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Current through July 1, 2025

You have searched **Finished drug products**

Search Results: '0641-0477'

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Proprietary Name	NDC Package Code	Strength	Dosage Form	Route	Application Number / Monograph ID	Labeler Name	Product NDC	Non Proprietary Name	Substance Name	Product Type
 Phenobarbital Sodium	0641-0477-25	130mg/mL	INJECTION	INTRAMUSCULAR; INTRAVENOUS	N/A	Hikma Pharmaceuticals USA Inc.	0641-0477	Phenobarbital Sodium	PHENOBARBITAL SODIUM	HUMAN PRESCRIPTION DRUG
Start Marketing Date 01/01/1971										
End Marketing Date N/A										
Market Category UNAPPROVED DRUG OTHER										
Package Description 25 VIAL in 1 CARTON (0641-0477-25) / 1 mL in 1 VIAL (0641-0477-21)										
Pharm Class N/A										
DEA CIV										
Sample Package No										
Listing Record Certified Through 12/31/2025										

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