Parecoxib prevents early postoperative cognitive dysfunction in elderly patients undergoing total knee arthroplasty

A double-blind, randomized clinical consort study

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Abstract

Background: Trial design neuroinflammation and postoperative pain after surgery are increasingly reported in association with postoperative cognitive dysfunction (POCD). Parecoxib, a selective cyclooxygenase (COX)-2 inhibitor, is used for postoperative analgesia for its potent anti-inflammatory and analgesic effects. This study aimed to evaluate parecoxib’s effects on POCD in elderly patients undergoing total knee arthroplasty.

Methods: Around 134 elderly patients undergoing total knee arthroplasty were randomly divided into parecoxib (group P) and control (group C) groups, and treated with parecoxib sodium and saline, respectively, shortly after induction of general anesthesia and 12-h post-surgery, respectively. Perioperative plasma IL-1β, IL-6, TNF-α, and C-reactive protein (CRP) levels were measured. Postoperative pain was assessed following surgery. Neuropsychological tests were performed before surgery, and 1 week and 3 months postoperation.

Results: POCD incidence in group P was significantly lower compared with that of group C at 1 week after surgery (16.7% vs 33.9%; P < 0.05); no significant difference was found between groups C and P at 3-month follow-up (9.7% vs 6.7%). Compared with group C values, visual analog pain scale (VAS) scores at 3, 6, and 12 hours after surgery were significantly lower in group P (P < 0.05). Plasma IL-1β, IL-6, and TNF-α levels were lower in group P than in group C after the operation (P < 0.05). No significant difference in the plasma CRP level was found between groups P and C.

Conclusions: Parecoxib sodium decreases POCD incidence after total knee arthroplasty in elderly patients and may explain how this drug suppresses inflammation and acute postoperative pain caused by surgical trauma.

Abbreviations: ASA = American Society of Anesthesiologists, BMI = body mass index, COX = cyclooxygenase, CRP = C-reactive protein, MMSE = Mini-Mental State Examination, POCD = postoperative cognitive dysfunction, VAS = visual analog pain scale.

Keywords: inflammation, parecoxib, postoperative cognitive dysfunction, postoperative pain

1. Introduction

Postoperative cognitive dysfunction (POCD) is a common complication after major surgery in the elderly, and associated with prolonged hospitalization, inability to cope independently, premature unemployment, and possible permanent dementia. Although POCD mechanisms remain elusive, mounting evidence has revealed that inflammation plays a key role in the disease process. Peripheral inflammation due to surgical trauma and the accompanying release of systemic inflammatory mediators have been shown to influence inflammatory processes in the central nervous system. Animal studies indicated that proinflammatory cytokines, such as interleukin 1β (IL-1β) and tumor necrosis factor-α (TNF-α), play a pivotal role in mediating surgery-induced neuroinflammation. Indeed, increased expression of proinflammatory cytokines results in performance deficits of hippocampus-dependent cognitive memory. It is well known that inflammation triggered by surgical trauma is also implicated in postoperative pain. Recent studies showed postoperative pain contributes to cognitive dysfunction.

Selective cyclooxygenase (COX)-2 inhibitor drugs are increasingly used as analgesics in postoperative analgesia. Such drugs attenuate inflammation by decreasing prostaglandin formation via inhibition of COX-2 activity in both peripheral and central tissues. Given their anti-inflammatory and analgesic activities, COX-2 inhibitors might potentially prevent the development of POCD. Animal studies have suggested COX-2 inhibitors to be promising candidates for the treatment of neuroinflammation and cognitive decline caused by surgical trauma. However, whether administration of COX-2 inhibitors in the perioperative period can decrease POCD incidence in clinical practice remains unclear.
Therefore, we assessed the effect of the COX-2 inhibitor parecoxib on POCD incidence in aged patients following total knee arthroplasty with general anesthesia; in addition, parecoxib’s effects on proinflammatory cytokine expression and visual analog pain scale (VAS) scores in these patients were investigated.

