NP THYROID- thyroid tablets tablet Calvin Scott & Co Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Take 1 tablet daily on an empty stomach or as directed

Do not exceed recommended dose. Keep out of reach of children.

Calcium stearate, dextrose, maltodextrin, mineral oil, modified wheat starch

Used as a thyroid hormone replacement for hypothyroidism.

Thyroid hormone replacement

Thyroid 90 mg

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center immediately.

NDC 17224-504-30

Package label for NP Thyroid 90 mg tablets (30 tablets per bottle). Contains product name, dosage strength,

PROVIDER Address Phone DEA#

NP Thyroid 90 mg

30 Brown Round Tablets

Directions: Take 1 tablet daily as directed.

CS NDC # 17224-504-30 MFG NDC #42192-331-01

Date: _____ EXP: _____

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

uted by: Calvin Scott & Co In erque, NM 87123 CIV RX Only No Refill

> F: 12/31 I: 12346597549451 IIN: 0030299383330I



, manufacturer/repacker name, and usage instructions as printed on the label.

NP THYROID

thyroid tablets tablet

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LEVOTHYROXINE (UNII: Q51BO43MG4) (LEVOTHYROXINE - UNII:Q51BO43MG4) LIOTHYRONINE (UNII: 06LU7C9H1V) (LIOTHYRONINE - UNII:06LU7C9H1V) LIOTHYRONINE (UNII: 06LU7C9H1V) (LIOTHYRONINE - UNII:06LU7C9H1V)

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
MINERAL OIL (UNII: T5L8T28FGP)		
CALCIUM STEARATE (UNII: 776XM7047L)		

Product Characteristics			
Color	brown (LIGHT BROWN, AS PER THE LABEL)	Score	no score
Shape	ROUND (STANDARD ROUND TABLET)	Size	8mm
Flavor		Imprint Code	AP;331
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17224-504- 30	30 in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/01/2025	

Labeler - Calvin Scott & Co Inc. (073404626)

Establishment

Name	Address	ID/FEI	Business Operations
Calvin Scott & Co Inc.		073404626	repack(17224-504)

Revised: 8/2025 Calvin Scott & Co Inc.