

[9261177](http://www.uniprot.org/citations/9261177), PubMed: [16373578](http://www.uniprot.org/citations/16373578), PubMed: [22942274](http://www.uniprot.org/citations/22942274), PubMed: [26859324](http://www.uniprot.org/citations/26859324), PubMed: [27226593](http://www.uniprot.org/citations/27226593)). This complex transformation is initiated by abstraction of hydrogen at carbon 13 (with S-stereochemistry), followed by insertion of molecular O₂ to form the endoperoxide bridge between carbon 9 and 11 that defines prostaglandins. The insertion of a second molecule of O₂ (bis-oxygenase activity) yields a hydroperoxy group in PGG₂ that is then reduced to PGH₂ by two electrons (PubMed: [7947975](http://www.uniprot.org/citations/7947975), PubMed: [7592599](http://www.uniprot.org/citations/7592599), PubMed: [9261177](http://www.uniprot.org/citations/9261177), PubMed: [16373578](http://www.uniprot.org/citations/16373578), PubMed: [22942274](http://www.uniprot.org/citations/22942274), PubMed: [26859324](http://www.uniprot.org/citations/26859324), PubMed: [27226593](http://www.uniprot.org/citations/27226593)). Similarly catalyzes successive cyclooxygenation and peroxidation of dihomo-gamma-linoleate (DGLA, C20:3(n-6)) and eicosapentaenoate (EPA, C20:5(n-3)) to corresponding PGH₁ and PGH₃, the precursors of 1- and 3- series prostaglandins (PubMed: [11939906](http://www.uniprot.org/citations/11939906), PubMed: [19540099](http://www.uniprot.org/citations/19540099)). In an alternative pathway of prostanoid biosynthesis, converts 2-arachidonoyl lysophospholipids to prostanoid lysophospholipids, which are then hydrolyzed by intracellular phospholipases to release free prostanoids (PubMed: [27642067](http://www.uniprot.org/citations/27642067)). Metabolizes 2-arachidonoyl glycerol yielding the glyceryl ester of PGH₂, a process that can contribute to pain response (PubMed: [22942274](http://www.uniprot.org/citations/22942274)). Generates lipid mediators from n-3 and n-6 polyunsaturated fatty acids (PUFAs) via a lipoxygenase-type mechanism. Oxygenates PUFAs to hydroperoxy compounds and then reduces them to corresponding alcohols (PubMed: [11034610](http://www.uniprot.org/citations/11034610), PubMed: [11192938](http://www.uniprot.org/citations/11192938), PubMed: [9048568](http://www.uniprot.org/citations/9048568), PubMed: [9261177](http://www.uniprot.org/citations/9261177)). Plays a role in the generation of resolution phase interaction products (resolvins) during both sterile and infectious inflammation (PubMed: [12391014](http://www.uniprot.org/citations/12391014)). Metabolizes docosahexaenoate (DHA, C22:6(n-3)) to 17R-HDHA, a precursor of the D- series resolvins (RvDs) (PubMed: [12391014](http://www.uniprot.org/citations/12391014)). As a component of the biosynthetic pathway of E-series resolvins (RvEs), converts eicosapentaenoate (EPA, C20:5(n-3)) primarily to 18S-HEPE that is further metabolized by ALOX5 and LTA4H to generate 18S-RvE1 and 18S- RvE2 (PubMed: [21206090](http://www.uniprot.org/citations/21206090)). In vascular endothelial cells, converts docosapentaenoate (DPA, C22:5(n-3)) to 13R-HDPA, a precursor for 13-series resolvins (RvTs) shown to activate macrophage phagocytosis during bacterial infection (PubMed: [26236990](http://www.uniprot.org/citations/26236990)). In activated leukocytes, contributes to oxygenation of hydroxyeicosatetraenoates (HETE) to diHETES (5,15-diHETE and 5,11-diHETE) (PubMed: [22068350](http://www.uniprot.org/citations/22068350), PubMed: [26282205](http://www.uniprot.org/citations/26282205)). Can also use linoleate (LA, (9Z,12Z)- octadecadienoate, C18:2(n-6)) as substrate and produce hydroxyoctadecadienoates (HODEs) in a regio- and stereospecific manner, being (9R)-HODE ((9R)-hydroxy-(10E,12Z)-octadecadienoate) and (13S)- HODE ((13S)-hydroxy-(9Z,11E)-octadecadienoate) its major products (By similarity). During neuroinflammation, plays a role in neuronal secretion of specialized preresolving mediators (SPMs) 15R-lipoxin A4 that regulates phagocytic microglia (By similarity).

Cellular Location

Microsome membrane; Peripheral membrane protein. Endoplasmic reticulum membrane; Peripheral membrane protein. Nucleus inner membrane; Peripheral membrane protein. Nucleus outer membrane; Peripheral membrane protein. Note=Detected on the luminal side of the endoplasmic reticulum and nuclear envelope

PTGS2 Antibody (Center P378) Blocking peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

PTGS2 Antibody (Center P378) Blocking peptide - Images

PTGS2 Antibody (Center P378) Blocking peptide - Background

Prostaglandin-endoperoxide synthase (PTGS), also known as cyclooxygenase, is the key enzyme in prostaglandin biosynthesis, and acts both as a dioxygenase and as a peroxidase. There are two isozymes of PTGS: a constitutive PTGS1 and an inducible PTGS2, which differ in their regulation of expression and tissue distribution. This gene encodes the inducible isozyme. It is regulated by specific stimulatory events, suggesting that it is responsible for the prostanoid biosynthesis involved in inflammation and mitogenesis.

PTGS2 Antibody (Center P378) Blocking peptide - References

Duggan, K.C., et al. J. Biol. Chem. 285(45):34950-34959(2010) Feher, A., et al. Am J Geriatr Psychiatry 18(11):983-987(2010) Wang, C.H., et al. Anticancer Res. 30(9):3649-3653(2010) Han, E.H., et al. J. Toxicol. Environ. Health Part A 73 (21-22), 1451-1464 (2010) :Cao, H., et al. Tohoku J. Exp. Med. 222(1):15-21(2010)