

SELECTIVE COX2 INHIBITORS%0A

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The cyclooxygenase active site is created by a long hydrophobic channel that is the site of non-steroidal anti-inflammatory drug binding. This active site extends from the membrane-binding domain (the lobby) to the core of the catalytic domain (20, 21).

<http://www.thegraphicdesign.co/Selective-COX-2-Inhibitors--A-Review-of-Their-Structure--.pdf>

COX 2 inhibitor Wikipedia

Selective COX-2 inhibitors are a type of nonsteroidal anti-inflammatory drug (NSAID) that directly targets cyclooxygenase-2, COX-2, an enzyme responsible for inflammation and pain. Targeting selectivity for COX-2 reduces the risk of peptic ulceration, and is the main feature of celecoxib, rofecoxib and other members of this drug class.

<http://www.thegraphicdesign.co/COX-2-inhibitor-Wikipedia.pdf>

THE STRUCTURE OF NS 398 BOUND TO PubMed Central PMC

NS-398 is a COX-2-selective inhibitor of the sulfonanilide class and as our present results establish, binds uniquely within the cyclooxygenase channel of COX-2. In contrast to that observed for the diarylheterocyclic coxibs, NS-398 does not penetrate the side pocket of COX-2 past the side chain of Val-523 or interact with the side chain of Arg-513 at the base of the pocket. Instead, the

<http://www.thegraphicdesign.co/THE-STRUCTURE-OF-NS-398-BOUND-TO---PubMed-Central--PMC-.pdf>

Discovery and development of cyclooxygenase 2 inhibitors

The impetus for development of selective COX-2 inhibitors was the adverse gastrointestinal side-effects of NSAIDs. Soon after the discovery of the mechanism of action of NSAIDs, strong indications emerged for alternative forms of COX, but little supporting evidence was found.

<http://www.thegraphicdesign.co/Discovery-and-development-of-cyclooxygenase-2-inhibitors--.pdf>

In silico molecular docking analysis of prodigiosin and

Introduction. Inflammation is the tissue reaction against infection, irritation or foreign substance. It is a part of the host defense mechanisms that is known to be involved in the inflammatory reactions associated with the release of histamine, bradykinin & prostaglandins.

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RESEARCH Open Access silico molecular docking analysis of

RESEARCH Open Access In silico molecular docking analysis of prodigiosin and cycloprodigiosin as COX-2 inhibitors Pabba Shiva Krishna¹, Kompally Vani², Metuku Ram Prasad¹, Burra Samatha¹,

<http://www.thegraphicdesign.co/RESEARCH-Open-Access-silico-molecular-docking-analysis-of--.pdf>

DOCKING STUDIES ON XANTHONES OF MANGOSTEEN AS COX 2 INHIBITORS

COX-2 were ranked according to the GOLD fitness function. The docking results showed that all The docking results showed that all the xanthone derivatives of mangosteen are active COX inhibitors with a significant preference

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