

Systemic antibiotics after incision and drainage of simple abscesses: a meta-analysis

Adam J Singer, Henry C Thode Jr

Department of Emergency Medicine, Stony Brook University, Stony Brook, New York, USA

Correspondence to

Dr Adam J Singer, Department of Emergency Medicine, Stony Brook University, Stony Brook, NY 11794-8300, USA; adam.singer@stonybrookmedicine.edu, adam.singer@stonybrook.edu

Received 7 March 2013

Accepted 12 April 2013

Published Online First

18 May 2013

ABSTRACT

Background Over the last decade, there has been a significant increase in the number of cutaneous abscesses. While there is general agreement that abscesses should be treated with incision and drainage, it is unclear whether systemic antibiotics should be routinely prescribed.

Objective To evaluate whether systemic antibiotics, when compared with a placebo, improve cure rates in patients with simple abscesses after incision and drainage.

Methods Design Systematic review and meta-analysis using RevMan5.

Patients and settings Children and adults with simple abscesses treated in outpatient clinics or emergency departments.

Data sources Cochrane Central, Medline, Embase and bibliographies.

Outcome measures Percentage of patients with complete resolution of abscess without the need for recurrent incision and drainage, additional antibiotics, or hospital admission within 7–10 days of treatment.

Results We included four trials, consisting of 589 patients in total (428 adults and 161 children). Patients were randomised to one of three antibiotics (cephridine (27), cephalexin (82), or trimethoprim sulfamethoxazole (161)) or to placebo (285), with 34 lost to follow-up or having incomplete data. When given in addition to incision and drainage, systemic antibiotics did not significantly improve the percentage of patients with complete resolution of their abscesses 7–10 days after treatment (88.1% vs 86.0%; OR 1.17 (95% CI 0.70 to 1.95)).

Conclusions When given in addition to incision and drainage, systemic antibiotics do not significantly improve the percentage of patients with complete resolution of their abscesses.

INTRODUCTION

A large number of patients present to the emergency department (ED) each year with a skin abscess, and this number is rising. A study of the national epidemiology of cutaneous abscess conducted between 1996 and 2005 found that the number of ED visits for abscesses rose from 1.2 to 3.3 million, which was considerably greater than the rise in total ED visits during the study period.¹ While incision and drainage of abscesses is considered the standard care, the role of adjunctive antibiotics in simple abscesses remains unclear.² Based on prior studies, antibiotics are prescribed in roughly half the ED patients with abscesses, with 51% of the antibiotic regimens including trimethoprim-sulfamethoxazole active against community-associated methicillin-resistant *Staphylococcus aureus*.^{1–3} There is a large degree of variability in use of antibiotics, with some

centres using antibiotics in all patients,⁴ while others use them only in a minority of patients. There have been several small observational studies and clinical trials that have evaluated the potential benefits of adjunctive systemic antibiotics in patients with simple abscess though the results have been inconclusive. Several large ongoing National Institutes of Health (NIH)-sponsored clinical trials are evaluating the need for adjunctive antibiotics in patients with simple abscess. However, it is unclear when these results will become available.

The lack of clear-cut evidence supporting routine administration of adjunctive systemic antibiotics in patients with simple abscesses prompted us to conduct a meta-analysis of interventional trials comparing the effects of antibiotics and placebo on outcome after incision and drainage of simple cutaneous abscesses.

METHODS

Data selection and extraction

We conducted a systematic search of the MEDLINE, Embase, Scopus databases and the Cochrane registry of clinical trials (last accessed 15 Dec 2012) using the terms ‘abscess’, ‘antibiotics’, ‘clinical trial’ and ‘outcome’. We performed an additional search using Google Scholar and a list of bibliographies. We also searched trial registries and abstracts of major emergency medicine (EM) congresses. We included all studies that were placebo-controlled, randomised trials that compared any oral antibiotic (intervention) with a placebo (comparator) in ED patients after incision and drainage of their abscesses (patients) and provided outcome data on clinical cure (outcomes). We excluded animal trials and crossover trials. Two investigators independently screened studies by title and abstract to evaluate whether the trial fitted the inclusion criteria.

