# Cox2 (D5H5) XP® Rabbit mAb

Small 100 µl (10 western blots)

Petite 40 ul (4 western blots)

Cell Signaling

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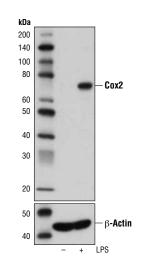
## For Research Use Only. Not For Use In Diagnostic Procedures.

Applications	Species Cross-Reactivity*	Molecular Wt.	Isotype	
W, IHC-P, IF-IC, F	H, M, R	74 kDa	Rabbit IgG**	

Background: Cyclooxygenase1 (Cox1) and cyclooxygenase2 (Cox2), family members with 60% homology in humans, catalyze prostaglandin production from arachidonic acid (1,2). While Cox1 expression is constitutive in most tissues, Cox2 expression is induced by lipopolysaccharide (LPS) and peptidoglycan (PGN) (3). PGN activates Ras, leading to phosphorylation of Raf at Ser338 and Erk1/2 at Tyr204. The activation of MAP kinase signaling results in subsequent activation of IKK $\alpha/\beta$ , phosphorylation of I $\kappa$ B $\alpha$ at Ser32/36, and NF-kB activation. Finally, activation of the transcription factor NF-κB is responsible for the induction of Cox2 expression (4). Investigators have shown that LPS and PGN induce the clinical manifestations of arthritis and bacterial infections, such as inflammation, fever, and septic shock (5). Research studies have indicated that Cox1 and Cox2 may also play a role in the neuropathology of Alzheimer's disease by potentiating  $\gamma$ -secretase activity and β-amyloid generation (6).

Specificity/Sensitivity: Cox2 (D5H5) XP® Rabbit mAb recognizes endogenous levels of total Cox2 protein.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding His108 of human Cox2 protein.



Western blot analysis of extracts from Raw 264.7 cells, untreated (-) or LPS-treated (1 μg/ml, 24 hr; +), using Cox2 (D5H5) XP®

Entrez-Gene ID #5743 Swiss-Prot Acc. #P35354

**Storage:** Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

\*Species cross-reactivity is determined by western blot.

\*\*Anti-rabbit secondary antibodies must be used to detect this antibody.

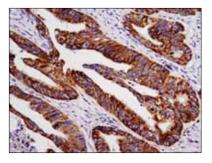
### **Recommended Antibody Dilutions:**

1:1000 Western blotting Immunohistochemistry (Paraffin) 1:600† Unmasking buffer: Citrate SignalStain® Antibody Diluent #8112 Antibody diluent: Detection reagent: SignalStain® Boost (HRP, Rabbit) #8114 †Optimal IHC dilutions determined using SignalStain® Boost IHC

Detection Reagent. Immunofluorescence (IF-IC) 1:800 1:200 Flow Cytometry

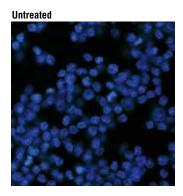
For product specific protocols please see the web page for this product at www.cellsignal.com.

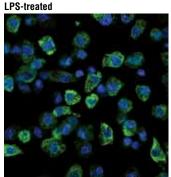
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Immunohistochemical analysis of paraffin-embedded human colon carcinoma using Cox2 (D5H5) XP® Rabbit mAb.

Rabbit mAb or β-Actin (D6A8) Rabbit mAb #8457.



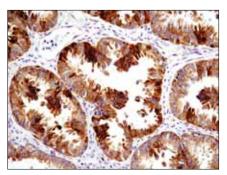


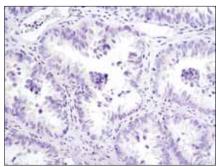
Confocal immunofluorescent analysis of Raw 264.7 cells, untreated (left; -) or LPS-treated (right; 1ug/ml, 24 hrs; +), using Cox2 (D5H5) XP® Rabbit mAb (green). Blue pseudocolor = DRAQ5® #4084 (fluorescent DNA dye).

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

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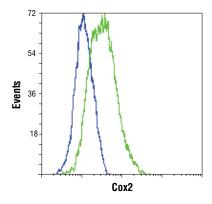
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Immunohistochemical analysis of paraffin-embedded human ovarian carcinoma using Cox2 (D5H5) XP® Rabbit mAb in the pres-

# ence of control peptide (left) or antigen-specific peptide (right).



Flow cytometric analysis of Raw 264.7 cells, untreated (blue) or LPS-treated (1µg/ml, 24 hrs; green), using Cox2 (D5H5) XP® Rabbit mAb. Anti-rabbit IgG (H+L), F(ab'), Fragment (Alexa Fluor® 488 Conjugate) #4412 was used as a secondary antibody.

## **Background References:**

- (1) Xie, W.L. et al. (1991) Proc Natl Acad Sci USA 88, 2692-6.
- (2) Vane, J.R. et al. (1998) Annu Rev Pharmacol Toxicol 38, 97-120.
- (3) O'Neill, G.P. et al. (1994) Mol Pharmacol 45, 245-54.
- (4) Chen, B.C. et al. (2004) J Biol Chem 279, 20889-97.
- (5) Wang, Q. et al. (2001) Infect Immun 69, 2270-6.
- (6) Qin, W. et al. (2003) J Biol Chem 278, 50970-7.