The implications of novel oral anticoagulation in ocular surgery

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Background

Anticoagulation drugs play a significant role in modern day surgery- to treat and prevent thromboembolic disease. Warfarin is one of the commonest anti-coagulation used. The dosing of warfarin can be accurately managed by checking international normalized ratio (INR). It is of vital importance when any surgery is planned for a patient so that the dose of warfarin can be adjusted to decrease complication related with bleeding. The advent of new oral anticoagulation therapy with improved safety profile has improved clinical outcomes. However it’s difficult to adjust the dosing of oral anticoagulants for any surgery due to lack of monitoring test like INR and subsequently difficult to reverse their anti-coagulant effect. Anti-coagulation needs to be very closely monitored and dose adjusted to avoid any complications of intra-ocular surgery. The clinician’s ability to understand appropriate dose management and follow-up decisions will have a deep impact, if such decisions are wrong. There is limited evidence and guideline for use and adjusting the dosage of newer anti-coagulant for intra-ocular surgery. The aim of this study is to provide evidence-based guidelines for the management of patients receiving novel oral anticoagulant therapy in Ophthalmology practice. A reproducible framework has been created with this guideline that may provide a process to synchronize outpatient anticoagulation supervision. This guideline aims to provide a practical, clinician focused approach to peri-procedural anticoagulant management, which is valuable to both ophthalmologists and anesthetists who play a major part in providing local blocks and sedation, monitoring patients and managing life threatening conditions, should the need arise.

New oral anticoagulants (NOACs)

Until recently, warfarin was the only oral anticoagulant available [1]. The problem is that it requires onerous laboratory monitoring and the list of food and drug interactions is quite wide. It also carries bleeding risk, making the drug unavailable to one-thirds of the patients who should get warfarin. The good news is that already three novel oral anticoagulants: dabigatran (Pradaxa), rivaroxaban (Xarelto) and apixaban (Eliquis) that have been approved recently. These drugs can be used as alternatives to warfarin. Figure 1 depicts the blood coagulation cascade and the role of anticoagulants [2]. During a disorder of coagulation, it is clinically practical to use of drugs for adjustment of this cascade (Figure 1).

The characteristics of commonly used NOAC drugs are compared in Table 1.

NOACs during anaesthesia

The European society of anesthesia (ESA) designed a standardized pre-operative questionnaire to identify anti-haemostatic drugs by using the patient’s bleeding history and medication history: unfractionated heparin and low-molecular-weight heparins, vitamin K-antagonists, heparinoids, fondaparinux, direct thrombin inhibitors, ADP receptor- antagonists, factor X inhibitors, cyclooxygenase I-inhibitors and glycoprotein IIb/IIIa-inhibitors. Assessment of the patient has to be carried out in advance of elective surgical operations in time for additional logistic, diagnostic and therapeutic outcomes (nervous blockade, individual surgery timing, optimization of coagulation and pro-coagulant therapy and performance of appropriate drug monitoring) [4]. All the patients should be thoroughly informed of their risks of bleeding or thrombosis, prior to consent. The recommendations of ESA on time intervals are chiefly founded on the pharmacology of the concerned anti-hemostatic agents instead of scarce prospective, randomized clinical studies.

NOACs in prevention of venous thromboembolism

Atrial fibrillation (AF) is a common condition that raises the risk of cerebral thromboembolism. Its prevalence increases with age, resulting in 8% of people over 80 years of age having AF [5]. Patients with AF are at significant risk of stroke and thromboembolic complications with associated mortality and morbidity. Anticoagulant therapy reduces the risk systemic thromboembolism that leads to transient ischaemic attacks or stroke, the most threatening complications of AF. NOACs (Table 2) are being recommended with increasing frequency as inhibitors of thrombin and activated factor X, and have a number of advantages over conventional warfarin, which includes predictable response, fewer drug and food interactions and little need for monitoring or dose change. Apixaban is an anticoagulant that results in blood coagulation by direct inhibition of activated factor X (factor Xa), thereby inhibiting thrombin formation [6]. Apixaban has a marketing authorisation for significant prevention of venous thromboembolic events. For the prevention of deep venous thrombosis
in patients undergoing hip or knee replacement surgery, another NOAC, Rivaroxaban is recommended (Table 2).

**NOACs in ophthalmic surgery**

There are no clear guidelines in the literature regarding the management of patients' anticoagulated with NOACs. These patients are often elderly and at major risk of life-threatening thromboembolic disorder if anticoagulation is discontinued [8]. Several reports suggest that ophthalmic surgical procedures (with an exception to major lid and orbital surgery) usually correspond to minimal blood loss.

