QC-01

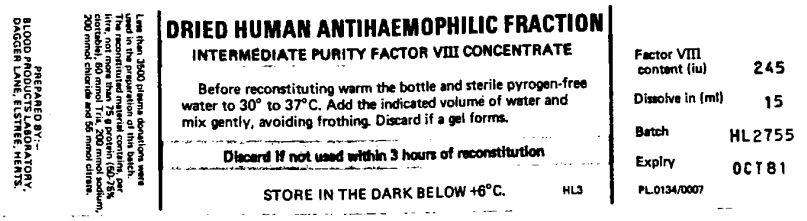
Final evaluation

Batch HL 2755

Name of product: HUMAN ANTIHAEMOPHILIC FRACTION
(Intermediate Purity Factor VIII Concentrate)

Product Licence No: 0134/0007.

Label: Completed sample label, (exactly as attached to vials)



Formulation of final product: See BP 1978 p.11 "Dried Antihaemophilic Fraction"

OCTOBER 80

QC-01

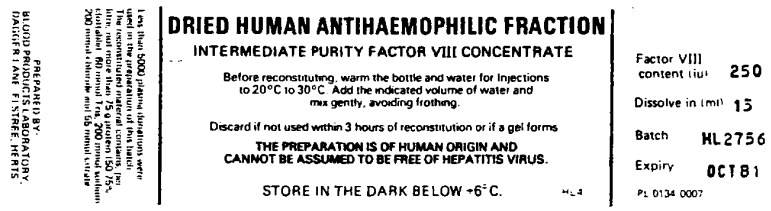
Final evaluation

Batch HL 2756

Name of product: HUMAN ANTIHAEMOPHILIC FRACTION
(Intermediate Purity Factor VIII Concentrate)

Product Licence No: 0134/0007.

Label: Completed sample label, (exactly as attached to vials)



Formulation of final product: See BP 1978 p.11 "Dried Antihaemophilic Fraction"
Label side-panels
Summary of Quality Control tests, QC-02 etc.

Fractionation date: 11/9/80

Sterilisation date: 11/9/80

Factor VIII assay date (for expiry): 3.10.80.

Sub-batches included in this batch:

540	2		
35 ml	35 ml	ml	ml

Vials filled at 35 ml before freeze-drying.

Contents recommended for re-solution in 15 ml Water For Injections