METALYSE® IS NOW AVAILABLE

for the fibrinolytic treatment of patients with acute ischaemic stroke[1].

Boehringer Ingelheim

metalyse® tenecteplase 25mg

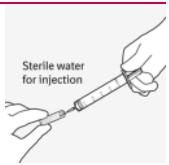
How to prepare Metalyse®

1. Reconstitute immediately before administration.

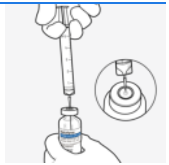
2. Remove the protective cap on the vial containing Metalyse® dry substance by flipping it up with a thumb.

3. Swab the rubber top of the vial with an alcohol wipe.

4. Aseptically withdraw 5 mL sterilised water for injection.

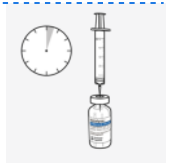


5. Transfer the 5 mL sterilised water for injection into the Metalyse® vial by introducing the needle vertically into the middle of the rubber stopper, directing the diluent stream into the powder.



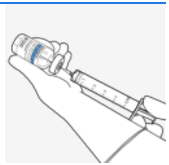
6. Take the vial with reconstituted Metalyse® and swirl gently to dissolve any remaining powder, but do not shake, as this will produce foam.

If there are bubbles, let the solution stand undisturbed for a few minutes to allow them to disappear.



7. The reconstituted solution consists of 5 mg/ml Metalyse®. It should be clear and colourless to pale yellow and it should not contain any particles.

8. Remove the amount required using a needle and a syringe.



9. Use immediately. Dispose of any unused solution.

Click on or scan the QR code to view the preparation video.

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How to dose & administer Metalyse®

Metalyse® is administered by a single IV bolus **over 5 to 10 seconds,** eliminating the need for a one-hour infusion, as required for Actilyse®.[2-4]

Calculate the dose based on the patient's body weight

Weight	Dose
<60 kg	3 ml (15 mg)
60 to <70 kg	3.5 ml (17.5 mg)
70 to <80 kg	4 ml (20 mg)
80 to <90 kg	4.5 ml (22.5 mg)
90+ kg	5 ml (25 mg)

Abbreviations
IV: intravenous.

References
1. Metalyse® European Summary of Product Characteristics. 2. Menon BK, et al. Lancet 2022; 400: 161-169. 3. Bivard A, et al. Lancet Neurol. 2022; 21: 520-27. 4. Actilyse® European Summary of Product Characteristics.
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