

# Datasheet for ABIN870790

# anti-Malondialdehyde antibody

2 Images



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### Overview

Quantity:	30 µg
Target:	Malondialdehyde (MDA)
Reactivity:	Please inquire
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This Malondialdehyde antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunohistochemistry (Frozen Sections) (IHC (fro)), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))

# **Product Details**

Immunogen:	Malondialdehyde (MDA)-modified KLH
Isotype:	IgG2 lambda
Specificity:	MDA-modified proteins, especially dihydropyridine (DHP)-lysine type derivative
Purification:	Protein A purified

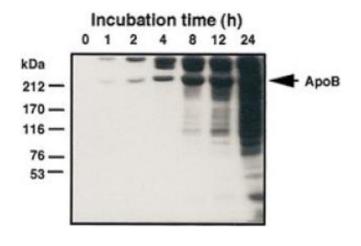
# Target Details

Target:	Malondialdehyde (MDA)
Alternative Name:	Malondialdehyde / MDA (MDA Products)
Target Type:	Chemical

## **Application Details**

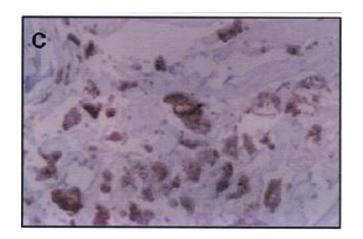
Application Notes:	Paraffin sections, $0.5 - 1 \mu\text{g/ml}$ , Cryosections. This antibody has not been tested for use in all applications. This does not necessarily exclude its use for non-tested procedures. The stated dilutions are recommendations only.est that the app
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Resuspend in aqua bidest.
Buffer:	10 mM PBS, 0.1 % NaN3, 0.1 % BSA
Preservative:	Sodium azide
Precaution of Use:	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C

## **Images**



#### **Western Blotting**

**Image 1.** Western Blot analysis of MDA-LDL adduct formation. LDL (0.5 mg) was incubated With 1 mM MDA in 1 ml of 50 r-nM PBS (pH 7.2) at 37°C. LDL was separated by SDS-PAGE and immunoblotted With A ABIN870790 (2 ug/ml). A ABIN870790 does not react with native LDL apoB, whereas the MDA-modified LDL apoB shows intense immunoreactivity. Yamada S et al. (2001) J Lipid Res 42(8): 1187-96.



#### **Immunohistochemistry**

**Image 2.** Immunohistochemistry image of dihydropyridine adduct staining in paraffn section of human atherosclerotic aorta. Endogenous peroxide activity was quenched by incubating the section for 10 min With 3% hydrogen peroxide. The section was incubated With ABIN870790 and detected using Vectastain ABC kit (Vector Laboratories). 3,3-diaminobenzidine tetrahydrochloride served as the chromogen. Sections were counterstained With hematoxylin. C. AAI 01 1 stains the cytoplasm of most foamy or spindle macrophages. Original magnification: XI 00. Yamada S et al. (2001) J Lipid Res 42(8): 1187-96.