

**LEVOTHYROXINE SODIUM- levothyroxine sodium tablet  
DirectRX**

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**LEVOTHYROXINE SODIUM**

**Description**

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**Clinical Pharmacology**

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**Indications and Usage**

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**Contraindications**

- 

**Warnings**

- 

**Precautions**

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**Precautions continued**

**Adverse Reactions**

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**Overdosage**

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**Dosage and Administration**

- 

**Package Label**

Mfg For: Lannett Co. Inc.  
Philadelphia, PA 19136  
NDC 0527-1342-01

**D**

**LEVOTHYROXINE SODIUM  
50mcg 30 Tabs**

Generic For: **SYNTHROID**  
Each tablet contains: Levothyroxine Sodium, USP  
50mcg (0.05mg)

Lot#  
Prod# 403-30

Discard After: 02/18

Packaged and Distributed By: **DIRECT B**

Alpharetta, GA 30005

**M**

NDC 61919-403-30

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

LEVOTHYROXINE SODIUM 50mcg  
NDC 61919-403-30 30 Tabs  
Lot Exp Date 02/18  
Mfg NDC 0527-1342-01

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Lot Exp Date 02/18  
Mfg NDC 0527-1342-01

Mfg Lot:  
12/2/2015

Revision | Effective Date

## LEVOTHYROXINE SODIUM

levothyroxine sodium tablet

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-403(NDC:0527-1342)
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEVOTHYROXINE SODIUM (UNII: 9J765S329G) (LEVOTHYROXINE - UNII:Q51BO43MG4)	LEVOTHYROXINE SODIUM ANHYDROUS	0.05 mg

### Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ACACIA (UNII: 5C5403N26O)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
LACTOSE (UNII: J2B2A4N98G)	
STARCH, CORN (UNII: O8232NY3SJ)	

### Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	7mm
Flavor		Imprint Code	JSP;514
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:6 19 19-403-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2015	

**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(6 19 19-403)

Revised: 12/2015

DirectRX