

Proof Delivery Form

CUP reference: 9872(JLO1731)

Date of delivery: 10.03.07

Journal and Article number: S0022215107007311jra

JOURNAL OF LARYNGOLOGY & OTOTOLOGY

Volume and Issue Number: 121 and 0

Number of pages (not including this page): 5

There follows a proof of the article you have written for publication in the *JOURNAL OF LARYNGOLOGY & OTOTOLOGY*. Please check the proofs carefully, make any corrections necessary and answer queries on the proofs. Queries raised by the sub-editor are listed below; the text to which the queries refer is flagged in the margins of the proof. Line numbers have been included on the proof to aid in listing any corrections.

Please return your corrections via email as soon as possible (but no later than 5 days after receipt) to both of the following addresses:

Paul Beaney: pbeaney@techset.co.uk;

Pravina Patel: ppatel@techset.co.uk

It is vitally important that any corrections we receive are easy to decipher, therefore please supply a separate typed sheet of all your corrections and answers to any queries. Please refer to the relevant line number and column when listing your corrections.

Please ensure you use the JLO identification number in all correspondence. If absolutely necessary, corrections can be returned to us via fax or courier, should you wish to return corrections in this way please contact us before doing so.

Techset Composition Limited
Chalke House, 3 Brunel Road, Churchfields,
Salisbury, SP2 7PU, UK

Direct Line: +44(0)1722 420689

Switchboard: +44(0)1722 332949

Facsimile: +44(0)1722 323159

Web address: www.techset.co.uk

- You are responsible for correcting your proofs! Errors not found may appear in the published journal.
- The proof is sent to you for correction of typographical errors only. Revision of the substance of the text is not permitted.
- Please answer carefully any queries raised from the sub-editor.
- A new copy of a figure must be provided if correction of anything other than a typographical error introduced by the printer is required.

Notes:

1. Your copyright and offprint forms should be returned directly to

Ms Susan Perkins
Production Editor
Cambridge University Press
The Edinburgh Building
CB2 8RU, UK
Tel: +44(0)1223 326163
Fax: +44(0)1223 325802
Email: sperkins@cambridge.org

2. The quality of half-tones will be checked by the Editorial Office.
 3. If you have any queries, please telephone the Editorial Office.
-

Author queries:

- Q1 Figure 1 and 2 were not cited in the text. Figure 1 and 2 have been swapped around. Position of citation acceptable?
- Q2 Table II was not cited in the text – is it OK here?
-

Typesetter queries:

Non-printed material:

Thank you in advance for your cooperation

Paul Beaney and Pravina Patel
Techset Composition Limited

On Behalf of
Ms Susan Perkins
Production Editor
Cambridge University Press
The Edinburgh Building
CB2 8RU
UK

transfer of copyright

Please read the notes overleaf and then complete, sign, and return this form to Sue Perkins, Cambridge University Press, The Edinburgh Building, Shaftesbury Road, Cambridge CB2 8RU, UK as soon as possible.

JOURNAL OF LARYNGOLOGY & OTOTOLOGY

In consideration of the publication in JOURNAL OF LARYNGOLOGY & OTOTOLOGY

of the contribution entitled:.....

.....

by (all authors' names):.....

.....

1 To be filled in if copyright belongs to you

Transfer of copyright

I/we hereby assign to JLO (1984) Ltd, full copyright in all formats and media in the said contribution.

I/we warrant that I am/we are the sole owner or co-owners of the material and have full power to make this agreement, and that the material does not contain any libellous matter or infringe any existing copyright.

I/we further warrant that permission has been obtained from the copyright holder for any material not in my/our copyright and the appropriate acknowledgement made to the original source. I/we attach copies of all permission correspondence.

I/we hereby assert my/our moral rights in accordance with the UK Copyrights Designs and Patents Act (1988).

Signed (tick one) ☐ the sole author(s)

☐ one author authorised to execute this transfer on behalf of all the authors of the above article

Name (block letters)

Institution/Company.....

Signature: Date:

(Additional authors should provide this information on a separate sheet.)

