Helsinki experience on nonvitamin K oral anticoagulants for treating cervical artery dissection

Mustanoja, Satu

2015-08

Mustanoja, S, Metso, T M, Putaala, J, Heikkinen, N, Haapaniemi, E, Salonen, O & Tatlisumak, T 2015, 'Helsinki experience on nonvitamin K oral anticoagulants for treating cervical artery dissection', Brain and Behavior, vol. 5, no. 8, 00349. https://doi.org/10.1002/brb3.349

http://hdl.handle.net/10138/166369
https://doi.org/10.1002/brb3.349

cc_by
publishedVersion

Downloaded from Helda, University of Helsinki institutional repository.
This is an electronic reprint of the original article.
This reprint may differ from the original in pagination and typographic detail.
Please cite the original version.
Helsinki experience on nonvitamin K oral anticoagulants for treating cervical artery dissection

Satu Mustanoja¹, Tiina M. Metso¹, Jukka Putaala¹, Noora Heikkinen¹, Elena Haapaniemi¹, Oili Salonen² & Turgut Tatlisumak¹

¹Department of Neurology, Helsinki University Central Hospital, Helsinki, Finland
²Department of Neuroradiology, Helsinki University Central Hospital, Helsinki, Finland

Keywords
Acute stroke, anticoagulation, cervical arterial dissection

Correspondence
Satu Mustanoja, Department of Neurology, Helsinki University Central Hospital, P.O. Box 340, FI-00029 HUS, Helsinki, Finland.
Tel: +35894711; Fax: +358947174089; E-mail: satu.mustanoja@hus.fi

Abstract

Background: Cervical arterial dissection (CeAD) patients with or without stroke are frequently treated with either antiplatelet agents or vitamin K antagonists (VKAs), but few data are reported on the use of nonvitamin K oral anticoagulants (NOACs). Methods: Between November 2011 and January 2014, we recorded data from patients with a stroke due to vertebral (VAD) or internal carotid artery dissection (ICAD). Patients using oral anticoagulants were included in the study and were divided into two treatment groups: patients using NOACs and those using VKAs. Excellent outcome was defined on modified Rankin Scale (mRS) ≤ 1 at 6 months. Results: Of 68 stroke patients (67% male; median age 45 [39–53]), six (8.8%; two with VAD and four with ICAD) were treated with NOACs: three with direct thrombin inhibitor dabigatran and three with direct factor Xa inhibitor rivaroxaban. National Institutes of Health Stroke Scale score at baseline was 4 (3–7) in the NOAC versus 2 (1–7) in the VKA groups. Complete recanalization at 6 months was seen in most patients in the NOAC (n = 5; 83%) and VKA (n = 34; 55%) groups. All the patients using NOACs had mRS ≤ 1 at 6 months and none had an intracerebral hemorrhage (ICH). In the VKA group most patients (n = 48; 77%) had mRS ≤ 1, one patient (1.7%) had an ICH and one died. Conclusions: In this small, consecutive single-center patient sample treating ischemic stroke patients with CeAD with NOACs did not bring up safety concerns and resulted in similar, good outcomes compared to patients using VKAs.

Background
Cervical arterial dissections (CeAD), that is, vertebral artery (VAD) and internal carotid artery (ICAD) dissections are common etiologies of ischemic stroke in the young (Yesilot Barlas et al. 2013). Early treatment is crucial as it can prevent vessel occlusion or embolic sequelae and avoid serious neurologic deficits. Most physicians prescribe anticoagulants for stroke prevention in acute CeAD, although there are no randomized trials comparing the safety and efficacy of anticoagulants with antiplatelets or placebo (Engelter et al. 2007; Sarikaya et al. 2013).

Anticoagulation with nonvitamin K oral anticoagulants (NOACs) is increasingly used for stroke prevention in patients with atrial fibrillation (AF), instead of vitamin K antagonists (VKAs), as both direct factor Xa (Granger et al. 2011; Patel et al. 2011) and direct thrombin (Connolly et al. 2009) inhibitors have been shown to have similar or better safety and efficacy profiles compared with warfarin. There is few data on their use in ischemic stroke patients with CeAD (Caprio et al. 2014); and only one report was found with 10 stroke patients using NOACs as the secondary prevention of ischemic stroke.

