

METFORMIN HYDROCHLORIDE- metformin hydrochloride tablet, film coated
Cadila Healthcare Limited

METFORMIN HYDROCHLORIDE TABLETS

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 65841-028-01 in bottle of 100 tablets

Metformin Hydrochloride Tablets USP, 500 mg

Rx only

100 tablets

ZyGenerics
NDC 65841-028-01
**METFORMIN
HYDROCHLORIDE**
Tablets, USP
500 mg
Rx only
100 TABLETS

PHARMACIST: Dispense the Patient Information provided separately to each patient.

Each film-coated tablet contains:
Metformin hydrochloride, USP ...500 mg

Usual Dosage: See package insert for complete prescribing information.

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

Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Lot:
Exp:
Rev: 11/14

NDC 65841-809-01 in bottle of 100 tablets

Metformin Hydrochloride Tablets USP, 500 mg

Rx only

100 tablets

No Varnished Area (Do Not Print)
(18 x 41 mm)

NDC 65841-809-01

Metformin Hydrochloride Tablets, USP

70 500 mg Z

PHARMACIST: Dispense the Patient Information provided separately to each patient.

100 TABLETS
Rx only

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Rev: 07/18

Each film-coated tablet contains:
Metformin hydrochloride, USP 500 mg

Usual Dosage: See package insert for complete prescribing information.

Store at 20° - 25°C (68° - 77°F)
[See USP Controlled Room Temperature].
Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

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NDC 65841-029-01 in bottle of 100 tablets
Metformin Hydrochloride Tablets USP, 850 mg
Rx only
100 tablets

ZyGenerics

NDC 65841-029-01

METFORMIN HYDROCHLORIDE
Tablets, USP

850 mg

PHARMACIST: Dispense the Patient Information provided separately to each patient.

Rx only
100 TABLETS

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Each film-coated tablet contains:
Metformin hydrochloride, USP. 850 mg

Usual Dosage: See package insert for complete prescribing information.

Store at 20° - 25°C (68° - 77°F)
[See USP Controlled Room Temperature].
Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

Lot:
Exp:
Rev: 11/14

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6584102901
7

NDC 65841-810-01 in bottle of 100 tablets
Metformin Hydrochloride Tablets USP, 850 mg

Rx only

100 tablets

No Varnished Area (Do Not Print)
(18 x 50.5 mm)

NDC 65841-810-01

Metformin Hydrochloride Tablets, USP

850 mg

PHARMACIST: Dispense the Patient Information provided separately to each patient.

zydus
pharmaceuticals

100 TABLETS
Rx only

Each film-coated tablet contains:
Metformin hydrochloride, USP 850 mg

Usual Dosage: See package insert for complete prescribing information.

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

Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India


Rev: 07/18

NDC 65841-030-01 in bottle of 100 tablets

Metformin Hydrochloride Tablets USP, 1000 mg

Rx only

100 tablets



Lot:
Exp:
Rev: 11/14

ZyGenerics

NDC 65841-030-01

METFORMIN HYDROCHLORIDE

Tablets, USP

1000 mg

PHARMACIST: Dispense the Patient Information provided separately to each patient.

Rx only

100 TABLETS

Each film-coated tablet contains:
Metformin hydrochloride, USP.. 1000 mg

Usual Dosage: See package insert for complete prescribing information.

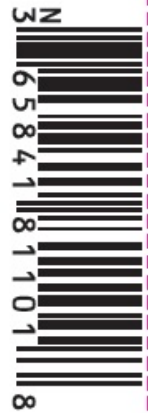
Store at 20° - 25°C (68° - 77°F)
[See USP Controlled Room Temperature].
Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

NDC 65841-811-01 in bottle of 100 tablets
 Metformin Hydrochloride Tablets USP, 1000 mg
 Rx only
 100 tablets



No Varnished Area (Do Not Print)
(18 x 50.5 mm)



NDC 65841-811-01


Metformin Hydrochloride Tablets, USP

1,000 mg

PHARMACIST: Dispense the Patient Information provided separately to each patient.

