A comparative study of aceclofenac versus etoricoxib in the management of acute low back pain in a tertiary care hospital

Hema Jagannathan¹, Amulya Thota¹, Ashok Kumar B. Kumarappa² and Githa Kishore¹

¹Department of Pain Management, Pharmacy Practice, Visveswarapura Institution of Pharmaceutical Sciences, Bangalore, India; ²Kempegowda Institute of Medical Sciences, Bangalore, India

ABSTRACT

Background: The aim of management of acute low back pain is to alleviate the pain quickly and improve functional ability. Non-steroidal anti-inflammatory drugs are the first line of treatment. The challenge lies in deciding which NSAIDs will provide greater symptomatic relief, while also being cost-effective.

Objective: To compare the effectiveness of aceclofenac and etoricoxib in the management of acute low back pain.

Methods: This prospective, open label, observational study was conducted at a tertiary care hospital. Patients over 18 years of age and presenting with low back pain of less than 6 weeks duration were enrolled in the study. Fifty patients with non-specific low back pain were randomized into two groups: Group A received aceclofenac (2 mg/kg) twice a day and Group B received etoricoxib (1 mg/kg) twice a day for 1 week. The Numerical Rating Scale (NRS) and Oswestry Low Back Disability Index (ODI) determined the clinically meaningful outcomes.

Results: The decrease in pain intensity in Group A was 52.27%, while in Group B it was 62.53%. However, the decrease in pain scores between the groups was not statistically significant (p = 0.3795). Improvement in functional ability in Group A and Group B was 57.01% and 61.48%, respectively. However, this improvement between the groups was not statistically significant (p > 0.999) at the end of 1 week. The average cost-effectiveness ratio indicated that etoricoxib was the dominant treatment over aceclofenac. Therefore, etoricoxib was found to be the cost-effective option for short-term pain relief in acute low back pain for 1 week.

Conclusion: Both aceclofenac and etoricoxib were clinically effective in reducing the pain intensity and in improving functional ability. However, etoricoxib was found to be the cost-effective intervention.

ARTICLE HISTORY
Received 10 June 2019
Accepted 7 February 2020

KEYWORDS
Acute low back pain; aceclofenac; etoricoxib; cost-effectiveness

Introduction

Low back pain (LBP) is defined as pain and discomfort localized below the costal margin and above the inferior gluteal folds, with or without radiating to the legs (sciatica). LBP is a symptomatic and a self-limiting condition. It includes pain, muscle spasm, or stiffness. Acute low back pain is defined as an episode which persists for less than 6 weeks.

Being a common musculoskeletal condition, the point prevalence among the world population ranges between 60–80%. The risk of LBP increases above 35 years of age, with prevalence being more among females. People involved in jobs requiring prolonged sitting and standing or handling of heavy loads are at greater risk. The etiology of LBP can vary from mechanical, systemic and non-specific causes. About 90% of patients are diagnosed with non-specific pain, which is defined as “low back pain not attributed to known, recognizable and specific pathology”. It is possibly from a sustained muscle spasm.

The aim of treatment of acute low back pain is to obviate the pain in the shortest duration and improve the functional ability. Non-steroidal anti-inflammatory drugs are the first choice of treatment, since they reduce pain, improve functional ability, and have an acceptable tolerability profile.

Aceclofenac is a preferential COX-2 inhibitor with anti-inflammatory and analgesic properties. Aceclofenac also targets the synthesis of glycosaminoglycan and mediates chondroprotective effects. It presents with more gastrointestinal side-effects like dyspepsia, abdominal pain, and nausea. Etoricoxib is a COX-2 selective inhibitor with anti-inflammatory, analgesic properties and potential antineoplastic properties. It presents with lesser incidence of gastrointestinal side-effects, but increased cardiovascular adverse events.
Along with efficacy, the cost of the drug plays a vital role in ensuring adherence to a therapy. Cost-effectiveness analysis (CEA) is a pharmacoeconomic method for assessing the health gains relative to the costs of different health interventions. It directly relates the financial and scientific implications of different interventions. This method helps to assess whether the additional cost paid is worth the additional benefit.