Two authors extracted data independently on a predefined data extraction form. A two-by-two summary table was completed for each outcome and for each trial, summarising the number of patients who experienced the event or outcome for each group. These data were double-checked. If there was any disagreement, then the source data were evaluated jointly. The primary outcome was clinical cure at 7 or 10 days, and a secondary outcome was abscess recurrence at 30 or 90 days.

For the randomised trials, two authors independently collected information from all studies to assess the risk of bias using the Cochrane risk of bias tool.⁵ We collected information on random sequence allocation (selection bias), allocation concealment (selection bias), blinding of participants and personnel (performance bias), blinding of



CrossMark

To cite: Singer AJ, Thode HC Jr. *Emerg Med J* 2014;**31**:576–578.

Table 1 Risk of bias

Bias	Liera 1985	Rajendran 2007	Duong 2010	Schmitz 2010
Random sequence generation (selection bias)	L	L	L	L
Allocation concealment (selection bias)	L	L	L	L
Blinding of participants and personnel (performance bias)	L	L	L	L
Blinding of outcome assessment (detection bias)	L	L	U	L
Incomplete outcome data (attrition bias)	L	L	L	L
Selective reporting (reporting bias)	L	L	L	L
Other bias	L	L	L	L

L, low; U, unclear.

outcome assessment (detection bias), incomplete outcome data (attrition bias), selective reporting (reporting bias), and any other obvious bias.

We assigned 'low risk', 'high risk' or 'unable to determine' to each item of the Cochrane risk of bias tool. Data are reported in accordance with the PRISMA guidelines for systematic reviews.⁶ Since this was a meta-analysis of published articles, it was exempt from institutional review board (IRB) review.

Statistical analysis

For each randomised trial and the aggregate analysis, we used the OR as the summary measure of association. We evaluated statistical heterogeneity among studies using Cochrane Q and I² statistics.⁷ We evaluated publication bias by means of visual inspection of the funnel plot.^{8–9} Data were analysed using REVMAN5 using weighted analysis where the weights were the inverse of the variance of the effects estimates.⁵ All reported p values are two-sided, and p < 0.05 was considered to be statistically significant.

RESULTS

A total of 106 studies were identified, of which 102 were excluded (retrospective, observational, complicated abscess, lack of placebo) so that four trials were kept for analysis. All these were randomised, placebo-controlled trials in ED patients with simple non-complicated abscesses. Risk of bias was determined to be low based on the criteria in REVMAN5 (table 1).

The four randomised trials included 589 patients in total (428 adults and 161 children). Patients were randomised to one of three antibiotics (cephridine (27),¹⁰ cephalexin (82),¹¹ or

trimethoprim sulfamethoxazole (161)^{12–13} or to placebo (285) with 34 lost to follow-up or having incomplete data. Three studies used clinical cure at 7 days,^{10–11–13} and one study used clinical cure at 10 days as their primary end point.¹² When given in addition to incision and drainage, systemic antibiotics did not significantly improve the percentage of patients with complete resolution of their abscesses 7–10 days after treatment (88.1% (95% CI 85.7 to 92.9) vs 86.0% (95% CI 79.6 to 88.1); difference, 2% (95% CI –4% to 7%); OR 1.17 (95% CI 0.70 to 1.95)) (figure 1). There was no significant heterogeneity among trials (p=0.30; I²=18%). The funnel plot did not reveal any publication bias. Limiting the analysis to only those studies with a 7-day endpoint had no impact on the results.

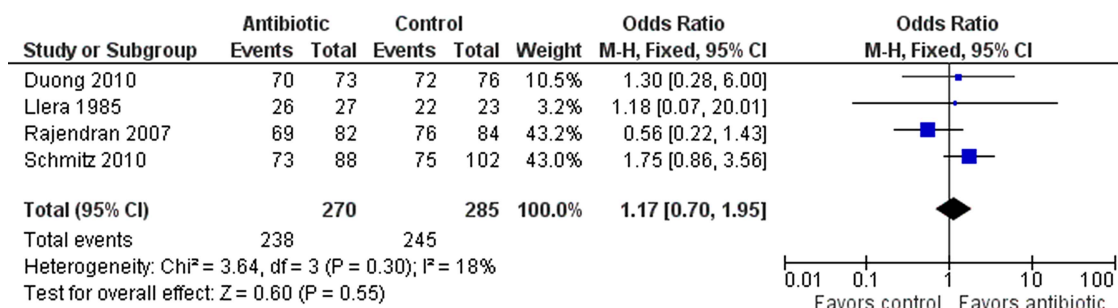
Follow-up beyond the first 7–10 days was performed in two of the trials, at 30 and at 90 days after incision and drainage.^{12–13} When given in addition to incision and drainage, systemic antibiotics did not significantly increase the percentage of patients without abscess recurrence 30–90 days after treatment (81.5% (95% CI 73.6 to 89.5) vs 71.6% (95% CI 62.8 to 80.3); difference 10% (95% CI –2 to 22), OR 1.74 (95% CI 0.88 to 3.45)) (figure 2).

