Abstract

Products of Non-steroidal anti-inflammatory drugs (NSAIDs) are principally and most widely utilized due to their multipurpose actions, i.e., analgesic anti pyretic and anti-inflammatory effect. Beside their therapeutic effect their toxic effect based on their selectivity of cyclooxygenase (COX) enzyme. While cardiovascular toxicities associated with NSAIDs are mainly related with COX-2 selective inhibitor and primarily dependent on their dose time and duration. Variability in therapeutic response and susceptibility to toxicity is well recognized and can be manageable if considerable precautions implement.

Keywords: Non-steroidal anti-inflammatory; Celecoxib; Cardiovascular

Introduction

In conditions like fever, pain, headache, arthritis, lupus disease and other inflammatory problems, there are numerous drugs which have frequently been recommended from the class of therapeutic agents known as non-steroidal anti-inflammatory drugs [1]. Aspirin (the proto type drug of this class) and other NSAIDs have become the cornerstone therapy to avert any thrombotic and cardiolesional episodes in majority of high risk sick population [2]. Although, NSAIDs are effective and beneficial in outline pathological conditions, prolonged use of NSAIDs are highly associated with severe side effects. However, many of these side effects can be prevented by careful management and by implementation of preventive strategies.

Mechanism of Action

The clinical effects of non-steroidal anti-inflammatory were first described by Vane and Piper. They suggest that NSAIDs inhibit the biosynthesis of prostanoids by inhibiting the enzyme cyclooxygenase (COX). Arachidonic acid is the substrate for COX enzyme lead to the formation of prostanoids including prostaglandins, prostacyclins, and thromboxanes [3]. Different isoform have been discovered [4] out of which COX-1 (PGHS-1) and COX-2 (PGHS-2) are the two allied COX enzymes which have been delineated. In sequences of amino acids, both of these isoforms have 60% homology that conserved for COX enzymes which have been delineated. In sequences of amino acids, both of these isoforms have 60% homology that conserved for the arachidonic acid catalysis [5-9]. Both of these isoforms serves as a catalyst in converting arachidonic acid to COX-2 which results in formation of prostaglandin, the biologically active form. Out of the two, the first isoform; prostaglandin H2 Synthase-1 (PGHS-1) is the major and exclusive loci for non-steroidal anti-inflammatory drug activity but the second isozyme; PGH Synthase-2 (PGHS-2) is also sensitive to actions of NSAIDs which is also recognized with it.

Anti-inflammatory glucocorticoids inhibit the expression of COX-2 which further explains the involvement of this enzyme in inflammation. One of the possible mechanisms for pinpointing the secure and more viable use of non-steroidal anti-inflammatory drugs arises from such findings that both the isoforms of COX enzyme can be segregate halted. Drugs which are inhibitors of PGHS-2 are the inflammation reducing agents and on the other hand they reserve the synthesis of PG by kidneys and stomach and therefore decrease unfavorable adverse reactions which are most frequently linked with NSAIDs [10]. Physiological stimuli including shear stress in vasculature and ovulation regulate both isoforms [11,12]. The enzyme COX-1 involve in various physiological conditions including mucosal protection and cardiovascular protection, while, certain processes of inflammation brings about COX-2 enzyme in turn the products of this isoform involve in growth factors, lymphokines and other inflammatory mediators. In Hypertension, diabetes mellitus (DM), bone fracture, heart failure and similar pathological conditions and even in non-pathological kidneys, there is enhanced interrelation of mRNA and proteins of COX-2 enzyme [13].

Therapeutic and harmful activities are linked with the suppression of COX enzyme. Studies suggested that actions of non-steroidal agents against inflammations are because of suppression of COX-2 whereas, the dwindling of COX-1 lead to stomach lining irritation [14].

Classification of NSAIDS

Based on the characteristics including chemical (Table 1) and pharmacological properties and (Table 2) show the classification of NSAIDs on the bases of selectivity toward COX enzymes [15,16].

