Pulmonary embolism four days after interruption of therapy with rivaroxaban

Gabor Göndör; Claudia Stöllberger

Keywords
Non-vitamin K-antagonist, oral anticoagulants, rivaroxaban rebound phenomenon, atrial fibrillation, pulmonary embolism

Summary
Thrombosis after cessation of anticoagulation, also named rebound thrombosis, is a matter of concern and controversy. There are only few published data about occurrence of rebound thrombosis associated with non-vitamin K-antagonist oral anticoagulant drugs (NOACs). We report on a 58-year-old male with paroxysmal atrial fibrillation (AF) who developed central pulmonary embolism four days after interruption of rivaroxaban because of parotid surgery. He had received 40 mg enoxaparin/d. The parotid gland was partially resected within 6 hours without blood loss. Pulmonary embolism and AF occurred on the first postoperative day. He recovered with low-molecular-weight heparin in therapeutic dosages and amiodarone and was discharged with phenprocoumon.

The relevance of a rivaroxaban rebound phenomenon, manifesting as arterial embolism, stroke or venous thromboembolism should be clarified. It should be assessed if rebound-phenomena also exist for the NOACs dabigatran, apixaban and edoxaban. Thus, the randomized trials and registries investigating patients with AF or venous thromboembolism should be re-analysed and, based on these data, recommendations should be developed for situations in which NOAC-therapy has to be interrupted or ceased.

Schlüsselwörter
Nicht Vitamin K-Antagonisten, orale Antikoagulantien, Rivaroxaban, Rebound-Phänomen, Vorhofflimmern, Pulmonalembolie

Zusammenfassung

Die Relevanz eines Rivaroxaban-Rebound-Phänomens, das als arterielle Embolie, Schlaganfall oder venöse Thromboembolie auftreten kann, sollte genauer erforscht werden. Es sollte festgestellt werden, ob ähnliche Phänomene auch bei den NOAKs Dabigatran, Apixaban und Edoxaban auftreten. Randomisierte Studien und Register, die Patienten mit AF oder venöser Thromboembolie einschlossen haben, sollten unter diesem Gesichtspunkt ausgewertet werden. Es sollten Empfehlungen ausgearbeitet werden für die Antikoagulation in Situationen, in denen die Therapie mit NOAKs unterbrochen wird.

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Introduction
Rivaroxaban is a direct anti-Xa oral anticoagulant and belongs to the group of non-Vitamin K-antagonist oral anticoagulants (NOACs), which are increasingly used in patients with atrial fibrillation for prevention of stroke or embolism, based on the results of a large randomized trial (1). Furthermore, rivaroxaban is also approved for the treatment of venous thromboembolism based on the results of two randomized trials (2, 3).

In comparison to the traditionally used vitamin-K-antagonists, the NOACs do not require any routine monitoring. On the ot-
her hand, in emergency situations it is more difficult to estimate their plasma concentration and there are no recommended specific antidotes so far.

Furthermore, rivaroxaban and other NOACs carry a black box warning regarding the risk of rebound thrombosis after discontinuation (4, 5).

Case report

A 58-year-old male patient was admitted to our department because of an acute pulmonary embolism. He had a history of paroxysmal atrial fibrillation since 3 years, currently under therapy with rivaroxaban 20 mg, arterial hypertension since many years, peripheral arterial disease stage Iib since 5 years, fatty liver disease, prostatic hyperplasia, cholecystectomy 5 years before, lateral parotidectomy of the left parotid gland 13 years before, substituted hypothyroidism since the thyroidectomy at the age of 41 years and appendectomy at the age of 14 years. His CHA2DS2-VASc score was 4 and the HAS-BLED score was 2. He was scheduled to an elective parotidectomy because of cystadenolymphoma of the right parotid gland.

