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Current through July 1, 2025

You have searched Finished drug products

Search Results: *'0641-0476'*

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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 Name
- Phenobarbital Sodium	0641-0476-25	65mg/mL	INJECTION	INTRAMUSCULAR; INTRAVENOUS	N/A	Hikma Pharmaceuticals USA Inc.	0641-0476	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 (0641-0476-25) / 1 mL in 1 VIAL (0641-0476-21)

Pharm Class N/A

DEA CIV

Sample Package No

Listing Record Certified Through 12/31/2025

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Nivagen Ex. 2025
Sun v. Nivagen
IPR2025-00893