Principal Display Panel

NDC 0173-0772-02

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Maintenance Pack

R_x only

25 mg

30 Tablets

This Maintenance Pack is intended for patients already taking LAMICTAL.

NOT for initial titration.

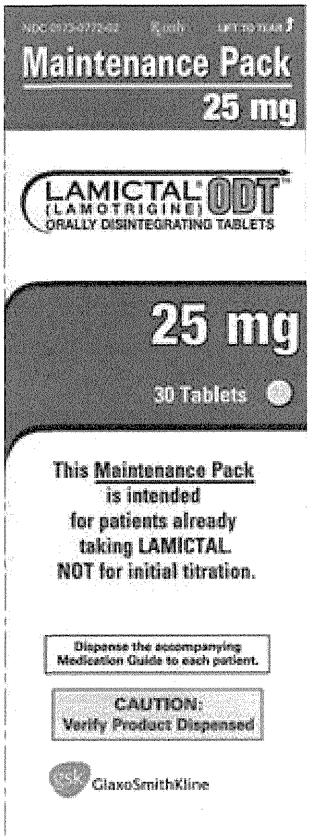
Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Made in India

10000000091910 Rev. 3/11



Principal Display Panel

NDC 0173-0774-02

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Maintenance Pack

R_x only

50 mg

30 Tablets

This Maintenance Pack is intended for patients already taking LAMICTAL.

NOT for initial titration.

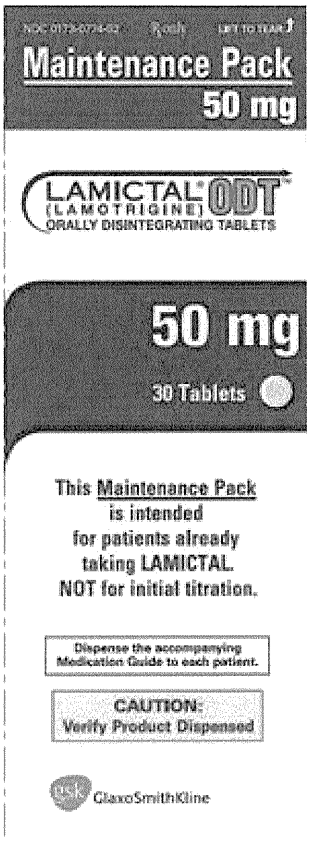
Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Made in India

10000000092032 Rev. 2/11



Principal Display Panel

NDC 0173-0776-02

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Maintenance Pack

R_x only

100 mg

30 Tablets

This Maintenance Pack is intended for patients already taking LAMICTAL.

NOT for initial titration.

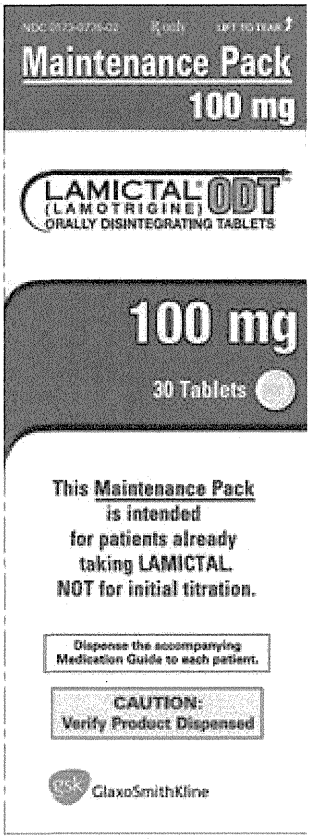
Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Made in India

10000000092031 Rev. 3/11



Principal Display Panel

NDC 0173-0777-02

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Maintenance Pack

R_x only

200 mg

30 Tablets

This Maintenance Pack is intended for patients already taking LAMICTAL.

NOT for initial titration.

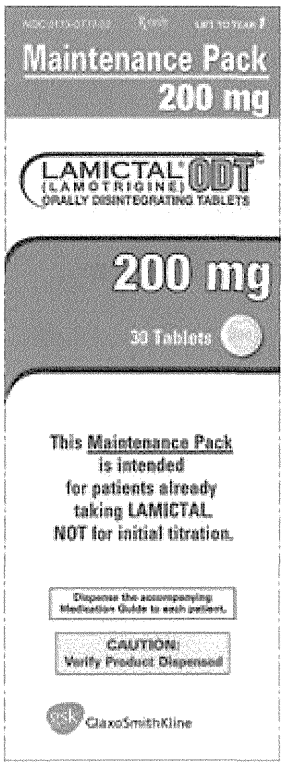
Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Made in India

10000000091909 Rev. 2/11



Principal Display Panel

NDC 0173-0779-00

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Patient Titration Kit

R_X ONLY

For patients TAKING valproate*

*See prescribing information for other drugs that may affect dosing of LAMICTAL ODT.