There is a paucity of cost-effectiveness analysis studies comparing NSAIDs in the management of acute low back pain. The objective of the study was to evaluate the effectiveness and determine the more cost-effective intervention between Aceclofenac and Etoricoxib in the management of acute non-specific low back pain. Data on the cost-effectiveness of drugs for acute low back pain will be invaluable to healthcare professionals for better informed decision-making when choosing treatments.

Materials and methods

Study design

A prospective, open label, comparative observational study was conducted in the Department of Pain Management and in the Department of Orthopedics of a tertiary care hospital, Bangalore on patients with acute low back pain prescribed with Aceclofenac (2 mg/kg BD) and Etoricoxib (1 mg/kg BD). The study was conducted in accordance to the permission granted by the institutional ethical committee [IR No: VIPS/IEC/2017-04].

Sample size calculation

We assumed a standard deviation change in the Oswestry Disability Index of 9.9 and a minimum clinically important difference on the Oswestry Disability Index of 8.0.

At 5% level of significance and 80% power of test, $\beta$ of 0.2, the sample size was calculated as 24 patients per group. An additional 10% was added to compensate for the patients lost to follow-up. Hence, the sample size was calculated to be 30 patients in each group.

Study criteria

Outpatients between 18–80 years of age presenting with low back pain of duration less than 6 weeks and prescribed with Aceclofenac (2 mg/kg BD) or Etoricoxib (1 mg/kg BD) were included in the study.

Patients with back pain caused by malignancy and/or infection, fractures, non-compliant patients, patients with renal and/or hepatic impairment, patients with rheumatological problems, patients with disc herniation, patients with cardiovascular disorders, patients on antidepressants and anticoagulants, and pregnant and lactating women were excluded.

Study procedure

A total of 60 patients participated in the study. All the patients were informed about the purpose and requirements of the study and details of the drugs. Written informed consent was obtained from the patients prior to their enrollment in the study. Details of the patient's demographic profiles, medication history, socio-economic status, and social history were recorded on a specially designed form.

Interventions

Patients prescribed with Aceclofenac (2 mg/kg BD) were assigned to Group A. Patients prescribed with Etoricoxib (1 mg/kg BD) were assigned to Group B by the physician. The patients were followed up for a period of 1 week.

Outcome measures

The primary outcome in this study was improvement in pain and functional disability. The severity of acute low back pain and the efficacy of the drugs in reducing the pain were assessed using the Numerical rating Scale. The Numerical rating Scale (NRS) is a segmented 11-point numeric scale, with 0 representing “no pain” and 10 representing “worst pain imaginable.”

Patients are required to self-report the pain intensity. Hence, to facilitate this, the Wong-Baker Faces Pain Rating Scale was used in our study as an aid so that the patient can report their pain intensity with ease by looking at the visual representation of various intensity of pain. The Oswestry Disability Index (ODI) questionnaire, which is considered as the gold standard to assess the degree of disability in acute low back pain, was used in our study to assess the functional disability.

The baseline NRS scores and ODI scores were recorded at the start of the study. After a follow-up period of 1 week, the scores were again recorded to analyze any clinically significant change in pain intensity and functional disability.

The average cost-effectiveness ratio (ACER) is the ratio of the cost to benefit of an intervention. ACER estimates average cost spent per effect. The analysis included the direct costs incurred by patients for drug acquisition, consultation costs, cost involved in the treatments of adverse events, and cost of co-prescribed drugs. Costs incurred were estimated for a period of 1 week.

Statistical analysis

The Shapiro-Wilk test and Jarque–Bera test were used to assess the normal distribution of the data. Normality tests were performed using XLSTAT package version 2018.2. Suitable parametric (like t-test) and non-parametric tests (like Wilcoxon sign ranked test and Mann Whitney U-test) were carried out for analyzing the data at 5% level of significance ($p < .05$) using Graphpad Prism software 7.04.
Results

Socio-demographic details

During the 6-month study period, 60 patients were enrolled in the study (Figure 1). Two patients did not give consent, three patients were not eligible, and five patients were lost to follow-up.

