Intravenous Thrombolysis With Recombinant Tissue-Type Plasminogen Activator in a Stroke Patient Receiving Dabigatran Anticoagulant After Antagonization With Idarucizumab

Jörg Berrouschot, MD, PhD; Anett Stoll, MD; Theresa Hogh, MD; Christoph Cyrill Eschenfelder, MD

Background and Purpose—Therapeutic options for acute ischemic stroke patients presenting on effective anticoagulation are limited. Idarucizumab, a humanized, monoclonal antibody fragment for immediate reversal of dabigatran, may allow this subgroup of orally anticoagulated patients to regain eligibility for thrombolysis. Methods—We report the first successful acute antagonization of dabigatran by idarucizumab before intravenous thrombolysis with recombinant tissue-type plasminogen activator.

Results—Idarucizumab was given to a 76-year-old male patient on dabigatran ≈3.5 hours after his last dose. Neurological status on admission was NIHSS (National Institutes of Health Stroke Scale) 11. Recombinant tissue-type plasminogen activator was initiated immediately after dabigatran reversal. The patient was discharged with a favorable outcome of NIHSS 1 on day 7. No complications were observed.

Conclusions—This case represents a new therapeutic paradigm. It is further supported by in vitro data showing no nonspecific interactions of idarucizumab with recombinant tissue-type plasminogen activator–induced thrombolysis. Thus, patients effectively anticoagulated with dabigatran who were previously contraindicated for thrombolytic therapy in this situation may now receive treatment because of the ability to rapidly reverse the anticoagulant activity of dabigatran with idarucizumab.

(Stroke. 2016;47:1936-1938. DOI: 10.1161/STROKEAHA.116.013550.)

Key Words: anticoagulants ■ antidotes ■ dabigatran ■ idarucizumab ■ stroke
Reversal of Dabigatran Prior to r-tPA Treatment

He presented in our emergency department 2 hours and 30 minutes after symptom onset. Neurological status showed a global aphasia, neglect to the right, and a slight brachiofacial hemiparesis pronounced in the right arm (NIHSS [National Institutes of Health Stroke Scale] 11 points). According to his wife, the patient took his last dabigatran dose around 6:00 pm, ≈1 hour before the start of symptoms.

The computed tomography (CT) scan showed the previous left-sided middle cerebral artery infarction and no early signs of infarction. The large, brain-supplying vessels were without relevant occlusion as analyzed by CT angiography.

The thrombin time was 218 seconds (normal range: 15–36 seconds), and the activated partial thromboplastin time (aPTT) was 73.3 seconds (normal range: 20–40 seconds) on admission. The remaining routine laboratory values, including creatinine clearance, were normal.

We treated the patient with an intravenous bolus of 5 g idarucizumab given for 5 minutes to antagonize the anticoagulant effect of dabigatran. Immediately afterward, intravenous thrombolysis with 69 mg r-tPA, equivalent to 0.9 mg r-tPA/kg body weight was initiated.

The next morning, the hemiparesis and neglect were almost fully regressed, and aphasia was significantly improved (NIH SS 3 points). The follow-up CT scan showed no new signs of cerebral infarction or hemorrhagic transformation (a magnetic resonance imaging was not possible because of the pacemaker). The thrombin time was 26.9 seconds and the aPTT was 26.6 seconds at follow-up.

A cardiovascular examination of the patient revealed slight atherosclerotic lesions in the brain-supplying arteries without relevant stenoses. Transthoracic and transesophageal echocardiography showed no additional evidence of a cardiac embolic source. Ultimately, nonvalvular atrial fibrillation remained the major determinant of the recurrent stroke in this patient. For this reason, we started oral anticoagulation again with an increased dabigatran dose (150 mg BID) beginning the following day.

After 7 days, the patient was discharged for early rehabilitation. Another CT scan showed no evidence of new cerebral infarction. At the time of transfer, the patient showed a slight fluid aphasia with primary anomia (NIH SS 1 point).

Discussion

For the first time, we report the successful use of idarucizumab in combination with subsequent intravenous thrombolysis using r-tPA, in an acute ischemic stroke patient receiving anticoagulation with 110 mg BID dabigatran.

Idarucizumab, a specific antidote for dabigatran, is a humanized antibody fragment (Fab fragment; Fab stands for fragment antigen binding) from the IgG1 isotype, which directly binds the thrombin inhibitor dabigatran and was approved in 2015 by the US Food and Drug Administration and European Medicines Agency for use in dabigatran-treated patients when rapid reversal of its anticoagulant effects is needed, such as in emergency surgery or urgent procedures or life-threatening or uncontrolled bleeding. These patients are also being enrolled in the currently ongoing, multicenter open-label, single-arm phase III study RE-VERSE AD (Reversal Effects of Idarucizumab on Active Dabigatran), where efficacy and tolerability of idarucizumab is being studied in clinical practice. Thus, patients requiring thrombolysis could also be enrolled in the RE-VERSE AD study as those requiring an urgent procedure after dabigatran reversal. However, information on this particular patient population has been limited to date.

