Intracameral vancomycin following cataract surgery: An eleven-year study

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Aim: To compare the incidences of endophthalmitis after cataract operations before and after introduction of intracameral vancomycin at the end of surgery.

Methods: A retrospective analysis was performed of presumed infectious endophthalmitis after cataract surgery from January 1, 1998 to December 31, 2008. From January 2001, the practice of using intracameral vancomycin at the end of cataract surgery was introduced. The period before introduction of intracameral vancomycin is considered as period A and that after as period B. The incidences of presumed or culture-proven endophthalmitis during periods A and B were compared.

Results: A total of 16,606 cataract surgeries were performed during the study period. The incidence of endophthalmitis per 1000 cataract surgeries was 3.0 during period A and 0.08 during period B. This reduction was statistically significant (Chi-squared test 36.6, P value < 0.0001). The relative risk of developing endophthalmitis without intracameral vancomycin prophylaxis was 38. The absolute risk reduction was 292 cases of endophthalmitis per 100,000 cataract surgeries.

Conclusions: Intracameral vancomycin significantly reduced the incidence of postoperative endophthalmitis after cataract surgery. There is a universal need to adopt this mode of microbial prophylaxis to reduce the burden of endophthalmitis after cataract surgery.

Keywords: endophthalmitis, intracameral vancomycin, cataract surgery

Introduction

Cataract surgery is the commonest operation performed in health care systems worldwide. The technique has evolved from extracapsular extraction to sutureless phacoemulsification surgery over the past 15 years. Intraocular infection following cataract surgery and lens implantation is a rare but dreaded complication that can have devastating consequences for sight. The Endophthalmitis Vitrectomy Study (EVS) estimated that up to half of patients developing endophthalmitis following cataract surgery had visual acuity worse than 20/40 and 15% had worse than 5/200.1 The incidence of endophthalmitis following cataract surgery varies between 0.04 and 0.27%.2–4 Even though several risk factors have been identified, this complication is often unexpected, with variable response to standard treatment. As the number of cataract surgeries is expected to rise with an increasing aging population, the burden of endophthalmitis is likely to rise. A robust preventative strategy is needed to contain this serious complication. The European Society of Cataract and Refractive Surgeons (ESCRS) in a multinational, partially-masked placebo-controlled trial have provided strong evidence for using intracameral antibiotics in preventing postoperative endophthalmitis following cataract surgery.
surgery. In this study we examined the effect of introduction of intracameral vancomycin at the end of cataract surgery on the incidence of postoperative endophthalmitis in a single eye unit in North West England.

**Methods**

The study included all cataract surgeries performed in the Ophthalmic Department at Warrington Hospital in North West England from January 1, 1998 to December 31, 2008. The standard practice of cataract surgery over this period was similar except for the introduction of intracameral vancomycin at the end of cataract surgery from January 2001. In the anesthetic room, 10 minutes prior to cataract surgery, each patient had periocular skin cleaning with 5% povidone iodine. Two drops of 5% povidone iodine were instilled into the conjunctival sac at this stage. The surgeon, after scrubbing and draping, used 5% povidone iodine to cleanse the eyelids and periorcular skin. The eyelids including the eye lashes were draped. All cataract surgeries were performed in a dedicated ophthalmic theatre assisted by ophthalmic specialist nurses. Phacoemulsification was performed either using a peristaltic or venturi system. Postoperatively, patients were prescribed prednisolone acetate 1% two-hourly for two weeks which was then tapered off over a further four weeks. Topical fusidic acid was used for 10 days.

However, in 2000, following a cluster of four cases of endophthalmitis after cataract surgery during a period of three months, the practice of microbial prophylaxis was reviewed. Based on evidence existing at the time, and following an external investigation and extensive discussions with the hospital infection control team, a decision was made to use vancomycin in a dose of 1 mg in 0.1 mL of normal saline, to be injected intracameral into the capsular bag as the final step of cataract surgery.

A case of endophthalmitis was defined as one which presented within six weeks of cataract surgery, with typical clinical findings of intraocular inflammation, reduced vision, and with or without culture-positive vitreous biopsy. These cases were entered prospectively into a data collection system. The detection and management of a case of endophthalmitis was under the direct supervision of a consultant ophthalmologist. All but two cases of suspected endophthalmitis had emergency vitreous sampling along with intravitreal injection of antibiotics. One case with a visual acuity of perception of light, had pars plana vitrectomy with intravitreal antibiotics as per EVS study. The vitreous samples were analyzed according to standard practice, with the microbiologist using gram stain and culture on blood agar, chocolate agar, and Sabouraud’s agar. Gram stain identified Gram-positive or Gram-negative organisms. Aerobic organisms were identified using incubation of blood and chocolate agar plates at 35°C for five days. Anaerobic organisms were identified by incubation of blood agar plate at 35°C with increased carbon dioxide in the atmosphere. Sabouraud’s media was incubated at 35°C for 30 days to identify fungal organisms. Identified organisms had antibiotic sensitivity tests performed to identify susceptibility.