Mechanism of Cardiovascular Toxicity of NSAIDS

It is believed that at therapeutic doses, COX-2 enzyme inhibitors halt COX-2 isozyme but it results in metabolic disturbances which leads to the over production of by products that are deleterious and damages arterial wall and produces arterial blood clotting [17,18]. Leukotriene B4 and thromboxane A2 (TXA2) will be formed in significant amounts and prostacyclin (PGI2) from arachidonic acid will be synthesized very less when COX-2 is inhibited therefore this balancing tip allows TXA2 to function unopposed that results in higher risk for CV adverse

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When inhibition of prostacyclin takes place which is also an atheroprotective prostanooid, cardio-toxicity occurs and this is due to the administration of traditional as well as COX-2 selective non-steroidal anti-inflammatory agents. The traditional non-steroidal agents along with coxibs frequently and selectively inhibits COX-2 enzyme which is also linked with the suppression of platelet COX-1 enzyme but it is not adequate to suppress the function of platelets and particularly this property of coxibs and traditional non-steroidal anti-inflammatory agents mainly elucidates their apportioned CVS toxicities [29]. Non-steroidal anti-inflammatory agents having actions on both of the isoforms of COX enzyme are less appealing to use as compared to the drugs which are selective for COX-2 enzyme and this is due to the GI side effects of non-steroidal anti-inflammatory drugs targeting non-selectively both enzymes, hence the drugs that are particular for COX-2 enzyme suppression have equal actions against inflammation like the non-selective agents but unlike non-selective drugs, they have negligible harm to GI system [30].

The safety of drugs particularly suppressing the COX-2 enzymes and non-selective traditional agents regarding cardiovascular system have been a considerable topic to discuss therefore the higher chances for such cardiovascular incidents associated with the use of different formulations of rofecoxib was highlighted following the influential trials like APPROV [31].

Hypertension and volume retention are also one of the multiple mechanisms like differences in endothelial functions and oxidative stress due to CV risks associated with NSAIDs [32]. The suppression of prostacyclin formation which is dependent on COX-2 enzyme inhibition without the dwindling of production of thromboxane due to COX-1 enzyme inhibition is due to the increased risks of thrombotic events caused by COX-2 selective inhibitors [32,33].

Ungrprasert et al. in 2015 investigated the association of non-aspirin non-steroidal anti-inflammatory drugs and risk of hemorrhagic stroke. A range of observational studies were systematically evaluated and meta-analysis of selected studies was performed. It was concluded that NSAIDs as particular group was not connected with a notably eminent threat of hemorrhagic stroke. But, an amplified risk was reported in numerous moieties of NSAID preferentially in meloxicam that NSAIDs as particular group was not connected with a notably eminent threat of hemorrhagic stroke. But, an amplified risk was reported in numerous moieties of NSAID preferentially in meloxicam.

Efficacy and Dosage

Individual drug in NSAIDs group possess own therapeutic

<table>
<thead>
<tr>
<th>Groups</th>
<th>Example(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Salicylates</td>
<td>Acetyl salicylic acid, Sulfasalazine</td>
</tr>
<tr>
<td>Propionic acid derivatives</td>
<td>Ibuprofen, naproxen, ketoprofen, flurbiprofen, fenoprofen, Oxaprozin.</td>
</tr>
<tr>
<td>Pyrocarboxylic acids</td>
<td>Etodolac, Tolmetin, diclofenac, ketorolac</td>
</tr>
<tr>
<td>Heteroaryl acetic acid</td>
<td>Nabumeton</td>
</tr>
<tr>
<td>Alkanones</td>
<td>Indomethacin, sulindac, etodolac</td>
</tr>
<tr>
<td>Indoleacetic, Indeneacetic acids</td>
<td>Piroxicam, meloxicam, Ketorolac</td>
</tr>
<tr>
<td>Fenamates</td>
<td>Mefenamic acid, meclofenamic acid</td>
</tr>
<tr>
<td>Diaryheterocycles</td>
<td>Rofecoxib, celecoxib, veldoxib, paracoxib, etoricoxib, lumaricoxib</td>
</tr>
</tbody>
</table>

Table 1: Chemical classification of NSAIDS.

