

Product	specification: Epirubicin pre-filled syringes
Name of product	Epirubicin injection. Aseptically prepared from licensed sterile starting materials.
Concentration	Concentration 2mg/ml in pre-filled syringes
Diluent	None
Volume	Dose specific volumes in accordance with the national dose banding tables
Final container	Polypropylene or polycarbonate sterile luer lock syringe, graduated in millilitres and sealed with sterile syringe cap/hub. Syringe and syringe cap/hub integrity to be valid. The validation should conform to the Protocols for the integrity testing of syringes published by the NHS Pharmaceutical QA Committee (2nd Edition, April 2013). Air bubbles within the final product must be minimised so the end user can use within need to remove air.
Starting materials	Licensed Epirubicin 2mg/ml solution for injection or powder for reconstitution for injection cells and syringes and syringe caps/hubs
Labelling	Labelling must be compliant with the principles of labelling for safety and the BP Gen specification on unlicensed medicines. Tall Man lettering must be used for the drug name.
Label sample	An example label is provided below stating the minimum requirements only (the labe format is not restrictive and suppliers can use their preferred layout):- EPIrubicin xxmg in xxml For Intravenous Injection Store in a Refrigerator at 2-8°C Protect From Light Expiry: dd/mm/yyyy BN: XXXXXXXXX Keep out of the reach and sight of children Manufacturer's details MS XXXXXX Caution Cytotoxic: Handle with care
Batch Number	All products will have a unique batch identification number
Latex status of - components - manufacturing process	All materials and manufacturing processes will be latex free or clearly labelled if not.
Stability	Stability studies should conform to the Standard Protocol for deriving and assessmer stability of Aseptic preparations (small molecules) published by the NHS Pharmaceu QA Committee (4th Edition, April 2017).