100 TABLETS
Rx only



Each film-coated tablet contains:
Metformin hydrochloride, USP 1,000 mg

Usual Dosage: See package insert for complete prescribing information.

Store at 20° - 25°C (68° - 77°F)
[See USP Controlled Room Temperature].
Dispense in light-resistant container.

KEEP THIS AND ALL THE DRUGS OUT OF THE REACH OF CHILDREN.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Rev: 07/18

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-028
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	500 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	ROUND (ROUND)	Size	13mm
Flavor		Imprint Code	70;Z
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-028-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
2	NDC:65841-028-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
3	NDC:65841-028-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
4	NDC:65841-028-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
5	NDC:65841-028-77	100 in 1 CARTON	09/28/2005	
5	NDC:65841-028-30	1 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	ANDA077064	09/28/2005	

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-029
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	850 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	OVAL (OVAL)	Size	19mm
Flavor		Imprint Code	69;Z
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-029-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
2	NDC:65841-029-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
3	NDC:65841-029-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
4	NDC:65841-029-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
5	NDC:65841-029-77	100 in 1 CARTON	09/28/2005	
5	NDC:65841-029-30	1 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	ANDA077064	09/28/2005	

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-030
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	1000 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	2 pieces
Shape	OVAL (OVAL)	Size	19mm
Flavor		Imprint Code	Z;71
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-030-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
2	NDC:65841-030-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
3	NDC:65841-030-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
4	NDC:65841-030-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2005	
5	NDC:65841-030-77	100 in 1 CARTON	09/28/2005	
5	NDC:65841-030-30	1 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	ANDA077064	09/28/2005	

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-809
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	500 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	ROUND (ROUND)	Size	13mm
Flavor		Imprint Code	70;Z
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-809-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
2	NDC:65841-809-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
3	NDC:65841-809-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
4	NDC:65841-809-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
5	NDC:65841-809-77	100 in 1 CARTON	12/09/2014	
5	NDC:65841-809-30	1 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	ANDA203686	12/09/2014	

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-810
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	850 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	OVAL (OVAL)	Size	19mm
Flavor		Imprint Code	69;Z
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-810-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
2	NDC:65841-810-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
3	NDC:65841-810-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
4	NDC:65841-810-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
5	NDC:65841-810-77	100 in 1 CARTON	12/09/2014	
5	NDC:65841-810-30	1 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	ANDA203686	12/09/2014	

METFORMIN HYDROCHLORIDE

metformin hydrochloride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-811
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METFORMIN HYDROCHLORIDE (UNII: 786Z46389E) (METFORMIN - UNII:9100L32L2N)	METFORMIN HYDROCHLORIDE	1000 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	2 pieces
Shape	OVAL (OVAL)	Size	19mm
Flavor		Imprint Code	Z;71
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-811-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
2	NDC:65841-811-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
3	NDC:65841-811-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
4	NDC:65841-811-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2014	
5	NDC:65841-811-77	100 in 1 CARTON	12/09/2014	
5	NDC:65841-811-30	1 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	ANDA203686	12/09/2014	

Labeler - Cadila Healthcare Limited (918596198)

Registrant - Cadila Healthcare Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Cadila Healthcare Limited		918596198	ANALYSIS(65841-028, 65841-029, 65841-030, 65841-809, 65841-810, 65841-811) , MANUFACTURE(65841-028, 65841-029, 65841-030, 65841-809, 65841-810, 65841-811)

Establishment

Name	Address	ID/FEI	Business Operations
Cadila Healthcare Limited		863362789	ANALYSIS(65841-028, 65841-029, 65841-030, 65841-809, 65841-810, 65841-811) , MANUFACTURE(65841-028, 65841-029, 65841-030, 65841-809, 65841-810, 65841-811)

Revised: 9/2020

Cadila Healthcare Limited