### WHO International Standard

#### 2nd International Standard 2016 Coagulation Factor XI, Plasma, Human

#### 1st International Standard 2017 Coagulation Factor XII, Plasma, Human

**NIBSC code: 15/180**

**Instructions for use**

*Version 2.0, Dated 20/12/2017*

### 1. INTENDED USE

This material is the 2nd International Standard for Factor XI, Plasma, Human and the 1st International Standard for Factor XII, Plasma, Human. It consists of ampoules, coded 15/180, containing approximately 1 mL aliquots of human normal plasma, freeze-dried. This preparation is intended for use in the measurement of FXI functional activity (FXI:C) and antigen (FXI:Ag) and FXII functional activity (FXII:C) and antigen (FXII:Ag) in plasma. In addition it can be used for potency assignment of FXI therapeutic concentrates.

The ECBS reports are available from the WHO [website](http://www.who.int/entity/biologicals/ECBS_2016_BS2281_FXI-2nd_IS_final.pdf)

http://www.who.int/biologicals/expert_committee/BS2326_FXII.pdf?ua=1

### 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 standard was assayed in an international collaborative study. The units assigned to this preparation are:

- Factor XI Functional activity (FXI:C): 0.71 IU/ampoule
- Factor XI Antigen (FXI:Ag): 0.78 IU/ampoule
- Factor XII Functional activity (FXII:C): 0.86 IU/ampoule
- Factor XII Antigen (FXII:Ag): 0.80 IU/ampoule

Results from 29 laboratories employing one-stage clotting assays were used to value assign functional activity (FXI:C) to the 1st IS relative to the 1st IS. The FXI antigen value (FXI:Ag) was assigned relative to local normal pooled plasma (total number of donors >20,000) by ELISA method, using data from 11 laboratories. For FXII, results from 20 laboratories using one-stage clotting assays were used to value assign functional activity (FXII:C) relative to local normal plasma pools (number of donors 566). The FXII antigen value (FXII:Ag) was also assigned relative to local normal plasma pools (216 donors), using results from 8 laboratories using ELISA methods.

Uncertainty: the assigned unitage does not carry an uncertainty associated with its calibration. The uncertainty may therefore be considered to be the variance of the ampoule content which was determined to be +/- 0.25 %.

### 4. CONTENTS

**Country of origin of biological material: United Kingdom.**

The material was prepared from a plasma pool derived from normal healthy donors (United Kingdom National Blood Transfusion Service). Blood was collected into CPD-adenine anticoagulant and subjected to two centrifugation steps after which the plasma was frozen rapidly and stored at -70°C until the day of ampoule filling. Individual donations were tested and found negative for HBsAg, anti-HIV-1/2 and anti-HCV. The material was formulated with glycine and a buffering agent HEPES (N-[2-Hydroxyethyl]piperazine-N'-[2-ethanesulfonic acid) at a final concentration of 1 % w/v and 40 mmol/L respectively. To avoid activation of FXI/FXII, polyethylene vessels were used for storage and transport of the pooled plasma. The frozen pooled plasma was thawed at 37°C and maintained at room temperature throughout the process. The material was filled into siliconised glass ampoules and freeze dried under conditions used for International Biologicals Standards (1).

**Activation status of the material:** The non-activated partial thromboplastin time (NAPTT) is known to be sensitive to activated clotting factor especially factor XIa and so it was used to assess the activation status of the finished product. The long mean clotting time of 300s (n = 9; sd ± 2.26) for 15/180 indicates the samples to be relatively unactivated.

### 5. STORAGE

Unopened ampoules should be stored in the dark below -20°C

**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.

- Tap the ampoule gently to collect the material at the bottom (labeled) end. Ensure that the disposable ampoule safety breaker provided is pushed down on the stem of the ampoule and against the shoulder of the ampoule body. Hold the body of the ampoule in one hand and the disposable ampoule breaker covering the ampoule stem between the thumb and first finger of the other hand. Apply a bending force to open the ampoule at the coloured stress point, primarily using the hand holding the plastic collar.

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. Take care that no material is lost from the ampoule and no glass falls into the ampoule. Within the ampoule is dry nitrogen gas at slightly less than atmospheric pressure. A new disposable ampoule breaker is provided with each DIN ampoule.

### 7. USE OF MATERIAL

No attempt should be made to weigh out any portion of the freeze-dried material prior to reconstitution

- Allow ampoules to warm to room temperature. Open ampoule, taking care to ensure that all material is in the lower part, and reconstitute with 1.0 mL of distilled water. After reconstitution please store the material as indicated in section 8 (On Bench Stability).

### 8. STABILITY

Referential 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 material. It is the policy of WHO not to assign expiry dates to international reference materials. They remain valid with the assigned potency and status until withdrawn or amended. Accelerated degradation studies, which involve potency estimation of ampoules stored at elevated temperatures relative to ampoules stored at below -150°C, has indicated that the Standard is stable when stored at -20° or below.
On Bench Stability: It is recommended that assays are to be performed as soon as possible after reconstitution. The stability of coagulation factors in plasma standards, after reconstitution, is mainly affected by two components - the surface of the container and the storage temperature. Unlike other WHO IS for blood coagulation factors it is recommended that upon reconstitution, the standard should either be transferred to a plastic tube or retained in the siliconised ampoule at room temperature (18-22 °C) in order to prevent cold activation of FXI or FXII. Results from NIBSC indicated no significant change in FXI or FXII clotting activity or antigen measurement when the reconstituted material was stored at room temperature in the siliconised ampoules for over 3 hours. However, users will be advised that local validation will be necessary if the reconstituted standard is stored under different conditions. The use of frozen aliquots of this International Standard cannot be recommended since the effect of freezing and thawing, under local conditions, on the FXI or FXII activity is unpredictable.

9. REFERENCES

10. ACKNOWLEDGEMENTS
The participants of the 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_stands.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
- **Stable:** Yes
- **Hygroscopic:** No
- **Flammable:** No
- **Corrosive:** No
- **Oxidising:** No
- **Irritant:** Yes
- **Handling:** See caution, Section 2
- **Contains material of human origin**

### Toxicological properties
- **Effects of inhalation:** Not established, avoid inhalation

### Suggested First Aid
**Inhalation:** Seek medical advice

### 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

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_Int_biol_esstandardsrev2004.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.