Thrombolytic Therapy for Stroke: Intravenous Tenecteplase Era

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Acute ischemic stroke (AIS) is a worldwide health burden and the leading cause of disability in adults. The management of AIS is developing rapidly. Currently, the two Food and Drug Administration (FDA)-approved treatment options for AIS are intravenous thrombolysis (IVT) with alteplase within the first 4.5 h of the AIS symptom onset and endovascular treatment (EVT) in patients with an occlusive clot in one of the intracranial large vessels within the first 24 h of symptom onset depending on the salvagability of the brain tissue [1-10]. Some studies have shown the potential benefit of EVT during the first 16 to 24 h from the AIS symptom onset if there is a favorable penumbra without a large infarct core [11, 12]. Another study revealed that EVT within 6 to 16 h after onset of AIS symptom could decrease the length hospital stay and improve patient functional outcome [13].

Alteplase is an intravenous recombinant tissue plasminogen activator (t-TPA) with a short half-life that needs continuous infusion for about 1 h [14]. Infusion of alteplase as a bridge prior to EVT is still a treatment option for eligible AIS patients with intracranial large vessel occlusions [15-17]. Thus far, alteplase is the only FDA-approved pharmacological treatment for AIS [1, 5, 18]; however, infusion of alteplase has a relatively low rate of recanalization and reperfusion of the large vessel occlusions prior to EVT; hence, it might not be the ideal treatment option [15, 19].

Tenecteplase is a modified form of alteplase with 14-fold more fibrin specificity and enhanced ability for thrombolysis. It also has a longer half-life and 80 times greater resistance to plasminogen activator inhibitor-1 compared to the alteplase; as such, it can be used as an intravenous bolus rather than a bolus followed by 1 h infusion as with alteplase [20]. Single bolus infusion of tenecteplase can give practical benefits to AIS patients who need transportation between hospitals to access EVT [15]. Single bolus infusion of tenecteplase can also reduce the time to receive EVT after the onset of stroke symptoms [16]. Furthermore, according to the information available on drugs.com, in the USA, tenecteplase would cost much less when compared with the alteplase [20].

To the best of our knowledge, at least five randomized clinical trials that compared alteplase with tenecteplase in the treatment of AIS have shown no significant differences in the mortality rate or in the chance of neurological recovery between these two options [15]. Parson et al [18], in a randomized clinical trial on AIS patients who received thrombolytic therapy, found that tenecteplase was superior to alteplase concerning the angiographic outcomes, rate of reperfusion, and also neurological improvement in 24 h. They also showed lower rate of severe disability at 90 days [18].

Campbell et al [15] studied tenecteplase 0.25 mg/kg versus the standard dose of alteplase for patients with AIS within 4.5 h of symptom onset prior to EVT. They reported significantly greater chance of reperfusion of the occluded vessel along with meaningfully better chance of 3-month functional outcome [15, 20].

Concerning the safety outcomes, two of the most recent meta-analyses comparing the efficacy and safety of the both treatment options for AIS, found no statistically significant difference in the rates of intracerebral hemorrhage between the two, but there were tendency toward lower rate of intracerebral hemorrhage with tenecteplase (odds ratio (OR): 0.81, 95% confidence interval (CI): 0.56 - 1.17; P = 0.26) [14, 21].

When it comes to the administration of the tenecteplase, current evidences advocate that 0.25 mg/kg (maximum 25 mg) is the most appropriate dose. The 0.1 mg/kg dose was not as effective as 0.25 mg/kg dose in the study by Parsons et al [18], and the 0.4 mg/kg dose may result in higher rates of intracerebral hemorrhage as shown by Haley et al [22].

In conclusion, tenecteplase is shown to be as effective as alteplase with respect to functional outcome after treatment of AIS. Furthermore, it can be used as an intravenous bolus rather than a bolus followed by 1 h infusion as with alteplase which makes it more convenient to administer and may have lower bleeding risks when compared to alteplase. It is also cheaper. Health care providers should contemplate using tenecteplase rather than alteplase for the treatment of AIS at the recommended dose if available in their centers.

Acknowledgments

None to declare.

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Financial Disclosure

None to declare.

Conflict of Interest

None to declare.

Author Contributions

All authors have contributed to the theoretical formalism, designing the study, data collection, data analysis and writing the manuscript.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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