SLEEPEZE EZE MELTS- diphenhydramine hydrochloride tablet
Medtech Products Inc.

SleepEze eze melts_Export only 63029-925-16

Drug Facts

Active ingredient (in each tablet)

Diphenhydramine Hydrochloride 25 mg

Purpose

Nighttime sleep aid

Uses

● relieves occasional sleeplessness ● helps to reduce difficulty falling asleep

Warnings

Do not use

●if you are elderly, as this drug may cause excitation rather than sedation ●with any other product containing diphenhydramine, even one used on skin ●in children under 12 years of age

Ask a doctor or pharmacist before use if you

●have ■a breathing problem such as emphysema or chronic bronchitis ■glaucoma ■difficulty urinating ●take sedatives or tranquilizers ●are pregnant or breastfeeding

When using this product

avoid drinking alcohol.

Stop use and ask a doctor if

sleeplessness continues for more than 2 weeks. Sleeplessness may be a symptom of a serious underlying medical illness.

Keep out of reach of children.

In case of overdose, call a poison control centre or get medical help right away

Directions

●Adults and children 12 years and over: ■take 1 or 2 tablets at bedtime if needed, or as directed by a doctor ■if you feel drowsy in the morning, reduce dose to 1 tablet

■do not take more than directed

Other information

■ Store at 20° to 25°C

Inactive ingredients

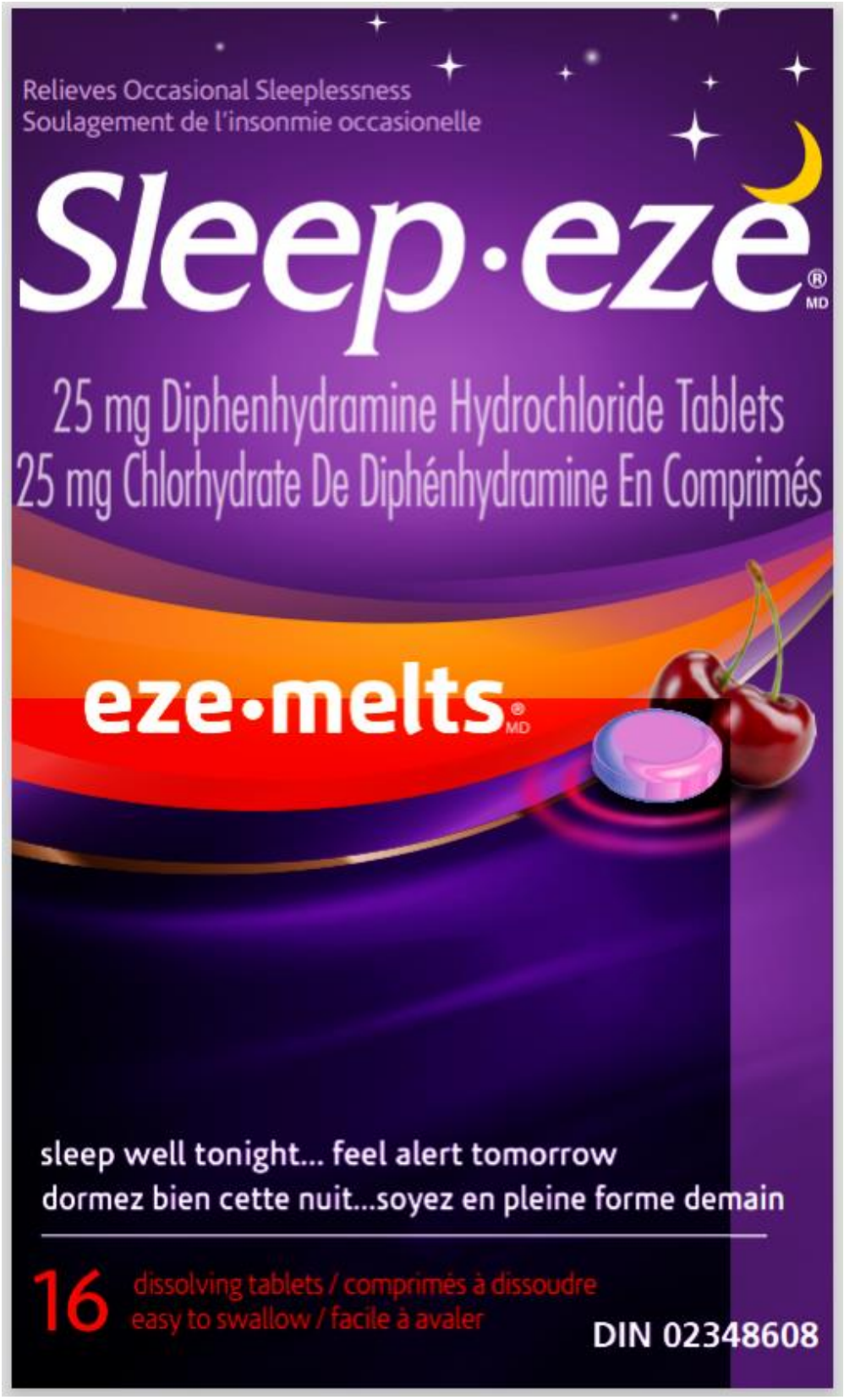
citric acid, d&c red 7 lake, ethylcellulose, flavour, hydroxypropyl cellulose, magnesium stearate, mannitol, microcrystalline cellulose, povidone, stearic acid, sucralose, sucrose

Questions?

1-800-443-4908

PRINCIPAL DISPLAY PANEL

Sleep.eze
25 mg Diphenhydramine Hydrochloride Tablets
16 dissolving tablets



SLEEPEZE EZE MELTS

diphenhydramine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63029-925

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ETHYLCELLULOSES (UNII: 7Z8S9VYZ4B)	
D&C RED NO. 7 (UNII: ECW0LZ41X8)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MINERAL OIL (UNII: T5L8T28FGP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE (UNII: FZ989GH94E)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	pink	**Score**	no score
Shape	ROUND	**Size**	13mm
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-925-16	1 in 1 CARTON	08/03/2010	
1		8 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
Export only		08/03/2010	

Labeler - Medtech Products Inc. (122715688)