Cases of endophthalmitis following cataract surgery were identified from the data collected on the postoperative complication reporting system in the Ophthalmic Department. The incidences of endophthalmitis before (period A) and after the introduction of intracameral vancomycin after cataract surgery (period B) were calculated. The relative risk, relative risk reduction, and absolute risk reduction were calculated from the data.

**Results**

A total of 16,606 cataract surgeries with intraocular lens implantations were performed between January 1, 1998 and December 31, 2008. The distribution of cataract surgeries over the study period is shown in Figure 1. Between January 1, 1998 and December 31, 2000 (period A), prior to introduction of intracameral vancomycin, 3904 cataract surgeries were performed. Thirteen patients developed endophthalmitis during this period. Eleven of the 13 cases of endophthalmitis had vitreous sampling and intravitreal antibiotics and two cases had aqueous sampling and intracameral antibiotics. From January 1, 2001 to December 31, 2008 (period B), 12,702 cataract surgeries were performed using intracameral vancomycin at the end of cataract surgery. One patient during this period developed presumed endophthalmitis with a negative vitreous biopsy.

The incidence of endophthalmitis prior to the introduction of intracameral antibiotic prophylaxis was 0.3%. The annual incidence of endophthalmitis during the study period is shown in Figure 2. After the introduction of intracameral antibiotics, the incidence of endophthalmitis dropped to 0.008% (Table 1). This reduction was statistically significantly (Chi-squared test, 36.6; \( P < 0.0001 \)). The relative risk of developing endophthalmitis without intracameral antibiotic prophylaxis was 38 (95% confidence interval [CI]: 7–252). With the use of intracameral vancomycin, the absolute risk reduction was 292 cases of endophthalmitis per 100,000 cataract surgeries. The relative risk reduction was 97%. Five patients with clinical endophthalmitis had a microorganism identified on gram stain or culture. Eight patients had
negative gram stain and culture. Three vitreous samples grew *Streptococcus pneumoniae* (Table 2).

The experience of surgeons ranged from consultants with more than eight years of experience in performing cataract surgery, to middle-grade surgeons with more than four years of experience, and to junior-grade surgeons with less than four years of experience in performing cataract surgery. The incidence of posterior capsular rupture with vitreous loss as a complication of cataract surgery was similar over the study period at 2.1% during period A and 2.2% during period B. Other microbial prophylactic measures including use of 5% povidone iodine, and postoperative antibiotics remained unchanged in the two periods.

**Discussion**

Dickey et al have shown that anterior segment intraocular surgery frequently introduces bacteria into the anterior chamber. In 65% to 100% of healthy subjects, conjunctiva and adnexa were found to be colonized by microorganisms with the potential to cause endophthalmitis. The EVS found that in 82% of cases of endophthalmitis, the organism isolated was the same species as that found in the conjunctiva and lids of the same patient. In the 420 cases of endophthalmitis in the EVS series, 70% of the isolates were coagulase-negative, Gram-positive staphylococci. As in the EVS study, our study also showed the majority of the culture-positive cases of endophthalmitis to be Gram-positive organisms (Table 2).
Prophylactic measures to reduce postoperative endophthalmitis after cataract surgery are wide and varied. Several perioperative and postoperative methods have been recognized. Preoperative iodine antisepsis combined with preoperative and postoperative topical antibiotic therapy is considered the standard of care for this purpose. The last decade has seen the evolution of intracameral antibiotics as a prophylactic measure to prevent postoperative endophthalmitis after cataract surgery, claiming superior effectiveness compared with standard care. Intracameral antibiotics can achieve high concentrations in the anterior chamber but have the potential to cause corneal endothelial toxicity. Yoeruek et al studied the toxic effects of cefuroxime and vancomycin on human corneal endothelial cells, and found them safe in clinically used concentrations. Higher concentrations could cause irreversible cell death. The choice of intracameral antibiotic prophylaxis appears to be empirical, based on the fact that the majority of isolates are Gram-positive organisms. The first report of intracameral antibiotic prophylaxis was by Peyman et al using gentamicin. Gimbel reported no case of endophthalmitis in a series of 12,000 patients after using an infusion fluid of gentamicin together with vancomycin during cataract surgery. Garat et al in their study used intracameral cefazolin because it has a bactericidal effect against Gram-positive organisms. Montan et al used intracameral cefuroxime for microbial prophylaxis after cataract surgery for similar reasons. They argued that intracameral vancomycin is probably inappropriate due to its poor efficacy against Gram-positive enterococci and its exclusive capacity to combat infections caused by multiresistant Gram-positive bacteria. For this reason, the Centers for Disease Control and Prevention in the US has cautioned against prophylactic vancomycin in cataract surgery. Garat et al argued that cefazolin might induce less drug resistance when compared with cefuroxime. Their choice of cefazolin