<table>
<thead>
<tr>
<th>COX-1 Isform</th>
<th>COX2/COX1 Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ketorolac</td>
<td>-</td>
</tr>
<tr>
<td>Flurbiprofen</td>
<td>0.078</td>
</tr>
<tr>
<td>Ketoprofen</td>
<td>-</td>
</tr>
<tr>
<td>Indomethacin</td>
<td>30</td>
</tr>
<tr>
<td>Acetyl Salicylic acid</td>
<td>2-167</td>
</tr>
<tr>
<td>Ibuprofen</td>
<td>0.15-1.7</td>
</tr>
<tr>
<td>Naproxen</td>
<td>0.6</td>
</tr>
<tr>
<td>Tenidap</td>
<td>122</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>COX-2 Isform</th>
<th>COX2/COX1 Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Celecoxib</td>
<td>30</td>
</tr>
<tr>
<td>Rofecoxib</td>
<td>4.0</td>
</tr>
<tr>
<td>Etodolac</td>
<td>1.9</td>
</tr>
<tr>
<td>Meloxicam</td>
<td>0.33</td>
</tr>
<tr>
<td>Diclofenac NaK</td>
<td>0.6 - 2.2</td>
</tr>
<tr>
<td>Sulindac</td>
<td>-</td>
</tr>
<tr>
<td>Sodium salicylate</td>
<td>-</td>
</tr>
<tr>
<td>Piroxicam</td>
<td>33</td>
</tr>
<tr>
<td>Tenoxicam</td>
<td>15</td>
</tr>
</tbody>
</table>

Table 2: Classification of NSAIDS based on their cox-1 or cox-2 selectivity [60].

Events since TXA2 is vasocostricitive and pro-aggregatory while PGI2 in vasodilatory and anti-aggregatory (Figure 1).

Side effects associated with cardiovascular system (CVS) and gastrointestinal (GI) mainly linked with the use of traditional non-steroidal anti-inflammatory drugs and COX-2 enzyme inhibitors [19]. And there is virtually higher chance of CVS pathologies in the population of patients who administered the drug formulations of COX-2 inhibitors as revealed through many recent clinical trials but it is quite evident that the hazards of GI issues are decreased by using the inhibitors of COX-2 enzymes [20-23].

In patients involving their cardiovascular system and having increased risk for such incidents, the main thing is the use of such agents that can give adequate hostage of toxicity prevention and fulfill provision of efficacy [24,25]. It was initially thought about COX-2 inhibitors that they have lesser analgesic activity as compared to traditional non-steroidal anti-inflammatory agents, moreover it was verified through the table constructed by the Oxford pain group [26-28].

![Cardiovascular toxicity mechanism](image)
effectiveness and strength (optimal dose) for different conditions. The most common doses of NSAIDs prescribe ranging 200 mg, 400 mg, 600 mg and 800 mg (Table 3). The products of Celecoxib having shorter half-lives and administered in doses of once daily are less intruding with the COX-enzyme system than the drugs having half-lives longer and administration is in dosages of once daily or the drugs possess shorter half-lives prescribed in twice daily dosing regimen. This shows the clinical importance of pharmacokinetics of the drug [33]. These drugs are normally available over-the-counter for the suppression of conditions such as headache, back and joint pain, and menstrual pain.

Numbers of studies have been found to be evaluating the GI and cardiovascular safety of alternative dosing strategies such as alternate day dosing, once daily verses twice daily dosing or periodic drug holidays.

Comparison and Facts of Different NSAIDS

Prolong and high doses of NSAIDS have a wide range of adverse effect, mainly on alteration in GI, renal functions and CVS system [34]. However, those patients who are taking therapeutic dose of NSAIDS for shorter time tolerates them well [35]. Number of studies have been reported that the risk associated with the NSAIDs depend upon the intensity, dose and duration [36]. It was consider that the use of NSAIDS are associated with CV side effects however later it was found that only non-selective NSAIDs accompanying with CV adverse effects such as increased risk of myocardial infarction (MI) [37,38]. Conclusively, on the basis of the results of randomized control trials (RCTs), at higher doses the effects of NSAIDS, except for naproxen, showing links to high hazards for MI development and in those patient who were previously diagnosed with coronary heart disease but for diclofenac and Rofecoxib risk associated is not dependent on doses [38].