**Suggested practice points**

- Patients with coronary stents and prosthetic heart valves should not discontinue anticoagulant agents for cataract surgery.
- A fresh international normalized ratio (INR) result is required to be obtained on the day of surgery, prior to anaesthetic/surgical intervention.
- The INR must be checked and be in the indicated therapeutic range.

**NOACs in cataract surgery**

Cataract operation is one of the most common ophthalmic surgical procedures; the vast majority of the patients are aged with a higher risk of associated systemic disease such as coronary artery disease [9]. Patients with coronary artery disease, hypertension and diabetes are at increased risk of suprachoroidal haemorrhage. In recent years there has been a steady rise in the use of local anaesthetic techniques, but many patients are on antiplatelet or anticoagulant drugs. Emerging data suggest that long-term use of these drugs reduces the risk of thromboembolic disorders in patients with AF and atherosclerotic disease. Several studies suggest that stopping anticoagulant medication, particularly in patients with prosthetic heart valves, AF or recent coronary disease may increase high risk of thromboembolic disorders. This may greatly outweigh the risk of intra or extraocular haemorrhage. Holbrook et al. [10] suggested frequent INR testing during the first couple of weeks of the onset or discontinuation of other medical treatment. Preoperative INR should be tested close to the time of surgery and if there have been any recent changes in the patient's routine medication or diet, then re-checking the INR on the day of surgery is suggested. Although it is reported that most ophthalmic surgical procedures can be safely performed in patients who are therapeutically anticoagulated but proper physician consultation is required, to maintain safe level of anticoagulation for the shortest possible time during surgery [8].

**NOACs in vitreoretinal surgery**

At present, there is conflicting evidence of bleeding risk in NOAC patients undergoing vitre-retinal surgery. Continuing of NOACs for patients undergoing vitreoretinal surgery is recommended at present, but the published literature does report certain complications such as subretinal haemorrhage associated with external drainage at the time of scleral buckling procedures. This may occur due to acute hypotony or trauma to choroidal vessels. Spontaneous massive subretinal bleeding in a patient with diabetic retinopathy and on treatment with warfarin would...

![Figure 1. Mechanism of action of oral anticoagulants.](image-url)
Table 1. Comparative analysis of the general characteristics of NOACs and current use in medicine [modified from Cupp, 2011] [3].

| Drug          | Dabigatran (Pradaxa) | Rivaroxaban (Xarelto) | Apixaban (Eliquis) |
|---------------|----------------------|-----------------------|-------------------|
| Mechanism of action | Direct inhibitor of thrombin | Inhibitor of Factor Xa | Inhibitor of Factor Xa |
| Pharmacokinetics | $T_{1/2} : 1 - 6$ hrs | $T_{1/2} : 2 - 4$ hrs | $T_{1/2} : 1 - 4$ hrs |
| $T_{1/2}$: 12 to 17 hrs | Half-life: 5-9 hrs | Half-life: 12 hrs |
| Dialyzable (68%): | Excretion: 66% renal | Excretion: 25% renal |
| Dosage | | 2.5 to 5 mg twice a day. |
| 150 mg twice a day | | Initiation 12 -24 hrs following surgery to prevent. |
| 75 mg twice a day in patients with renal function impairment | | Treatment for 10 to 14 days following knee surgery and 32 to 38 days following hip surgery. |
| Advantages | | Unchanged excreted in urine. |
| Regular monitoring is not required | | Use in patients with moderate to severe (Child-Pugh class B or C) hepatic disorders should be avoided. |
| Disadvantages | | Use in patients with mild to moderate (Child-Pugh class A or B) hepatic disorders should be avoided. |
| Irreversible | | For CrCl ≥30 mL/min: 2.5 mg twice daily (no dosage adjustment). For CrCl 15-29 mL/min: Use with caution. For CrCl 15-29 mL/min: DO NOT use. |
| Drug must be used within 4 months of opening and stored in original container | | Apixaban prolongs aPTT, modified prothrombin time, and HepTest |
| Monitoring | | APTT can provide an approximation of anticoagulant activity. |
| Bleeding risk can be assessed by the ecarin clotting time (ECT); mainly used in research | | Without food 6 to 10 hrs following surgery. |
| No specific monitoring test is recommended. Rivaroxaban prolongs aPTT, anti-factor Xa activity, Neoplastinprothrombin time, and the HepTest. | | Initiation 12 -24 hrs following surgery to prevent. |
| Bioavailability | | Axapaxaban prolongs aPTT, modified prothrombin time, and HepTest |
| 6.5% | | APTT can provide an approximation of anticoagulant activity. |
| Monitoring | | Without food 6 to 10 hrs following surgery. |
| Conversion to/from other oral/ parenteral anticoagulants | | APTT can provide an approximation of anticoagulant activity. |
| MPM: Start dabigatran 0-2 hours before the next dose of parenteral anticoagulant would have been due or at the time of discontinuation of intravenous heparin. | | Without food 6 to 10 hrs following surgery. |
| Management prior to surgery | | APTT can provide an approximation of anticoagulant activity. |
| $CrCl ≥ 50$ mL/min: Discontinue dabigatran 1 to 2 days before procedure | | APTT can provide an approximation of anticoagulant activity. |
| $CrCl < 50$ mL/min: Discontinue dabigatran 3 to 5 days before procedure. Major procedures, spinal puncture, spinal or epidural catheter placement may require discontinuation for a longer period of time. | | APTT can provide an approximation of anticoagulant activity. |
| NOACs in glaucoma surgery |