2. Methods

2.1. Patients and allocation

This was a double-blind, randomized clinical consort study. It was approved by the Ethics Committee of Xuzhou Central Hospital. Written informed consent was obtained from all patients. The sample size of the study was calculated using a free software (http://www.statpages.org) based on a pilot study. Eligible subjects were ASA I or II patients between 65 and 80 years of age, scheduled for total knee arthroplasty. Exclusion criteria were: ASA > II; peptic ulcer disease, cardiac-cerebral vascular disease, diabetes mellitus; neurological or psychiatric disorders; history of allergic reactions to NSAIDs; history of drug and alcohol abuse; hepatic and/or kidney dysfunction; BMI > 35; patients on analgesics or antidepressants; Mini-Mental State Examination (MMSE) score < 23; inability to comply with the study protocol or procedures. Using computer-generated randomized table, patients meeting the eligibility criteria were randomized to receive an intravenous bolus of parecoxib sodium 40 mg or placebo (normal saline), shortly after induction of general anesthesia and 12 hours postsurgery, respectively. Patients, anesthetists, and investigators were blinded for treatment allocation.

2.2. Anaesthesia and postoperative treatment

All participants received general anesthesia according to a standardized protocol. Anesthesia was induced with 0.1 mg/kg midazolam, 0.2 mg/kg cisatracurium, 2 mg/kg propofol, and 0.6 µg/kg sufentanil, and maintained with remifentanil and propofol. Bispectral Index Score was maintained at 40–60 by adjusting the propofol infusion rate. Heart rate, arterial pressure, respiratory rate, PETCO2 and SpO2 were recorded continuously. Patients revived spontaneously without administration of any anesthetic antagonists. Fentanyl was used for patient-controlled analgesia (PCA) and tropisetron was administered for nausea and vomiting.

2.3. Neuropsychological tests

Neuropsychological tests were administered before surgery, and at 1 week and 3 months after surgery. An experienced neurologist, blinded to treatment group assignments, carried out the neuropsychological tests at both times in tranquil surroundings. The test battery, which included 7 tests with 9 subscales, was designed to measure memory, attention and concentration, and psychomotor skills. The tests included: the Mental Control and Digit Span (forward and backward) subtests of the Wechsler Memory Scale (attention and concentration), Visual Retention and Paired Associate Verbal Learning subtests of the Wechsler Memory Scale (figural memory and verbal learning/memory), Digit Symbol subtest of the Wechsler Adult Intelligence Scale-Revised (psychomotor speed), Halstead-Reitan Trail Making Test (Part A) (hand-eye coordination, attention, and concentration), and Grooved Pegboard Test (favored and unfavored hand) (manual dexterity). A decline of 1 or more SDs in 2 or more tests was considered to reflect POCD.\[13\]

3. Results

3.1. Patient characteristic

From February 2014 to May 2015, 134 patients were included in the trial. The flow of patients through the study and detailed reasons for exclusion are provided in Fig. 1. A total of 5 and 7 patients were lost to follow-up at 1-week follow-up in groups C (control) and P (parecoxib treatment), respectively. In addition, 17 and 13 patients were lost to follow-up at 3-month follow-up in control and parecoxib groups, respectively. The basic demographic and clinical characteristics of patients in both groups are presented in Table 1. A significant difference was obtained in smoker number between both groups (P < 0.05), but the other characteristics were similar.

3.2. Neuropsychological test results and incidence of POCD

There were significant differences in values obtained for mental control, Digit symbol, and Pegboard favored hand between both groups. The control group showed a statistically significant downward trend in values obtained for the most neuropsychological tests compared with the parecoxib group (Table 2). Importantly, the parecoxib group showed lower POCD incidence compared with controls at 1-week follow-up (16.7% vs 33.9%, P < 0.05). However, no statistically significant difference in POCD incidence between both groups was observed at 3-month follow-up (Table 3).

3.3. Postoperative pain and plasma levels of pro-inflammatory cytokines

Significantly lower VAS scores were found at 3, 6, and 12 hours in group P, compared to group C values (Table 4). ELISA data showed that plasma levels of IL-1β, IL-6, TNF-α, and CRP were higher after surgery compared with baseline levels in both groups.
Figure 1. Consort diagram of patients’ randomization, intervention, and analysis.

