Influenza Reagent
Influenza Anti N3 Neuraminidase Serum SH503
NIBSC code: 09/100
Instructions for use
(Version 2.0, Dated 22/08/2011)

1. INTENDED USE
Influenza antiserum reagent 09/100 is prepared in sheep for neuraminidase identity tests.

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

3. UNITAGE
No unitage is assigned to this material.

4. CONTENTS
Country of origin of biological material: United Kingdom.
The antiserum reagent was prepared in sheep SH503 to NIBRG-107 (H2N2) virus . NIBRG-107, derived from A/mallard/England/727/2006 and prepared by reverse genetics at NIBSC. The antigen was extracted from purified virus by treatment with bromelain and purified by sedimentation on sucrose gradients (Brand, CN and Skehel, JJ, Nature, New Biology, 1972, 238, 145-147.

The immunisation schedule for sheep SH503 was as follows: One dose of approximately 50 micrograms of virus protein with Freund’s Complete Adjuvant was given intramuscularly, followed two weeks later with a 10 microgram dose of virus protein with Freund's Incomplete Adjuvant (FIA), four further 10 microgram doses of virus protein with FIA were given at weekly intervals. Seven weeks after the initial immunisation, serum was collected and sodium azide (0.05% w/v) was added.

The serum was treated by maintenance of pH5.49 for 40 minutes followed by restoration of the original pH and filled into vials in 1ml volumes.

5. STORAGE
+2-8°C
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 ‘flip-up’ circular cap. Either on the cap or the collar of the vial, there is an indication of the point at which to lever off the cap. This exposes an area of the stopper through which reconstitution and withdrawal of the preparation can be made using a hypodermic needle and syringe. If use of a pipette is preferred, then fully remove the metal collar using, for example, forceps, taking care to avoid cuts by wearing appropriate gloves. Remove the stopper for access. Care should be taken to prevent loss of the contents.

7. USE OF MATERIAL
Reagent 09/100 should be used in tests of neuraminidase identity, such as the neuraminidase inhibition (NI) test of Aymard-Henry M, Coleman MT, Dowdle WR, Laver WG, Schild GC and Webster RG. Bull WHO, 1973, 48, 199-202.

Although reagent 09/100 is not assigned a unitage, it is issued before final testing has been completed, NI test titres will be advised as soon as testing is complete.

8. STABILITY
Reference materials are held at NIBSC within assured, temperature-controlled storage facilities. Reference Materials should be stored on receipt as indicated on the label, please also see attached storage information sheet.

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

9. REFERENCES
N/A

10. ACKNOWLEDGEMENTS
This reagent was produced on behalf of the FLUSECURE project with funding from the EU.

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

<table>
<thead>
<tr>
<th>Physical and Chemical properties</th>
<th>Corrosive</th>
<th>No</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stable: Yes Oxidising: No</td>
<td>No Handling: See caution, Section 2</td>
<td></td>
</tr>
<tr>
<td>Flammable: No Other (specify): Contains sheep serum and sodium azide</td>
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<table>
<thead>
<tr>
<th>Toxicological properties</th>
<th></th>
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<tbody>
<tr>
<td>Effects of inhalation: Not established, avoid inhalation</td>
<td></td>
</tr>
<tr>
<td>Effects of ingestion: Not established, avoid ingestion</td>
<td></td>
</tr>
<tr>
<td>Effects of skin absorption: Not established, avoid contact with skin</td>
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</tbody>
</table>

<table>
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<tr>
<th>Suggested First Aid</th>
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<tbody>
<tr>
<td>Inhalation: Seek medical advice</td>
</tr>
<tr>
<td>Ingestion: Seek medical advice</td>
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</tbody>
</table>
Contact with eyes:  Wash with copious amounts of water. Seek medical advice
Contact with skin:  Wash thoroughly with water.

<table>
<thead>
<tr>
<th>Action on Spillage and Method of Disposal</th>
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</thead>
<tbody>
<tr>
<td>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.</td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Country of origin for customs purposes*: United Kingdom</th>
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<tbody>
<tr>
<td>* 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.</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Net weight: 1g</th>
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<tr>
<td>Toxicity Statement: Non-toxic</td>
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Veterinary certificate or other statement if applicable.
STORAGE OF REAGENT 09/100

NIBSC has prepared a number of reagents for single radial diffusion assay of influenza subtypes of pandemic potential.

Since it is not known when these reagents may be required, it is desirable that they have an indefinite shelf life and they are stored at NIBSC in colder conditions than reagents prepared for the assay of epidemic strains. Therefore the recommended storage temperature marked on the label for reagent 09/100 is -20°C.

However it is assumed that a customer ordering this reagent, will use it within a short period similar to that for a conventional reagent. Consequently, this reagent is not normally shipped frozen and the recommended storage temperature is +4°C.
VETERINARY CERTIFICATE OF ANIMAL HEALTH

This is to certify that I have today examined a Sheep with ear tag number: UK 281 038 2010 [Virology no. SH503], which has been used in the production of blood antiserum between 12th February 2009 and 7th April 2009. Both the ear tag number and the animals’ record show that it is of UK origin.

This animal was a breeding Ewe which became surplus to requirements. In my opinion at the time of clinical examination, the ewe was in good health and showed no clinical signs of infectious disease.

Kath Hardcastle BVetMed Cert LAS MRCVS
Named Veterinary Surgeon

Kath Hardcastle BVetMed Cert LAS MRCVS
Office: 020 7468 5333 Fax: 020 7468 5378 Mobile: 07809199998
Email: khardcastle@rvc.ac.uk
Foot and Mouth Disease Virus Inactivation Certificate

This is to certify that serum collected from Sheep no.UK 281 038 2010 [Virology no.SH503 ] has been treated by an APHIS approved method for inactivation of Foot and Mouth Disease Virus. The treatment method used was maintenance of pH5.5 or lower for a minimum of 40 minutes.

Dr Philip Minor
Deputy Director
National Institute for Biological Standards and Control