### Table 1 Incidence of endophthalmitis before and after introduction of intracameral vancomycin prophylaxis

<table>
<thead>
<tr>
<th>Cataract surgeries</th>
<th>Endophthalmitis cases</th>
<th>Incidence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Period A 1998–2000 (Preintracameral vancomycin)</td>
<td>3904</td>
<td>13</td>
</tr>
<tr>
<td>Period B 2000–2008 (Postintracameral vancomycin)</td>
<td>12702</td>
<td>1</td>
</tr>
</tbody>
</table>

### Table 2 Characteristics of cases of presumed endophthalmitis after cataract surgery over the 11-year study period

<table>
<thead>
<tr>
<th>Case</th>
<th>Year</th>
<th>Age</th>
<th>Section</th>
<th>Intraoperative complication</th>
<th>Suture</th>
<th>Onset of symptoms</th>
<th>Intraocular antibiotic (Intravitreal or intracameral)</th>
<th>Organism on gram stain or culture</th>
<th>Visual acuity at last visit</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Period A</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case 1</td>
<td>1998</td>
<td>85</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Nil</td>
<td>Negative</td>
<td>HM</td>
</tr>
<tr>
<td>Case 2</td>
<td>1998</td>
<td>80</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>10.0 nylon</td>
<td>9 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/12</td>
</tr>
<tr>
<td>Case 3</td>
<td>1998</td>
<td>76</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/12</td>
</tr>
<tr>
<td>Case 4</td>
<td>1998</td>
<td>85</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>1 day</td>
<td>Gentamicin, Vancomycin</td>
<td>Gram-positive cocci</td>
<td>No PL</td>
</tr>
<tr>
<td>Case 5</td>
<td>1999</td>
<td>91</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>10 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/60</td>
</tr>
<tr>
<td>Case 6</td>
<td>1999</td>
<td>65</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Cefuroxime, Vancomycin</td>
<td>Psuedomonas aerginosa</td>
<td>6/9</td>
</tr>
<tr>
<td>Case 7</td>
<td>1999</td>
<td>52</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Teicoplanin, Ciprofloxacin</td>
<td>Nil</td>
<td>6/12</td>
</tr>
<tr>
<td>Case 8</td>
<td>2000</td>
<td>80</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>1 day</td>
<td>Vancomycin, Cefazidime</td>
<td>Streptococcus pneumoniae</td>
<td>No PL</td>
</tr>
<tr>
<td>Case 9</td>
<td>2000</td>
<td>69</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>10.0 nylon</td>
<td>5 days</td>
<td>Teicoplanin, Ciprofloxacin</td>
<td>Streptococcus pneumoniae</td>
<td>6/18</td>
</tr>
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<td>Case 10</td>
<td>2000</td>
<td>75</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>6 days</td>
<td>Cefuroxime, Vancomycin</td>
<td>Negative</td>
<td>6/36</td>
</tr>
<tr>
<td>Case 11</td>
<td>2000</td>
<td>60</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>3 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/9</td>
</tr>
<tr>
<td>Case 12</td>
<td>2000</td>
<td>78</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>14 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/12</td>
</tr>
<tr>
<td>Case 13</td>
<td>2000</td>
<td>93</td>
<td>Scleral tunnel</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>No PL</td>
</tr>
<tr>
<td><strong>Period B</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case 14</td>
<td>2008</td>
<td>73</td>
<td>Clear corneal</td>
<td>Nil</td>
<td>Nil</td>
<td>2 days</td>
<td>Vancomycin, Cefazidime</td>
<td>Negative</td>
<td>6/6</td>
</tr>
</tbody>
</table>
Intracameral vancomycin following cataract surgery

was also based on its cost being 41% less than that of
cefuroxime.17 The preliminary report of principal results
from a randomized, partially masked, multicentre study by
the ESCRS1 provided evidence that the risk of contracting
depthalmitis is significantly reduced by an intracameral
injection of cefuroxime at the end of surgery. The choice
of intracameral antibiotic in the ESCRS study was based
on the study by Montan PG et al of the effectiveness and
safety of cefuroxime.18,19

