Results at seven years after the use of intracameral cefazolin as an endophthalmitis prophylaxis in cataract surgery

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Abstract
Background: To evaluate results after seven years using prophylactic intracameral cefazolin for the prevention of endophthalmitis in cataract surgery.

Methods: A prospective, observational study of all patients submitted to cataract surgery over the period January 1996 to December 2009. All cases of postoperative endophthalmitis over that period were reviewed. The patients were classified in two groups: Group 1 (11,696 patients) operated on between January 1996 and December 2002, Group 2 (13,305 patients) between January 2003 and December 2009 (in whom a 1 mg/0.1 bolus of intracameral cefazolin was instilled).

Results: During the study period, 76 cases of endophthalmitis were observed in Group 1, and seven in Group 2. The rate of postoperative endophthalmitis reduced from 0.63% to 0.05% with a cefazolin injection. The relative risk (RR) for endophthalmitis in Group 1 against group 2 was 11.45 [95% CI 5.72-22.84, p < 0.001].

Conclusions: An intracameral bolus injection of cefazolin (1 mg in 0.1 ml solution) at the conclusion of the cataract surgery significantly reduced the rate of postoperative endophthalmitis.

Keywords: endophthalmitis, cataract surgery, cefazoline, prophylactic ophthalmic surgery, cephalosporines

Background
Endophthalmitis remains a serious complication after cataract surgery, although prophylactic measures introduced in recent years have reduced the number of patients with this complication.

Currently there are two streams of opinion towards endophthalmitis prophylaxis, the use of fourth-generation quinolones (gatifloxacin and moxifloxacin) topically [1-4], or the introduction of intracameral cephalosporins, the latter being cefuroxime (a second generation cephalosporin), which is the most widely used and accepted [5-10]. However, our study group [11], as well as Garat et al [12,13], prefers the use of cefazolin (a first generation cephalosporin). Having previously studied the bacteria that cause endophthalmitis in our environment most frequently, we prefer cefazolin because of its higher frequency of gram-positive bacteria in our medium, and because it best covers infections by such bacteria. Furthermore, cefazolin shows no corneal toxicity at doses of 1 mg or 2 mg; its toxicity was established when doses of 5 mg or more were injected. We consider the risk of an infection caused by a cefazolin-resistant bacterium low, based on the bacteria cultured since 1994 and their antibiogram. We consider there not to be any risk of coverage of gram-negative bacteria by cefazolin, and its incidence in endophthalmitis was low in our Health Care District.

Our two groups are included in the Barcelona Endophthalmitis Group (GEB), formed by 38 public and private Hospitals in Catalonia (Spain), who have been studying the epidemiological factors of postoperative endophthalmitis and assessing the prophylaxis and treatment of endophthalmitis in our country since 2000. Appendix 1 gives a list of GEB participating hospitals and ophthalmologists.

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This study will present the results obtained after seven years of using intracameral cefazolin after cataract surgery at doses of 1 mg in 0.1 ml.

**Methods**

Since 1996, there has been an ongoing registration of all postoperative endophthalmitis patients at Hospital St Joan (Table 1). All cases of postoperative endophthalmitis were studied in the epidemiological unit of the hospital in order to determine their origin.

**Design**

A prospective observational study. The population comprised all patients submitted to un-combined cataract surgery in the period from January 1996 to December 2009. All cases of postoperative endophthalmitis were related. Only patients with phacoemulsification were included in the study in order to reduce any possible bias introduced by the technical cataract surgery. Technical surgery includes a sutureless, clear corneal incision of 3.2 mm using a venturi (Millenium, Bausch Lomb®) phaco unit.

For all patients included in our hospital, the prophylactic measures for reducing the bacterial flora of the conjunctiva since 1996 were:

- Topical 10% povidone-iodine in the skin of peri-orbital region
- Topical 5% povidone-iodine on the conjunctiva and eyelashes for a minimum of 1 minute
- Draping of the peri-orbital region and eyelashes
- Postoperative use of eye drops with tobramycin 3.00 mg/mL associated to dexametasone 1.00 mg/mL instilled every 4 hours (these eye drops were gradually tapered after 1 week), and diclofenac every 6 hours. These drops were continued at week 3, and the topical diclofenac was used for 1 month after surgery.

The patients were classified in two groups:

- Group 1 included patients operated on from January 1996 to December 2002 (11696 patients), during which time intracameral cefazolin was not used.
- Group 2 included patients operated on from January 2003 to December 2009 (13305 patients) when we used 1.00 mg/0.1 mL intracameral cefazolin was used at the end of cataract surgery.