The sociodemographic details of the patients are detailed in Table 1.

The complaint of acute low back pain is common in patients in the age group of 40–50 years (36%). Acute LBP is more prevalent among females (62%) when compared to males (38%).

The majority (52%) of the patients were homemakers. Of the study subjects, 88% were found to work for 8–12 h a day; 6% consumed alcohol (60–100 ml per day) and 8% were smokers. The mean BMI among females was around 26.6 and among males was around 29.4.

| Gender, n (%)   | Age (mean ± SD) years | Working hours, n (%) |
|----------------|----------------------|---------------------|
| Female         | 45.04 ± 4.24         | 4–8 h               |
| Male           | 39.68 ± 19.09        | 8–12 h              |
|                |                      | 12–16 h             |

Primary outcomes

Pain intensity—Numerical Rating Scale

In Group A, there was about a 52.27% reduction in pain intensity post-treatment with Aceclofenac. The average decrease in pain score was found to be 3.36, which was statistically significant at $p < .0001$ (Wilcoxon signed ranked test) (Table 2). In Group B, there was an about 62.53% decrease in the pain intensity. The average decrease in pain score was found to be 2.32, which was statistically significant at $p < .0001$ (Wilcoxon signed ranked test). However, the decrease in pain intensity between the two groups treated with Aceclofenac and Etoricoxib was not statistically significant at $p = .3795$ (Mann Whitney U-test).

Functional disability—ODI

In patients prescribed with Aceclofenac (Group A), the functional ability was improved around 57.01%. There was an average decrease in ODI score by 15.08, which was statistically significant at $p < .0001$ (Paired t-test) (Table 3). In patients prescribed with Etoricoxib (Group B), there was an about 61.48% reduction in ODI scores. There was an average decrease in ODI score by 18.24, which was statistically significant at $p < .0001$ (paired t-test). Though clinically significant, the improvement in functional disability between the two groups treated with Aceclofenac and Etoricoxib was not statistically significant ($p > .999$) (unpaired t-test).

Apart from the drugs under study, the participants were co-prescribed with other drugs like proton pump inhibitors, neuroprotectants, muscle relaxants and drugs for their co-morbid conditions which is been presented in Table 4.

Cost-effectiveness analysis

Cost-effectiveness analysis identifies the intervention which has the potential to yield the greatest improvement in health

Table 2. Pre- and post-treatment numerical rating scale scores in both groups.

| Parameters       | Group A (Aceclofenac) | Group B (Etoricoxib) |
|------------------|-----------------------|----------------------|
| Mean ± SD        | 4.52 ± 0.82           | 5.44 ± 1.41          |
| Median           | 4                     | 5                    |
| $p$-value        | <.0001                | <.0001               |
| Average decrease in pain score | 2.32 (52.27%) | 3.36 (62.53%) |

Table 3. Pre- and post-treatment Oswestry Disability Index scores in both groups.

| Parameters       | Group A (Aceclofenac) | Group B (Etoricoxib) |
|------------------|-----------------------|----------------------|
| Mean ± SD        | 26.96 ± 5.69          | 29.84 ± 5.35         |
| $p$-value        | <.0001                | <.0001               |
| Average decrease in ODI score | 15.08 (57.01%) | 18.24 (61.48%) |
for the least resources. The costs incurred for the drugs including co-prescribed drugs, diagnostic methods used and physician consultation costs were estimated. The cost of the drugs were obtained from CIMS, January–April 2018.

The average decrease in NRS and ODI scores were used as primary outcomes.

**Average cost-effectiveness ratio (ACER)**

The average cost-effectiveness ratio (ACER) is the ratio of the cost to benefit of an intervention. There was a greater decrease in the pain intensity and better functional ability in patients receiving Etoricoxib when compared with patients receiving Aceclofenac.

Upon calculation it was found that the ACER of Etoricoxib was less when compared to Aceclofenac, indicating Etoricoxib is the cost-effective intervention. Hence, it is evident that Etoricoxib is the dominant treatment over Aceclofenac for a duration of 1 week, making it the cost-effective option for short-term pain relief in acute low back pain.