2 To be filled in if copyright does not belong to you

a Name and address of copyright holder.....

.....

.....

.....

b The copyright holder hereby grants to JLO (1984) Ltd the non-exclusive right to publish the contribution in the journal and to deal with requests from third parties in the manner specified in paragraphs 3 and 5 overleaf.

(Signature of copyright holder or authorised agent)

3 US Government exemption

I/we certify that the paper above was written in the course of employment by the United States Government so that no copyright exists.

Signature: Name (Block letters):

4 Requests received by Cambridge University Press for permission to reprint this article should be sent to (see para. 4 overleaf)

Name and address (block letters)

.....

.....

Notes for contributors

- 1 The Journal's policy is to acquire copyright in all contributions. There are two reasons for this: (a) ownership of copyright by one central organisation tends to ensure maximum international protection against unauthorised use; (b) it also ensures that requests by third parties to reprint or reproduce a contribution, or part of it, are handled efficiently and in accordance with a general policy that is sensitive both to any relevant changes in international copyright legislation and to the general desirability of encouraging the dissemination of knowledge.
- 2 Two 'moral rights' were conferred on authors by the UK Copyright Act in 1988. In the UK an author's 'right of paternity', the right to be properly credited whenever the work is published (or performed or broadcast), requires that this right is asserted in writing.
- 3 Notwithstanding the assignment of copyright in their contribution, all contributors retain the following **non-transferable** rights:
 - The right to post *either* their own version of their contribution as submitted to the journal (prior to revision arising from peer review and prior to editorial input by Cambridge University Press) *or* their own final version of their contribution as accepted for publication (subsequent to revision arising from peer review but still prior to editorial input by Cambridge University Press) on their **personal or departmental web page**, or in the **Institutional Repository** of the institution in which they worked at the time the paper was first submitted, provided the posting is accompanied by a prominent statement that the paper has been accepted for publication and will appear in a revised form, subsequent to peer review and/or editorial input by Cambridge University Press, in Journal of Laryngology & Otology published by Cambridge University Press, together with a copyright notice in the name of the copyright holder (Cambridge University Press or the sponsoring Society, as appropriate). On publication the full bibliographical details of the paper (volume: issue number (date), page numbers) must be inserted after the journal title, along with a link to the Cambridge website address for the journal. Inclusion of this version of the paper in Institutional Repositories outside of the institution in which the contributor worked at the time the paper was first submitted will be subject to the additional permission of Cambridge University Press (not to be unreasonably withheld).
 - The right to post the definitive version of the contribution as published at Cambridge Journals Online (in PDF or HTML form) on their **personal or departmental web page**, no sooner than upon its appearance at Cambridge Journals Online, subject to file availability and provided the posting includes a prominent statement of the full bibliographical details, a copyright notice in the name of the copyright holder (Cambridge University Press or the sponsoring Society, as appropriate), and a link to the online edition of the journal at Cambridge Journals Online.
 - The right to post the definitive version of the contribution as published at Cambridge Journals Online (in PDF or HTML form) in the **Institutional Repository** of the institution in which they worked at the time the paper was first submitted, no sooner than **one year** after first publication of the paper in the journal, subject to file availability and provided the posting includes a prominent statement of the full bibliographical details, a copyright notice in the name of the copyright holder (Cambridge University Press or the sponsoring Society, as appropriate), and a link to the online edition of the journal at Cambridge Journals Online. Inclusion of this definitive version after one year in Institutional Repositories outside of the institution in which the contributor worked at the time the paper was first submitted will be subject to the additional permission of Cambridge University Press (not to be unreasonably withheld).
 - The right to make hard copies of the contribution or an adapted version for their own purposes, including the right to make multiple copies for course use by their students, provided no sale is involved.
 - The right to reproduce the paper or an adapted version of it in any volume of which they are editor or author. Permission will automatically be given to the publisher of such a volume, subject to normal acknowledgement.
- 4 We shall use our best endeavours to ensure that any direct request we receive to reproduce your contribution, or a substantial part of it, in another publication (which may be an electronic publication) is approved by you before permission is given.
- 5 Cambridge University Press co-operates in various licensing schemes that allow material to be photocopied within agreed restraints (e.g. the CCC in the USA and the CLA in the UK). Any proceeds received from such licenses, together with any proceeds from sales of subsidiary rights in the Journal, directly support its continuing publication.
- 6 It is understood that in some cases copyright will be held by the contributor's employer. If so, JLO (1984) Ltd requires non-exclusive permission to deal with requests from third parties, on the understanding that any requests it receives from third parties will be handled in accordance with paragraphs 4 and 5 above (note that your approval and not that of your employer will be sought for the proposed use).
- 7 Permission to include material not in your copyright
If your contribution includes textual or illustrative material not in your copyright and not covered by fair use / fair dealing, permission must be obtained from the relevant copyright owner (usually the publisher or via the publisher) for the non-exclusive right to reproduce the material worldwide in all forms and media, including electronic publication. The relevant permission correspondence should be attached to this form.

