

WHO Reference Reagent WHO International Reference Reagent for Antibodies to Q Fever (Coxiella burnetii) Antigens (human plasma) NIBSC code: 21/304 Instructions for use (Version 1.0, Dated 07/11/2023)

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1. INTENDED USE

The intended use of this reference reagent is to allow cross comparison, standisation and harmonisation of immunological assays used in biological research as well as in preclinical and clinical studies of vaccine development. Can be used as a standard or reference material in relation to manufacturing or quality control testing of biological products or in the field of in vitro diagnostics.

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

The content of this preparation has been asses by a collaborative study to have an activity of 100 100 U/ampoule for Phase I antigens 16 U/ampoule for Phase II antigens

4. CONTENTS

Country of origin of biological material: The Netherlands. The preparation contains material of human origin, which has been tested and found negative for HBsAg, HIV antibody, HCV antibody and HCV RNA by PCR. Each ampoule contains 0.5 mL of freeze-dried material of pooled convalescent plasma.

5. STORAGE

This preparation should be stored at -20C

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 freezedried material prior to reconstitution

The material should be reconstituted in 0.5 mL distilled water or millipore water and stored at -20C until needed

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8. STABILITY

Reference materials are held at MHRA South Mimms Site within assured, temperature-controlled storage facilities. Reference Materials should be stored on receipt as indicated on the label.

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

9. REFERENCES

N/A

10. ACKNOWLEDGEMENTS

Special thanks are due to Dr Boris Hogema and Dr B Tomson of the Sanquin Blood Supply Foundation (The Netherlands) for donating the material used in this preparation. Special thanks are also due to all the participating laboratories and scientist involved in the collaborative study.

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: Freeze dried powder		Corrosive:	No	
Stable: Yes	r	Oxidising:	No	
Hygroscopi No c:		Irritant:	Unknown	
Flammable: No		Handling: Se	e caution, Section 2	
Other Contains human plasma (specify):				
Toxicological properties				
Effects of inhalation: Not		established, avoid inhalation		
Effects of ingestion: Not		established, avoid ingestion		
Effects of sl absorption:	kin Not skin		avoid contact with	



Medicines & Healthcare products Regulatory Agency



Suggested First Aid			
Inhalation:	Seek medical advice		
Ingestion:	Seek medical advice		
Contact with	Wash with copious amounts of water. Seek		
eyes:	medical advice		
Contact with	Wash thoroughly with water.		
skin:			

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: 0.5175g per ampoule

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_l nter_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.

