

INSTRUCTIONS FOR USE

	Perform appropriate hand hygiene when preparing any medication.
	Check the order and confirm the right solution and the right patient.
50 mL	Remove the aluminum overwrap.
DIN 02485532 FBCA4005E STERILE SOLUTION/SOLUTION STÉRILE [®] HYDROmorphone Hydrochloride Injection USP [®] Chlorhydrate d'HYDROmorphone injectable USP 20 mg / 50 mL (0.4 mg / mL)	 Firmly squeeze the bag and check for leaks; if leaks found, do not use the bag as sterility may be impaired.
Opioid Analgesic – Isotonic / Single dose for IV Use 0thy Analgesique opioide – Isotonique / Dose unique pour usage IV seulement Each mt. contins: Hr090cmpother bydrochtied 0. Ang. unic acid 0.104 mg (as anhydroxis, sodum ottabe 0.1357 mg (as Brightath), sodum ottahet 0.9% and vatar for injection. Presentate Fields: Enterbanets 1 and 2577. Protect from freeding.	9 Perform visual inspection to check for particles; only use clear solution.
Chapter mit, content: chapter de docum. 1374 mit de docum. 144 mit de docum. 1374 et au pour injection. Stars agent de conservation. Conserver entre 15 et 2572. Profuge du get. ESTERCIMENTE: L'Altre de docum. 1374 et au pour injection. Stars agent de conservation. Conserver entre 15 et 2572. Profuge du get. Later-free / Nan-BitP / Nan-PVC Container contentient stars Contentient stars append to conserve documents? / B4-4:282-2838 St-Hyacimithe Oc Canada J25 0.39 (01) 0 72:40176 00016 3	6 Open the puncture port of the bag by twisting the two flaps in opposite directions to break the first flap (see illustration). NOTE : The puncture port septum remains closed and sterile right after removal of the tip.
	Follow the infusion set instructions to properly connect to the puncture port.
ADDITIVE PORT	Follow the infusion pump instructions to properly connect the infusion set.
PUNCTURE PORT	
The	 Verify the infusion pump programing and make sure vou follow your institution core rights for safe medication administration before starting the infusion.
	Patients should be monitored for respiratory depression, especially during initiation of ®HYDROmorphone Hydrochloride Injection USP or following a dose increase.
The Additive port may be covered with a tamper-proof cap. Do not remove.	The ready to use ®HYDROmorphone Hydrochloride Injection USP is for single use only. No preservative.
	Discard any unused portion as per your hospital protocol. * Please refer to the product monograph for detailed information.

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