If you are in doubt about whether or not permission is required, please consult the Permissions Controller, Cambridge University Press, The Edinburgh Building, Shaftesbury Road, Cambridge CB2 8RU, UK. Fax: +44 (0)1223 315052.

Email: lnicol@cambridge.org.

The information provided on this form will be held in perpetuity for record purposes. The name(s) and address(es) of the author(s) of the contribution may be reproduced in the journal and provided to print and online indexing and abstracting services and bibliographic databases

Please make a duplicate of this form for your own records

Offprint order form



CAMBRIDGE
UNIVERSITY PRESS

PLEASE COMPLETE AND RETURN THIS FORM. WE WILL BE UNABLE TO SEND OFFPRINTS UNLESS A RETURN ADDRESS AND ARTICLE DETAILS ARE PROVIDED.

VAT REG NO. GB 823 8476 09

The Journal of Laryngology & Otology (JLO)

Volume:

no:

Offprints

To order offprints, please complete this form and send it to **the publisher** (address below). Please give the address to which your offprints should be sent. They will be despatched by surface mail within one month of publication.

Number of offprints required:

Email:

Offprints to be sent to (print in BLOCK CAPITALS):

Post/Zip Code:

Telephone:

Date (dd/mm/yy):

Author(s):

Article Title:

All enquiries about offprints should be addressed to **the publisher**: Journals Production Department, Cambridge University Press, The Edinburgh Building, Shaftesbury Road, Cambridge CB2 8RU, UK.

Charges for offprints (excluding VAT) Please circle the appropriate charge:

Number of copies	25	50	100	150	200	per 50 extra
1-4 pages	£68	£109	£174	£239	£309	£68
5-8 pages	£109	£163	£239	£321	£399	£109
9-16 pages	£120	£181	£285	£381	£494	£120
17-24 pages	£131	£201	£331	£451	£599	£131
Each Additional 1-8 pages	£20	£31	£50	£70	£104	£20

Methods of payment

If you live in Belgium, France, Germany, Ireland, Italy, Portugal, Spain or Sweden and are not registered for VAT we are required to charge VAT at the rate applicable in your country of residence. If you live in any other country in the EU and are not registered for VAT you will be charged VAT at the UK rate.

If registered, please quote your VAT number, or the VAT number of any agency paying on your behalf if it is registered.

VAT Number:

Payment **must** be included with your order, please tick which method you are using:

- ☐ Cheques should be made out to Cambridge University Press.
- ☐ Payment by someone else. Please enclose the official order when returning this form and ensure that when the order is sent it mentions the name of the journal and the article title.
- ☐ Payment may be made by any credit card bearing the Interbank Symbol.

Card Number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Expiry Date (mm/yy):

/

Card Verification Number:

--	--	--	--

The card verification number is a 3 digit number printed on the **back** of your **Visa** or **Master card**, it appears after and to the right of your card number. For **American Express** the verification number is 4 digits, and printed on the **front** of your card, after and to the right of your card number.

