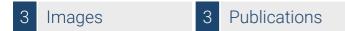


## Datasheet for ABIN3020064

# anti-MAPK14 antibody (pThr180, pTyr182)





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Quantity:	100 μL	
Target:	MAPK14	
Binding Specificity:	pThr180, pTyr182	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This MAPK14 antibody is un-conjugated	
Application:	Western Blotting (WB)	
Product Details		
Immunogen:	Synthetic peptide	
Cross-Reactivity:	Mouse (Murine), Rat (Rattus)	
Target Details		
Target:	MAPK14	
Alternative Name:	MAPK14 (MAPK14 Products)	
Molecular Weight:	360 kDa	
Gene ID:	1432	
UniProt:	Q16539	
Pathways:	MAPK Signaling, Neurotrophin Signaling Pathway, Activation of Innate immune Response,	

Cellular Response to Molecule of Bacterial Origin, Regulation of Muscle Cell Differentiation,
Regulation of Cell Size, Hepatitis C, Toll-Like Receptors Cascades, Autophagy, Thromboxane A2
Receptor Signaling, BCR Signaling, S100 Proteins

## **Application Details**

Restrictions:

For Research Use only

#### **Publications**

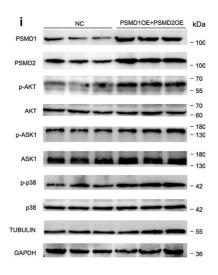
Product cited in:

Kumar, Wang, Liu, Ding, Dong, Zheng, Ye, Liu: "Hypoxia-Induced Mitogenic Factor Promotes Cardiac Hypertrophy via Calcium-Dependent and Hypoxia-Inducible Factor-1α Mechanisms." in: **Hypertension (Dallas, Tex.: 1979)**, Vol. 72, Issue 2, pp. 331-342, (2018) (PubMed).

Dong, Liu, Meng, Liu, Bi, Wu, Jin, Yao, Tang, Wang, Li, Zhang, Yu, Zhan, Chen, Ge, Yang, Li: "Keratin 8 limits TLR-triggered inflammatory responses through inhibiting TRAF6 polyubiquitination." in: **Scientific reports**, Vol. 6, pp. 32710, (2018) (PubMed).

Xiao, Deng, Lv, Kang, Ma, Yan, Song, Gao, Zhang, Xu: "Hydrogen Peroxide Induce Human Cytomegalovirus Replication through the Activation of p38-MAPK Signaling Pathway." in: **Viruses**, Vol. 7, Issue 6, pp. 2816-33, (2016) (PubMed).

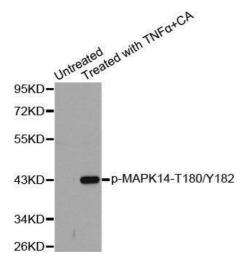
## Images

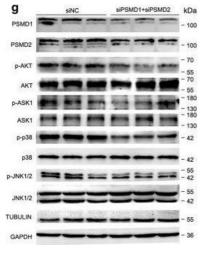


#### **Western Blotting**

**Image 1.** PSMD1 and PSMD2 regulate the expression level of fatty acids (FAs) and lipid synthesis-related genes. a Interference efficiency detection by qRT-PCR. b The expression level of fatty acid synthesis-related genes was detected by qRT-PCR. c The expression level of lipid synthesis-related genes was detected by qRT-PCR. d Overexpression efficiency detection by qRT-PCR. e The expression level of fatty acid synthesis-related genes was detected by qRT-PCR. f The expression level of lipid synthesis-related genes was detected by qRT-PCR. g The

ASK1-p38-JNK and AKT signaling in groups of interfered cells and control cells was detected by Western Blot experiments. TUBULIN and GAPDH were the reference proteins. h Grey value analysis of g. ImageJ software was used for this analysis, according to the instructions. i The ASK1-p38-JNK and AKT signaling in the overexpression cell group and control cells was detected by Western Blot experiments. TUBULIN and GAPDH were the reference proteins. j Grey value analysis of i. ImageJ software was used for this analysis, according to the instructions. The statistical significance of differences between means was assessed using an unpaired Student's t-test (n=3, \*p<0.05, \*\*p<0.01) vs. NC - figure provided by CiteAb. Source: PMID31703613





### **Western Blotting**

**Image 2.** Western blot analysis of extracts from Hela cells using Phospho-MAPK14-T180/Y182 antibody.

#### **Western Blotting**

**Image 3.** PSMD1 and PSMD2 regulate the expression level of fatty acids (FAs) and lipid synthesis-related genes. a Interference efficiency detection by qRT-PCR. b The expression level of fatty acid synthesis-related genes was detected by qRT-PCR. c The expression level of lipid synthesis-related genes was detected by qRT-PCR. d Overexpression efficiency detection by qRT-PCR. e The expression level of fatty acid synthesis-related genes was detected by qRT-PCR. f The expression level of lipid synthesis-related genes was detected by qRT-PCR. g The

ASK1-p38-JNK and AKT signaling in groups of interfered cells and control cells was detected by Western Blot experiments. TUBULIN and GAPDH were the reference proteins. h Grey value analysis of g. ImageJ software was used for this analysis, according to the instructions. i The ASK1-p38-JNK and AKT signaling in the overexpression cell group and control cells was detected by Western Blot experiments. TUBULIN and GAPDH were the reference proteins. j Grey value analysis of i. ImageJ software was used for this analysis, according to the instructions. The statistical significance of differences between means was assessed using an unpaired Student's t-test (n=3, \*p<0.05, \*\*p<0.01) vs. NC - figure provided by CiteAb. Source: PMID31703613