NHS England

Product specification: Vinblastine 50ml infusion bags

	Name of product	Vinblastine infusion. Aseptically prepared from licensed sterile starting materials.
	Concentration	Concentration range 0.1mg/ml to 0.3mg/ml in 50ml infusion bags in accordance with the national dose banding tables. The Vinblastine concentrate should be added to the bag without withdrawal of equivalent volume of Sodium Chloride 0.9% w/v Infusion unless withdrawal of some infusion fluid is required to accomodate the total dose.
	Diluent	Sodium Chloride 0.9% w/v
	Volume	50ml
	Final container	Non-PVC e.g. polyolefin infusion bag with additive port cover. *Ideally infusion bag design will incorporate self sealing giving port to minimise H&S risks associated with accidental spillage during administration.*
	Starting materials	Licensed Vinblastine Injection Licensed Sodium Chloride 0.9% w/v Infusion bags
	Labelling	Labelling must be compliant with the principles of labelling for safety (in particular must comply with HSC 2008/001 Updated national guidance on the safe administration of intrathecal chemotherapy and NPSA/2008/RRR004 i.e. must be labelled 'For Intravenous Use Only – Fatal If Given by Other Routes') and the BP General 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 label format is not restrictive and suppliers can use their preferred layout):- vinBLAStine xxmg in xxml Sodium Chloride 0.9% w/v FOR INTRAVENOUS USE ONLY FATAL IF GIVEN BY OTHER ROUTES Infuse the entire contents of the bag Store in a Refrigerator at 2-8^{O}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 assessment of stability of Aseptic preparations (small molecules) published by the NHS Pharmaceutical QA Committee (4th Edition, April 2017).