Corticosteroid treatment reduces headache in eosinophilic meningitis: a systematic review

Sittichai Khamsai1, Kittisak Sawanyawisuth1, Vichai Senthong1, Panita Limpawattana1, Jarin Chindaprasit1, Pewpan M. Intapan1, Wanchai Maleewong2, Somsak Tiamkao2, Verajit Chotmongkol1, Chetta Ngamjarus3

1Department of Medicine, Faculty of Medicine, Khon Kaen University, Khon Kaen - Thailand
2Department of Parasitology, Faculty of Medicine, Khon Kaen University, Khon Kaen - Thailand
3Department of Epidemiology and Biostatistics, Faculty of Public Health, Khon Kaen University, Khon Kaen - Thailand

ABSTRACT

Background: Eosinophilic meningitis (EOM) is an emerging parasitic disease that can be found worldwide, of which acute severe headache is a presenting symptom. Although such headaches may persist for up to 2 months, studies have found corticosteroid to be effective in reducing this symptom. As the most recent systematic review was published in 2015, the aim of this study was to provide a more up-to-date examination of the role of corticosteroids in EOM.

Methods: We included randomized controlled trials of corticosteroid treatment for EOM regardless of comparators. Research articles published in five databases were searched and evaluated. The primary outcome was headache, which was compared among various treatment regimens.

Results: We found a total of 257 articles after duplication removal. Of those, two met the study criteria. According to these studies, oral prednisolone alone or in a combination of albendazole resulted in fewer patients with headache after a 2-week course of treatment compared with placebo (maximum of 9.1% vs. 45.5%). The duration of headache was also shorter in the prednisolone arm vs. placebo (maximum of 5 vs. 13 days). There were no serious side effects reported.

Conclusion: A 2-week course of treatment with oral corticosteroid with or without albendazole reduced headaches in patients with EOM.

Keywords: Angiostrongylus cantonensis, Headache, Prednisolone
with placebo. It is recommended that systematic reviews be updated every 2 years to include any newly published articles (13). We thus updated and examined studies of corticosteroid treatment in EOM regardless of comparators.

Materials and methods

This study is a systematic review focused on the effects of corticosteroids in human EOM. We included randomized controlled trials of corticosteroid treatment for EOM regardless of comparators. The types of interventions were corticosteroids vs. placebo or other active treatments. The participants in the studies examined were EOM patients aged 15 years or older (12). Those who had already taken corticosteroids prior to study participation were excluded.

We searched five databases in this review: PubMed, Central database, Scopus, CINAHL Plus, and Web of Science. The search terms used were “meningit*”, “eosinophil*”, “Angiostrongylus cantonensis”, and “randomized controlled trials”. The full list of search terms are shown in Appendix 1 (See supplementary material). The final search was performed on November 22, 2019.

After duplication removal, initial screening was carried out for nonrelevant articles. Studies were considered relevant if they had been conducted to evaluate the role of corticosteroids in any form of headache caused by EOM. The full-text reports were subsequently reviewed by two independent authors (SK, KS). Of these, any randomized controlled trials were included in the final analysis. The primary outcomes were number of patients with persistent headache at 2 weeks after treatment and duration of headache by corticosteroid treatment regimen. Other outcomes included number of patients who underwent repeated lumbar puncture and side effects of treatment. We presented frequency with percentage for categorical variables. Continuous variables were described using median and range. Risk ratios (RR) with 95% confidence intervals (CI) were used to compare to compare a risk between two groups from randomized controlled trials (RCTs).

Biases of eligible studies were evaluated across six domains (sequence generation, allocation concealment, blinding of participants/personnel and outcome assessors, incomplete outcome data, selective outcome reporting, and other potential sources of bias) by two authors independently (SK, KS). Biases were categorized as low risk, high risk, or unclear according to the guidelines specified in the Cochrane Handbook for Systematic Reviews of Interventions (12). Disagreements were reviewed and reported by a third reviewer (CN).

Results

We found a total of 288 articles in the following five databases: Scopus (254 articles), PubMed (21 articles), Web of Science (9 articles), Central database (3 articles), and CINAHL Plus (1 article). Two hundred fifty-seven remained after duplication removal (Fig. 1), of which 252 were excluded as nonrelevant. In total, five articles were selected for full-text review (11,14-17). Of these, only two met the study criteria. With regard to the excluded studies, one did not evaluate outcomes or headache, and the other two were not randomized controlled trials.

