bs-6679R

[Primary Antibody]

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MAOA Rabbit pAb

DATASHEET —

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GenelD: 4128 **SWISS:** P21397

Target: MAOA

Immunogen: KLH conjugated synthetic peptide derived from human MAOA:

281-380/527.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

Background: Monoamine oxidase A and B catalyzes the oxidative deamination of biogenic and xenobiotic amines and has important functions in the metabolism of neuroactive and vasoactive amines in the central nervous system and peripheral tissues. MAOA preferentially oxidizes biogenic amines such as 5-hydroxytryptamine (5-HT), norepinephrine and epinephrine. MAOB preferentially degrades

benzylamine and phenylethylamine

Applications: WB (1:500-2000)

IHC-P (1:100-500) **IHC-F** (1:100-500) **IF** (1:100-500) **ELISA** (1:5000-10000)

Reactivity: Human (predicted: Mouse,

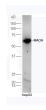
Rat, Rabbit, Pig, Cow, Dog,

Horse)

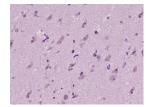
Predicted 60 kDa

Subcellular Location: Cell membrane ,Cytoplasm

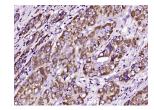
VALIDATION IMAGES



Sample: HepG2 Cell (Human) Lysate at 40 ug Primary: Anti-MAOA (bs-6679R) at 1/300 dilution Secondary: HRP conjugated Goat-Anti-rabbit IgG (bs-0295G-HRP) at 1/5000 dilution Predicted band size: 60 kD Observed band size: 60 kD



Paraformaldehyde-fixed, paraffin embedded (human brain glioma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (MAOA) Polyclonal Antibody, Unconjugated (bs-6679R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human liver carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (MAOA) Polyclonal Antibody, Unconjugated (bs-6679R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.

- SELECTED CITATIONS -

• [IF=4.8] Caiyao Han. et al. Study on the synergistic mechanism of fermented Yaomu on Huafengdan in the treatment of ischemic stroke. JOURNAL OF ETHNOPHARMACOLOGY. 2025 Mar 13:343:119438. Western blot; 39909116