VENLAFAXINE HYDROCHLORIDE- venlafaxine hydrochloride tablet
Zydus Lifesciences Limited

Venlafaxine Hydrochloride Tablets

SPL MEDGUIDE

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 65841-671-01 in bottle of 100 Tablets

Venlafaxine Hydrochloride Tablets, 25 mg

Rx only

100 TABLETS

NDC 65841-672-01 in bottle of 100 Tablets

Venlafaxine Hydrochloride Tablets, 37.5 mg

Rx only

100 TABLETS

NDC 65841-673-01 in bottle of 100 Tablets

Venlafaxine Hydrochloride Tablets, 50 mg

Rx only

100 TABLETS

NDC 65841-674-01 in bottle of 100 Tablets

Venlafaxine Hydrochloride Tablets, 75 mg

Rx only

100 TABLETS

NDC 65841-675-01 in bottle of 100 Tablets

Venlafaxine Hydrochloride Tablets, 100 mg

Rx only

100 TABLETS

VENLAFAXINE HYDROCHLORIDE

venlafaxine hydrochloride tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:65841-671
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VENLAFAXINE HYDROCHLORIDE (UNII: 7D7RX5A8MO) (VENLAFAXINE - UNII:GRZ5RCB1QG)	VENLAFAXINE	25 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	ORANGE (PEACH)	**Score**	2 pieces
Shape	ROUND (ROUND)	**Size**	6mm
Flavor		**Imprint Code**	ZC;64
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-671-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
2	NDC:65841-671-14	60 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
3	NDC:65841-671-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
4	NDC:65841-671-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
5	NDC:65841-671-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
6	NDC:65841-671-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077653	06/13/2008	

VENLAFAXINE HYDROCHLORIDE

venlafaxine hydrochloride tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:65841-672
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VENLAFAXINE HYDROCHLORIDE (UNII: 7D7RX5A8MO) (VENLAFAXINE - UNII:GRZ5RCB1QG)	VENLAFAXINE	37.5 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	ORANGE (PEACH)	**Score**	2 pieces
Shape	ROUND (ROUND)	**Size**	7mm
Flavor		**Imprint Code**	ZC;65
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-672-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
2	NDC:65841-672-14	60 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
3	NDC:65841-672-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
4	NDC:65841-672-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
5	NDC:65841-672-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
6	NDC:65841-672-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077653	06/13/2008	

VENLAFAXINE HYDROCHLORIDE

venlafaxine hydrochloride tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:65841-673
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
VENLAFAXINE HYDROCHLORIDE (UNII: 7D7RX5A8MO) (VENLAFAXINE - UNII:GRZ5RCB1QG)	VENLAFAXINE	50 mg

Inactive Ingredients	
Ingredient Name	**Strength**
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics			
Color	ORANGE (PEACH)	**Score**	2 pieces
Shape	ROUND (ROUND)	**Size**	8mm
Flavor		**Imprint Code**	ZC;66
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65841-673-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
2	NDC:65841-673-14	60 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
3	NDC:65841-673-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
4	NDC:65841-673-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
5	NDC:65841-673-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
6	NDC:65841-673-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077653	06/13/2008	

VENLAFAXINE HYDROCHLORIDE

venlafaxine hydrochloride tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:65841-674
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VENLAFAXINE HYDROCHLORIDE (UNII: 7D7RX5A8MO) (VENLAFAXINE - UNII:GRZ5RCB1QG)	VENLAFAXINE	75 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	ORANGE (PEACH)	**Score**	2 pieces
Shape	ROUND (ROUND)	**Size**	10mm
Flavor		**Imprint Code**	ZC;67
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-674-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
2	NDC:65841-674-14	60 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
3	NDC:65841-674-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

4	NDC:65841-674-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
5	NDC:65841-674-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
6	NDC:65841-674-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077653	06/13/2008	

VENLAFAXINE HYDROCHLORIDE

venlafaxine hydrochloride tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:65841-675
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VENLAFAXINE HYDROCHLORIDE (UNII: 7D7RX5A8MO) (VENLAFAXINE - UNII:GRZ5RCB1QG)	VENLAFAXINE	100 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	ORANGE (PEACH)	**Score**	2 pieces
Shape	ROUND (ROUND)	**Size**	10mm
Flavor		**Imprint Code**	ZC;68
Contains			

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:65841-675-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
2	NDC:65841-675-14	60 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
3	NDC:65841-675-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
4	NDC:65841-675-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
5	NDC:65841-675-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	
6	NDC:65841-675-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077653	06/13/2008	

Labeler - Zydus Lifesciences Limited (918596198)

Revised: 9/2023 Zydus Lifesciences Limited