

Retrospective study in 608 cases on the rate of surgical site infections after orbital surgery without prophylactic systemic antibiotics

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ABSTRACT

Objective To determine postoperative surgical site infection (SSI) rates in three separate patient groups who underwent orbital surgery without prophylactic systemic antibiotics.

Study design Single-centre retrospective descriptive case series.

Study population We studied the notes of 639 consecutive patients who had undergone orbital surgery in our hospital from 2009 through 2013. All patients belonged to either of three groups: (1) clean orbital surgery (n=226); (2) clean orbital surgery with implant (n=290); (3) clean-contaminated surgery (n=92). Thirty-one patients were excluded.

Results Of the total of 608 patients, without systemic antibiotic prophylaxis, only five were diagnosed with SSI 5/608 (0.82%): 1/226 in the 'clean' group, 3/290 in the 'clean-with-implant' group and 1/92 in the 'clean-contaminated' group. All five patients with SSI were effectively treated with antibiotics.

Conclusion In this study 'clean', 'clean-with-implant' and 'clean-contaminated' orbital surgery was safely performed without prophylactic antibiotics. Where postoperative infection did occur, the patients were effectively treated with systemic antibiotics. We suggest to restrict the administration of systemic antibiotic prophylaxis in orbital surgery.

INTRODUCTION

Recently, WHO advised to minimise the use of antibiotics because of increasing resistance and lack of new generations of antibiotics.¹ However, antibiotic drugs continue to be widely used both for prophylaxis and treatment of infection before, during or after surgical procedures, while standards of perioperative care regarding treatment regimens vary between countries. The literature about the use of antibiotics in oculoplastic and orbital surgery is scarce.²

Recent retrospective studies about eviscerations and enucleations show that withholding antibiotic prophylaxis from these patients did not increase the number of surgical site infections (SSIs).²⁻⁴ Similarly, a recent retrospective study from our group did not show any SSI in 186 enucleations (without prophylactic antibiotics) by a single surgeon.⁵ We are not aware of studies about the prophylactic antibiotics use in other intraorbital surgical procedures.

In the National Institute for Health and Care Excellence update (February 2017) on (general)

SSIs, it was advised to provide prophylactic antibiotics in clean-contaminated and in clean surgery with implants or prosthesis, by giving a single dose of intravenous antibiotics prior to incision on starting anaesthesia.⁴ In our hospital, however, we have not routinely used prophylactic systemic antibiotics in orbital surgery for clean, clean with implant or clean-contaminated surgery. The only indications in our institute for preoperative or perioperative systemic antibiotics are severely immunocompromised patients and/or patients with an active infection such as endophthalmitis with corneal perforation, orbital cellulitis or patients with increased risk of endocarditis.

In this study, we evaluate the postoperative infection rate of orbital surgery without prophylactic systemic antibiotics in a retrospective case series.

METHODS

We retrospectively analysed the records of all patients who underwent orbital surgery in our hospital between 2009 and 2013. We included patients who underwent orbital decompression, orbitotomy for tumour excision, enucleation, evisceration, exenteration, orbital biopsy and other orbital (reconstructive) surgery. Patients required a follow-up of at least 30 days after the procedure. Patients were categorised in three groups according to the type of orbital surgery: 'clean' orbital surgery, 'clean-with-implant' orbital surgery and 'clean-contaminated' orbital surgery. Of each patient we recorded the age, gender, American Society of Anesthesiologists (ASA) classification, type of surgery, use of systemic antibiotics and occurrence of postoperative SSI.

DEFINITIONS

- ▶ *Clean*: an uninfected operative wound in which no inflammation is encountered, and the respiratory tracts are not entered.
- ▶ *Clean with implant*: as stated, but with implant of an allogenic material.
- ▶ *Clean-contaminated*: operative wounds in which the respiratory tracts are entered under controlled conditions and without unusual contamination, provided no evidence of infection or major break in technique is encountered.
- ▶ SSI is defined as an event within 30 days after the orbital surgical procedure. We retrospectively categorised to the Centers for Disease Control and Prevention (CDC) definitions. This



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Table 1 Characteristics of the patient groups studied

	Clean N	Clean with implant N	Clean- contaminated N	Total
Mean age at surgery (range)	54 (1–87)	58 (4–98)	50 (3–87)	
Gender				
Male	108	152	20	280
Female	118	138	72	328
ASA classification				
No of patients ≤ 2	198	233	82	513
No of patients > 2	28	57	10	95
Total	226	290	92	608
Surgery				
Biopsy	71			
Debulking	18			
Tumour excision	96			
Decompression*	11		92	
Exenteration	6			
Socket reconstruction†	17	57		
Other	7			
Evisceration		159		
Enucleation		74		
Total	226	290	92	608
Excluded patients				
Due to prophylaxis	3	7	3	13
Due to preoperative infection	4	8	6	18
Total	7	15	9	31

*Eleven patients (6 bilateral) had solely lateral wall decompression and 93 (76 bilateral) had decompression with an opening to the nose or one of the sinuses.