Signature of
card holder:

Amount
(Including VAT
if appropriate):

£

Please advise if address registered with card company is different from above

Is bacterial colonisation of the tonsillar fossa a factor in post-tonsillectomy haemorrhage?

J C STEPHENS, CHRISTOS GEORGALAS*, M KYI†, K GHUFOOR‡

Abstract

Objectives: To identify if there is a link between bacterial colonisation of the tonsillar fossa and post-tonsillectomy haemorrhage.

Study design and setting: Prospective non interventional study of 105 patients who underwent tonsillectomy during a seven-month-period. The study took place in a secondary care centre, the West Middlesex University Hospital.

Participants: The participants were 105 patients who consecutively underwent tonsillectomy. The exclusion criteria were any patients with suspected or known malignancy, or known bleeding dyscrasias. The participants underwent microbiological sampling of the tonsil pre-operatively.

Main outcomes measures: The outcome measures were primary or secondary bleeding, defined as any evidence of haemorrhage in the tonsillar fossae.

Results: Twenty four percent of patients undergoing tonsillectomy had positive cultures from their tonsils pre-operatively. Patients with bacterial colonisation of the tonsillar fossa pre-operatively had an increased rate of post-tonsillectomy haemorrhage (odds ratio: 3.8, 1.1–12.1, 95 per cent confidence intervals $p = 0.04$).

Conclusion: This prospective study has found a relationship between bacterial colonisation of the tonsillar fossa and post-tonsillectomy haemorrhage. This suggests that there may be an argument for the use of antibiotics in those cases with positive pre-operative cultures. In view of the types of pathogens isolated, we feel that the management of a post-tonsillectomy bleed should include a beta lactamase inhibiting antibiotic.

Key words: Tonsillectomy; Haemorrhage; Microbiology; Infection

Introduction

Tonsillectomy is one of the most common surgical procedures undertaken in the UK, and post-operative haemorrhage is the most significant complication, with reported rates ranging between 3 to 20 per cent.¹ The severity of bleeding can range from very minor to fatal haemorrhage so a thorough analysis of the risk factors is extremely useful. Consensus opinion suggests that the cause of secondary haemorrhage is infection in the tonsillar bed, and it has also been demonstrated that the use of hot techniques for tonsillectomy are associated with higher rates of secondary haemorrhage. The management of mild to moderate bleeds is usually conservative and includes hospital bed rest, fluid resuscitation, intravenous antibiotics and close monitoring.² However, this practice has little evidence base, and there is no data to show any association between bacterial colonisation of the tonsillar bed and the rates of post-operative haemorrhage.

Several studies have examined tonsillar micro flora, both during episodes of acute tonsillitis and in periods without any clinical evidence of infection.^{3–5} It has been shown that even in the absence of inflammation, polymicrobial flora are present in tonsillar tissue and during the inflammatory process these increase significantly in number.⁴ The commonest bacteria isolated have been *Haemophilus influenzae*, *Staphylococcus aureus*, and mixed anaerobes.^{3,5}

Post-tonsillectomy infection is a recognised complication, and can result in pyrexia, increased pain and analgesia requirements, nausea and vomiting, otalgia, halitosis and general malaise.^{6,7} Opinion over whether the routine use of antibiotics post-operatively is justified is divided and several studies have examined this. Results have varied, earlier trials showing a decrease in the symptoms of post-tonsillectomy infection with antibiotics,⁶ but later studies finding no justification for the routine use of antimicrobials.^{7,8}

From the Department of ENT, Charing Cross Hospital, the *Department of ENT, St Mary's Hospital, the †Department of Microbiology, West Middlesex Hospital and the ‡Department of ENT, St Bartholomew's Hospital, London.
Accepted for publication: 20 January 2007.

The aim of this prospective study is to examine whether there is a relationship between bacterial colonisation and post-operative bleeding, and to identify the organisms involved.