**Discussion**

Low back pain is a common self-limiting musculoskeletal condition, mostly with a non-specific etiology which presents with pain, muscle tension, and stiffness. In our study population, acute low back pain was more common in patients in the age group of 40–50 years. The mean age in all the study subjects was found to be 41.67 years.

In a study conducted by Gupta et al.\textsuperscript{17} it was found that low back pain was common in the third and fourth decades of life. The average age of patients was found to be 38.39 years.

In our study, 62% of the total study subjects were females. The female preponderance can be attributed to Spinal osteoarthritis, joint degeneration, psychological factors, female hormone fluctuation, and menstrual history.

Around 52% of the total subjects were homemakers and 24% were professionals. A similar epidemiological study conducted by Nazeer et al.\textsuperscript{18} reported that housewives formed the majority of cases (66%).

The various reasons attributed could be the unduly working hours, working posture, and physical exhaustion. The majority of the patients reported to be working for 8–12 h a day.

The majority (70%) reported to have a sedentary lifestyle. In our study about 80% of the total study subjects were literate (attended more than primary education).

The mainstay of the management for acute low back pain is to alleviate the pain in the shortest duration with least side-effects. Analgesics are the first line of drugs, since they provide symptomatic relief and have an acceptable tolerability profile.

The challenge lies in selecting the most effective, safest, and cost-effective analgesic.

Aceclofenac, a preferential COX-2 inhibitor and Etoricoxib, a selective COX-2 inhibitor both have analgesic, anti-inflammatory effects. However they differ in their adverse effect profile.

Aceclofenac is associated with more GI harm, while Etoricoxib is associated with less GI adverse effects and more cardiovascular adverse effects\textsuperscript{19}. The information about their adverse effect profile is based on the literature search.

Based on our study results, Etoricoxib demonstrated a greater reduction in pain (62%) when compared with Aceclofenac (52%). Also the improvement in the functional disability was more in Etoricoxib (61%) when compared with Aceclofenac (57%). None of the study participants discontinued the therapy and no adverse effects were reported.

Proton pump inhibitors were co-prescribed with NSAIDs to prevent GI discomfort.

28% in group A and 20% in group B were prescribed with Pantoprazole.

Proteolytic enzymes play a key role in reducing inflammation by causing the lysis of the peptide bonds\textsuperscript{20}. Enzymes like serratiopeptidase, trypsin were co-prescribed in the present study. About 44% of the study population were given proteolytic enzymes.

Low back pain is generally associated with muscle spasms. Hence, muscle relaxants are frequently co-prescribed with NSAIDS.

---

**Table 4. Drugs co-prescribed with the interventional drugs.**

| Drugs administered | Total no. of patients | Percentage |
|--------------------|-----------------------|------------|
| GROUP A (n = 25)   |                       |            |
| Aceclofenac + Muscle Relaxant (Combination) | 22 | 44% |
| Aceclofenac + Paracetamol (Combination) | 1 | 2% |
| Aceclofenac + Paracetamol + serratiopeptidase | 2 | 4% |
| GROUP B (n = 25)   |                       |            |
| Etoricoxib + Muscle Relaxant (Combination) | 23 | 46% |
| Etoricoxib 90mg alone | 2 | 4% |

**Drugs co-prescribed in both the groups**

| Name of the drugs | Group A | Group B |
|-------------------|---------|---------|
|                   | No. of patients | Percentage | No. of patients | Percentage |
| Pantoprazole       | 7 | 28% | 5 | 20% |
| Gabapentine        | 9 | 36% | 8 | 32% |
| Proteolytic enzymes | 6 | 24% | 15 | 60% |
| Vitamin B12 + Pregabalin | 3 | 12% | 4 | 16% |
| Inj. Vitamin B12  | 1 | 4% | 1 | 4% |
In our study, Thiocolchicoside (4 mg) was the most commonly prescribed muscle relaxant. It is a GABA agonist and acts on the muscular contracture by activating the GABA-ergic inhibitory pathways, thereby acting as a potent muscle relaxant\(^\text{21}\). In total, 84% of the total patients enrolled in the study were co-administered with muscle relaxant (Table 4).