Originally, he took phenprocoumon since 3 years, but the oral anticoagulation was changed to rivaroxaban six weeks before the actual surgery by his general practitioner. The cause for the readjustment was a transient ischemic attack of the right medial cerebral artery with sensory abnormality of the left arm for 15 minutes. The International Normalized Ratio at the time of the TIA was 2.75.

Preoperatively, the Caprini DVT risk score (a risk stratification score for periprocedural venous thromboembolism) was 4, which corresponds to high risk and, in case of inclusion of TIA, the score would add up to 9 (6). He stopped taking rivaroxaban four days before the surgery and received enoxaparin sodium 40 mg once daily. Besides, he took amiodipine/valsartan/hydrochlorothiazide 5/160/25 mg once, bisoprolol 5 mg twice and levothyroxine sodium 100 µg once daily. The surgery took 6 hours and succeeded without any complication. The patient had no blood loss and he did not need any erythrocyte concentrates. The histological result showed a partial resection of the parotid gland with two benign cystadenolymphomas.

Postoperatively the patient complained of pain in the right popliteal fossa and 15 hours after the end of the surgery, he started to suffer from sudden onset of severe dyspnea why he was transferred to the medical department. At admission, the blood pressure was 140/85 mmHg, the heart rate 150/min and the O2-saturation 88%. After supplemental oxygen therapy with 5 l/min, the oxygen saturation increased to 92%. The D-dimer was 7.60 mg/l (Table 1). The electrocardiogram showed tachycardious atrial fibrillation.

The CT-angiography of the pulmonary arteries showed a central pulmonary embolism on both sides of the lung reaching into the segmental arteries (Figure 1).

Concomitantly, the veins of the pelvis and lower extremities were investigated by CT and did not show any thrombus. A bedside-echocardiography showed a slightly reduced left ventricular systolic function, a normally sized right ventricle with normal function, no pericardial effusion and no thrombus. Tachycardious atrial fibrillation was converted into sinus rhythm after the application of amiodarone, potassium and magnesium. Thrombolysis was considered but refused because of the recent surgery. Thus, enoxaparin sodium 100 mg was injected as a bolus and prescribed two times daily. An oral anticoagulation therapy with phenprocoumon was initiated after 7 days and the patient was discharged without any complaints after 12 days.

Tab. 1 Laboratory findings

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Parameter abbreviations: PT = prothrombin time; INR = international normalized ratio; NM = not measured; PSA = prostate-specific antigen; PT = prothrombin time; PS = prostate-specific antigen; CK = creatine phosphokinase; CRP = C-reactive protein; GGT = gamma-glutamyl transferase; ALT = alanine-aminotransferase; CK = creatine phosphokinase; CRP = C-reactive protein; GGT = gamma-glutamyl transferase; INR = international normalized ratio; NM = not measured; PSA = prostate-specific antigen; PT = prothrombin time; PS = prostate-specific antigen; CK = creatine phosphokinase; CRP = C-reactive protein; GGT = gamma-glutamyl transferase; INR = international normalized ratio; NM = not measured; PSA = prostate-specific antigen; PT = prothrombin time
Discussion

Rivaroxaban is a NOAC, which takes its effect by inhibiting the prothrombinase-complex and the free factor Xa directly. The prothrombinase-complex is an enzymatic complex of the blood coagulation cascade consisting of the activated factors X and V, calcium and phospholipids (7). The advantage of rivaroxaban compared to the vitamin-K-antagonists is that it does not require any routine monitoring. Based on randomized trials in patients with venous thromboembolism (2, 3), the guidelines of acute pulmonary embolism mention rivaroxaban 20mg once daily as an alternative to vitamin-K-antagonists if extended anticoagulation treatment is necessary as a class IIa, level B recommendation (8).