Patients undergoing glaucoma surgery especially trabeculectomy are at increased risk of haemorrhagic complications with anticoagulation, including suprachoroidal haemorrhage, intrableb bleeding and hyphaema [13]. Patients on anticoagulation throughout glaucoma surgery had the maximum rate of hemorrhagic complications (31.8%) compared to patients who discontinued treatment prior to surgery (P = .001). At present there is no authoritative guideline available for
management of patients on NOACs undergoing glaucoma surgery, although complications occur more frequently on NOACs. In a questionnaire survey of glaucoma surgeons in England [14], diversity in clinical practice was observed with regard to continuation of anticoagulation therapy with majority of the surgeons not stopping anticoagulation drugs prior to glaucoma surgery.

Other ophthalmic surgeries

To date there are no known published studies reporting an increased risk of bleeding on NOAC patients undergoing oculoplastic, corneal or squint surgeries.

Guidelines for proper use of NOACs

The use of newly developed oral anticoagulants (apixaban, dabigatran and rivaroxaban) is recommended to follow the guidelines mentioned below:

- Since all these anticoagulants (NOACs) need dose reduction based on renal function, monitoring of renal function is necessary.
- Carrying out proper adjustments by taking in consideration the metabolic changes that take place with age and other factors that may increase bleeding tendency eg personal history or family history of bleeding disorders, liver failure etc.
- More intensive monitoring of patients during the first months of treatment when control of anticoagulant tends to be less stable.
- The presence of the direct thrombin inhibitor, daebigatran is qualitatively indicated by the activated partial thromboplastin time (APTT).
- Similarly the existence of apixaban and rivaroxaban (both inhibitors of factor Xa) are indicated by the prothrombin time (PT).
- Rivaroxaban should always be taken with food while the other NOACs can be taken with or without food.
- In case of patients receiving dronedarone, itraconazole or ketoconazole, the dabigatran dose should be reduced to 75 mg twice daily.
- Patients taking rifampin, ritonavir, or phenytoin should avoid NOACs if possible.
- Warfarin may be preferable over NOAC if patients assumed with low compliance as the anticoagulant effects of NOACs dissipate rapidly 12-24 hours after a dose.
- If there is confirmation of good compliance to NOAC (prescribed treatment for ≥3 weeks preceding cardioversion).

- The NOACs should be withheld for ≥24 hours in patients with normal renal function prior to an elective surgical procedure coupled with a minor bleeding risk and ≥48 hours before surgical procedures associated with a major bleeding risk.
- Apixaban should be discontinued at least 24 hours before surgery in low risk patients and at least 48 hours in moderate to high risk patients and recommenced as soon as possible after the patient’s surgery.
- Rivaroxaban should be discontinued at least 24 hours before surgery and recommenced as soon as possible after the patient’s surgery.
- Surgery may be required to reverse bleeding due to anticoagulation by NOACs. Intravenous tranexamic acid may be used with haematology input.

Table 2. Oral anticoagulants for prevention of stroke in atrial fibrillation [Khan, 2014] [7].

|                | Apixaban | Dabigatran | Rivaroxaban | Warfarin |
|----------------|----------|------------|-------------|----------|
| Dose           | 2.5 mg-5 mg | 110 mg to 150 mg | 10 mg | variable |
| Target         | Inhibition of Factor Xa | Inhibition of IIa (thrombin) | Inhibition of Factor Xa | Inhibition of Vit K dependent factor II,V,VI,IX and X |
| Frequency      | 2 times daily | 2 times daily | Daily | Daily |
| Metabolism     | Metabolism of liver (70%) | Metabolism of liver (20%) | Metabolism of liver (60%) | Hepatic cytochrome P₄₅₀isoenzyme |

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