Neuromodulators are also given to treat neuropathic pain. Methylcobalamin and Pregabalin, Gabapentin were co-prescribed along with NSAIDs in patients with neurological deficits. Methylcobalamin has an important role in the regeneration of myelin sheath and helps to restore the function of the nerve in neuropathy\(^\text{22}\).

Gabapentin and Pregabalin have a high affinity for the auxiliary \(\alpha_{2}\beta\) sub-units of the voltage-gated calcium channel and thus blocks Ca\(^{2+}\) influx into nerve terminals, which leads to reduced transmitter release\(^\text{23}\). Of the patients with LBP, 34% were prescribed with Gabapentin alone; 14% of the subjects were prescribed with Methylcobalamin and Pregabalin as a combination.

Cost-effectiveness analysis highlights the interventions that are relatively inexpensive, yet have the potential to reduce the disease burden substantially. Costs are measured in a common monetary value and the effectiveness in terms of physical units. The average cost-effectiveness ratio calculated for 1 week indicated Etoricoxib to be dominant over Aceclofenac, indicating Etoricoxib is the cost-effective intervention.

Our study is in agreement with a study conducted in Norway by Jansen et al.\(^\text{24}\) where they evaluated the cost-effectiveness of Etoricoxib versus Celecoxib and non-selective NSAIDs in the treatment of ankylosing spondylitis.

Their economic evaluation suggested that Etoricoxib was the most cost-effective initial NSAID treatment for Ankylosing Spondylitis patients since there was a > 98% probability that treatment with Etoricoxib resulted in greater Quality Adjusted Life Years than the other interventions\(^\text{24}\).

There are certain limitations in our study. The above study was a single center, open label study. Also the shorter duration of the study period and the smaller size is another limitation.

Low back pain is a common complaint and patients prefer speedy recovery. Also there are a number of NSAIDs available to reduce pain. Therefore, further studies need to be carried on large populations and at different centers to extrapolate the findings of the safety and efficacy of NSAIDs.

**Conclusion**

According to the results of the present prospective observational study, both Etoricoxib and Aceclofenac are equally effective in reducing the pain intensity and improving the functional ability in acute low back pain. However, cost-effectiveness analysis indicated Etoricoxib to be a more cost-effective intervention when compared with Aceclofenac. Hence, both Etoricoxib and Aceclofenac are effective analgesics in acute low back pain, nonetheless Etoricoxib was estimated to be a cost-effective intervention.

**Transparency**

**Declaration of funding**

There is no funding to disclose.

**Declaration of financial/other relationships**

The authors and peer reviewers on this manuscript have no relevant financial or other relationships to disclose.

**Acknowledgements**

We express our gratitude to Dr. Shivananda S., Professor and Head of the Department of Orthopedics KIMS Hospital for his support. We extend our thankfulness to Dr. G. Y. Narmada, Honorable Principal, for her support.

**References**

[1] Verhagen AP, Downie A, Popal N, et al. Red flags presented in current low back pain guidelines: a review. Eur Spine J. 2016; 25(9):2788–2802.

[2] Soumya RV, Babu L, Shafi K M, et al. A prospective comparative study on the efficacy of aceclofenac and diclofenac in low back pain. EJPMR. 2018;5(1):200–203.

[3] Bhattacharj S, Paudel Chhetri H, Alam K, et al. A study on factors affecting low back pain and safety and efficacy of NSAIDs in acute low back pain in a tertiary care hospital of western Nepal. JCDR. 2013;7(12):2752–2758.

[4] Bindra S, Sinha AGK, Benjamin AI. Epidemiology of low back pain in Indian population: a review. Int J Basic Appl Med Sci 2015;5(1):166–179.

