

Faculty of Pain Medicine Safety Checklist:

*Intrathecal Drug Delivery Device - In Hospital/Secondary Care Refill Checklist

Sign in to Theatre/Refill Room. (To be read out loud)

Place addressograph label here

I. Preparing for an Intrathecal Drug Delivery Device refill:

Please tick if appropriate:-

- Initial team brief undertaken and staff members have introduced themselves?
- Two personnel present for the pump refill and independent check?
(note - in exceptional circumstances one person could do as per local hospital guidance and SOP)
- Patient identity confirmed as per local protocol?
- Is the necessary equipment available for pump refill (Prescription, IT programmer, Drugs and Refill kit, Ultrasound/Radiology guidance if required)?
- Have personnel confirmed drugs in refill syringe match the physician prescription as per local prescribing policy?

II. Refilling the Intrathecal Drug Delivery Device:

Please tick or cross. If a cross, kindly address:-

- IT pump programme read and expected residual volume documented?
- IT pump reservoir accessed easily?
Record difficulties here:.....
- Does the actual reservoir volume (aspirated) correlate with the expected reservoir volume?
If (X), confirm the access needle is placed correctly?
If still unexplained, is this discrepancy within the manufacturer recommended tolerance limits?
If discrepancies are outside the manufacturer recommended tolerance limits or if there is failure to aspirate the reservoir, then involve a senior colleague and consider technical guidance from the manufacturer. If you are unsure always seek help.
- Aspirated drug discarded as per local drug policy?
- Refill drug easy to aspirate fully from the reservoir after delivery of the first 5mls
If (X), check needle position

III. Following Intrathecal Drug Delivery Device Refill:

- ITDD programmed with new reservoir volume and doses cross checked?
- Pump alarm date documented and communicated to patient and refill date organised as per local practice?
- ITDD residual battery life over 18 months?
If under 18 months confirm date documented and communicated to the patient
- Patient to remain in the clinical area for 30 minutes following refill, in order to detect any early adverse effects of the pump refill.
NB: Side effects from an inadvertent pocket refill can be delayed and in cases of doubt the patient should be admitted to a HDU area

*This document is a checklist and appropriate assessment of the patient should be done as per local protocol.

Signatures

A.....

B.....