Table 1
Demographic and clinical characteristics.

| Admission characteristics | Group C (n=62) | Group P (n=60) | P   |
|---------------------------|---------------|---------------|-----|
| Age, y                    | 74.3±7.6      | 75.1±8.2      | 0.56|
| BMI, kg/m²                | 27.3±5.2      | 27.8±4.9      | 0.58|
| Education, y              | 9.3±2.4       | 8.9±3.2       | 0.47|
| Hypertension              | 22 (35.5%)    | 24 (40%)      | 0.75|
| Diabetes mellitus         | 18 (29%)      | 14 (23.3%)    | 0.83|
| COPD                      | 7 (11.3%)     | 8 (13.3%)     | 0.88|
| Smoker                    | 13 (20.9%)    | 6 (10%)       | 0.04|
| MMSE scores               | 27.9±2.1      | 27.5±2.4      | 0.61|
| Length of surgery, min    | 68±7          | 71±10         | 0.48|
| Estimated blood loss, mL  | 212±35        | 230±44        | 0.36|
| Preoperative hospital stay, days | 2.8±0.4 | 2.7±0.5 | 0.55|
| Postoperative hospital stay, days | 6.6±1.2 | 7.1±1.0 | 0.41|
| Postoperative nausea, vomiting | 6 (9.7%) | 5 (8.3%) | 0.81|
| Postoperative itchiness   | 6 (9.7%)      | 4 (6.7%)      | 0.72|

Values are mean±SD or number (percentages).
BMI = body mass index, COPD = chronic obstructive pulmonary disease, MMSE = Mini-Mental State Examination.
Table 2
Neuropsychological test results at baseline, 7 days, and 90 days follow-up in patients.

| Variables                        | Group | Baseline   | After operation 7th day | After operation 90th day | F Statistic |
|----------------------------------|-------|------------|-------------------------|--------------------------|-------------|
|                                  |       |            |                         |                          |             |
| Mental control                   | C     | 82.4±13.5  | 71.4±10.5               | 76.8±14.3                | 0.54        |
|                                  | P     | 81.8±15.1  | 79.5±8.2                | 81.1±11.3                | 0.48        |
| Visional rational                | C     | 9.5±3.3    | 8.7±2.6                 | 9.4±3.8                  | 0.36        |
|                                  | P     | 9.8±4.0    | 9.6±3.4                 | 9.7±4.1                  | 0.23        |
| Paired associate verbal learning | C     | 17.8±2.3   | 15.8±3.1                | 17.2±3.2                 | 0.18        |
|                                  | P     | 17.2±2.9   | 16.4±3.3                | 17.4±3.8                 | 0.19        |
| Digit span forward               | C     | 7.2±1.3    | 7.0±1.1                 | 7.3±1.4                  | 0.23        |
|                                  | P     | 7.5±1.1    | 7.1±1.2                 | 7.4±1.6                  | 0.23        |
| Digit span backward              | C     | 4.2±1.4    | 4.1±1.9                 | 4.1±1.1                  | 0.23        |
|                                  | P     | 4.2±1.1    | 4.1±1.4                 | 4.2±1.8                  | 0.23        |
| Digit symbol                     | C     | 28.2±10.3  | 17.3±11.3               | 25.2±18.4                | 0.59        |
|                                  | P     | 27.7±9.6   | 26.2±13.4               | 26.9±15.4                | 0.74        |
| Trails A                         | C     | 135.8±44.6 | 116.7±48.2              | 131.8±45.2               | 0.74        |
|                                  | P     | 137.6±38.4 | 128.4±45.2              | 135.8±46.8               | 0.74        |
| Pegboard favored hand            | C     | 83.6±12.7  | 70.1±12.2               | 75.5±14.2                | 0.76        |
|                                  | P     | 84.5±11.4  | 82.5±10.1               | 83.8±11.1                | 0.76        |
| Pegboard unfavored hand          | C     | 85.5±13.4  | 80.1±14.8               | 83.2±14.7                | 1.17        |
|                                  | P     | 86.3±15.2  | 85.2±10.1               | 85.8±16.1                | 1.17        |