[5] Shah JM, Patel K, Shah MR, et al. Comparison of efficacy and safety of thiocolchicoside and pregabalin in the treatment of acute non-specific low back pain: an open label randomized prospective study. Int J Basic Clin Pharmacol. 2016;5(5):1733–1738.

[6] Kumar S, Rani S, Siwach R, et al. To compare the efficacy and safety of fixed dose combination of thiocolchicoside and aceclofenac versus chloroxazone, aceclofenac and paracetamol in patients with acute lower backache associated with muscle spasm. Int J App Basic Med Res. 2014;4:101–105.

[7] Roloefs PD, Deyo RA, Koes BW, et al. Nonsteroidal anti-inflammatory drugs for low back pain. Spine. 2008;33(16):1766–1774.

[8] Tripathi KD. Essentials of medical pharmacology. 7th ed. New Delhi: Jaypee Brothers Medical Publishers (P) LTD; 2003.

[9] National Center for Biotechnology Information. PubChem Compound Database; CID = 71771. [accessed 6 Feb 2018]. Available from: https://pubchem.ncbi.nlm.nih.gov/compound/71771.

[10] National Center for Biotechnology Information. PubChem Compound Database; CID = 123619. [accessed 6 Feb 2018]. Available from: https://pubchem.ncbi.nlm.nih.gov/compound/123619.

[11] Jamison DT, Breman JG, Measham AR, editors, et al. Priorities in health. Washington (DC): The International Bank for Reconstruction and Development/the World Bank; 2006. Chapter 3, Cost-effectiveness analysis. [accessed 15 march 2018]. Available from: https://www.ncbi.nlm.nih.gov/books/NBK10253/.

[12] Brzeziński K, Wordliczek J. Comparison of the efficacy of dexketoprofen and diclofenac in treatment of non-specific low back pain. Ann Agric Environ Med. 2013;1:52–56.

[13] Michener LA, Snyder AR, Leggin BG. Responsiveness of the Numeric Pain Rating Scale in patients with shoulder pain and the effect of surgical status. J Sport Rehabil. 2011;20(1):115–128.
[14] Kumar P, Tripathi L. Challenges in pain assessment: pain intensity scales. Indian J Pain. 2014;28:61–70.

[15] Mehra A, Baker D, Disney S, et al. Oswestry Disability Index Scoring made easy. Ann R Coll Surg Engl. 2008;90(6):497–499.

[16] Bang H, Zhao H. Average cost-effectiveness ratio with censored data. J Biopharm Stat. 2012;22(2):401–415.

[17] Gupta R, Mahajan S, Dewan D, et al. An epidemiological study of low back pain in a tertiary care hospital of Jammu, Jammu and Kashmir, India. Int J Res Med Sci. 2017;5(3):835–839.

[18] Nazeer M, Rao SM, Soni S, et al. Low back pain in South Indians: causative factors and preventive measures. Sch J App Med Sci. 2015;3(1D):234–243.

[19] Walker C. Are all oral COX-2 selective inhibitors the same? A consideration of celecoxib, etoricoxib, and diclofenac. Int J Rheum. 2018;2018:1–12.

[20] Viswanatha Swamy AHM, Patil PA. Effect of some clinically used proteolytic enzymes on inflammation in rats. Indian J Pharm Sci. 2008;70(1):114–117.

[21] Umekar AR, Bavaskar SR, Yewale PN. Thiocolchicoside as muscle relaxant: a review. Int J Pharm Bio Sci. 2011;1(3):364–371.

[22] Onkar YU, Swami DC. Sustained-release pregabalin with methylcobalamin in neuropathic pain: an Indian real-life experience. Int J Gen Med. 2013;6:413–417.

[23] Taguchigarashi T, Watt AS, et al. Effectiveness of pregabalin for the treatment of chronic low back pain with accompanying lower limb pain (neuropathic component): a non-interventional study in Japan. J Pain Res. 2015;8:487–497.

[24] Jansen JP, Gaugris S, Choy EH, et al. Cost effectiveness of etoricoxib versus celecoxib and non-selective NSAIDS in the treatment of ankylosing spondylitis. Pharmacoeconomics. 2010;28(4):323–344.