DISCUSSION

This systematic review and meta-analysis reveals that use of adjunctive oral antibiotics after incision and drainage of simple cutaneous abscesses in the ED does not increase the early 7–10-day cure rate, nor does it reduce the rate of abscess recurrence 30–90 days after incision and drainage. Individually and collectively, the clinical trials included in this systematic review showed no advantage of antibiotics over placebo.

The results of this meta-analysis are in agreement with the recent clinical practice guidelines by the Infectious Diseases Society of America for the treatment of methicillin-resistant *Staphylococcus aureus* infections in adults and children. These guidelines recommend incision and drainage alone for simple abscesses with systemic antibiotics only for severe or extensive disease, rapid progression in the presence of associated cellulitis, signs and symptoms of systemic illness, associated comorbidities or immunosuppression, extremes of age, abscess in an area difficult to drain, associated septic phlebitis, and lack of response to incision and drainage.²

Only two of the included trials involved an antibiotic with specific activity against most strains of community associated methicillin resistant staphylococcus aureus (CA-MRSA), which is responsible for more than half of all skin and soft tissue infections in a recent multicenter ED-based study.¹⁴ Even when the meta-analysis is limited to the latter two studies, there is no statistically significant difference in 7–10 cure rates (mean difference 6.2%, 95% CI –1.3 to 13.7). Empiric antibiotics, if prescribed, should demonstrate in vitro activity against CA-MRSA.

**Figure 1** Forest plots for 7–10-day cure rates.

Original article

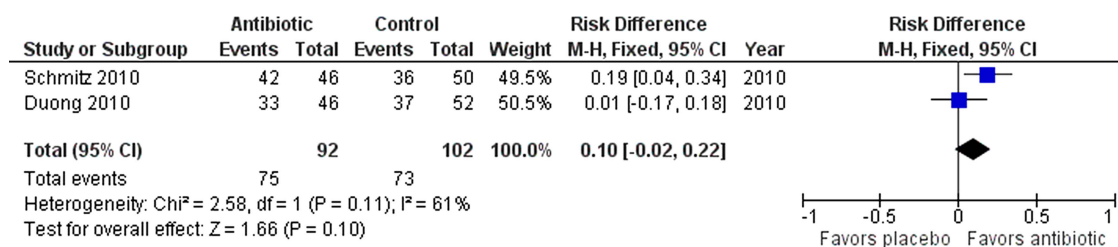


Figure 2 Forest plots for 30–90-day recurrence rates.

Trimethoprim sulfamethoxazole, clindamycin, and tetracycline have all demonstrated in vitro activity against >90% of over 300 isolates tested in the 2008 US ED-based surveillance study.¹⁵ If the abscess is associated with significant cellulitis, or cannot be distinguished from cellulitis, consideration should be given to adding an antibiotic active against *Streptococcus pyogenes*, which is responsible for most cases of cellulitis without abscess.

LIMITATIONS

Our study has several limitations. First, it is limited to four randomised clinical trials including less than 600 patients. Thus, it may have been underpowered to detect small yet clinically significant differences in outcomes. Second, longer-term follow-up was only performed in two of the studies which further increased the risk of a type II error. Third, most of the studies lacked enough detail to determine the effect of confounding variables, such as adequacy of drainage and compliance with assigned treatments.

In conclusion, this meta-analysis of ED patients with simple abscesses indicates that adjunctive oral antibiotics do not increase the cure rate or recurrence rates after incision and drainage.

Contributors The study was conceived by AJS, and data analysis was by HCT and AJS. AJS wrote the first draft and all authors reviewed and approved the final manuscript.

Competing interests None.