**Antibiotic selection**

The choice of cefazolin as an intracameral antibiotic was based on bacterial and antibiogram studies in endophthalmitis cases since 1996 (Table 2), and the antibiogram (Table 3).

Table 3 was created from the analysis of an antibiogram of 14,626 cultures obtained from samples of patients with community- and hospital- acquired infections (urinary infections, pneumonia, sepsis, etc.), which occurred during 2002 in the population serviced by our Health Care District. In the table we presented only the results obtained with the bacteria cultured in endophthalmitis observed in our area, and compared with the prophylactic antibiotics most frequently used for endophthalmitis in Spain (cefazolin, cefuroxime, vancomycin, and levofloxacin). At this point we should point out that in our Hospital in cases of allergy to cephalosporines we used vancomycin.

The dose of cefazolin (1 mg/0.1 ml) was based on our calculations that this anterior chamber concentration of cefazolin exceeded the minimum inhibitory concentration (MIC) for susceptible bacteria.

**Inclusion criteria**

Patients in our dependent Health care district who were submitted to cataract surgery by phacoemulsification by clear cornea incision.

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**Table 1 Description of cases of endophthalmitis since 1996**

| Group 1 1996-2002 | 1996 | 1997 | 1998 | 1999 | 2000 | 2001 | 2002 | Total |
|-------------------|------|------|------|------|------|------|------|-------|
| Cases*            | 8    | 9    | 9    | 11   | 12   | 13   | 14   | 76    |
| Percentage        | 0.637% | 0.613% | 0.629% | 0.664% | 0.634% | 0.687% | 0.692% | 0.649% |
| Number of operations** | 1256 | 1467 | 1430 | 1656 | 1892 | 1972 | 2023 | 11696 |

| Group 2 2003-2009 | 2003 | 2004 | 2005 | Total |
|-------------------|------|------|------|-------|
| Cases*            | 1    | 1    | 1    | 7     |
| Percentage        | 0.058% | 0.052% | 0.055% | 0.052% |
| Number of operations** | 1707 | 1911 | 1812 | 13305 |

Cases* = number of cases of endophthalmitis (proven + unproven) occurring each year

Patients** = Number of cataract interventions carried out each year
Table 2: Bacteria cultured in cases of endophthalmitis in the two groups.

| Bacterial species                  | Number of cases (percentage) 1996-2002 | Number of cases (percentage) 2003-2009 |
|------------------------------------|----------------------------------------|----------------------------------------|
| Negative cultures                  | 26 (34.21%)                            | 2 (28.57%)                             |
| Gram positive (96.00%)             |                                        |                                        |
| Staphylococcus epidermidis         | 37 (74.00%)                            |                                        |
| *Staphylococcus aureus* meticillin| 5 (10.00%)                              |                                        |
| sensitive                          |                                        |                                        |
| Streptococcus pneumoniae           | 2 (4.00%)                               |                                        |
| Streptococcus viridans             | 1 (2.00%)                               |                                        |
| Bacillus spp                       | 3 (6.00%)                               |                                        |
| Corynebacterium                    | 1 (2.00%)                               |                                        |
| Klebsiella pneumoniae              | 1 (2.00%)                               |                                        |
| Pseudomonas aeruginosa             | 1 (14.28%)                              |                                        |
| Proteus mirabilis                  | 1 (2.00%)                               |                                        |
| E Colli                            | 1 (2.00%)                               |                                        |
| Staphylococcus epidermidis         | 37 (74.00%)                             |                                        |
| Serratia marcesens                 | 1 (2.00%)                               |                                        |
| Pseudomonas aeruginosa             | 1 (14.28%)                              |                                        |
| Proteus mirabilis                  | 2 (28.57%)                              |                                        |

Table 3: Sensitivity to antibiotics of the most frequent bacteria in our Area in 2002

