

**LEVOTHYROXINE SODIUM- levothyroxine sodium tablet
DirectRX**

LEVOTHYROXINE SODIUM

Description

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Clinical Pharmacology

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Indications and Usage

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Contraindications

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Warnings

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Precautions

-

Precautions section continued

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Adverse Reactions

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Overdosage

-

Dosage and Administration

-

Package Label

D

LEVOTHYROXINE SODIUM
88mcg 30 Tabs

Generic For: **SYNTHROID**
Each tablet contains: Levothyroxine Sodium, USP
88mcg (0.088mg)

Lot# 976-30 Discard After: 02/18

Produced and Distributed By: **DIRECT**

Alpharetta, GA 30005

Mfg For: Lannett Co., Inc.
Philadelphia, PA 19136
NDC 0527-1344-01

Mfg Lot: 12/2015

AFFECT
Caution: Federal law prohibits transfer of this drug to any person other than the patient for whom it was prescribed.

RX ONLY-KEEP OUT OF REACH OF CHILDREN
Dosage: See package insert. Store between 68-77 degrees F.

M

NDC 61919-976-30

LEVOTHYROXINE SODIUM 88mcg
NDC 61919-976-30 30 Tabs
Lot Exp Date 02/18
Mfg NDC 0527-1344-01

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LEVOTHYROXINE SODIUM

levothyroxine sodium tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-976(NDC:0527-1344)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEVOTHYROXINE SODIUM (UNII: 9J765S329G) (LEVOTHYROXINE - UNII:Q51BO43MG4)	LEVOTHYROXINE SODIUM ANHYDROUS	0.088 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ACACIA (UNII: 5C5403N26O)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	green (Olive)	Score	2 pieces
Shape	ROUND	Size	7mm
Flavor		Imprint Code	JSP;561
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-976-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/02/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA authorized generic	NDA021210	12/02/2015	

Labeler - DirectRX (079254320)**Establishment**

Name	Address	ID/FEI	Business Operations
DirectRX		079254320	repack(61919-976)

Revised: 12/2015

DirectRX