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# anti-Monoamine Oxidase A antibody (Center)





Publication



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Quantity:	100 μL	
Target:	Monoamine Oxidase A (MAOA)	
Binding Specificity:	Center	
Reactivity:	Human, Mouse	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This Monoamine Oxidase A antibody is un-conjugated	
Application:	Western Blotting (WB)	

### **Product Details**

Immunogen:	Recombinant protein encompassing a sequence within the center region of human monoamine oxidase A. The exact sequence is proprietary.	
Isotype:	IgG	
Cross-Reactivity:	Mouse (Murine), Dog (Canine), Pig (Porcine), Rat (Rattus), Cow (Bovine)	
Cross-Reactivity (Details):	Mouse (89 %), Dog (90 %), Pig (90 %), Rat (90 %), Bovine (89 %)	
Characteristics:	Rabbit polyclonal antibody to monoamine oxidase A (monoamine oxidase A) monoamine oxidase A antibody [N2C3]	
Purification:	Purified by antigen-affinity chromatography.	

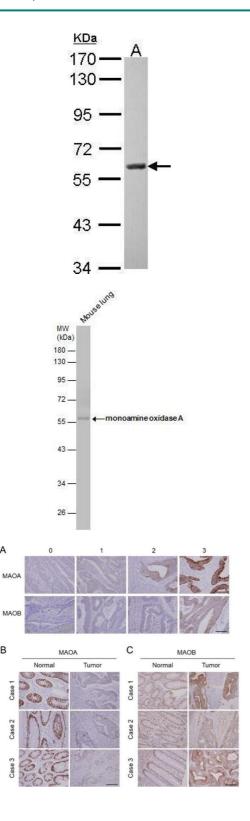
# **Target Details**

Target:	Monoamine Oxidase A (MAOA)	
Alternative Name:	Monoamine Oxidase A (MAOA Products)	
Background:	This gene encodes monoamine oxidase A, an enzyme that degrades amine neurotransmitters, such as dopamine, norepinephrine, and serotonin. The protein localizes to the mitochondrial outer membrane. The gene is adjacent to a related gene on the opposite strand of chromosome X. Mutation in this gene results in monoamine oxidase deficiency, or Brunner syndrome.	
	Cellular Localization: Mitochondrion outer membrane	
Molecular Weight:	60 kDa	
Gene ID:	4128	
Application Details		
Application Notes:	Suggested dilution Reference Western blot 1:500-1:3000* Not tested in other applications.  *Optimal dilutions/concentrations should be determined by the researcher.Suggested dilutionReferenceWestern blot1:500-1:3000*	
Comment:	Positive Control: HepG2	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	0.33 mg/mL	
Buffer:	0.1M Tris, 0.1M Glycine, 10 % Glycerol (pH 7). 0.01 % Thimerosal was added as a preservative.	
Preservative:	Thimerosal (Merthiolate)	
Precaution of Use:	This product contains Thimerosal (Merthiolate): a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	-20 °C	
Storage Comment:	Keep as concentrated solution. Aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.	

Product cited in:

Wu, Yen, Kou, Wu: "Luteolin and Apigenin Attenuate 4-Hydroxy-2-Nonenal-Mediated Cell Death through Modulation of UPR, Nrf2-ARE and MAPK Pathways in PC12 Cells." in: **PLoS ONE**, Vol. 10, Issue 6, pp. e0130599, (2016) (PubMed).

## Validation report #104426 for Immunohistochemistry (IHC)



#### **Western Blotting**

**Image 1.** WB Image Sample (30 ug of whole cell lysate) A: Hep G2 , 7.5% SDS PAGE monoamine oxidase A antibody antibody diluted at 1:1000

#### **Western Blotting**

**Image 2.** WB Image monoamine oxidase A antibody [N2C3] detects monoamine oxidase A protein by western blot analysis. Mouse tissue extracts (50  $\mu$ g) was separated by 10% SDS-PAGE, and the membrane was blotted with monoamine oxidase A antibody [N2C3], diluted at 1:1000.

#### **Immunohistochemistry**

**Image 3.** Immunohistochemical (IHC) results of monoamine oxidase A (MAOA) and MAOB expressions in a Taiwanese colorectal cancer cohort. (A) Representative pictures of expressions of MAOA and MAOB by IHC staining. An intensity score of 0 was defined as negative cytoplasmic staining, of 1 was defined as weak cytoplasmic staining, of 2 was defined as moderate cytoplasmic staining, and of 3 was defined as strong cytoplasmic

staining. Scale bar indicated 100  $\mu$ m. (B,C) Representative IHC staining images for MAOA (B) and MAOB (C) levels in paired normal (N) and tumor tissues (T) from selected colorectal cancer patients. The magnifying factor used in these representative pictures is x400, and the intensity score of MAOA in the N part was 3. Scale bar indicated 100  $\mu$ m. (D,E) Quantified results of cytoplasmic levels of MAOA (D) and MAOB (E) from IHC staining in primary colorectal cancer and corresponding normal colon mucosa. A total of 59 N/T paired data were included. The scores were calculated as the staining intensity score x percentage of stained cells. - figure provided by CiteAb. Source: PMID32316576