This manufacturers declaration is not part of USDA APHIS Certification. It is intended solely for use with consigments entering the European Union through Border Inspection Posts in England.

ANNEX I

MANUFACTURER'S DECLARATION

IMPORT OF BLOOD PRODUCTS UNDER MODEL HEALTH CERTIFICATES 4(C) AND 4(D) AS LAID DOWN IN ANNEX XV OF REGULATION (EU) NO 142/2011 (AS AMENDED) INTENDED FOR IMPORT TO THE UK VIA AN ENGLISH BORDER INSPECTION POST

NOTES

1. THE DECLARATION ONLY APPLIES TO BLOOD PRODUCTS IMPORTED USING MODEL HEALTH CERTIFICATES 4(C) AND 4(D) AS LAID DOWN IN ANNEX XV OF REGULATION (EU) NO 142/2011 (AS AMENDED) FOR IMPORTS INTO THE UK WHICH ARE TO BE LANDED IN A BORDER INSPECTION POST IN ENGLAND.

THIS DECLARATION DOES NOT APPLY TO ANY OTHER MEMBER STATE OR FOR THOSE CONSIGNMENTS LANDING AT A BIP IN WALES OR SCOTLAND.

2. THE DECLARATION MUST BE SIGNED AND STAMPED BY THE SENIOR MANAGER/ VETERINARY SUPERVISOR OF THE APPROVED THIRD COUNTRY ESTABLISHMENT WHICH HAS PRODUCED THE BLOOD PRODUCT INTENDED FOR IMPORT ON THEIR COMPANY HEADED PAPER.

3. THE DECLARATION MUST BE THE ORIGINAL SIGNED COPY AND MUST ACCOMPANY THE ORIGINAL HEALTH CERTIFICATE TO THE BORDER INSPECTION POST IN ENGLAND.

4. PHOTOCOPIES WILL NOT BE ACCEPTED AND THE DECLARATION WILL NOT BE TAKEN INTO ACCOUNT.

5. EACH HEALTH CERTIFICATE MUST BE ACCOMPANIED BY A NEW DECLARATION EACH TIME.

6. THE DECLARATION MUST BE WORDED AND SET OUT AS PER THE MODEL BELOW. FAILURE TO FOLLOW CORRECT LAYOUT AND WORDING WILL RESULT IN THE DECLARATION NOT BEING CONSIDERED AND THE CONSIGNMENT BEING CLASSED AS CATEGORY 1 MATERIAL.

<table>
<tr><td rowspan="2">1. Name and address of approved establishment where the blood product was produced (in full)

Approval number as per EU third country approval list</td><td>MANUFACTURER'S DECLARATION
Imports of blood products not intended for food or feed use –
To accompany the original health certificate as per model 4(C) or 4(D) of Annex XV of Regulation (EU) No 142/2011</td></tr>
<tr><td>Certificate reference number to which this declaration refers : (refer to certificate that has been issued)</td></tr>
<tr><td rowspan="3">2. Name and address of consignee (in full)</td><td>3. Country of origin
US-UNITED STATES</td></tr>
<tr><td></td></tr>
<tr><td colspan="2">4. Identification of the consignment (must match health certificate)</td></tr>
</table>

5. DECLARATION

I, the undersigned senior manager/ veterinary supervisor of the establishment which has produced the blood products intended for export to the UK/England hereby confirm:

1. The blood products are not intended for human or animal consumption;
2. The blood products comply with the requirements laid down in the health certificate 4(C)* (for untreated blood products) and/ or 4(D)***** (for treated blood products);
3. The blood product has been produced from blood that did not originate from Category 1 material as defined in Article 8(c) or 8(d) of Regulation (EC) No 1069/2009;
4. The blood products intended for export has been produced from Category 3 material as listed in the health certificate number: (insert Certificate Reference Number)
   …….....

Place Date

Signature of senior manager/ veterinary supervisor of manufacturing facility

……

Name and position in capital letters

Address

Signature must be in a different colour to that in the printed declaration