

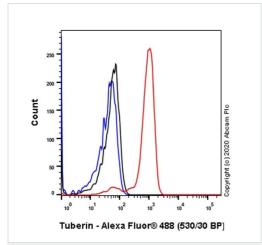
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab32554</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Tuberin antibody [Y320] -BSA and Azide free (ab178441)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human lung carcinoma tissue sections labeling Tuberin with purified <u>ab32554</u> at 1/100 dilution (2.97 µg/mL). Heat mediated antigen retrieval using Bond[™] Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

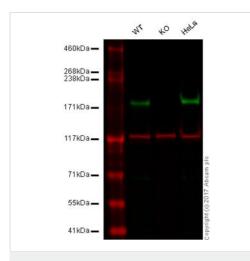
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab32554**).



Flow Cytometry (Intracellular) - Anti-Tuberin antibody [Y320] - BSA and Azide free (ab178441)

Intracellular Flow Cytometry analysis of Jurkat (Human T cell leukemia T lymphocyte) cells labeling Tuberin with purified **ab32554** at 1/30 dilution (10 μg/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit IgG (Alexa Fluor[®] 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab32554</u>).



Western blot - Anti-Tuberin antibody [Y320] - BSA and Azide free (ab178441)

This WB data was generated using the same anti-Tuberin antibody clone, Y320, in a different buffer formulation (cat# <u>ab32554</u> - unpurified).

Lane 1: Wild-type HAP1 whole cell lysate (20 µg) Lane 2: Tuberin knockout HAP1 whole cell lysate (20 µg) Lane 3: HeLa whole cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab32554</u> observed at 180 kDa. Red - loading control, <u>ab18058</u>, observed at 130 kDa.

<u>ab32554</u> was shown to specifically react with tuberin in wild-type HAP1 cells. No band was observed when tuberin knockout samples were used. Wild-type and tuberin knockout samples were subjected to SDS-PAGE. Ab32554 and <u>ab18058</u> (Mouse anti Vinculin loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/10000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye[®] 800CW) preabsorbed <u>ab216773</u> and Goat anti-Mouse IgG H&L (IRDye[®] 680RD) preabsorbed <u>ab216776</u> secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



(ab178441)

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