In our case, the significantly elevated aPTT and thrombin time values before the start of thrombolysis were seen ≈4 hours after the last intake and demonstrated an anticoagulant effect consistent with peak dabigatran values. Because CT angiography showed no proximal vascular occlusion, mechanical thrombectomy was not indicated in this patient. Although not specifically listed as an indication for use of idarucizumab as per labeling information, we first antagonized the anticoagulant effect of dabigatran with 5 g idarucizumab before immediately starting intravenous thrombolysis with r-tPA (see Table for details).

Idarucizumab rapidly reverses dabigatran anticoagulation and returns the patient coagulation status to their normal or baseline state. Thus, if they are undergoing a thrombotic event, which is procoagulant in nature, then thrombolysis should be initiated rapidly after dabigatran reversal. This is supported by recent data demonstrating that there are no interactions between idarucizumab and r-tPA–induced thrombolysis in human plasma in vitro. In addition, idarucizumab has no procoagulant effect, in contrast to coagulation factor concentrates, and as an antibody fragment is neutral regarding any pro- or anticoagulant effect. It only directly inhibits dabigatran, thus will not in itself further potentiate coagulation.

Table. Extract From Prescribing Information of r-tPA and Idarucizumab

<table>
<thead>
<tr>
<th></th>
<th>Alteplase³</th>
<th>Idarucizumab⁴</th>
</tr>
</thead>
<tbody>
<tr>
<td>4.4 Special warnings and precautions for use</td>
<td>Indications and usage</td>
<td></td>
</tr>
<tr>
<td>Patients receiving oral anticoagulant treatment:</td>
<td>Praxbind is a humanized monoclonal antibody fragment (Fab) indicated in patients treated with Pradaxa when reversal of the anticoagulant effects of dabigatran is needed:</td>
<td></td>
</tr>
<tr>
<td>The use of Actilyse may be considered when dosing or time since the last intake of anticoagulant treatment makes residual efficacy unlikely confirmed by appropriate test(s) of anticoagulant activity for the product(s) concerned showing no clinically relevant activity on the coagulation system (eg, INR ≤1.3 for vitamin K antagonists or other relevant test(s) for other oral anticoagulants are within the respective upper limit of normal).</td>
<td>• For emergency surgery/urgent procedures.³</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• In life-threatening or uncontrolled bleeding.³</td>
<td></td>
</tr>
</tbody>
</table>

INR indicates international normalized ratio; and r-tPA, recombinant tissue-type plasminogen activator.

*Urgent procedures: any urgent procedure that cannot be postponed, any diagnostic, therapeutic, interventional, or pharmacological intervention (such as thrombolysis), and where rapid removal of an anticoagulatory effect of dabigatran is required, may be considered. In the RE-VERSE AD trial (Reversal Effects of Idarucizumab on Active Dabigatran; group B), no clinical restrictions were prespecified.³,¹² Fab indicates fragment antigen binding.
Thus, this represents a new therapeutic paradigm. Patients effectively anticoagulated with dabigatran who were previously contraindicated for thrombolytic therapy in this situation may now receive treatment because of the ability to rapidly reverse the anticoagulant activity of dabigatran with idarucizumab. Particularly since both vitamin K antagonists, such as warfarin, and also other non–vitamin K antagonist oral anticoagulants currently have no specific reversal agent.

Although there is to date limited clinical evidence and no specific labeling information for ischemic stroke use available, our case illustrates a potential beneficial use of idarucizumab in this specific situation.

Conclusions

Because of the increasing use of non–vitamin K antagonist oral anticoagulants, more anticoagulated acute stroke patients presenting within the time window for intravenous thrombolytic therapy will become more frequent in stroke departments. When these patients are treated with dabigatran, the anticoagulation can be antagonized with idarucizumab, thereby abruptly returning them to a status of normal hemostasis with respect to dabigatran anticoagulation. Afterward—presumably with a comparable bleeding risk to a nonanticoagulated patient—they may undergo intravenous thrombolysis with rt-PA. Further studies on this emergency condition are needed to increase the level of evidence from an individual treatment to a generalized recommendation.

Acknowledgments

Boehringer Ingelheim provided assistance for preparation of this article.

Disclosures

C.C. Eschenfelder is an employee of Boehringer Ingelheim. The other authors report no conflicts.

References


Intravenous Thrombolysis With Recombinant Tissue-Type Plasminogen Activator in a Stroke Patient Receiving Dabigatran Anticoagulant After Antagonization With Idarucizumab

Jörg Berrouschot, Anett Stoll, Theresa Hogh and Christoph Cyrill Eschenfelder

*Stroke*. 2016;47:1936-1938; originally published online June 14, 2016;
doi: 10.1161/STROKEAHA.116.013550

*Stroke* is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2016 American Heart Association, Inc. All rights reserved.
Print ISSN: 0039-2499. Online ISSN: 1524-4628

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://stroke.ahajournals.org/content/47/7/1936

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in *Stroke* can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to *Stroke* is online at:
http://stroke.ahajournals.org//subscriptions/