High risk patient population including patients having congestive heart failure (CHF), hepatic and kidney disorders, patients administering diuretics and angiotensin converting enzyme inhibitors, are more prone to develop hypertension by using the drugs selectively targeting COX-2 enzyme [39]. Results of various RCTs emphasizes on evidence of the hazards which are associated with the use of non-steroidal anti-inflammatory agents and inhibitors of COX-2 enzymes, explaining highest risk of MI was associated with Rofecoxib and lumiracoxib, stroke occurrence was found to be hooked up with the use of Ibuprofen and Diclofenac, while many of the cardiovascular events associated with use of Etoricoxin and Diclofenac. When the Diclofenac used in doses higher than needed for therapeutic response, risk of death was remarked through various RCTs [29].

Chronic utilization of non-steroidal anti-inflammatory drugs linked with various deleterious health issues. Summary of the results of a study showed that the relative risk of Rofecoxib in doses of less than and equals to 25 mg with 95% Confidence Interval was found to be 1.33 (1.00-1.79) while on the other hand when the dose was increased and greater than 25 mg with similar confidence intervals, the relative risk of CVS events associated with Rofecoxib was found to be 2.19 (1.64-2.91). Similarly, Celecoxib showed to have relative risk of 1.06 (0.91-1.23) with 95% confidence interval while Diclofenac showed highest values of relative risks 1.40 (1.16-1.70) with 95% confidence interval. Moreover Meloxicam and Piroxicam showed relative risks of 1.25 (1.00-1.55) and 1.06 (0.70-1.59) respectively. Ibuprofen showed 1.07 (0.97-1.18) and Indomethacin was found to have relative risk of 1.30 (1.07-1.60). Naproxen among all NSAIDs included in the study showed lowest relative risks of 0.97 (0.87-1.07) with 95% confidence interval. It showed that Rofecoxib was found to be most harmful in either dosage regimen and Naproxen was least dangerous contrary to which Diclofenac was on the top of the list for increasing hazards of CV toxicity [40-42].

Likewise, with various other trials including adenoma prevention with Celecoxib (APC) trial and the prevention of spontaneous adenomatous polyps (PreSAP) trial for evaluating the safety and efficacy of Celecoxib in cardiovascular events and blood pressure divulge, the hazard ratio was found to be 2.6% (95% CI; 1.1-6.1) in the patients taking 200 mg Celecoxib twice daily. The hazard ratio was found to be 3.4% (95% CI; 1.5-7.9) in patients with Celecoxib dose of 400 mg twice daily in APC trial and 1.3 (95% CI, 0.6-2.6) in patients who had taken Celecoxib 400 mg OD in PreSAP trial. Highly significant elevation in the systolic blood pressure was also shown by both dosage groups in APC trial while in PreSAP group no blood pressure changes were seen. Cardiovascular events and blood pressure is possibly raised up to two-folds by increasing doses of Celecoxib and there is meaningful reduction in CV events by administering lower doses or other dose intervals [43-46].

Management of CVS Toxicities by NSAIDS

In the presence of partial suppression of platelet COX-1, there seems to be a deep-seated suppression of COX-2 dependent PGI-2 and it put one's finger on the fact that this is the most credible process of cardiovascular risks and toxicities which are associated with the use of non-steroidal anti-inflammatory agents both selective and non-selective [47,48]. The use of a biomarker strategy of COX inhibition has permitted the understanding that the extent on how much patients are exposed (magnitude and duration) is substantial determinant of a much raised risk of nonfatal MI [49]. Different study outcomes make a strong foundation to suggest the possible strategy for the treatment and prevention of cardiovascular hazards in the patient population with different pathologies of vascular system along with various inflammatory conditions [50].