Ethics approval IRB.

Provenance and peer review Not commissioned; internally peer reviewed.

REFERENCES

- 1 Taira BR, Singer AJ, Thode HA, *et al.* National epidemiology of cutaneous abscesses: 1996 to 2005. *Am J Emerg Med* 2009;27:289–92.
- 2 Liu C, Bayer A, Cosgrove SE, *et al.* Infectious Diseases Society of America. Clinical practice guidelines by the Infectious Diseases Society of America for the treatment of methicillin-resistant *Staphylococcus aureus* infections in adults and children. *Clin Infect Dis* 2011;52:e18–55.
- 3 Pallin DJ, Egan DJ, Pelletier JA, *et al.* Increased US department visits for skin and soft tissue infections, and changes in antibiotic choices, during the emergency of community-associated methicillin-resistant *Staphylococcus aureus*. *Ann Emerg Med* 2008;51:291–8.
- 4 Lee MC, Rios AM, Aten MF, *et al.* Management and outcome of children with skin and soft tissue abscess caused by community-acquired methicillin-resistant *Staphylococcus aureus*. *Pediatr Infect Dis J* 2004;23:123–7.
- 5 Review Manager (REVMAN) v5.1, Copenhagen: The Nordic Cochrane Centre, The Cochrane Collaboration, 2011.
- 6 Liberati A, Altman DG, Tetzlaff J, *et al.* The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. *PLoS Med* 2009;6:e1000100.
- 7 Higgins JP, Thompson SG. Quantifying heterogeneity in a meta-analysis. *Stat Med* 2002;21:1539–58.
- 8 Egger M, Davey Smith G, Schneider M, *et al.* Bias in meta-analysis detected by a simple, graphical test. *BMJ* 1997;315:629–34.
- 9 DerSimonian R, Laird N. Meta-analysis in clinical trials. *Control Clin Trials* 1986;7:177–88.
- 10 Ller JL, Levy RC. Treatment of cutaneous abscess: a double-blind clinical study. *Ann Emerg Med* 1985;14:15–19.
- 11 Rajendran PM, Young D, Maurer T, *et al.* Randomized, double-blind, placebo-controlled trial of cephalexin for treatment of uncomplicated skin abscesses in a population at risk for community-acquired methicillin-resistant *Staphylococcus aureus* infection. *Antimicrob Agents Chemother* 2007;51:4044–8.
- 12 Duong M, Markwell S, Peter J, *et al.* Randomized, controlled trial of antibiotics in the management of community-acquired skin abscesses in the pediatric patient. *Ann Emerg Med* 2010;55:401–7.
- 13 Schmitz GR, Bruner D, Pitotti R, *et al.* Randomized controlled trial of trimethoprim-sulfamethoxazole for uncomplicated skin abscesses in patients at risk for community-associated methicillin-resistant *Staphylococcus aureus* infection. *Ann Emerg Med* 2010;56:283–7.
- 14 Moran GJ, Krishnadasan A, Gorwitz RJ, *et al.* Methicillin-resistant *S. aureus* infections among patients in the emergency department. *N Engl J Med* 2006;355:666–74.
- 15 Talan DA, Krishnadasan A, Gorwitz RJ, *et al.* Comparison of *Staphylococcus aureus* from skin and soft tissue infections in U.S. emergency department patients, 2004 and 2008. *Clin Infect Dis* 2011;53:144–9.



Systemic antibiotics after incision and drainage of simple abscesses: a meta-analysis

Adam J Singer and Henry C Thode, Jr

Emerg Med J 2014 31: 576-578 originally published online May 18, 2013

doi: 10.1136/emmermed-2013-202571

Updated information and services can be found at:

<http://emj.bmj.com/content/31/7/576.full.html>

These include:

References

This article cites 14 articles, 4 of which can be accessed free at:

<http://emj.bmj.com/content/31/7/576.full.html#ref-list-1>

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections

Articles on similar topics can be found in the following collections

[Drugs: infectious diseases](#) (255 articles)

[Child health](#) (210 articles)

Notes

To request permissions go to:

<http://group.bmj.com/group/rights-licensing/permissions>

To order reprints go to:

<http://journals.bmj.com/cgi/reprintform>

To subscribe to BMJ go to:

<http://group.bmj.com/subscribe/>