Materials and methods

There were 105 consecutive patients included in the study. The median age was seven with a range from two to 38. There were 79 paediatric cases and 26 adults, 61 males and 44 females. All patients undergoing tonsillectomy were added to the database, excluding those with suspected or proven malignancy, or known bleeding tendency. All patients underwent tonsillectomy in the operating theatres at the West Middlesex University Hospital, Middlesex, under general anaesthesia. The surgical technique used for each patient was bipolar diathermy. This study predated the National Prospective Tonsillectomy Audit, and the interim guidance issued in March 2004, which found that the use of diathermy for tonsillectomy dissection and haemostasis is associated with higher rates of post-tonsillectomy haemorrhage and suggesting that diathermy be used with caution.⁹ Seventy-seven per cent of procedures were carried out by registrars, 17 per cent by senior house officers and 6 per cent by consultants. None of the patients received pre- intra- or post-operative antibiotics or steroids. Demographic data were collected, including the age, gender, indication for surgery and any additional procedure performed, as well as the grade of surgeon and outcome in terms of post-operative bleeding. Patients were given standard advice sheets on post-operative care and dealing with most post-discharge questions.

Microbiological sampling was taken from the tonsillar surface pre-operatively immediately before removing the tonsils.

These swabs were placed in Stuart's transport medium and transported to the microbiology laboratory. The swabs were cultured directly onto four types of culture media plates to identify all bacterial organisms in the upper aerodigestive tract. The plates were incubated at 37°C in appropriate conditions and re-examined after 24 hours. They were re-incubated for a further 24 hours and re-examined before the plates were discarded. Any likely significant pathogens from the first and second readings were followed up for complete identification and sensitivity testing.

Patients with post-operative problems were advised to follow the instructions on the advice sheet – namely to return to hospital or the nearest emergency department or call the ENT ward. All patients requiring re-admission to hospital with an episode of bleeding were also re-swabbed at the time of admission. Bleeding was defined as an episode of fresh (red) or altered (brown) blood expectoration of any volume in the post-operative phase, and bleeding episodes were identified by an interview with the patient which took place in the hospital at seven to 10 days post-operatively. All results were entered into an Excel data file, and subsequently transferred and analysed in SPSS 12.0. All comparisons in proportions were performed using 2-sided

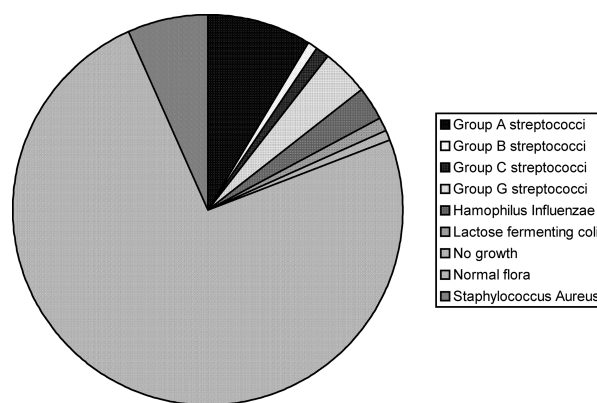


FIG. 1

Details of microbiology – percentage of microbials identified.

Pearson's chi-square and Fisher's exact test as appropriate. Continuous variables parametrically distributed were compared using two-sided *t*-tests.

Results and analysis

One hundred and five patients were included in the study. Twenty-four point eight percent of patients had positive pre-operative cultures, with the organisms identified detailed in Figure 1. The full microbiology results are summarised in the Table I. The total number of patients who experienced post-tonsillectomy haemorrhage (defined as any amount of fresh or altered blood from the tonsillar fossae) was 14 (13.3 per cent). The median time to bleeding episode was seven days, mean 8.1 days, range 0–16 days (95 per cent confidence intervals [CI] 5.3–9.4 days) (Figure 2). We found that males had a higher bleeding rate than females, 16.2 per cent of males versus 9.1 per cent of females, although this difference was not statistically significant. Operative procedures performed by senior house officers had the highest rates of bleeding at 23.5 per cent, followed by registrars with a bleeding rate of 9 per cent while none of the five patients who were operated on by consultants bled again. Statistically this difference was significant ($p = 0.009$). Similarly, analysis of the patient's ages showed that those who bled tended to be older (median age 17 in the haemorrhage group versus six in the non

