

METHOCARBAMOL- methocarbamol tablet, film coated

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DirectRX

METHOCARBAMOL

Description

-

Clinical Pharmacology

-

Pharmacokinetics

-

Elderly

-

Special Populations

The clearance of methocarbamol in 8 renally-impaired patients on maintenance hemodialysis was reduced about 40% compared to 17 normal subjects, although the mean (\pm SD) elimination half-life in these two groups was similar: 1.2 (\pm 0.6) versus 1.1 (\pm 0.3) hours, respectively.

Indications and Usage

-

Contraindications

-

Warnings

-

Precautions

-

Adverse Reactions

Overdosage

-

Dosage and Administration

-

How supplied

SPL Unclassified

Package Label

D

Dir. By: Virtus Pharmaceuticals, LLC
Tampa, FL 33619
NDC 76439-136-50

Mfg Lot:
7/17/2016

**METHOCARBAMOL
750mg 90 Tabs**

Generic For: **ROBAXIN**
Each tablet contains 750 mg of Methocarbamol Tablets, USP

Lot#
Prod# 627-90

Discard After: 04/17

Packaged and Distributed By: Alpharetta, GA 30005

M

NDC 61919-627-90

May cause DROWSINESS. ALCOHOL may INTENSIFY this effect. Use care when operating dangerous machinery.

METHOCARBAMOL 750mg
NDC 61919-627-90 90 Tabs
Lot Exp Date 04/17
Mfg NDC 76439-136-50

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D

Mfg For: Granules Pharmaceuticals Inc.
Charlottesville, VA 20151
NDC 70010-754-01

Mfg Lot:
7/27/2018

**METHOCARBAMOL
500mg 60 Tabs**

Generic For: **ROBAXIN**
Each Film-Coated Tablet Contains: Methocarbamol USP, 500mg

Lot#
Prod# 610-60

Discard After: 04/20

Packaged and Distributed By: Alpharetta, GA 30005

M

NDC 61919-610-60

May cause DROWSINESS. ALCOHOL may INTENSIFY this effect. Use care when operating dangerous machinery.

METHOCARBAMOL 500mg
NDC 61919-610-60 60 Tabs
Lot Exp Date 04/20
Mfg NDC 70010-754-01

METHOCARBAMOL 500mg
NDC 61919-610-60 60 Tabs
Lot Exp Date 04/20
Mfg NDC 70010-754-01

METHOCARBAMOL 500mg
NDC 61919-610-60 60 Tabs
Lot Exp Date 04/20
Mfg NDC 70010-754-01

METHOCARBAMOL 500mg
NDC 61919-610-60 60 Tabs
Lot Exp Date 04/20
Mfg NDC 70010-754-01

D

Mfg For: Camber Pharm., Inc.
Piscataway, NJ 08854
NDC 31722-534-05

Mfg Lot:
BW 9/8/2017 916575

**METHOCARBAMOL
750mg 90 Tabs**

Generic For: **ROBAXIN**
*Each uncoated tablet contains Methocarbamol USP 750 mg

Lot#
Prod# 903-90

Discard After: 04/17

Packaged and Distributed By: Alpharetta, GA 30005

M

NDC 61919-903-90

May cause DROWSINESS. ALCOHOL may INTENSIFY this effect. Use care when operating dangerous machinery.

METHOCARBAMOL 750mg
NDC 61919-903-90 90 Tabs
Lot Exp Date 04/17
Mfg NDC 31722-534-05

METHOCARBAMOL 750mg
NDC 61919-903-90 90 Tabs
Lot Exp Date 04/17
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METHOCARBAMOL 750mg
NDC 61919-903-90 90 Tabs
Lot Exp Date 04/17
Mfg NDC 31722-534-05

METHOCARBAMOL 750mg
NDC 61919-903-90 90 Tabs
Lot Exp Date 04/17
Mfg NDC 31722-534-05

METHOCARBAMOL

methocarbamol tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-627(NDC:76439-135)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHOCARBAMOL (UNII: 125OD7737X) (METHOCARBAMOL - UNII:125OD7737X)	METHOCARBAMOL	750 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TRIACETIN (UNII: XHX3C3X673)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	yellow	Score	no score
Shape	CAPSULE	Size	19mm
Flavor		Imprint Code	AP211
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-627-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA200958	01/01/2015	

METHOCARBAMOL

methocarbamol tablet, coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:6 19 19-6 10 (NDC:700 10-754)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHOCARBAMOL (UNII: 125OD7737X) (METHOCARBAMOL - UNII:125OD7737X)	METHOCARBAMOL	500 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE (UNII: FZ989GH94E)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSE 2910 (6 MPAS) (UNII: 0WZ8WG20P6)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color	orange	Score	2 pieces
Shape	ROUND	Size	13mm
Flavor		Imprint Code	G;500
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:6 19 19-6 10-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	04/18/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209312	04/18/2019	

METHOCARBAMOL

methocarbamol tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-903(NDC:31722-534)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHOCARBAMOL (UNII: 125OD7737X) (METHOCARBAMOL - UNII:125OD7737X)	METHOCARBAMOL	750 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white (White to Offwhite)	Score	no score
Shape	OVAL (Capsule shaped)	Size	19mm
Flavor		Imprint Code	115;H
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-903-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/23/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090200	04/23/2019	

Labeler - DirectRX (079254320)**Establishment**

Name	Address	ID/FEI	Business Operations
DirectRX		079254320	repack(61919-627, 61919-903) , relabel(61919-610)