

LACOSAMIDE - lacosamide tablet, film coated
MSN LABORATORIES PRIVATE LIMITED

LACOSAMIDE film coated tablet, for oral use, CV

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 69539-025-60




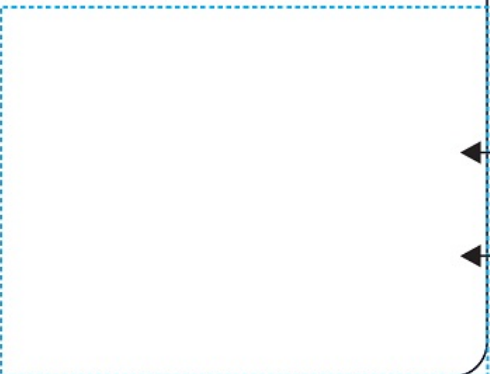

Lacosamide Tablet, USP

50 mg

Rx Only

PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

60 Tablets

Rx Only	NDC 69539-025-60	Each tablet contains 50 mg lacosamide, USP.	Issued: 03/2022 DXXXXXX-XX	
Lacosamide  Tablets, USP		USUAL DOSAGE: See package insert for dosage information.	 N 3 69539102560 2	
50 mg	PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT	Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [See USP Controlled Room Temperature]. M.L. No.: 5/MN/TS/2014/F/G		
	60 Tablets	Manufactured by: MSN Laboratories Private Limited Telangana- 509228, INDIA		
		Distributed by: MSN Pharmaceuticals Inc. Piscataway, NJ 08854-3714		

NDC 69539-026-60

Lacosamide Tablet, USP

100 mg

Rx Only


PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

60 Tablets

Rx Only NDC 69539-026-60


Each tablet contains 100 mg lacosamide, USP.

Issued: 03/2022
DXXXXXX-XX

Lacosamide 
Tablets, USP

100 mg

PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

 60 Tablets



USUAL DOSAGE:
See package insert for dosage information.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [See USP Controlled Room Temperature].

M.L. No.: 5/MN/TS/2014/F/G

Manufactured by:
MSN Laboratories Private Limited
Telangana- 509228,
INDIA

Distributed by:
MSN Pharmaceuticals Inc.
Piscataway, NJ 08854-3714

NDC 69539-027-60

Lacosamide Tablet, USP

150 mg

Rx Only


PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

60 Tablets

Rx Only NDC 69539-027-60


Each tablet contains 150 mg lacosamide, USP.

Issued: 03/2022
DXXXXXX-XX

Lacosamide 
Tablets, USP

150 mg

PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

 60 Tablets



USUAL DOSAGE:
See package insert for dosage information.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [See USP Controlled Room Temperature].

M.L. No.: 5/MN/TS/2014/F/G

Manufactured by:
MSN Laboratories Private Limited
Telangana- 509228,
INDIA

Distributed by:
MSN Pharmaceuticals Inc.
Piscataway, NJ 08854-3714

NDC 69539-028-60

Lacosamide Tablet, USP

200 mg

Rx Only

PHARMACIST: DISPENSE WITH THE ACCOMPANYING MEDICATION GUIDE TO EACH PATIENT

60 Tablets

Rx Only

NDC 69539-028-60

Each tablet contains 200 mg lacosamide, USP.

Issued: 03/2022
DXXXXXX-XX



Lacosamide 
Tablets, USP

200 mg

PHARMACIST: DISPENSE WITH THE
ACCOMPANYING MEDICATION
GUIDE TO EACH PATIENT



60 Tablets

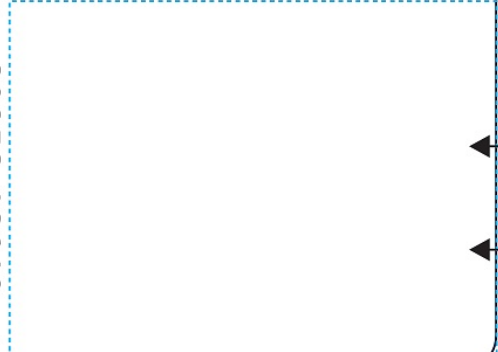
USUAL DOSAGE:
See package insert for dosage information.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C to 30°C (59°F to 86°F) [See USP Controlled Room Temperature].

M.L. No.: 5/MN/TS/2014/F/G

Manufactured by:
MSN Laboratories Private Limited
Telangana- 509228,
INDIA

Distributed by:
MSN Pharmaceuticals Inc.
Piscataway, NJ 08854-3714



LACOSAMIDE

lacosamide tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69539-025
Route of Administration	ORAL	DEA Schedule	CV

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LACOSAMIDE (UNII: 563KS2PQY5) (LACOSAMIDE - UNII:563KS2PQY5)	LACOSAMIDE	50 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPVIDONE (120 .MU.M) (UNII: 68401960MK)	
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
HYDROXYPROPYL CELLULOSE (160000 WAMW) (UNII: RFW2ET671P)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	PINK	Score	no score
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Shape	OVAL	Size	10mm
Flavor		Imprint Code	11
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69539-025-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	
2	NDC:69539-025-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204921	03/19/2022	

LACOSAMIDE

lacosamide tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69539-026
Route of Administration	ORAL	DEA Schedule	CV

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LACOSAMIDE (UNII: 563KS2PQY5) (LACOSAMIDE - UNII:563KS2PQY5)	LACOSAMIDE	100 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPROVIDONE (120 .MU.M) (UNII: 68401960MK)	
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	YELLOW	Score	no score
Shape	OVAL	Size	13mm
Flavor		Imprint Code	12
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69539-026-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	
2	NDC:69539-026-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204921	03/19/2022	

LACOSAMIDE

lacosamide tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69539-027
Route of Administration	ORAL	DEA Schedule	CV

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LACOSAMIDE (UNII: 563KS2PQY5) (LACOSAMIDE - UNII:563KS2PQY5)	LACOSAMIDE	150 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPROVIDONE (120 .MU.M) (UNII: 68401960MK)	
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	BROWN (tan)	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	113
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69539-027-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	
2	NDC:69539-027-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204921	03/19/2022	

LACOSAMIDE

lacosamide tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:69539-028
Route of Administration	ORAL	DEA Schedule	CV

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LACOSAMIDE (UNII: 563KS2PQY5) (LACOSAMIDE - UNII:563KS2PQY5)	LACOSAMIDE	200 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPROVIDONE (120 .MU.M) (UNII: 68401960MK)	
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	

HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	BLUE	Score	no score
Shape	OVAL	Size	17mm
Flavor		Imprint Code	114
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69539-028-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	
2	NDC:69539-028-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204921	03/19/2022	

Labeler - MSN LABORATORIES PRIVATE LIMITED (650786952)

Establishment

Name	Address	ID/FEI	Business Operations
MSN LABORATORIES PRIVATE LIMITED		650786952	ANALYSIS(69539-025, 69539-026, 69539-027, 69539-028) , MANUFACTURE(69539-025, 69539-026, 69539-027, 69539-028)

Revised: 4/2022

MSN LABORATORIES PRIVATE LIMITED