Rivaroxaban was also found to be non-inferior to warfarin for the prevention of stroke and systemic embolism in patients with moderate- to high-risk nonvalvular atrial fibrillation and concerns about a potential rebound effect after cessation of rivaroxaban have been raised in association with the ROCKET AF (Rivaroxaban Once Daily Oral Direct Factor Xa Inhibition Compared with Vitamin K Antagonism for Prevention of Stroke and Embolism Trial in Atrial Fibrillation) study, a multicenter, randomized, double-blind trial with the participation of 14,264 patients (1). After the ROCKET AF study, an end of study (EOS) visit was performed and the participants were transitioned to open-label vitamin-K-antagonist-therapy. Between days 3 and 30, an excess of stroke and systemic embolic events were observed in participants assigned to rivaroxaban, and the median time to reach a therapeutic INR value was 13 days in the rivaroxaban-group compared to 3 days in the warfarin-group (5, 9). Unfortunately, there are no data about venous thromboembolic events occurring after the ROCKET AF study since only stroke and arterial embolism were registered.

In our case, we considered three possible triggers that could have led to the thrombus formation and the subsequent pulmonary embolism.

First, the phenomenon of rebound thrombosis could have been a reason. Rivaroxaban carries a black box warning regarding the risk of thrombosis after discontinuation: „discontinuing rivaroxaban places patients at an increased risk of thrombotic events” and „an increased risk of stroke was observed following rivaroxaban discontinuation in clinical trials in atrial fibrillation patients” (4, 5). There are only limited data about the prevalence and clinical relevance of a rivaroxaban rebound phenomenon. Bakhit et al. reported a case with rebound thrombosis. Twenty-four hours after the cessation of rivaroxaban, an elective radiofrequency catheter ablation was performed but it had to be interrupted because a large septal thrombus in the left atrium was found (10).

The black box warning, however, exists for other oral anticoagulants as well. "Premature discontinuation of any oral anticoagulant in the absence of adequate alternative anticoagulation increases the risk of thrombotic events” (4). The direct oral anti Xa-inhibitors apixaban and edoxaban carry the same black box warning (11, 12). Regarding the direct oral thrombin-inhibitor dabigatran, the RE-MODEL trial suggested that dabigatran does not cause rebound thrombosis (13), other studies suggested the opposite (14). Thorne et al reported three arterial or venous thromboembolism cases within one month after cessation of dabigatran (15).

Data from a recently published non-industry-sponsored registry show that the NOACs rivaroxaban and dabigatran offer good protection against thromboembolic events in real-life patients with nonvalvular atrial fibrillation, but that interruptions of treatment represent a vulnerable period (16). In that registry, 866 patients with non-valvular atrial fibrillation with an average CHA2DS2-VASc-Score of 2.1, who were started on dabigatran or rivaroxaban, were analysed for thromboembolic events and survival. Patients who had temporary or permanent discontinuation of NOACs were compared to patients on continuous NOAC treatment. The incidence of thromboembolic events during treatment interruptions was more than 6-times higher than the “natural course” incidence predicted by the CHA2DS2-VASc-score. On the other hand, the incidence of thromboembolism was significantly lower in patients with uninterrupted NOAC treatment than the predicted incidence for untreated patients (16). Also in that registry, only arterial embolism and strokes, and no venous thromboembolic events, were counted.

The possibility of rebound thrombosis after cessation of therapy with vitamin K-antagonists has been subject of discussion for a long time, e.g. by Genewein et al. in 1996 (17).

The pathophysiologic mechanism of a rivaroxaban rebound phenomenon is uncertain. Haynes et al., based on an in vitro study, suggested that the discontinuation of rivaroxaban leads to prothrombotic activity. The reason of this phenomenon may be that when rivaroxaban plasma concentrations decrease after cessation of therapy, there is an unmasking of thrombus-associated prothrombinase (18).

A second pathogenetic possibility for the thrombotic event in our case are the 6 hour-long surgery and the following hospitalization which are well known risk factors for thrombosis (19). However, the incidence of thrombosis and pulmonary embolism are not as common in patients after
References

16. Vene N et al. Risk of Thromboembolic Events in Patients with Non-Valvular Atrial Fibrillation