Q1

TABLE I
PRE-OPERATIVE MICROBIOLOGY RESULTS

	Frequency	Per cent
Group A beta haemolytic streptococcus	9	8.6
Group B streptococcus	1	1
Group C haemolytic streptococcus	1	1
Group G beta haem streptococcus	4	3.8
<i>Haemophilus influenzae</i>	3	2.9
Lactose fermenting coli	1	1
No growth	1	1
Normal flora	78	74.3
<i>Staphylococcus aureus</i>	7	6.7
Total	105	100

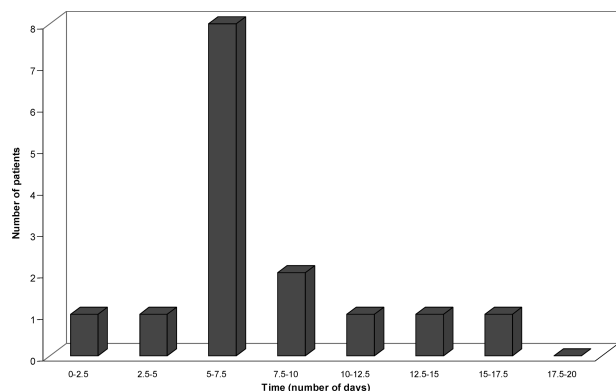


FIG. 2

Time interval to post-tonsillectomy haemorrhage.

haemorrhage group), although the difference was not statistically significant.

A comparison of all the patients with growth of pathogenic bacteria, versus those with normal flora or no bacteria showed that there is a consistent pattern of increased bleeding associated with colonisation of the tonsillar fossae (Table II).

We found that a pre-operative swab which showed normal flora or no bacterial growth was associated with an 8.8 per cent chance of bleeding versus a 26.9 per cent chance of bleeding with pathogen growth, and this finding was statistically significant (odds ratio; 3.8, 1.1 to 12.1, 95 per cent CI, $p = 0.04$). The different bleeding rates varied depending which bacteria had been cultured: four out of the nine patients with Group A streptococci returned with a post-tonsillectomy bleed, while one of the four patients with Group G streptococci bled, compared with only seven out of 72 patients with no growth or normal flora. The risk of bleeding was highest when the tonsillar fossa was colonised with haemolytic streptococci. The range of flora isolated from the oropharynx included Lancefield Groups A, B, C and G streptococci, *Staphylococcus aureus* and *Haemophilus influenzae*.

Discussion

Many potential risk factors in post-tonsillectomy haemorrhage have been assessed including the use

of local anaesthesia, pre-operative abnormalities in blood pressure or clotting,¹⁰ the role of age, gender and indication for surgery,¹¹ and even whether red-heads are more commonly affected, or higher rates seen on Friday the 13th.¹² None of these associations were shown to be positive. It has been shown that the use of bipolar diathermy can increase the rate of post-tonsillectomy haemorrhage, with rates three times higher observed than when using traditional cold steel techniques. The observation of a dose-response relation suggests that the extent to which diathermy is used is linked with the amount of damage to surrounding tissues, and therefore to the rate of secondary haemorrhage.¹³ Although no clear link has been shown between positive tonsillar microbiology and post-operative tonsillectomy haemorrhage, infection is widely accepted to play a role in the pathogenesis. Patients presenting to hospital with post-tonsillectomy bleeds are usually admitted and treated with intravenous antibiotics and this combined with a period of observation is frequently adequate intervention. Previous studies have shown that tonsillar tissue contains pathogenic bacteria,¹⁴ and that during tonsillectomy a transient bacteraemia occurs in as many as 27 per cent via the breach in oropharyngeal mucosa,^{3,15} but as this bacteraemia is short lived the use of prophylactic antibiotics is unnecessary unless the risk of metastatic infection is high.^{3,15}

