

Non WHO Reference Material Reteplase Reference Material NIBSC code: 10/104 Instructions for use (Version 1.0, Dated 16/11/2011)

This material is not for in vitro diagnostic use.

#### 1. INTENDED USE

The potency of the standard was determined in a collaborative study in comparison with the previous Reteplase reference preparation 93/726. The agreed potency is 24.3 milliunits per ampoule. This reference material is intended for use as a standard to determine the fibrinolytic activity of Reteplase test solutions in enzymatic bioassay systems.

## 2. CAUTION

# This preparation is not for administration to humans or animals in the human food chain.

The preparation contains material of human origin, and either the final product or the source materials, from which it is derived, have been tested and found negative for HBsAg, anti-HIV and HCV RNA. As with all materials of biological origin, this preparation should be regarded as potentially hazardous to health. It should be used and discarded according to your own laboratory's safety procedures. Such safety procedures should include the wearing of protective gloves and avoiding the generation of aerosols. Care should be exercised in opening ampoules or vials, to avoid cuts.

## 3. UNITAGE

24.3 milliunits per ampoule

#### 4. CONTENTS

Country of origin of biological material: EU.

# 5. STORAGE

Please store at -20 °C or below, away from direct light

Please note: because of the inherent stability of lyophilized material, NIBSC may ship these materials at ambient temperature.

## 6. DIRECTIONS FOR OPENING

DIN ampoules have an 'easy-open' coloured stress point, where the narrow ampoule stem joins the wider ampoule body. Various types of ampoule breaker are available commercially. To open the ampoule, tap the ampoule gently to collect material at the bottom (labelled) end and follow manufactures instructions provided with the ampoule breaker.

# 7. USE OF MATERIAL

No attempt should be made to weigh out any portion of the freeze-dried material prior to reconstitution

# 8. STABILITY

Reference materials are held at NIBSC within assured, temperaturecontrolled storage facilities. Reference Materials should be stored on receipt as indicated on the label.

# 9. REFERENCES

None

# 10. ACKNOWLEDGEMENTS

None

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## 11. FURTHER INFORMATION

Further information can be obtained as follows;

This material: enquiries@nibsc.org

WHO Biological Standards:

http://www.who.int/biologicals/en/

JCTLM Higher order reference materials: http://www.bipm.org/en/committees/jc/jctlm/

Derivation of International Units:

http://www.nibsc.org/standardisation/international standards.aspx

http://www.nibsc.org/standardisa Ordering standards from NIBSC:

http://www.nibsc.org/products/ordering.aspx

NIBSC Terms & Conditions:

http://www.nibsc.org/terms\_and\_conditions.aspx

#### 12. CUSTOMER FEEDBACK

Customers are encouraged to provide feedback on the suitability or use of the material provided or other aspects of our service. Please send any comments to enquiries@nibsc.org

## 13. CITATION

In all publications, including data sheets, in which this material is referenced, it is important that the preparation's title, its status, the NIBSC code number, and the name and address of NIBSC are cited and cited correctly.

#### 14. MATERIAL SAFETY SHEET

Classification in accordance with Directive 2000/54/EC, Regulation (EC) No 1272/2008: Not applicable or not classified

Physical and Chemical properties			
Physical appearance:		Corrosive:	No
Freeze dried white powder			
Stable: Yes		Oxidising:	No
Hygroscopic: Yes		Irritant:	Yes
Flammable: No		Handling:See	e caution, Section 2
Other (specify): Contains material of human origin			
Toxicological properties			
Effects of inhalation: Not		established, avoid inhalation	
Effects of ingestion: Not		established, avoid ingestion	
Effects of skin absorption: Not		established, avoid contact with skin	
Suggested First Aid			
Inhalation: See	Seek medical advice		
Ingestion: Seek medical advice			
Contact with eyes: Was			
medical advice			
Contact with skin: Was	sh thoroughly with water.		
Action on Spillage and Method of Disposal			

# Action on Spillage and Method of Disposal

Spillage of ampoule contents should be taken up with absorbent material wetted with an appropriate disinfectant. Rinse area with an appropriate disinfectant followed by water.

Absorbent materials used to treat spillage should be treated as biological waste.

# 15. LIABILITY AND LOSS

In the event that this document is translated into another language, the English language version shall prevail in the event of any inconsistencies between the documents.

Unless expressly stated otherwise by NIBSC, NIBSC's Standard Terms and Conditions for the Supply of Materials (available at http://www.nibsc.org/About\_Us/Terms\_and\_Conditions.aspx or upon request by the Recipient) ("Conditions") apply to the exclusion of all other





terms and are hereby incorporated into this document by reference. The Recipient's attention is drawn in particular to the provisions of clause 11 of the Conditions.

# 16. INFORMATION FOR CUSTOMS USE ONLY

Country of origin for customs purposes\*: United Kingdom
\* Defined as the country where the goods have been produced and/or sufficiently processed to be classed as originating from the country of supply, for example a change of state such as freeze-drying.

Net weight: 1.0 g

Toxicity Statement: Toxicity not assessed

Veterinary certificate or other statement if applicable.

Attached: No