The first of the remaining studies was published in 2000 and enrolled 110 EOM patients (11). This study found that treatment with prednisolone resulted in fewer patients experiencing headaches at 2 weeks after treatment and shorter duration of headache by corticosteroid treatment regimen. Other outcomes included number of patients who underwent repeated lumbar puncture and side effects of treatment. We presented frequency with percentage for categorical variables. Continuous variables were described using median and range. Risk ratios (RR) with 95% confidence intervals (CI) were used to compare to compare a risk between two groups from randomized controlled trials (RCTs).

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The first of the remaining studies was published in 2000 and enrolled 110 EOM patients (11). This study found that treatment with prednisolone resulted in fewer patients experiencing headaches at 2 weeks after treatment and shorter duration of headache when compared with placebo (9.1% vs 45.5%; RR 0.20 with 95% CI: 0.08 to 0.48 and 5 days vs. 13 days, respectively). The number of cases of repeated lumbar puncture was also significantly higher in the placebo group than in the prednisolone group (22 times vs. 7 times; RR 0.32 with 95% CI: 0.15 to 0.68), as shown in Table I.

Fig. 1 - Study flow of the systematic review on corticosteroid treatment in eosinophilic meningitis. RCT = randomized controlled trial.
The second study was published in 2009 and conducted in 104 EOM patients (17). It compared prednisolone plus albendazole with prednisolone alone and found no significant difference in terms of the primary outcome. No data on repeated lumbar puncture were reported (Tab. I).

Both studies were conducted in Thailand, used only oral prednisolone, and reported no serious side effects from either prednisolone or prednisolone plus albendazole. Allocation, concealment, and stratification were not reported in either study. The second study was not blinded (Fig. 2).

### TABLE I - Two randomized controlled trials of corticosteroid treatment in eosinophilic meningitis

| Factors                        | Study 1: 2000 | Study 2: 2009 |
|--------------------------------|---------------|---------------|
|                                | Active Control | Active Control |
| Treatment                      | Pred Placebo  | Pred + albendazole | Pred |
| No. of patients                | 55            | 53            | 51 |
| No. of patients with headache at 2 weeks | 5 (9.1%) | 25 (45.5%) | 0 (2.0%) |
| Median (range) duration of headache, days | 5 (1-60) | 13 (1-56) | 3 (1-14) |
| No. of patients with repeated LP (n, %) | 7 (12.8%) | 22 (40.0%) | NA |

LP = lumbar puncture, NA = not available, Pred: prednisolone.

### TABLE II - Previous single-arm studies of corticosteroid treatment in eosinophilic meningitis

| Factors                        | Study 1: 2004 | Study 2: 2004 | Study 3: 2006 |
|--------------------------------|---------------|---------------|---------------|
|                                | Prednisolone 1 week | Prednisolone + albendazole | Prednisolone + mebendazole |
| No. of patients                | 52            | 26            | 41            |
| No. of patients with headache at 2 weeks (n, %) | 6 (11.6%) | 3 (11.6%) | 4 (9.8%) |
| Median (range) duration of headache, days | NA | 4 (1-17) | 3 (1-20) |
| No. of patients with repeated LP (n, %) | 1 (2.0%) | 0 | 3 (7.4%) |

LP = lumbar puncture, NA = not available.

### Discussion

Although there were two eligible studies, we were unable to perform a conventional meta-analysis or network meta-analysis due to the inconsistent treatment arms and incomplete cycle, respectively. As a result, this report was defined as a systematic review.

In both studies, oral prednisolone with or without albendazole yielded favorable outcomes in terms of both numbers of EOM patients with headache and duration of headache after 2 weeks of treatments. These findings were compatible with those of other single-arm studies examining the effects of prednisolone alone vs. prednisolone plus anthelmintics (15,18,19), as shown in Table II. The corticosteroid-based regimen with albendazole and that with mebendazole yielded comparable headache durations in the two studies in this review (3-4 days vs. 3-5 days), but the 1-week prednisolone treatment had higher numbers of patients with headache at 2 weeks (11.6%). There were no serious side effects of corticosteroid found in any of the five studies examined.

There were some limitations in these studies. First, both studies were conducted in northeast Thailand, an area endemic for EOM. Second, some information was not reported in the randomized controlled trials such as concealment or blinding. These items were reported as possible biases (Fig. 2). Finally, only oral prednisolone was administered for a duration of 2 weeks.

### Conclusion

Two weeks of treatment with oral corticosteroid with or without albendazole reduced headaches in EOM patients.

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