

GABAPENTIN- gabapentin tablet, film coated
Zydus Lifesciences Limited

GABAPENTIN TABLETS

Manufactured by:

Cadila Heathcare Ltd.

India

SPL MEDGUIDE

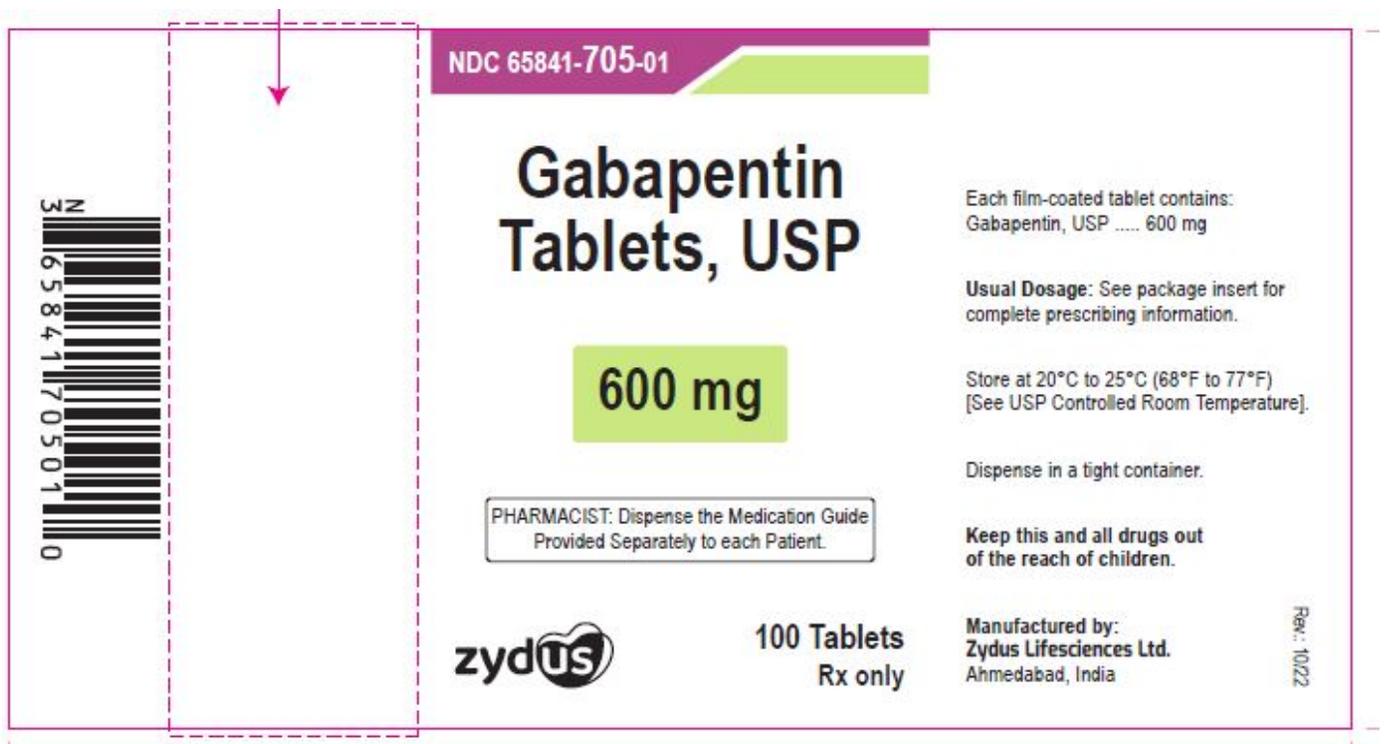
PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 65841-705-01 in bottle of 100 tablets

Gabapentin Tablets USP, 600 mg

R_x only

100 tablets



NDC 65841-706-01 in bottle of 100 tablets

Gabapentin Tablets USP, 800 mg

R_x only

100 tablets

NDC 65841-706-01

Gabapentin Tablets, USP

800 mg

PHARMACIST: Dispense the Medication Guide Provided Separately to each Patient.

zydus 100 Tablets Rx only

Each film-coated tablet contains: Gabapentin, USP 800 mg

Usual Dosage: See package insert for complete prescribing information.

Store at 20°C to 25°C (68°F to 77°F) [See USP Controlled Room Temperature].

Dispense in a tight container.

Keep this and all drugs out of the reach of children.

Manufactured by: Zydus Lifesciences Ltd. Ahmedabad, India

Rev: 10/22

GABAPENTIN

gabapentin tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-705
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GABAPENTIN (UNII: 6CW7F3G59X) (GABAPENTIN - UNII:6CW7F3G59X)	GABAPENTIN	600 mg

Inactive Ingredients

Ingredient Name	Strength
COPOVIDONE K25-31 (UNII: D9C330MD8B)	
HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED (UNII: 2165RE0K14)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
POVIDONE (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	2 pieces
Shape	OVAL (OVAL)	Size	17mm
Flavor		Imprint Code	ZE72
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-705-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	
2	NDC:65841-705-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	
3	NDC:65841-705-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078926	10/16/2012	

GABAPENTIN

gabapentin tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-706
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GABAPENTIN (UNII: 6CW7F3G59X) (GABAPENTIN - UNII:6CW7F3G59X)	GABAPENTIN	800 mg

Inactive Ingredients

Ingredient Name	Strength
COPOVIDONE K25-31 (UNII: D9C330MD8B)	
HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED (UNII: 2165RE0K14)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
POVIDONE (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	2 pieces
Shape	OVAL (OVAL)	Size	20mm
Flavor		Imprint Code	ZE71
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-706-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	
2	NDC:65841-706-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	
3	NDC:65841-706-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2012	
4	NDC:65841-706-77	10 in 1 CARTON	10/16/2012	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078926	10/16/2012	

Labeler - Zydus Lifesciences Limited (918596198)

Registrant - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		918596198	ANALYSIS(65841-705, 65841-706) , MANUFACTURE(65841-705, 65841-706)

Revised: 11/2024

Zydus Lifesciences Limited