

WHO International Standard WHO 2nd International Standard for Streptodornase NIBSC code: 08/230 Instructions for use (Version 2.0, Dated 13/04/2010)

1. INTENDED USE

This standard is intended for use in the determination of streptodornase activity in preparations of streptokinase.

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

Each ampoule when reconstituted with 1 ml of water contains 3200 IU of streptodornase

4. CONTENTS

Country of origin of biological material: United Kingdom.

The active ingredient in this refrence preparation is streptodornase which is prepared from Streptococcus bacteria. Each ampoule also contains approximately 5 mg of human albumin included as a stabiliser. Albumin is prepared from human plasma, which has been tested and found negative for HBsAg, HIV antibody, HCV antibody and HCV RNA by PCR.

5. STORAGE

Ampoules should be stored at -20 °C upon receipt. The product is stable for shorter periods during shipping at ambient temperatures

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

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.

International Standards are intended to be used to calibrate other, working, standards and not for use directly in assays as references or reagents.

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.

NIBSC follows the policy of WHO with respect to its reference materials.

9. REFERENCES

A report of the collaborative study is available from WHO, reference number WHO/BS/09.2112

10. ACKNOWLEDGEMENTS

We are grateful to all laboratories who took part in the collaborative study to calibrate this standards (see WHO/BS/09.2112 for complete list).

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

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

No 1272/2008: Not applicable or not classified				
Physical and Chemical properties				
Physical appearance: white solid		Corrosive:	No	
Stable: Yes		Oxidising:	No	
Hygroscopic: Yes	ygroscopic: Yes		No	
Flammable: No	ammable: No		Handling:See caution, Section 2	
Other (specify): contains material of human and bacterial origin				
Toxicological properties				
Effects of inhalation: Not		established, avoid inhalation		
Effects of ingestion: N		ot established, avoid ingestion		
Effects of skin absorption: No		t established, avoid contact with skin		
Suggested First Aid				
Inhalation: Seek medical advice				
Ingestion: Seek medical advice				
Contact with eyes: Wasl				
medical advice				
Contact with skin: Wasl	Wash thoroughly with water.			
Action on Spillage and Method of Disposal				

Spillage of 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.





NIBSC Confidence in Biological Medicines

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: 20 mg

Toxicity Statement: Toxicity not assessed

Veterinary certificate or other statement if applicable.

Attached: No

17. CERTIFICATE OF ANALYSIS

NIBSC does not provide a Certificate of Analysis for WHO Biological Reference Materials because they are internationally recognised primary reference materials fully described in the instructions for use. The reference materials are established according to the WHO Recommendations for the preparation, characterization and establishment of international and other biological reference standards http://www.who.int/bloodproducts/publications/TRS932Annex2_Inter_bi olefstandardsrev2004.pdf (revised 2004). They are officially endorsed by the WHO Expert Committee on Biological Standardization (ECBS) based on the report of the international collaborative study which established their suitability for the intended use.