WHO Reference Reagent
WHO REFERENCE REAGENT FOR HEPATITIS E VIRUS
ANTIBODY, human serum
NIBSC code: 95/584
Instructions for use
(Version 4.0, Dated 26/04/2013)

1. INTENDED USE
The WHO Reference Reagent for antibodies to hepatitis E virus was established in 1997 by the Expert Committee for Biological Standardisation and will serve as a biological reference preparation for antibodies to hepatitis E virus (Ferguson et al., 2002).

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 an eye shield. Take care that no material is lost from the ampoule and no material might enter the eyes, for example, by the use of suitable gloves and an eye shield. Take care that no material is lost from the ampoule and no material might enter the eyes, for example, by the use of suitable gloves and an eye shield. Care should be taken to avoid cuts and projectile glass fragments that might enter the eyes, for example, by the use of suitable gloves and an eye shield.

Disassemble the disposable ampoule breaker covering the ampoule by tapping it gently on a hard surface or a firm table to break the neck of the ampoule. Ensure that the disposable ampoule safety breaker provided is correctly fitted. Refer to Section 2 for a listing of hazardous materials that may cause injury.

Physical appearance: Freeze-dried

Corrosive: No
Oxidising: No
Irritant: No
Handling: See caution, Section 2

Contains material of human origin

3. UNITAGE
This material has an assigned unitage of 50 units per ampoule i.e. 100 units per ml when reconstituted as directed in 0.5 ml distilled water.

4. CONTENTS
Country of origin of biological material: United Kingdom.

Each ampoule contains a freeze-dried residue comprising, under an atmosphere of nitrogen: human serum containing antibodies to hepatitis E virus.

5. STORAGE
Unopened ampoules should be stored at -20°C or below.

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

6. DIRECTIONS FOR OPENING
Disassemble the disposable ampoule breaker covering the ampoule by tapping it gently on a hard surface or a firm table to break the neck of the ampoule. Ensure that the disposable ampoule safety breaker provided is correctly fitted. Refer to Section 2 for a listing of hazardous materials that may cause injury.

Physical appearance: Freeze-dried

Corrosive: No
Oxidising: No
Irritant: No
Handling: See caution, Section 2

Contains material of human origin

Other (specify): No

7. USE OF MATERIAL
No attempt should be made to weigh out any portion of the freeze-dried material prior to reconstitution.

Dissolve the total contents of the ampoule with 0.5ml of sterile distilled water. Ensure that the entire freeze dried residue is dissolved in this solution.

For economy of use, it is recommended that the solution be sub-divided into several small aliquots, stored at -20°C or below. Avoid repeated thawing/freezing.

8. STABILITY
Reference materials are held at NIBSC within assured, temperature-controlled storage facilities and they should be stored on receipt as indicated on the label. It is the policy of WHO not to assign an expiry date to their international reference materials. They remain valid with the assigned potency and status until withdrawn or amended.

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

9. REFERENCES

10. ACKNOWLEDGEMENTS

11. FURTHER INFORMATION
Further information can be obtained as follows;

This material: enquiries@nibsc.org
WHO Biological Standards:
http://www.who.int/biological/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 appearance: Freeze dried | Corrosive: | No |
| Stable: Yes | Oxidising: | No |
| Hygroscopic: No | Irritant: | No |
| Flammable: No | Handling: | See caution, Section 2 |
| Other (specify): | Contains material of human origin |

Other (specify): No
### 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 |
| Contact with eyes: | Wash with copious amounts of water. Seek medical advice |
| Contact with skin: | Wash thoroughly with water. |

### 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: | 0.5g |
| Toxicity Statement: | Non-toxic |
| 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_biologicalstandardsrev2004.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.