

WHO Reference Reagent Poliovirus Reference Sabin Type 1 NIBSC code: 01/528 Instructions for use (Version 2.0, Dated 08/04/2010)

1. INTENDED USE

Preparation $01/\overline{5}28$ was established thorough an International collaborative study involving the six WHO Global Specialized Polio Laboratories. It is intended to be used in reference laboratories to measure the sensitivity of cell cultures for poliovirus infection, or other laboratory assays, see 7 below.

CAUTION

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

The material is of bovine origin. The material is certified to be obtained from animals taken from a closed herd in the female line since 1980, in which no animal has been clinically suspected of having BSE & which has not been fed rations containing ruminant derived protein during that period. 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. should be exercised in opening ampoules or vials, to avoid cuts.

UNITAGE

The expected virus titre for 01/528 reference standard is 5.1 log 10 CCID50/0.1 ml in RD cells and 4.9 log 10 CCID50/0.1 ml in L20B cells

4. CONTENTS

Country of origin of biological material: Belgium. Each vial contains approximately 800 µl of liquid containing: authenticated Poliovirus type 1 Sabin strain [LS-c, 2ab strain]. Minimal essential medium with foetal calf serum Thermal stabilisers were not added to this preparation

5. STORAGE

Unopened ampoules should be stored at -20°C or colder. Repeated freeze-thawing should be avoided.

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

6. DIRECTIONS FOR OPENING

Vials have a screw cap; an internal stopper may also be present. The cap should be removed by turning anti-clockwise. Care should be taken to prevent loss of the contents. Please note: If a stopper is present on removal of the cap, the stopper should remain in the vial or be removed with the cap.

7. USE OF MATERIAL

The 01/528 reference standard should be used to set up a laboratory procedure to evaluate cell-line sensitivity for poliovirus infection following WHO guidelines (see corresponding reference in section 9). It can also be used by laboratories seeking authenicated Sabin strains for other laboratory assays, such as virucidal tests for disinfectants. This material is supplied for use in its final form and must not be further diluted other than as required for individual assay procedures.

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

World Health Organization. 2004. Evaluation of cell-line sensitivity, in Polio laboratory manual, 4th Edition. WHO/IVB04.10. Pages 73-80.

10. ACKNOWLEDGEMENTS

We acknowledge all participants in the collaborative study from France, Holland, UK, USA, India and Japan.

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

Physical and Chemical properties			
Physical appearance:		Corrosive:	No
Liquid			
Stable: Yes		Oxidising:	No
Hygroscopic: No		Irritant:	No
Flammable: No		Handling:See caution, Section 2	
Other (specify):			
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: Seek medical advice			
Ingestion: Seek medical advice			
medical advice			
Contact with skin: Was	Wash thoroughly with water.		



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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.

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: 0.5 g

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_biolefstandardsrev2004.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.