KIT CONTAINS:

21	25 mg Tablets
7	50 mg Tablets

WEEKS	**TABLETS PER DAY**
1&2	Take 1 (25 mg) tablet every OTHER day
3&4	Take 1 (25 mg) tablet ONCE a day
5	Take 1 (50 mg) tablet ONCE a day

Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Visit www.LAMICTALODT.com for more information

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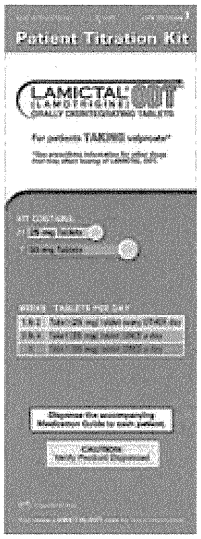

Principal Display Panel

NDC 0173-0780-00

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Patient Titration Kit

R_X ONLY

For patients TAKING carbamazepine, phenytoin, Phenobarbital, or primidone, and NOT TAKING valproate*

*See prescribing information for other drugs that may affect dosing of LAMICTAL ODT.

KIT CONTAINS:

42	50 mg Tablets
14	100 mg Tablets

WEEKS	**TABLETS PER DAY**
1&2	Take 1 (50 mg) tablet ONCE a day
3&4	Take 1 (50 mg) tablet TWICE a day
5	Take 1 (100 mg) tablet TWICE a day

Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Visit www.LAMICTALODT.com for more information\

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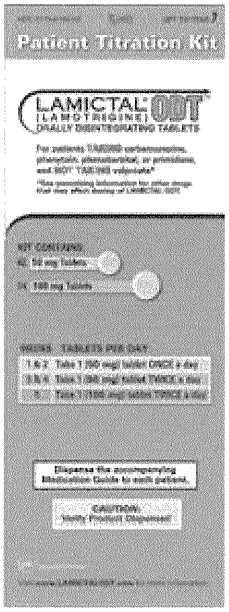

Principal Display Panel

NDC 0173-0778-00

LAMICTAL® ODT™

(LAMOTRIGINE)

ORALLY DISINTEGRATING TABLETS

Patient Titration Kit

R_X ONLY

For patients NOT TAKING carbamazepine, phenytoin, phenobarbital, primidone, or valproate*

*See prescribing information for other drugs that may affect dosing of LAMICTAL ODT.

KIT CONTAINS:

14	25 mg Tablets
14	50 mg Tablets
7	100 mg Tablets

WEEKS	**TABLETS PER DAY**
1&2	Take 1 (25 mg) tablet ONCE a day
3&4	Take 1 (50 mg) tablet ONCE a day
5	Take 1 (100 mg) tablet ONCE a day

Dispense the accompanying Medication Guide to each patient.

CAUTION: Verify Product Dispensed

GlaxoSmithKline

Visit www.LAMICTALODT.com for more information

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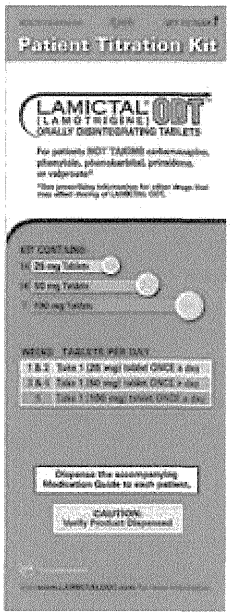

LAMICTAL

lamotrigine tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG LABEL	**Item Code (Source)**	NDC:0173-0633
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LAMOTRIGINE (LAMOTRIGINE)	LAMOTRIGINE	25 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE	
MAGNESIUM STEARATE	
CELLULOSE, MICROCRYSTALLINE	
POVIDONES	
SODIUM STARCH GLYCOLATE TYPE A POTATO	

Product Characteristics

Color	WHITE	**Score**	2 pieces
Shape	HEXAGON (6 sided)	**Size**	6mm
Flavor		**Imprint Code**	LAMICTAL;25
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0173-0633-02	100 in 1 BOTTLE		
2	NDC:0173-0633-10	35 in 1 DOSE PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020241	08/15/1996	

LAMICTAL

lamotrigine tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG LABEL	**Item Code (Source)**	NDC:0173-0642
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LAMOTRIGINE (LAMOTRIGINE)	LAMOTRIGINE	100 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6	
LACTOSE	
MAGNESIUM STEARATE	
CELLULOSE, MICROCRYSTALLINE	
POVIDONES	
SODIUM STARCH GLYCOLATE TYPE A POTATO	

Product Characteristics

Color	ORANGE (peach)	**Score**	2 pieces
Shape	HEXAGON (6 sided)	**Size**	11mm
Flavor		**Imprint Code**	LAMICTAL;100
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0173-0642-55	100 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020241	01/17/1995	

LAMICTAL

lamotrigine tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG LABEL	**Item Code (Source)**	NDC:0173-0643
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LAMOTRIGINE (LAMOTRIGINE)	LAMOTRIGINE	150 mg

Inactive Ingredients

Ingredient Name	Strength
FERRIC OXIDE YELLOW	
LACTOSE	
MAGNESIUM STEARATE	
CELLULOSE, MICROCRYSTALLINE	
POVIDONES	
SODIUM STARCH GLYCOLATE TYPE A POTATO	

Product Characteristics

Color	WHITE (cream)	**Score**	2 pieces
Shape	HEXAGON (6 sided)	**Size**	12mm
Flavor		**Imprint Code**	LAMICTAL;150
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0173-0643-60	60 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020241	01/17/1995	