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ROYAL FREE HOSPITAL  
POND STREET  
LONDON NW3 2QG  
TELEPHONE 0171 794 0500



31st October

Nicholas Ryness HIRSCH

This is to confirm that the enclosed patients received the batch factor 8y FHB4419. This was in 1995 and we have no further bottles of this batch in use.

INTERNATIONAL TRAINING CENTRE OF THE WORLD FEDERATION OF HAEMOPHILIA



## Facsimile Transmission

To: *F.A.O Prof C. Lee, Hemophilia*

Customer: Royal Free Hospital  
Of: Pond Street London

Fax Number:

GRO-C

From:

Customer Services Department

*Jane Martin*

Total pages:

1

Date:

30/11/1997

Subject:

Product Recall Incident No. PR 97/205/19

Bio Products Laboratory

Dagger Lane

Elstree

Herts. WD6 3DX

Telephone: 0181 258 2200

Fax: 0181 258 2604

### Product Recall

In accordance with instructions received from the MCA, based upon new advice received from the CPMP, Bio Products Laboratory have initiated a recall of the following product which your organisation has received. The recall is a precautionary measure only related to post donation information. Subsequent to donation the donor was found not to have met the current health requirements for C-JD. The advice from the Lothian Ethical committee is that the recipients (patients) should not be informed that product that they have received has been recalled for this reason.

Product Names:	batch	dose	expiry
Factor VIII type 8Y	FHB4419	500iu	06/06/98
Human Albumin Solution Zenalb 4.5	ADA0529	500ml	14/11/97

Quantity, Batch number(s) and date(s) of dispatch to you:  
278 of FHB4419 on 14/9/95

You are kindly requested to return any remaining stock that conforms to the above batch details. If you have no stock left you should confirm that in writing. Please recall any of the above material that you have supplied. Action under the recall should be completed within 48 hours.

Please note that before returning any stock to BPL you must contact our sales office on 0181 258 2251 or 0181 258 2267 to arrange return and delivery of replacements. We will, of course, replace such stock free of charge.

Product is to be returned to our Dispatch Department, labelled with Incident Number PR 97/205/19 and containing the senders name and organisation.

We apologise for any inconvenience that this recall may cause you.  
Yours sincerely,

Pam Hurd  
Customer Services Manager

SHOULD TRANSMISSION BE UNCLEAR PLEASE CONTACT THE SENDER IMMEDIATELY

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BIO PRODUCTS LABORATORY



A part of the National Blood Authority.  
A Special Health Authority within the NHS.