Aspirin and warfarin therapy in oculoplastic surgery

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Abstract

Background/aims—There are no nationally agreed guidelines on preoperative management of patients who are on aspirin or warfarin therapy. There is considerable evidence that complication rates in anticoagulated patients are low whereas there are higher rates of thromboembolic complications in those whose therapy is manipulated. This survey aimed to establish oculoplastic specialist and non-specialist ophthalmic surgeons’ current management practice of patients before oculoplastic surgery who are taking aspirin or warfarin and to assess the rate of complications in these patients.

Method—An anonymous postal questionnaire survey of all ophthalmic consultants and specialist registrars in the Wessex region along with oculoplastic specialists in the Southern region.

Results—The overall response rate was 92%. Preoperative management was influenced both by type of operation and by type of surgeon. A statistically significant higher proportion of surgeons would consider altering warfarin compared with aspirin treatment. For all procedures, non-specialists are unlikely to stop aspirin therapy, and are less likely to stop warfarin before all procedures apart from dacrocystorhinostomy. A significant proportion of surgeons (18%) would allow insufficient time for the coagulation status of the patient to change after altering treatment. In those who gave an indication of their practice, aspirin was stopped an average of 10 days before surgery (range 2–30, SD 7 days). Warfarin was stopped an average of 3 days before surgery (range 1–10 days, SD 2 days). However, 18% of surgeons volunteer that they would allow insufficient time for the coagulation status of the patient to change.

Conclusions—In this survey, at least half the surgeons questioned would consider stopping warfarin before oculoplastic procedures. Over half of all surgeons have seen complications related to aspirin or warfarin, some of which were serious. A suggested approach to minimising patient risk is given.

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There are no nationally agreed guidelines on the appropriate action to take preoperatively with patients on aspirin or warfarin therapy. In cataract surgery, the perioperative complication rate in such patients is low (0–1%) and outcome is not adversely affected.6–9 Despite this, a significant proportion of surgeons discontinue such therapy before cataract surgery resulting in serious or fatal systemic thromboembolic complications in 4.5%.1–3

There are few data available, however, with respect to oculoplastic surgery. This survey was designed to establish how oculoplastic specialist and non-specialist ophthalmic surgeons manage patients on aspirin or warfarin therapy undergoing oculoplastic surgical procedures. It also aimed to establish the frequency of local and systemic complications in order to assess whether these patients are being put at risk by this practice.

Methods

A questionnaire was sent to all ophthalmic consultants and specialist registrars in the Wessex region (47) along with 15 oculoplastic specialists in the Southern region.

Results—The overall response rate was 92% and 100% of the oculoplastic specialists responded. For each question, the results indicate the percentage of the total replies received.

Overall, 93% of surgeons did not mention that they would obtain advice before altering treatment and 74% did not specify that the indication for aspirin or warfarin would influence the decision to alter treatment. The minimum period required in order to alter coagulation status before surgery is 7 days for aspirin and 3 days for warfarin. A significant proportion of surgeons (18%) would allow insufficient time for the coagulation status of the patient to change after altering treatment. A considerable proportion of surgeons (54%) reported that they had seen complications as a result either of stopping or continuing anticoagulation therapy.

Conclusions—In this survey, at least half the surgeons questioned would consider stopping warfarin before oculoplastic procedures. Over half of all surgeons have seen complications related to aspirin or warfarin, some of which were serious. A suggested approach to minimising patient risk is given.

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Table 1 Adjustment of treatment by oculoplastic specialists and non-specialists for different procedures. Numbers of surgeons who sometimes or always stop treatment with percentage of total replies for each question.

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Non-oculoplastic specialists</th>
<th>Oculoplastic specialists</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Stop aspirin</td>
<td>Stop warfarin</td>
</tr>
<tr>
<td>Ectropion repair</td>
<td>9 (21%)</td>
<td>20 (50%)</td>
</tr>
<tr>
<td>Entropion repair</td>
<td>8 (20%)</td>
<td>19 (48%)</td>
</tr>
<tr>
<td>Ptosis procedures</td>
<td>13 (35%)</td>
<td>25 (69%)</td>
</tr>
<tr>
<td>DCR</td>
<td>23 (57%)</td>
<td>34 (87%)</td>
</tr>
</tbody>
</table>

Table 2 Risk management for patients on aspirin and warfarin treatment undergoing oculoplastic procedures.

<table>
<thead>
<tr>
<th>Indication</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heparinise</td>
<td>Stop aspirin, stop warfarin</td>
</tr>
</tbody>
</table>

For all procedures, non-specialists are unlikely to stop aspirin therapy (Table 1). Similar proportions of non-specialist and oculoplastic surgeons are likely to stop warfarin before a DCR (87% and 84% respectively). In contrast, fewer non-specialist than oculoplastic specialist surgeons sometimes or always stop warfarin before ptosis procedures (69% and 78% respectively), entropion repair (48% and 60% respectively), and ectropion repair (50% and 67% respectively).

In those patients in whom warfarin therapy had been discontinued, around 40% of all surgeons would consider heparinisation of patients in those undergoing ectropion and entropion repair. This figure increases to approximately 50% before ptosis procedures and DCR. For oculoplastic specialists, the target maximum international normalised ratio (INR) before all procedures was approximately 3. For non-specialists, the target was similar prior to entropion and ectropion repair, but was 2.5 before DCR.

A considerable proportion of surgeons (54%) reported that they had seen complications as a result of either stopping or continuance of anticoagulation therapy.

Discussion

The preoperative management of oculoplastic patients on aspirin or particularly warfarin therapy presents a difficult clinical problem. There were no statistically significant differences in the approach taken by non-specialists compared with oculoplastic specialists.

The actual incidence of haemorrhagic complications compromising surgery in oculoplastic patients is unknown. In a study by Bartley on anticoagulated oculoplastic patients whose anticoagulation was not altered, no significant complications occurred. However, there are significant risks involved in discontinuing anticoagulation. In this survey, 54% of the surgeons reported that they had seen both haemorrhagic surgical complications and, in common with other surveys, serious systemic embolic complications including one death due to cerebrovascular accident and one brachial artery embolus. It is possible that such embolic complications were underreported.

In the case of cataract surgery, the available evidence suggests the practice of reducing aspirin and warfarin preoperatively is unnecessary and is putting patients at risk. Until data on haemorrhagic complication rates in oculoplastic patients on aspirin and warfarin are available, it is hard to justify a similar increase in the risk of serious life threatening complications. In view of this, a suggested approach for the management of such patients is given in Table 2.

Ophthalmologists must not lose sight of known risks when trying to minimise perceived risks.

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