Methods
Between November 2011 and January 2014 we recorded data from consecutive patients with a stroke due to VAD or ICAD. This study was approved by our institutional authorities. Our institutional guidelines recommend the use of anticoagulants in all CeAD patients for 6 months, and the selection of the anticoagulant is decided by the treating neurologist together with the patient.

Patients using oral anticoagulation were included in the study and were divided into two groups: patients using NOACs, and those using VKAs. Patients who underwent endovascular stenting followed by antiplatelet therapy,
and patients treated with only heparin or LMWH were excluded. We excluded two patients with multiple traumatic injuries not receiving oral anticoagulation to keep the study population homogenous.

Recurrent ischemic stroke, or intracerebral hemorrhagic (ICH) stroke events, recanalization rate, and functional outcome on the modified Rankin Scale (mRS) within six months were evaluated and compared between the NOAC and VKA-treated groups. An excellent outcome was defined as mRS≤1 at 6 months.

### Statistical analyses

Statistical significance for intergroup differences was assessed by Chi-square test for categorical variables, and Mann–Whitney U-test for noncontinuous variables or skewed numerical variables. Statistical significance was set at <0.05. Analyses were performed with SPSS 19 (SPSS Inc., Chicago, IL, USA).

### Results

Of 68 stroke patients included (Table 1), six were treated with NOACs (three with dabigatran and three with rivaroxaban) and 62 with warfarin. There were slightly more men and VAD was seen in most patients (Table 2). There were no statistical differences between the two groups in stroke severity, recanalization rate, or outcome (P > 0.05). All the patients using NOACs had mRS ≤1 with no bleeding complications, or recurrent dissections.

### Discussion

Stroke is the most common complication of CeAD, occurring in two thirds of the cases (Lichy et al. 2012). Although the molecular mechanisms are still poorly understood in CeAD (Debette 2014), imaging studies and transcranial ultrasound have suggested that arterial embolism is the main mechanism of stroke (Benninger et al. 2004), explain-
ing why anticoagulation is frequently used. ICH is the most feared and serious complication of the use of oral anticoagulation in the prevention and treatment of strokes. In a meta-analysis it was recently concluded that instead of anticoagulants antiplatelets should be used because they cause less bleeding events (Sarikaya et al. 2013), despite a trend was seen during the first week toward more deaths in the antiplatelet group (Sarikaya et al. 2013).

An excellent outcome was seen in all of our patients in the NOAC group and 77% in the VKA group. In a recent report on NOACs and CeAD, there were more recurrent strokes and radiographic progression of dissection while bleeding complications were less common (Caprio et al. 2014). In our study group, only one death and one serious ICH occurred during the 6-month treatment period, both in the VKA group. Fewer bleeding complications have been seen with the NOACs in stroke prevention in AF in three randomized trials (Connolly et al. 2009; Granger et al. 2011; Patel et al. 2011) and the European Society of Cardiology recommends NOACs in preference to VKA therapy for stroke prevention in patients with AF. In the first report with NOACs and CeAD, there were no major bleeds and 5% minor hemorrhagic complications being equal to the rate in the antiplatelet group (Caprio et al. 2014). We anticipate that the indications for the use of NOACs will be extended over time, when new data on their use in different conditions have accumulated. Recently, another off-label indication for using NOACs was reported, as factor Xa inhibitors showed a similar clinical benefit as VKAs in the treatment of cerebral venous thrombosis in a small study cohort of seven patients (Geisbusch et al. 2014).

CeAD etiology dominates in the younger age groups (Metso et al. 2012), unlike AF with a higher risk for bleeding complications associated with older age (Pancholy et al. 2014). The NOAC plasma concentrations achieved with a given dose vary, depending on absorption, renal function, and other factors that can be problematic with the elderly (Reilly et al. 2014). In the young and socially active CeAD patients, at least those with less severe strokes, many could benefit of NOACs given as a fixed dose without laboratory monitoring. Currently it remains unknown whether there is a single concentration range, where the balance between thrombo-embolic events and bleeding events is optimal for CeAD patients. It could be, however, that in more stable CeAD stroke patients the concentration range can be wider, and that NOACs could serve as a first-line treatment for the relatively short treatment period used in CeAD.