†Seventeen patients had socket reconstruction without an alloplastic implant, 57 with alloplastic implant.

ASA, American Society of Anesthesiologists.

resulted in wound infections as a superficial SSI, preseptal cellulitis as deep incisional SSI and orbital cellulitis as organ/space SSI.⁶

- ▶ **ASA physical status:** assessment by the anaesthesiologist of the patient's preoperative physical condition using the ASA Classification of Physical Status.⁶

All patients with a contaminated, dirty or infected orbital disease were excluded. All patients having antibiotic prophylaxis in any form were excluded. We examined the records of 639 patients; 608 of these did not have systemic antibiotics prior to surgery, while 31 had received antibiotic treatment or had a dirty wound and were excluded.

RESULTS

Table 1 shows the patient characteristics, including age, gender, ASA classification and type of surgery in each study group.

Table 2 shows the incidence of SSI in the three patient groups and the ASA classification.

Infection of the surgical site occurred in one patient in the 'clean' group, three in the 'clean with implant' group and one in the 'clean-contaminated' group. SSI occurred more frequently in ASA ≤ 2 patients (n=5/508) than in ASA > 2 patients (0/95). A group of 31 patients was treated with antibiotics preoperatively: 13 with systemic antibiotic prophylaxis and 18 with antibiotics as therapy for underlying disease; some had prolonged therapy after endophthalmitis.

The patients with a SSI are shown in **table 3**.

Table 2 Outcome of SSIs found in the three studied patient groups

	Clean N	Clean with implant N	Clean- conta- minated N	Total
SSI in ASA groups				
No of patients ASA ≤ 2 without SSI	197	230	81	508
No of patients ASA ≤ 2 with SSI	1	3	1	5
No of patients > 2 without SSI	28	57	10	95
No of patients ASA > 2 with SSI	0	0	0	0
Total	226	290	92	608

ASA, American Society of Anesthesiologists; SSI, surgical site infection.

Of the five patients with clinical signs suggestive of infection, two had been clinically diagnosed with wound infection, two with preseptal orbital cellulitis and one with all the signs of an orbital cellulitis.

Two other patients from the group of 608 patients did not show signs of infection, but their general physicians had started oral antibiotics without consulting us because of 'bloody discharge' after a medial wall decompression and because of 'conjunctival swelling' after orbital surgery, respectively. We therefore considered these two cases as not having a surgical site infection.

DISCUSSION

This study is limited by the retrospective design, the low number of patients per treatment group and the heterogeneity of the procedures. We cannot exclude the possibility that, due to the low rate of SSI as observed in the study, this is explained by chance alone, but our results indicate that orbital surgery without the use of prophylactic antibiotic therapy is rarely complicated by SSIs. All patients who exhibited SSI responded well to subsequent administration of antibiotics.

Since the late 1980s, no new antibiotics have been discovered (the 'Discovery Void') while drug resistance is increasing.¹⁷ Therefore, the need to control the use of antibiotics becomes more and more important. Apart from resistance increase, antibiotic use is associated with a wide variety of side effects and complications such as anaphylaxis, Stephens-Johnson syndrome, gastrointestinal distress, nephrotoxicity, ototoxicity, hepatic toxicity, cardiac toxicity, renal insufficiency, neutropenia, neuritis, myelosuppression, vaginitis, sun sensitivity, and allergic response. Antibiotics may also interact unfavourably with other medication.^{3,8}

In 2005, results were published of a prospective randomised trial regarding the efficacy of adjuvant oral antibiotics preoperative next to one dose of systemic perioperative prophylactic antibiotics in colorectal surgery.⁹ No difference in SSIs was found between groups, while antibiotic use was associated with an increase in side effects, such as postoperative vomiting, nausea and abdominal pain. The side effects were significantly lower in the group of patients who did not receive any adjuvant oral preoperative antibiotics.⁹

