## Recurrent ischemic stroke and ventricular hemorrhage after IV thrombolysis: is early anticoagulation with Rivaroxaban safe? A case report

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Introduction: Direct Oral Anticoagulants (DOACs) have lower hemorrhagic complications rates, especially brain hemorrhage, compared to vitamin K antagonists<sup>1</sup>. Clinical trials investigating DOACs excluded patients with recent (=i.e. less than 7 days) ischemic stroke to avoid the risk for symptomatic Hemorrhagic Transformation (sHT)<sup>2</sup>. Despite the lack of clinical data, DOACs are perceived as safe and increasingly used in clinical practice even in the earlier days after stroke onset. We describe a case in which Rivaroxaban, an anti Xa coagulation factor, had been used for early ischemic stroke recurrence despite the presence of ventricular bleeding after thrombolysis.

Case report: A 88 years old gentleman with permanent AF developed acute onset aphasia and right side weakness. Head CT was unremarkable for hemorrhagic and ischemic lesions while showing Middle Cerebral Artery (MCA) M1 vascular



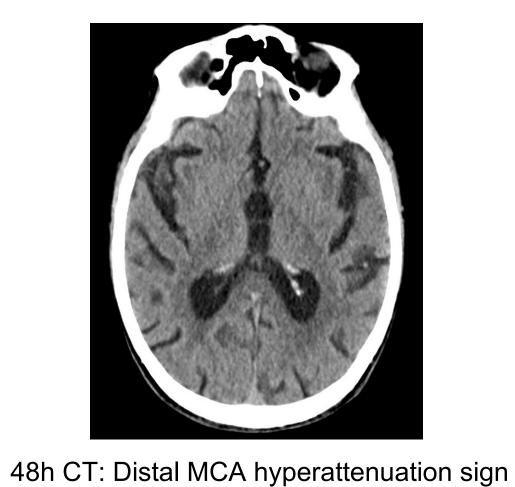


hyperattenuation sign. Intravenous thrombolysis within 3 hours from symptom onset was administered. On the following day, we observed an improvement of neurological status with milder right motor symptoms and minimal motor aphasia. 24 hours CT was negative for both parenchymal ischemia and vascular hyperattenuation signs while showing asymptomatic bilateral bleeding in occipital horns of lateral ventricles. At 48 hours CT, ischemic signs were evident in the terminal territory of left MCA inferior branch with vascular hyperattenuation spot on distal left MCA branch. On the fifth day, aphasia worsened. A brain CT showed a new ischemic lesion in left MCA inferior branch with distal MCA hyperattenuation sign with the persistence of intraventricular bleeding. Since an early ischemic event recurred, we decided to start anticoagulant therapy despite the bleeding. As the patient GFR was 34 ml/min, we used the reduced dose of Rivaroxaban 15 mg. During the following days the patient exhibited an improvement of neurological conditions. Eleventh day head CT demonstrated a reduction of intraventricular bleeding and the presence of small petechial HT within one ischemic lesion. Rivaroxaban was then continued after discharge.

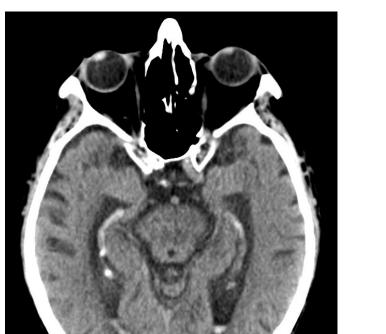
Before thrombolysis

24h CT: intraventricular bleeding



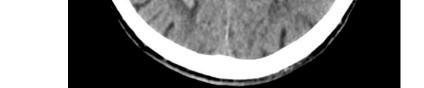


48h CT: intraventricular bleeding









5° day CT: intraventricular bledding



5° day CT new ischemic lesion

11° day CT:: reduction of intraventricular bleeding

**Discussion and Conclusion:** In patients with cardioembolic stroke the timing for starting anticoagulant therapy is still a controversial issue as the risk of sHT. Due to DOACs lower bleeding risk, a rule of thumb of 1 (TIA) - 3 (non disabling stroke) - 6 (moderate stroke) - 12 (severe stroke) days had been advocated even in absence of clinical data<sup>3</sup>. No data are available regarding the timing for starting DOACs therapy in the presence of bleeding complications. In our case, the use of Rivaroxaban was associated with a clinical improvement in the absence of radiological worsening of ventricular bleeding and sHT. Clinical data are required to determine the appropriate timing for DOACs after an ischemic stroke.

## References:

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## XLVII CONGRESSO NAZIONALE 22-25 OTTOBRE 2016 – VENEZIA