| Gram negative | Cefazolin (s2 ≥ 4)* | Cefuroxime (s2 ≥ 4) | Levofloxacin (s2 ≥ 4) | Vancomycin (s4 ≥ 32) |
|---------------|---------------------|---------------------|-----------------------|-----------------------|
| Staphylococcus aureus | 100% (s2 ≥ 4) | 96% (s2 ≥ 4) | 65% (s2 ≥ 4) | 66% (s2 ≥ 4) |
| Staphylococcus pneumoniae | 96% (s2 ≥ 4) | 97% (s2 ≥ 4) | 89% (s2 ≥ 4) | 100% (s4 ≥ 32) |
| E Colli       | 85% (s4 ≥ 32)       | 87% (s4 ≥ 32)       | 71% (s4 ≥ 32)       | 90% (s4 ≥ 32)       |
| Pseudomonas aeruginosa | 18% (s4 ≥ 32) | 15% (s4 ≥ 32) | 65% (s2 ≥ 4) | 65% (s4 ≥ 32) |
| Proteus mirabilis | 79% (s4 ≥ 32) | 82% (s4 ≥ 32) | 70% (s4 ≥ 32) | 100% (s4 ≥ 32) |
| Klebsiella pneumoniae | 75% (s4 ≥ 32) | 79% (s4 ≥ 32) | 79% (s4 ≥ 32) | 95% (s4 ≥ 32) |

* ≤ 0.60 = MIC (minimum inhibitory concentration) = Critical concentrations in µg/ml (critical concentration of susceptibility, critical concentration of resistance), according to the recommendations of MENSURA to the interpretation of antibiogram.

** = Unknown, we do not know the mechanisms of resistance, or the measure for resistance.

Exclusion criteria
Patients admitted for extracapsular cataract surgery. Patients allergic to cephalosporins and to whom we gave intracameral vancomycin.

Ethics
Local ethics committee approval the study CEIC HUSJ-2002-DEC-1022

Definition of acute postoperative endophthalmitis
Following the criteria [14] in the Endophthalmitis Vitrectomy Study (EVS), the ophthalmologist diagnosed presumed acute endophthalmitis. If a positive culture of a vitreous sample was obtained, we defined the case as a proven acute endophthalmitis. In all proven and unproven cases, the patients had swollen lids, pain and an opaque vitreous.

Microbiological method
Vitreous samples obtained by the ophthalmologist were immediately processed, then a Gram stain was carried out and the sample cultivated in petri dishes. Antibiogram susceptibility was measured according to the criteria laid down by the Spanish Committee for the Standardization of the Sensitivity and Resistance to Antimicrobials (MENSURA) [15-17] which includes members of the Spanish Society of Chemotherapy and the Spanish Society for Infectious Diseases and Microbiology [15]. This is a body with responsibilities similar to those of the National Committee for Clinical Laboratory Standards [18].

Statistical analyses for descriptive statistics were performed using SPSS statistical software (version 17.0). The data obtained was analysed with frequency and descriptive statistics. Values are expressed as mean ± SEM, and statistical significance was determined using the Student’s t-test for paired data.

Results
Demographic results in the two groups of patients
Group 1. Formed by patients without intracameral instillation of cefazolin. This group included a total of 11,696 patients, with a median age of 69.8 ± 7.55 years (53-89 years); a total of 6785 (58.01%) were females. The difference between groups was not statistically significant in the Student’s t-test.

Group 2. Formed by patients with intracameral instillation of cefazolin at doses of 1 mg/0.1 mL. This group included a total of 13,305 patients, with a median age of 66.17 ± 7.83 years (53-81 years); a total of 7717 (58.00%) were females. The difference between groups was not statistically significant in the Student’s t-test.

Endophthalmitis cases
In Group 1 there were 76 postoperative endophthalmitis cases, at a median elapsed time of 5.37 ± 2.33 days after...
surgery. In 16 (69.77%) cases, cultures were positive for the following sub-classified gram-positive cultured bacteria: 9 cases (39.13%) of Staphylococcus epidermidis, 4 cases (17.39%) of staphylococcus aureus and 2 cultures (8.70%) positive for Streptococcus spp. The gram-negative bacteria gave a culture positive for Klebsiella pneumoniae (4.35%). Group 2, there were seven postoperative endophthalmitis cases, out a median elapsed time of 5.41 ± 2.29 days after surgery. The statistical study of differences in respect to first group were significant at p < 0.001, RR: 11.45 [95% CI 5.72-22.84].

The bacteria cultured were:

- The first case involved patients with diabetes mellitus type II, treated with insulin for 30 years, with macro vascular disease with symptoms of intermittent claudication. The culture was positive for a Gram-negative bacterium (Klebsiella pneumoniae).
- The second case was a patient who lived alone and who had serious social problems; the patient also had domestic animals at home. The culture was positive for a Gram-positive anaerobic bacterium (Corynebacterium).
- The third case, positive for Proteus mirabilis, occurred in a 68-year-old woman with type II diabetes mellitus of long evolution (22 years), with poor glycaemic control and peripheral vascular macroangiopathy.
- The fourth case, positive for Proteus mirabilis, occurred in a 77-year-old man with type II diabetes mellitus and poor glycaemic control.
- The fifth case, positive for Pseudomonas aeruginosa, occurred in a 73-year-old man with viral C-hepatitis.
- Negative cultures: two cases of negative cultures (28.57%) were observed in this period of time.

