



## Background:

- 1) New anticoagulants, dabigatran (Pradax) and rivaroxaban (Xarelto), are being used more and more in James Bay for prevention of strokes in patients with non-valvular A. fib.
- 2) A significant concern is that there is no antidote in the case of major bleeding for these new medications.

## Summary of new anticoagulants:

- 1) Higher dose dabigatran (150mg bid) seems to prevent CVA's better than warfarin. Rivaroxaban and lower dose dabigatran (110mg bid) is non-inferior to warfarin.
- 2) Warfarin, dabigatran and rivaroxaban all cause similar rates of major bleeding.
- 3) There is a higher rate of GI bleeding for patients on dabigatran and rivaroxaban compared to warfarin but lower rates of intracranial bleeding.
- 4) Bleeding in patients on dabigatran and rivaroxaban may be more problematic given that there is no antidote.
- 5) If someone on dabigatran or rivaroxaban bleeds in our region it is even more problematic as we have no timely access to G/C scopes, CT, neurosurgeons, etc.

\*\*refer to INESSS document: “Fibrillation auriculaire chez l’adulte – Choix de l’anticoagulothérapie” for more detailed discussion

### Conclusion:

Warfarin will continue to be first line for CVA prevention in patients with atrial fibrillation (non-valvular) however there is still a role for the newer anticoagulants in James Bay. The indications for starting dabigatran or rivaroxaban are

- 1) patients with very labile INRs and/or
- 2) where following their INR would be difficult (same criteria as for medicaments d'exception). This may be especially true in the case of patients who may spend a significant time in the bush and returning to the clinic regularly for INR checks poses a challenge.

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The risks and benefits of anticoagulation needs to be clearly explained to the patients. Patients on dabigatran or rivaroxaban should have their renal function monitored closely.

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