The results in this study showed that the rate of bleeding throughout the entire cohort appears to be high at 13.3 per cent. This is due to the fact that all bleeding episodes were reported, including those which were minor and did not require hospital admission. This result is in keeping with the published series using a similar definition of post-operative bleeding, where bleeding rates as high as 20 per cent have been reported.¹ Our study identified an increased rate of bleeding in both males and with increasing age. Although neither of these findings were statistically significant, they reflect the findings of the National Prospective Tonsillectomy Audit, which showed a statistically significant difference in both categories.⁹ The increased rate of bleeding in tonsillectomies performed by trainees has been considered, and was thought to be due to increased use of diathermy during the procedure. Although the numbers were small, this has been useful in re-assessing the training and supervision required during the performance of this common procedure.

The second point of interest is the diverse range of bacteria which were identified from the tonsils and tonsillar beds. Previous research has found the commonest bacteria isolated were *Haemophilus influenzae*, *Staphylococcus aureus* and mixed anaerobes. Our data isolated Lancefield groups A B C & G streptococci and lactose fermenting coliforms as well as those mentioned above. The organisms seen most frequently were streptococci, including the beta haemolytic streptococci which comprised 63 per cent of the total. This was helpful in guiding the choice of antibiotics in the treatment of post-tonsillectomy infection or bleeding.

TABLE II

CORRELATION OF PRE-OPERATIVE MICROBIOLOGY WITH BLEEDING

Microbiology	Pre-operative microbiology	
	Pathogens grown	Normal flora/No growth
Bleed	7	7
No bleeding	19	72
Total	26	79
	105	

$p = 0.04$ Fisher's exact test

Trials assessing the effect of antibiotics post-tonsillectomy have not demonstrated a significant reduction in complications. This may be due to the diverse range of bacteria present and resistance to the commonly prescribed antimicrobials, which in most studies tended to be amoxicillin or erythromycin. This data shows the frequent presence of bacteria which are resistant to amoxicillin via the virulence factor beta lactamase, i.e. the beta haemolytic streptococci, and suggests it would be more appropriate to use an antibiotic which contains clavulanic acid such as co-amoxiclav (Augmentin).

- **Treatment of post-tonsillectomy haemorrhage often involves hospital admission and intravenous antibiotics**
- **However, there is no published data to show that infection is related to post-tonsillectomy haemorrhage**
- **One hundred and five patients underwent tonsillectomy and had microbiological sampling of their tonsils pre-operatively**
- **Twenty-four per cent of these patients had positive cultures from their tonsils**
- **Bacterial colonisation of the tonsil post-operatively increases the post-tonsillectomy bleeding rate by more than three times ($p = 0.04$)**

However, the most important finding was that bacterial colonisation of the tonsil in the pre-operative period increases the bleeding rate by more than three times, an increase that was statistically significant despite our small sample. Although it would be tempting to assume that eradication of pathogens would be associated with reduced bleeding rates, it remains to be proven. However, our study could potentially explain the divergent results of studies which assessed the efficacy of prophylactic antibiotics in preventing post-tonsillectomy bleeding. If one assumed that antibiotics were effective prophylactically only in patients with positive cultures, then their overall effectiveness would be dependent on the incidence of colonisation. In studies containing only a small number of patients with colonised tonsillar fossae their effectiveness would be diluted and not demonstrable. Thus, we do not feel that one could suggest routinely sterilising the tonsillar fossae pre-operatively as the number needed to treat to avoid an episode of bleeding would be extremely large. However, it may be a viable solution to prescribe antibiotics for patients with positive microbiology pre-operatively, although this would require all patients to undergo microbiological sampling, and this is also a significant undertaking. Further trials may be useful in assessing the viability of this proposition as well as its cost effectiveness.

Potential limitations of the study

We were not able to control for potential sources of bias within the confines of this study, for example use of other prescribed medications, including medications which predispose to bleeding – although all patients with a bleeding tendency were excluded from the study. We were also unable to control for exercise regimes and activities post-operatively, and had we done so it may have strengthened our findings.