Since Ibuprofen proves to interrupt the Aspirin effects which are anti-platelet therefore shunning the concomitant use of Ibuprofen is recommended [51,52]. If Aspirin is prescribed to a high risks patient of vascular incidents, Naproxen along with a proton pump inhibitor must be administered before two hours of aspirin. When choosing appropriate treatment in different patient traits, NSAIDs must be considered cautiously. Diclofenac showed highest mortality and morbidity in patients with increased cardiovascular risk while on the other hand naproxen serves to have least toxicity against cardiovascular system. Hence, there must be a rational prescribing pattern for non-steroidal anti-inflammatory agents [53].

Current NSAIDS Utilization Pattern

Rational Therapeutic utilization and patient education plays indispensable role when it comes to use of NSAIDs. Guidelines for the use of NSAIDs have been published by many different organizations.
and all these recommendations reflects a proper regime that can not only provide an effective pain control but optimal cardiovascular and GI protection along with the evaluation of CV and GI risks factors in patients prior to initiation of COX-2 or NSAIDs [54,55].

Following are the recommendations of FDA regarding NSAIDs that:

- When COX-2 inhibitors and traditional non-steroidal anti-inflammatory drugs (NSAIDs) are to be used for the management of individual patients; they should be prescribed with the lowest effective dose and for the shortest duration.
- They should not be prescribed for high risk patients, e.g., patients with a history of ischemic heart disease, stroke or congestive heart failure, or in patients who have recently undergone coronary artery bypass grafting (CABG).
- All prescription-strength NSAIDs will now display “black box” label warnings for the potential risk of cardiovascular and gastrointestinal adverse effects.
- Treatment with NSAIDs alone in patients aged less than 65 years who do not have gastrointestinal risk factors is considered appropriate. Co-therapy with a proton pump inhibitors (PPI) or treatment with a COX-2 inhibitor was considered unnecessary in these patients.
- The use of a NSAID alone was considered inappropriate in any patient with a previous gastrointestinal event and in those who concurrently receive aspirin, steroids or warfarin. These patients should receive tNSAID plus a PPI or a COX-2 inhibitor.
- Use of a COX-2 inhibitor with PPI co-therapy is appropriate only in patients at very high risk, such as those with a previous gastrointestinal event who are taking aspirin, and those who are taking aspirin plus steroids or warfarin [56].

Significant Considerations

Blood pressure, plasma half-life, dose dependency, interaction with ASA and COX-2 selectivity are few of the many variable factors that have affected CV hazards and risks which are coupled when non-steroidal anti-inflammatory agents are used [57]. It has been revealed through many observational studies, systematic reviews, meta-analysis and clinical trials that in comparison with other non-selective and COX-2 selective NSAIDs, Naproxen has consistently shown lower risk of CV events. Among non-selective NSAIDs, highest CV risk has been linked with use of Diclofenac. At twice daily regimens and at higher doses, there is evidence showing high risk of CV events associated with celecoxib. When Acetylsalicylic acid is used along with either diclofenac or naproxen, it is proved that there is negligible abatement for protective cardiovascular system however, there founds an interaction between ibuprofen and ASA. Therefore to enhance its anti-platelet activity, Acetylsalicylic acid must get to the patient 30 minutes before or 8 hours after ibuprofen use [58-63].

Conclusion

Due to anti-natriuretic effects as well as the harmful actions of non-steroidal anti-inflammatory agents on vascular activity of heart, it is found to delegitimize cardiovascular system, along with ineffectiveness of therapeutic agents. Selectivity in COX enzyme inhibition is major factor to be considered when prescribing individualized therapy to patient. So it can be concluded since inhibitors of COX-2 enzyme are associated with various cardiovascular pathologies like CHF, HTN and cardiac arrhythmias, therefore in patients with any of these existing conditions, they must be administered cautiously, by selecting drugs rationally in low doses and less frequently.

References


