NEO-SYNEPHRINE HYDROCHLORIDE- phenylephrine hydrochloride injection, solution
Hospira, Inc.

Neo-Synephrine® HCl

PRINCIPAL DISPLAY PANEL - 1 mL Ampule Label

1 mL
NDC 0409-1800-01

Neo-Synephrine®
HCl

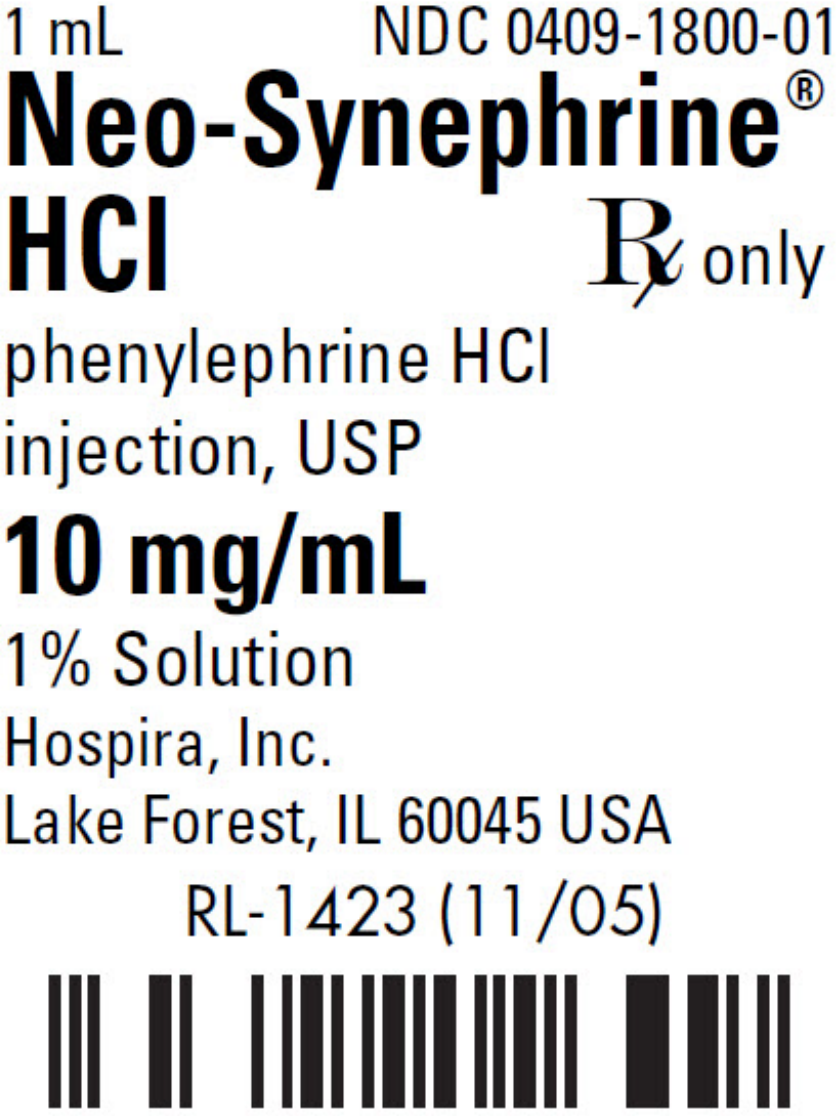
Rx only

phenylephrine HCl
injection, USP

10 mg/mL

1% Solution
Hospira, Inc.
Lake Forest, IL 60045 USA

RL-1423 (11/05)



PRINCIPAL DISPLAY PANEL - 1 mL Ampule Cello Pack Label

1 mL (10 mg)
Single-dose
5 Ampuls
NDC 0409-1800-01

Neo-Synephrine® HCl
phenylephrine hydrochloride injection, USP
1% Sterile Aqueous Injection

FOR PARENTERAL USE
Rx only

PROTECT FROM LIGHT.

Each mL contains 10 mg NEO-SYNEPHRINE hydrochloride, brand of phenylephrine hydrochloride, 3.5 mg sodium chloride, 4 mg sodium citrate, 1 mg citric acid monohydrate, and not more than 2 mg **sodium metabisulfite** as preservative.

For usual dosage and route of administration, see package insert. Store at 20 to 25°C (68 to 77°F). [See USP Controlled Room Temperature.]
WARNING: CONTAINS SULFITES.

Printed in USA
Hospira, Inc., Lake Forest, IL 60045 USA
Hospira

RL-2067 (3/07)

1 mL (10 mg) Single-dose 5 Ampuls NDC 0409-1800-01

Neo-Synephrine® HCl

phenylephrine hydrochloride injection, USP
1% Sterile Aqueous Injection

FOR PARENTERAL USE ℞ only

PROTECT FROM LIGHT.

Each mL contains 10 mg NEO-SYNEPHRINE hydrochloride, brand of phenylephrine hydrochloride, 3.5 mg sodium chloride, 4 mg sodium citrate, 1 mg citric acid monohydrate, and not more than 2 mg **sodium metabisulfite** as preservative. For usual dosage and route of administration, see package insert. Store at 20 to 25°C (68 to 77°F). [See USP Controlled Room Temperature.]

WARNING: CONTAINS SULFITES.

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Hospira, Inc., Lake Forest, IL 60045 USA

Hospira

(01) 1 030409 180001 1

RL-2067 (3/07)

NEO-SYNEPHRINE HYDROCHLORIDE

phenylephrine hydrochloride injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:0409-1800
Route of Administration	SUBCUTANEOUS, INTRAMUSCULAR, INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -	PHENYLEPHRINE	10 mg

UNII:1WS297W6MV)	HYDROCHLORIDE	in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X)	3.5 mg in 1 mL
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	4 mg in 1 mL
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	1 mg in 1 mL
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0409-1800-01	5 in 1 CELLO PACK	03/14/2006	
1		1 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
EXPORT ONLY		03/14/2006	

Labeler - Hospira, Inc. (141588017)

Establishment

Name	Address	ID/FEI	Business Operations
Hospira, Inc.		093132819	ANALYSIS(0409-1800) , LABEL(0409-1800) , MANUFACTURE(0409-1800) , PACK(0409-1800)

Revised: 1/2021 Hospira, Inc.