The effect of postoperative antibiotic prophylaxis has been studied in a retrospective, multicentre, comparative case series.³ Of 644 eye eviscerations or enucleations, 381 had been treated with postoperative antibiotic prophylaxis and 263 had not. In the first group, one presumed case of orbital cellulitis had occurred (0.26%) and also one (0.38%) in the second group. Both groups received one dose of perioperative antibiotic prophylaxis. A recent study of Pariseau compared 480 cases of enucleation and evisceration: in the group of 70 patients with

Tabel 3 SSI in 608 patients after orbital surgery without antibiotic prophylaxis

Patient	Age	Gender	ASA	Surgery	Group	SSI	Days after surgery	Treatment	Culture
1	39	F	1	Lateral orbitotomy for cavernous hemangioma	cl	Wound infection	2	Oral amoxicillin/clavulanic acid	None
2	59	F	1	Evisceration for painful blind eye; acrylic implant	c+i	Orbital cellulitis	3	Intravenous amoxicillin/clavulanic acid; oral clindamycin	<i>Staphylococcus aureus</i>
3	34	M	1	Evisceration after trauma; acrylic implant	c+i	Preseptal cellulitis	4	Oral amoxicillin/clavulanic acid	Culture negative
4	65	M	2	Evisceration after complications of intraocular surgery; acrylic implant	c+i	Preseptal cellulitis	5	Oral azithromycin	None
5	80	F	2	Orbital decompression medial wall for dysthyroid optic neuropathy	cc	Wound infection	7	Oral amoxicillin/clavulanic acid	None

Patient 6 got preoperative antibiotic prophylaxis, amoxicillin/clavulanic acid and was excluded from the study. ASA, American Society of Anesthesiologists; SSI, surgical site infection; cc, clean-contaminated; c+i, clean with implant; cl, clean.

perioperative intravenous antibiotics, the SSI rate was 1.4%; and in the group of 411 cases not given perioperative intravenous antibiotics, the SSI rate was 1.1% for eviscerations with implants and 0.7% for enucleations with implants. In the systematic review in the same paper, the SSI rate in other studies about evisceration was between 0.6% and 7%, and between 0% and 13% for enucleation.²

Several other studies on prophylactic (preoperative or postoperative) antibiotics regarding surgery of the jaws, ears and dental surgery also failed to show evidence for antibiotic prophylaxis to prevent SSIs.^{10 11}

In our single-centre retrospective case series without systemic prophylactic antibiotic treatment, patients with almost all types of orbital surgery were included. The rate of SSI proved comparable with those reported by other studies, with and without preoperative and/or perioperative prophylactic antibiotics.^{2 3}

We prescribe postoperative systemic antibiotics (usually a combination of oral amoxicillin and clavulanate) only in case of evident or suspected wound infection (deep or superficial) or evident or suspected orbital cellulitis (organ space as defined by the CDC).¹² We feel there is no conclusive evidence that demonstrates the usefulness of systemic antibiotics as a prophylaxis in orbital surgery, while it is known that adverse side effects occur relatively frequently.

In our series, 'clean', 'clean surgery with implant' and 'clean-contaminated' surgery in the absence of prophylactic systemic antibiotic treatment could be compared. No substantial differences with respect to postoperative infection rate were found, although numbers per group were low, and the groups itself are heterogenous and therefore limiting the conclusions to be drawn.

In conclusion, our retrospective series shows that orbital surgery may be performed safely without prophylactic systemic antibiotics. The small proportion of cases (0.82%) with presumed signs of postoperative infection could be treated effectively with standard antibiotics. We have restricted the use of preoperative and perioperative systemic prophylactic antibiotics to severely immunocompromised patients. This, however, also requires further study, preferably through a prospective trial.

Contributors All authors substantially contributed to the conception or design of the work, or the acquisition, analysis or interpretation of data; drafted the work or revised it critically for important intellectual content; and approved the final version published.

Competing interests None declared.

Patient consent for publication Not required.

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Data sharing statement Individual participant data that underlie the results reported in this article, after deidentification. This is including the data of the excluded patients. Interested orbital surgeons can request the anonymised data by mail, when a methodologically sound proposal is provided and has been approved by our independent review committee. Proposals should be directed to r.dekeizer@oogziekenhuis.nl. To gain access, data requestors will need to sign a data access agreement.

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