**Table 3**
Patients with postoperative neuropsychological deficit in the test battery at baseline, 7 days, and 90 days follow-up.

| Number of deficits | After operation 7th day | After operation 90th day |
|--------------------|-------------------------|--------------------------|
|                    | Group C | Group P | Group C | Group P |
| 1                  | 20      | 13      | 10      | 8       |
| 2                  | 16      | 6       | 5       | 3       |
| 3                  | 4       | 0       | 1       | 1       |
| 4                  | 1       | 0       | 0       | 0       |
| ≥5                 | 0       | 0       | 0       | 0       |
| **POCD patients**  | 22 (33.9%) | 10 (16.7%) | 6 (9.7%) | 4 (6.7%) |

POCD = postoperative cognitive dysfunction.

* Statistically significant, *P* < 0.05.

relish and astrocyte activation are associated with declined cognitive performance in humans and animals. Indeed, proinflammatory cytokines, such as TNF-α, IL-1β, and IL-6, can be released by activated astrocytes, triggering neuroinflammation and leading to cognitive dysfunction. High concentrations of proinflammatory cytokines inhibit long-term potentiation and impair memory. Neuroinflammation has been implicated in cognitive impairment; this may provide a viable target to prevent the development of POCD. Animal studies suggested COX-2 inhibitors to be promising candidates for the treatment of neuroinflammation and cognitive decline caused by surgical trauma. In the current study, the plasma levels of proinflammatory cytokines were significantly elevated after surgery in both parecoxib and control groups, but the degree of increase in the parecoxib group was markedly lower compared with that of controls. We speculate that the prophylactic effect of parecoxib on POCD likely results from its anti-inflammatory activity. However, more studies are needed to further understand parecoxib’s effects.

Postoperative acute pain is another potential risk factor for cognitive dysfunction. Animal studies demonstrated that postoperative acute- or inflammatory pain could exacerbate memory deficits. Available evidence indicates that ketoprofen, a nonsteroidal anti-inflammatory drug (NSAID), can prevent the

Table 4
Postoperative pain.

| Time-point of the VAS assessment | Group C | Group P | *P*
|----------------------------------|---------|---------|------|
| 1 h                              | 2.2±1.3 | 2.1±1.5 | 0.256|
| 3 h                              | 3.7±1.1 | 2.6±1.0 | 0.006|
| 6 h                              | 3.4±1.2 | 2.5±0.8 | 0.022|
| 12 h                             | 3.0±0.9 | 2.4±1.1 | 0.037|
| 24 h                             | 2.1±1.2 | 1.8±0.8 | 0.124|
| 48 h                             | 1.1±0.6 | 1.0±0.8 | 0.304|

Data presented as mean ± SD. VAS = visual analog scale.
development of surgery-associated cognitive dysfunction via its pain-relieving effects in aged rats. Parecoxib sodium is frequently used for postoperative analgesia because of less side effects. In the current study, parecoxib decreased postoperative pain scores in patients after surgery. Furthermore, parecoxib reduced POCD incidence in aged patients. Therefore, the effectiveness of parecoxib in reducing POCD incidence might also be explained by its analgesic activity, which alleviates cognitive deficits.

There are several limitations of this study that should be mentioned. First, sample size was relatively small to fully demonstrate the effectiveness of parecoxib. No statistically significant difference was found in POCD incidence between both groups at 3-month follow-up, which may result from the reduced sample size. In addition, POCD is a common neurological complication of surgery; therefore, cerebrospinal fluid samples may be more suitable than plasma specimens for assessing proinflammatory cytokines levels. Furthermore, despite randomization, fewer smokers were found in the control group compared with the parecoxib group. It is unknown whether smoking would affect such studies, but it was recently shown that nicotine, the main specific alkaloid in tobacco smoke, does not alter cognitive function. Ongoing work in our laboratory aims at addressing these issues.

In conclusion, parecoxib decreased POCD incidence after total knee arthroplasty in elderly patients. Anti-inflammatory therapy and effective postoperative pain control with parecoxib may have benefits in preventing POCD in elderly patients. The current study provides a strong basis for parecoxib use as a prophylactic drug for POCD prevention.

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