PrNALOXONE HYDROCHLORIDE INJECTION

AHFS 28:10.00 • OPIATE ANTAGONISTS

PRODUCT INFORMATION

THESE PRODUCTS ARE AVAILABLE AT YOUR LOCAL DISTRIBUTOR

GENERIC NAME	NALOXONE HYDROCHLORIDE INJECTION		
ADMINISTRATION	For intravenous, intramuscular or subcutaneous administration.		
STORAGE CONDITIONS	Store between 15 and 30°C. • Protect from light. • Discard 28 days after initial use.		
INDICATION	Indicated for the complete or partial reversal of opioid depression, including respiratory depression induced by opioids, including natural and synthetic opioids, propoxyphene, methadone and the agonist-antagonist nalbuphine, pentazocine and butorphanol. Naloxone is also indicated for the diagnosis of suspected acute opioid overdosage.		
STRENGTH	**0.4 mg/mL**		**1 mg/mL**
DIN	02393034		02393042
FORMAT	10 x **1 mL** Sterile multidose vial	1 x **10 mL** Sterile multidose vial	10 x **2 mL** Sterile multidose vial
PACKAGING	2 mL vial Latex-free stopper White Flip-off cap	10 mL vial Latex-free stopper White Flip-off cap	2 mL vial Latex-free stopper Blue Flip-off cap
UPC	8 01500 11222 0	8 01500 11223 7	8 01500 11224 4
GS1 DataBar (vial)	(01)10801500112227	(01)10801500112234	(01)10801500112241
OMEGA PRODUCT CODE	L0010222	L0010223	L0010224
MCKESSON PRODUCT CODE	045770	045744	045772
CPDN PRODUCT CODE	912220	912237	912244

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