PRODUCT INFORMATION



COX-1 (ovine) Polyclonal Antibody

Item No. 160108

Overview and Properties

Contents: This vial contains 200 µl of lyophilized antiserum.

Cyclooxygenase-1, PGHS-1, Prostaglandin Endoperoxide Synthase 1, Synonyms:

Prostaglandin G/H Synthase 1, Prostaglandin H2 Synthase 1

Immunogen: Peptide from an internal region of ovine COX-1

Cross Reactivity: (-) Ovine, human, and mouse COX-2

Species Reactivity: (+) Human, bovine, ovine, and porcine; (-) Mouse and rat COX-1

Uniprot No.: P05979 Form: Solid

-20°C (as supplied) Storage:

Storage Buffer: Lyophilized from serum, resuspend in 200 μ l double distilled water.

≥3 years Stability: Rabbit Host:

Applications: Western blot (WB); the recommended starting dilution for WB is 1:500. Suitable

for immunofluorescence and immunohistochemistry working dilution should be

determined empirically.

Description

Cyclooxygenase 1 (COX-1) is a bifunctional enzyme that exhibits both COX and peroxidase activities. 1,2 It is composed of an N-terminal signal peptide, an EGF-like domain, a membrane binding domain, a catalytic domain, and a C-terminal tail.3 COX-1 is constitutively expressed in the gastrointestinal tract, kidney, spleen, liver, and lung and localizes to the endoplasmic reticulum.^{4,5} The COX component converts arachidonic acid (Item Nos. 90010 | 90010.1 | 10006607) to a hydroperoxyl endoperoxide prostaglandin G₂ (PGG₂; Item No. 17010) and the peroxidase component reduces the endoperoxide to the corresponding alcohol PGH₂ (Item No. 17020), the precursor of PGs, thromboxanes, and prostacyclins. ^{1,2} COX-1 is the target of many non-steroidal anti-inflammatory drugs (NSAIDs) and is responsible for the undesirable gastrointestinal and renal side effects, such as ulcer formation and reductions in the glomerular filtration rate, respectively.^{6,7} Cayman's COX-1 (ovine) Polyclonal Antiserum can be used for Western blot (WB). The antibody recognizes COX-1 at 70 kDa from human, porcine, bovine endothelial, and ovine seminal vesicle samples.

References

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- 3. Smith, W.L. and DeWitt, D.L. Prostaglandin endoperoxide H synthases-1 and -2. Adv. Immunol. 62, 167-215 (1995).
- 4. Seibert, K., Zhang, Y., Leahy, K., et al. Pharmacological and biochemical demonstration of the role of cyclooxygenase 2 in inflammation and pain. Proc. Natl. Acad. Sci. USA 91(25), 12013-12017 (1994).
- 5. Morita, I., Schindler, M., Regier, M.K., et al. Different intracellular locations for prostaglandin endoperoxide H synthase-1 and -2. The Journal of Biological Chemisty 270(18), 10902-10908 (1995).
- Gierse, J.K., Hauser, S.D., Creely, D.P., et al. Expression and selective inhibition of the constitutive and inducible forms of human cyclo-oxygenase. Biochem. J. 305(Pt. 2), 379-484 (1995).
- Frölich, J.C. A classification of NSAIDs according to the relative inhibition of cyclooxygenase isoenzymes. Trends Pharmacol. Sci. 18(1), 30-34 (1997).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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