In the groups of patients that received intracameral cefazolin, there were no cases observed of toxic effect at the corneal or retina levels, nor was there any hypersensitivity reaction.

**Statistical analysis**

The relative risk for presenting with endophthalmitis in Group 1 compared with Group 2 was 11.45 [95% CI 5.72-22.84, p < 0.001]. When limiting the analysis to proven cases (50 cases in Group 1, against 5 cases in Group 2), the estimators of the relative risk were 14.07 [95% CI 7.68 - 24.48, p < 0.001].

**Visual acuity**

1. In Group 1 (period 1996-2002) 38 over 76 patients (50%) had a final visual acuity (VA) over 0.1 and six patients (7.9%) had VA > 0.4 on the Snellen charts. There were ten patients with no light perception (four cases with negative culture, one case produced by Seratia marcesens, and five cases of Streptococcus pneumoniae). Staphilococcus epidermidis in 18 cases caused a final VA between 0.1 and 0.4, and in 14 cases a final VA between light perception and 0.1; the Staphilococcus aureus predominately produced a final VA between light perception and 0.1 (four cases) and only one case with final VA > 0.4 on the Snellen charts (Table 4).

2. In Group 2 (period 2003-2009) four patients had final VA of no light perception (two cases with gram-negative bacteria, one case with negative culture and one case produced by gram-positive corynebacterium). In the other three cases, the final VA was inferior to 0.1 on the Snellen charts (two cases with gram-negative bacteria and one with a negative culture) (Table 4).

**Discussion**

In our health care district we had a high level of endophthalmitis. Previously, the intracameral use of antibiotics, as we point out in the present study in group 1, lead to the incidence of endophthalmitis at a rate of 0.649%. This value is higher than that reported in other countries [2,6,8,14], but the incidence in Spain was also higher than other study groups, Garcia-Saenz et al [19] found an incidence of 0.59% (95% CI, 0.50%-0.70%) between January 1999 and September 2005, and Garat et al showed an incidence of 0.422% (95% CI 0.279-0.613) in their studies [12,13]. Because our centre had an excessive number of endophthalmitis cases despite using all means of regular prophylaxis (a sterile ophthalmology operating room, povidone-iodine in skin (at 10%) and conjunctival sac (at 5%) with few surgical intraoperative complications etc.), we decided to use intracameral antibiotics after cataract surgery.

Peyman et al. published the first report of successful prophylactic bolus injections of antibiotics into the anterior chamber in 1977 [20]. Despite the efficacy of the injections, the technique later becomes forgotten about. It is well-established that the source of most infecting agents is the patients’ ocular flora; the most frequently reported being bacteria gram-positive, coagulase-negative, or positive staphylococcus.

Swedish physicians have pioneered the use of intracameral cefuroxime since 2002, with excellent outcomes in 400,000 surgical interventions. Montan et al. published the efficacy of cefuroxime 1 mg intracameral [5,6], a practice that has lowered the rate of postoperative endophthalmitis from 0.26% to 0.06%. The large, prospective, multi-centred study, the European Society of Cataract Refractive Surgeons (ESCR) confirmed the Swedish experience, finding that an injection of cefuroxime at the end of the surgery reduced endophthalmitis rates to just 0.05% [8-10].

We chose to use cefazolin, which is recommended by the Department of Microbiology and the Infectious
Diseases Committee of our Hospital, rather than cefuroxime because it is a first generation cephalosporin and has a wide range of activity against gram-positive bacteria in our Health Care District, rather than a second-generation cephalosporin, such as cefuroxime. A study in our Health Care District by Vila-Corcoles et al [21], showed an increased resistance of Streptococcus pneumoniae to cefuroxime but not to cefazolin.

It is interesting to observe in Table 3 that cefazolin and cefuroxime present a similar antibiotic resistance pattern, with regard to both gram-positive and gram-negative bacteria. Therefore, we still believe that cefazolin is a good option as a prophylaxis for endophthalmitis, and its substitution for cefuroxime is not important for reducing the number of cases of endophthalmitis. However, by using cefazolin, there is still a risk of infection by gram-negative bacteria, which is why the majority of proven cases of endophthalmitis in Group 2 (4/5 cases were gram-negative) were caused by this type of bacteria. This is clearly a limitation in the use of cefazolin as a prophylaxis for endophthalmitis.

