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anti-PTGS2 antibody (AA 368-604)

Images



Publications



Overview

Quantity:	150 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
Isotype:	lgG1
Cross-Reactivity:	Mouse (Murine), Human, Chicken
Characteristics:	1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
	2. Please refer to us for technical protocols.
	3. 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. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

Product Details

Purification:

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

Target Details

Buffer:

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	

Aqueous buffered solution containing BSA, glycerol, and \leq 0.09 % sodium azide.

Handling

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:

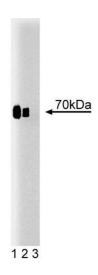
Shiotani, Denda, Yamamoto, Kitayama, Endoh, Sasaki, Tsutsumi, Sugimura, Konishi: "Increased expression of cyclooxygenase-2 protein in 4-nitroquinoline-1-oxide-induced rat tongue carcinomas and chemopreventive efficacy of a specific inhibitor, nimesulide." in: **Cancer research**, Vol. 61, Issue 4, pp. 1451-6, (2001) (PubMed).

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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).

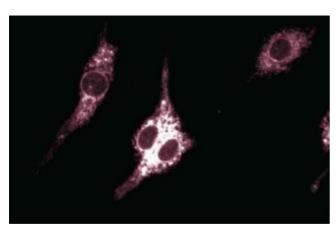
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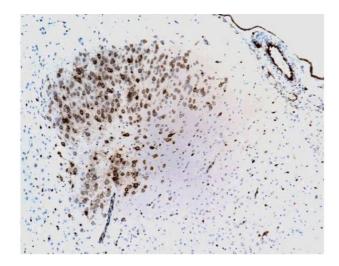
Western Blotting

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



Immunofluorescence

Image 2. Immunofluorescent staining of Mouse Macrophages.



Immunohistochemistry

Image 3. Staining of the neurons and endothelial cells of the blood vessels, formalin-fixed, citrate buffer pre-treatment, 10X