Our study has limitations. It is retrospective, and the number of patients treated with NOACs is small. As there are no randomized controlled trials going on, it adds new information on safety issues on secondary prevention with NOACs in stroke patients with CeAD.

**Conclusion**

In this small, consecutive single-center patient sample treating ischemic stroke patients with CeAD with NOACs did not bring up safety concerns and resulted in similar, good outcomes compared to patients using VKAs.

**Acknowledgments**

None.

**Conflict of Interest**

The authors declare that there is no conflict of interest.

**References**

Benninger, D. H., D. Georgiadis, C. Kremer, A. Studer, K. Nedeltchev, and R. W. Baumgartner. 2004. Mechanism of ischemic infarct in spontaneous carotid dissection. Stroke 35:482–485.

Caprio, F. Z., R. A. Bernstein, M. J. Alberts, Y. Curran, D. Bergman, A. W. Korutz, et al. 2014. Efficacy and safety of novel oral anticoagulants in patients with cerebral artery dissections. Cerebrovasc. Dis. 38:247–253.

Connolly, S. J., M. D. Ezekowitiz, S. Yusuf, D. Phil, J. Eikelboom, J. Oldgren, et al. 2009. Dabigatran versus warfarin in patients with atrial fibrillation. N. Engl. J. Med. 361:1139–1151.

Debette, S. 2014. Pathophysiology and risk factors of cerebral artery dissection: what have we learnt from large hospital-based cohorts? Curr. Opin. Neurol. 27:20–28.

Engelter, S. T., T. Brandt, S. Debette, V. Caso, C. Lichy, A. Pezzini, et al. 2007. Antiplatelets versus anticoagulation in cerebral artery dissection. Stroke 38:2605–2611.

Geisbusch, C., D. Richter, C. Herweh, P. A. Ringleb, and S. Nagel. 2014. Novel factor Xa inhibitor for the treatment of cerebral venous and sinus thrombosis: first experience in 7 patients. Stroke 45:2469–2471.

Granger, C. B., J. H. Alexander, J. J. McMurray, R. D. Lopes, E. M. Hylek, M. Hanna, et al. 2011. Apixaban versus warfarin in patients with atrial fibrillation. N. Engl. J. Med. 365:981–992.

Lichy, C., A. Metso, A. Pezzini, D. Leys, T. Metso, P. Lyrer, et al. 2012. Predictors of delayed stroke in patients with cerebral artery dissection. Int. J. Stroke. 10:360–363.

Metso, T. M., S. Debette, C. Grond-Ginsbach, S. T. Engelter, D. Leys, T. Brandt, et al. 2012. Age-dependent differences in cerebral artery dissection. J. Neurovi. 259:2202–2210.

Pancholy, S. B., P. S. Sharma, D. S. Pancholy, T. M. Patel, D. J. Callans, and F. E. Marchlinski. 2014. Meta-analysis of
gender differences in residual stroke risk and major bleeding in patients with nonvalvular atrial fibrillation treated with oral anticoagulants. Am. J. Cardiol. 113:485–490.
Patel, M. R., K. W. Mahaffey, J. Garg, P. Guohua, D. E. Singer, W. Hacke, et al. 2011. Rivaroxaban versus warfarin in nonvalvular atrial fibrillation. N. Engl. J. Med. 365:883–891.
Reilly, P. A., T. Lehr, S. Haertter, S. J. Connolly, S. Yusuf, J. W. Eikelboom, et al. 2014. The effect of dabigatran plasma concentrations and patient characteristics on the frequency of ischemic stroke and major bleeding in atrial fibrillation patients: the RE-LY Trial (Randomized Evaluation of Long-Term Anticoagulation Therapy). J. Am. Coll. Cardiol. 63:321–328.
Sarikaya, H., B. R. da Costa, R. W. Baumgartner, K. Duclos, E. Touze, J. M. de Bray, et al. 2013. Antiplatelets versus anticoagulants for the treatment of cervical artery dissection: Bayesian meta-analysis. PLoS One 8:e72697.
Yesilot Barlas, N., J. Putaala, U. Waje-Andreassen, S. Vassilopoulou, K. Nardi, C. Odier, et al. 2013. Etiology of first-ever ischaemic stroke in European young adults: the 15 cities young stroke study. Eur. J. Neurol. 20:1431–1439.