WHO International Standard
Timothy Pollen Extract
NIBSC code: 82/520
Instructions for use
(Version 5.0, Dated 24/07/2013)

1. INTENDED USE
The first International Standard for Timothy Grass (*Phleum pratense*)
pollen extract consists of ampoules, coded 82/520, containing the
freeze dried residue of 1ml aliquots of an extract of Timothy Grass
pollen. This preparation was established as the 1st International
standard for Timothy Grass (*Phleum pratense*) pollen extract by the
Expert Committee on Biological standardisation

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

The material is not of human or bovine origin. 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
A potency of 100,000 International units has been assigned to each
ampoule.

4. CONTENTS
Country of origin of biological material: United Kingdom.
The bulk material for the 1st International Standard for Timothy Grass
(*Phleum pratense*) pollen extract consisted of a freeze dried extract
prepared from a mixture of Timothy Grass pollen from 2 different lots
commissioned from a commercial allergen manufacturer. The pollen
used was more than 99% pure Timothy Grass pollen and there was
less than 5% non-pollen particles by weight. The pollen was extracted
using a 0.125M ammonium bicarbonate solution. The extract was
centrifuged, filtered and donated by the company as 4.5 litres of liquid.
It was stored at the National Institute for Biological Standards and
Control (NIBSC) at 4°C prior to filling.

At NIBSC the material was re-filtered ending with a Millipore
membrane of pore size of 0.22μm, homogenised by stirring and
distributed in 1ml volumes at +4°C into 4,00 ampoules, coded 82/520.
The mean weight of liquid content 68 checkweight ampoules taken at
intervals during the fill was 1.0021 +/- 0.049%. The contents of the
ampoules were then freeze dried and secondarily desiccated under the
conditions normally used for international biological standards. The
mean dry weight was 31.66mg +/- 0.47% (n=5) (of which
approximately 6mg was protein) and the moisture content was
0.3413% (n=5).

5. STORAGE
Unopened ampoules should be stored at -20°C.
Please note: because of the inherent stability of lyophilized
material, NIBSC may ship these materials at ambient temperature.

6. DIRECTIONS FOR OPENING
Tap the ampoule gently to collect the material at the bottom (labelled)
end. Ensure ampoule is scored all round at the narrow part of the
neck, with a diamond or tungsten carbide tipped glass knife file or
other suitable implement before attempting to open. Place the
ampoule in the ampoule opener, positioning the score at position 'A';
shown in the diagram below. Surround the ampoule with cloth or layers
of tissue paper. Grip the ampoule and holder in the hand and squeeze at
point 'B'. The ampoule will snap open. Take care to avoid cuts and
projectile glass fragments that enter eyes. Take care that no material is
lost from the ampoule and that no glass falls into the ampoule.

7. USE OF MATERIAL
No attempt should be made to weigh out any portion of the freeze-dried
material prior to reconstitution
The total contents of the ampoule should be reconstituted in 1.0ml of distilled
water and dissolved by gently swirling to avoid froth. The reconstituted
standard should be used as soon as possible after reconstitution.

8. STABILITY
Accelerated degradation studies have shown that the 1st International
Standard is very stable in unopened ampoules stored at -20°C.
No significant loss of activity was found in ampoules stored for 6 months
at temperatures up to 34°C.
Reference materials are held at NIBSC within assured, temperature-
controlled 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
collaborative study establishing the first international standard for Timothy
(*Phleum pratense*) grass pollen allergenic extract. J. Allergy Clin.
2. Campbell, P.J International Biological Standards and Reference
Preparations. II. Procedures used for the production of biological

10. ACKNOWLEDGEMENTS
All participants in the collaborative study.
We would like to express our gratitude to the following, who by provision
of finance, and in some case
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Copenhagen, Denmark; Antigen Laboratories, Liberty, Mo., USA;
Beecham, Betchworth, Surrey, UK; Berkeley Biologicals, Berkeley, Calif.
USA; Center Laboratories, Port Washington, N.Y, USA; Diephuis
Pharmacia, Groningen, The Netherlands; Greer Laboratories, Lenoir,
N.C, USA; HAL Allergen Laboratories, Haarlem, The Netherlands;
Hollister-Stein, Spokane, Wash., USA; Laboratories Hamon, Montreal,
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:
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>
</tr>
</thead>
<tbody>
<tr>
<td>Physical appearance: Freeze-dried powder</td>
</tr>
<tr>
<td>Stable: Yes</td>
</tr>
<tr>
<td>Hygroscopic: No</td>
</tr>
<tr>
<td>Flammable: No</td>
</tr>
<tr>
<td>Other (specify):</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Toxicalogical properties</th>
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<tbody>
<tr>
<td>Effects of inhalation:</td>
</tr>
<tr>
<td>Effects of ingestion: Not established, avoid ingestion</td>
</tr>
<tr>
<td>Effects of skin absorption:</td>
</tr>
</tbody>
</table>

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 be classed as originating from the country of supply, for example a change of state such as freeze-drying.

Net weight: 0.01g
Toxicity Statement: Non-toxic
Veterinary certificate

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 materials.
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.

Attached: No