The lack of availability of eye-drops of fourth-generation quinolones in Spain is the most obvious reason for the preference for intracameral antibiotics in the prophylaxis of endophthalmitis. Since June 2010, a moxifloxacin eye-drop has been available in Spain, with more possibilities of prophylaxis.

Results show that final VA was worse in Group 2 than in group 1. The explanation for this can be found in the type of bacteria that cause endophthalmitis. We found Corynebacterium, Klebsiella pneumoniae, Pseudomonas aeruginosa, and Proteus mirabilis in group 2. All these bacteria were poorly sensitive to cefazolin. Negative culture was observed in a 34.21% of cases in group 1 and 28.57% in Group 2, similar to other studies, such as the Endophthalmitis Vitrectomy Study (EVS), with values near to 30%.

Conclusions
Intracameral bolus injection of cefazolin in cataract surgery demonstrated prophylactic efficacy in diminishing the rate of postoperative endophthalmitis in our hospital, at two doses of 1 mg in 0.1 ml solution.

Furthermore, a close relationship between with the Department of Microbiology, and the Infectious Diseases Committee of our Hospital is essential for developing a proper antibiotic prophylaxis.

Further studies are needed with a larger number of patients in order to fully determine the effectiveness of cefazolin and its non-toxicity at corneal endothelium level.

Appendix 1
Hospitals that form the Barcelona Endophthalmitis Group (GEB). http://gebcn.org/geb.html
01. Hospital Sant Pau: Dr. Jesus Díaz
02. Hospital Clinic: Dra. Socorro Alforja - Dr. Joan Giralt
03. Hospital Bellvitge: Dr. Marc Rubio
04. Hospital Vall d’Hebron: Dr. Alex Fonollosa
05. Hospital General d’Hospitalet: Dra. Magela Garat
06. Hospital Germans Trias i Pujol (Can Ruti): Dr. Anglada
07. Hospital Parc Taulí: Dr. D. Muntaner - Dra. MªT. Sellares - Dra. C. Guardia
08. Mutua de Terrassa: Dra. Silvia Freixes
09. IMAS (H. Esperança i Mar): Dr. Daniel Vilaplana - Dr. Poposki
10. Hospital St. Rafael: Dra. Ciprés
11. Hospital Viladecans: Dr. Sergi Sedó
12. Institut Català de la Retina: Dra. Sust
13. Institut Oftalmic de Barcelona: Dra. Laura Sararols
14. Instituto de Microcirugía Ocular: Dra. Isabel Nieto
- Dra. Elena Arrondo
15. Hospital General de Cataluña: Dr. Rouras
16. Clínica Barraquer: Dr. Remberto Escoto
17. Hospital Espirit Sant: Dr. Rouras - Dr. Amias
18. Hospital Granollers: Dr. Jarel Hernecki
19. Hospital de Terrassa: Dr. Jose Juan Escobar - Dr. Jose Pradas
20. Institut Condal d’Oftalmologia (ICO): Dra. Sararols
21. Hospital Municipal de Badalona: Dra. Estela Barnola
22. OFTALMO: Dr. J. Cabot
23. Clínica de Vic i Hospital General de Vic: Dr. Manuel Amén
24. Consulta GMOftalmo: Dr. Gallofré
25. Hospital de Nens de Barcelona: Dr. J. Flors Despranell
26. Clínica Quiron: Dr. Aureli Rilo
27. Institut Oftalmològic Tres Torres: Dr. Emili Juarez
28. Hospital de Mollet: Dr. Francisco Goñi
29. Hosp. Universitari Josep Trueta: Dra. Flor Escalada
30. Hosp. Universitari Sant Joan de Reus: Dr. Pere Romero Dra. Isabel Mendez
31. Hosp. Universitari Joan XXIII Tarragona: Drs. Perez i Pardo
32. Hosp Verge de la Cinta de Tortosa: Dr. J Colome
33. Hosp Son Dureta, Palma de Mallorca: Dr. José Luis Olea
34. Clínica Sagrada Familia i Corachan: Dr. Carlos Martin
35. Clínica Plató: Dr. Joan Pujol - Dr. Sergio Miserais
36. Institut Oftalmològic de Menorca: Dr. Pere Vilallonga
37. Hospital Sant Joan de Deu de Martorell: Dra. Mª Carmen del Águila
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Authors’ contributions
PRA contributed to the study design, the researched data interpretation, and contributed to the interpretation of the study findings. All authors read and approved the final manuscript.

Competing interests
The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper. No financial support was received.

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