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Datasheet for ABIN967822 anti-PTGS2 antibody (AA 368-604)

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3	Images

Publications



Overview

Quantity:	50 µg
Target:	PTGS2
Binding Specificity:	AA 368-604
Reactivity:	Human, Mouse, Chicken
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This PTGS2 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF), Immunoprecipitation (IP)

Product Details

Immunogen:	Rat Cox-2 aa. 368-604
Clone:	33-Cox
lsotype:	lgG1
Cross-Reactivity:	Mouse (Murine), Human, Chicken
Characteristics:	 Since applications vary, each investigator should titrate the reagent to obtain optimal results. Source of all serum proteins is from USDA inspected abattoirs located in the United States. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive
	deposits in plumbing. 4. Please refer to us for technical protocols.

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Product Details

Purification:

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

Target Details

Target:	PTGS2
Alternative Name:	Cox-2 (PTGS2 Products)
Background:	Cyclooxygenase (Cox) is also known as prostaglandin H synthase or PGH synthase (E.C.
	1.14.99.1). It catalyzes the conversion of arachidonate to prostaglandin H2 (PGH2), the
	precursor of PGE2, PGF2alpha, PGD2, prostacyclin, and thromboxane A2. Cox actually has two
	different enzymatic activities: a cyclooxygenase that mediates the formation of PGG2 from
	oxygen and arachidonate and a hydroperoxidase that catalyzes a reduction of PGG2 yielding
	PGH2. Two Cox genes, Cox-1 and Cox-2, have been isolated in several species. A 4kb mRNA
	encodes the 604 amino acid Cox-2 protein. The two human Cox isoenzymes are 61% identical
	in amino acid composition with the active sites being highly conserved. Cox-2 mRNA and
	protein levels are induced by serum, lipopolysaccharides, growth factors, human chorionic
	gonadotropin and phorbol testers in various mammalian cell types. It has been shown that
	interleukin-1alpha (IL-1alpha) induces increased levels of Cox-2 mRNA and protein in human
	endothelial cells. The sustained increase in Cox-2 is apparently due (at least in part) to IL-1alpha
	increasing the stability of Cox-2 mRNA. This type of regulatory mechanism may play an
	important role in chronic inflammatory conditions.
	Synonyms: PGHS-2, Cyclooxygenase-2
Molecular Weight:	70 kDa
Pathways:	Brown Fat Cell Differentiation, Positive Regulation of fat Cell Differentiation
Application Details	
Comment:	Related Products: ABIN968550, ABIN967389
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	250 μg/mL
Buffer:	Aqueous buffered solution containing BSA, glycerol, and ≤0.09 % sodium azide.

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Handling	
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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
Storage Comment:	Store undiluted at -20°C.
Publications	
Product cited in:	Shiotani, Denda, Yamamoto, Kitayama, Endoh, Sasaki, Tsutsumi, Sugimura, Konishi: "Increased
	expression of cyclooxygenase-2 protein in 4-nitroquinoline-1-oxide-induced rat tongue
	carcinomas and chemopreventive efficacy of a specific inhibitor, nimesulide." in: Cancer
	research, Vol. 61, Issue 4, pp. 1451-6, (2001) (PubMed).
	Marrogi, Pass, Khan, Metheny-Barlow, Harris, Gerwin: "Human mesothelioma samples
	overexpress both cyclooxygenase-2 (COX-2) and inducible nitric oxide synthase (NOS2): in vitro
	antiproliferative effects of a COX-2 inhibitor." in: Cancer research, Vol. 60, Issue 14, pp. 3696-
	700, (2000) (PubMed).
	Giroux, Descoteaux: "Cyclooxygenase-2 expression in macrophages: modulation by protein
	kinase C-alpha." in: Journal of immunology (Baltimore, Md. : 1950), Vol. 165, Issue 7, pp. 3985
	91, (2000) (PubMed).
	Xie, Cho, Calaycay, Mumford, Swiderek, Lee, Ding, Troso, Nathan: "Cloning and characterizatior
	of inducible nitric oxide synthase from mouse macrophages." in: Science (New York, N.Y.), Vo
	256, Issue 5054, pp. 225-8, (1992) (PubMed).



Western Blotting

Image 1. Western blot analysis of Cox-2 on a lysate from mouse macrophages treated with IFNgamma and LPS. Lane 1: 1:250, lane 2: 1:500, lane 3: 1:1000 dilution of the Mouse Anti-Cox-2 antibody.



Immunofluorescence

Image 2. Immunofluoresence staining of mouse macrophages.



Immunohistochemistry

Image 3. Immunohistochemical staining of neurons and endothelial cells from blood vessels (formalin-fixed, citrate buffer pre-treatment, 10X).

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