Our choice of vancomycin in 2000 was guided by the EVS
and local consultations. Intracameral vancomycin has been
implicated as a cause of cystoid macular edema. Axer-Siegel
et al in a randomized controlled trial investigated fluorescein
angiogram evidence of cystoid macular edema associated
with use of intracameral vancomycin during extracapsular
cataract extraction.20 They found a significant increase in
cidence of cystoid macular edema in the vancomycin
group compared with the control group. However, design
flaws, long surgical time, and attrition of patients have raised
questions about the accuracy of their study results. A more
recent randomized controlled trial examined the effects of
intracameral vancomycin and gentamicin on macular thick-
ness as measured by ocular coherence tomography.21 They
found no statistically significant increase in macular thick-
ness in the group that received intracameral vancomycin
and gentamicin. In view of the strong evidence in favor of
intracameral cefuroxime and concerns about the development
of vancomycin resistance, we are reviewing our choice of
intracameral antibiotic. With the advent of fourth-generation
fluoroquinolones, with their superior antibacterial activity, the
choice of antibiotics in the ESCRS study might be outdated.
O’Brien et al advocate moxifloxacin as a better choice for
intracameral antibiotic prophylaxis.22 However, safety studies
are available only in animal models.

A recent survey23 on antibiotic prophylaxis practice after
cataract surgery in the UK raises some interesting issues,
given that 37% of the respondents did not use intracameral
antibiotic prophylaxis. The practice of the 61% who did
not respond is unclear. The main criticism of the ESCRS
study was the lack of a subconjunctival cefuroxime arm. An
uncontrolled but important study in this context demonstrated
a threefold reduction in the rate of presumed infectious
depthalmitis when prophylactic practice was changed
from subconjunctival to intracameral cefuroxime injection
after cataract surgery.24 A 2007 survey of American Society
of Cataract and Refractive Surgery members found that 77%
were not using intracameral antibiotics, with the majority
opting instead for preoperative and postoperative topical
antibiotics.25 Specifically, fourth-generation fluoroquinolones
seemed to be a popular option.

We acknowledge that our study has several limitations
due to its retrospective nature. Wound placement, type of
intraocular lens, and suture placement are variables that can
possibly affect development of endophthalmitis. The above
details were not recorded in most of our patient records. Lack
of data on these variables could have introduced bias which
could potential affect the results. A prospective study would
be required to eliminate such bias.

In summary, there is significant research evidence that
intracameral antibiotic prophylaxis reduces the incidence of
depthalmitis after cataract surgery. Our study, although
retrospective, with an inherent risk of bias, adds support to this
body of evidence. The visual, emotional, and financial costs
dealing with endophthalmitis following phacoemulsification are
huge. All ophthalmic surgeons should recognize this potential
complication and adopt intracameral antibiotics as an essential
part of intraocular surgery. Until a commercially available, pre-
diluted, single-dose, licensed antibiotic formulation is available,
arrangements with local hospital pharmacies should be made.

Disclosures
This study was not funded, and the authors report no conflict
of interest, either financial or proprietary, in this work.

References
1. Endophthalmitis Vitrectomy Study Group. Results of the Endophthalmitis
Vitrectomy Study: A randomized trial of immediate vitrectomy and
of intravenous antibiotics for the treatment of postoperative bacterial
acute endophthalmitis after cataract surgery: A systematic review of the literature.
4. Miller JJ, Scott IU, Flynn HW Jr, Smiddy WE, Miller D.
Acute onset endophthalmitis after cataract surgery: A population-based study.
study of prophlaxis of postoperative endophthalmitis after cataract
surgery: Preliminary report of principal results from a European mul-
6. Dickey JB, Thompson KD, Jay WN. Anterior chamber aspirate cultures
after uncomplicated cataract surgery. Am J Ophthalmol. 1991;112:
278–282.
7. Speaker MG, Milch FA, Shah MK, Eisner W, Kreiswirth BN. Role
of external bacterial flora in the pathogenesis of acute postoperative
8. Boes DA, Lindquist TD, Firtsche TR, Kalima RE. Effects of povidone
iodine chemical preparation and saline irrigation on the perilimal flora.