Conclusion

This prospective study has found a statistically significant relationship between bacterial colonisation of the tonsillar fossa and post-tonsillectomy haemorrhage. This suggests there may be an argument for the use of antibiotics in those cases with positive pre-operative cultures. In view of the types of pathogens isolated, we feel that the management of a post-tonsillectomy bleed should include a beta lactamase inhibiting antibiotic.

References

- 1 Myssiorek D, Alvi A. Post-tonsillectomy haemorrhage: an assessment of risk factors. *Int J Paediatr Otorhinolaryngol* 1996;**37**:35–43
- 2 Shah G, Ghani R, Ayub J. Frequency of post tonsillectomy haemorrhage following tonsillectomy with bipolar diathermy- an experience at a teaching hospital, Abbottabad. *Med Coll Abbottabad* 2004;**16**:38–9
- 3 Gaffney RJ, Freeman DJ, Walshy MA, Cafferkey MT. Differences in tonsil core bacteriology in adults and children: a prospective study of 262 patients. *Respir Med* 1991;**85**:383–8
- 4 Brook I, Foote PA Jr. Microbiology of “normal” tonsils. Department of Paediatrics, National Naval Medical Centre, Bethesda Maryland. *Ann Otol Rhinol Laryngol* 1990; **99**:980–3
- 5 Surrow JB, Handler SD, Telian SA, Fleisher GR, Baranak CC. Bacteriology of tonsil surface and core in children. Department of Otorhinolaryngology and Human Communication, Children’s Hospital of Philadelphia, PA 19104. *Laryngoscope* 1989;**99**:261–6
- 6 Telian SA, Handler SD, Fleisher GR, Baranak CC, Wetmore RF. The effect of antibiotic therapy on recovery after tonsillectomy in children. A controlled study. *Arch Otolaryngol Head Neck Surg* 1986;**112**:610–5
- 7 O’Reilly BJ, Black S, Fernandes J, Panesar J. Is the routine use of antibiotics justified in adult tonsillectomy? *J Laryngol Otol* 2003;**117**:382–5
- 8 Burkart CM, Steward DL. Antibiotics for reduction of post-tonsillectomy morbidity: a meta-analysis. *Laryngoscope* 2005;**115**:997–1002
- 9 British Association of Otorhinolaryngologists – Head and Neck Surgeons Comparative Audit Group, Clinical Effectiveness Unit, The Royal College of Surgeons of England. *National Prospective Tonsillectomy Audit: Final Report of an audit carried out in England and Northern Ireland between July 2003 and September 2004*. London: The Royal College of Surgeons of England, 2005.
- 10 Tami TA, Parker GS, Taylor RE. Post tonsillectomy bleeding: an evaluation of risk factors. *Laryngoscope* 1987;**11**:1307–11
- 11 Bhattacharyya N. Evaluation of post tonsillectomy bleeding in the adult population. *Ear Nose Throat J* 2001;**80**:544–9
- 12 Kumar VV, Kumar NV, Isaacson G. Superstition and post-tonsillectomy haemorrhage. *Laryngoscope* 2004;**114**:2031–3
- 13 Van der Muelen J. Tonsillectomy technique as a risk factor for post operative haemorrhage. *Lancet* 2004;**364**:697–703

- 14 Stjernquist-Desatnik A, Holst E. Tonsillar microbial flora:
comparison of recurrent tonsillitis and normal tonsils. *Acta*
Otolaryngol 1999;**119**:102–6
- 15 Yuldririm I, Okur E, Ciragil P, Aral M, Kilic MA, Gul M
et al. Bacteraemia during tonsillectomy. *J Laryngol Otol*
2003;**117**:619–23

West Hampstead
London, NW6 4NE, UK.

Fax: 44 (0)208 321 5904
E-mail: jstephens@doctors.org.uk

Address for correspondence:
Miss Joanna C Stephens,
68b Gascony Avenue,

Miss J C Stephens takes responsibility for the integrity
of the content of the paper.
Competing interests: None declared
