WAL-DRAM 2 QUICK-DISSOLVING- meclizine hydrochloride tablet, orally disintegrating
Walgreen Co.

Wal-Dram 2 Quick Dissolving

Active ingredient (in each tablet)
Meclizine hydrochloride 25mg

Purpose
Meclizine hydrochloride Antiemetic

Uses
• for the prevention and treatment of the nausea, vomiting, or dizziness associated with motion sickness

Warnings

Do not use in children under 12 years of age unless directed by a physician

Ask a doctor before use if you have
■ glaucoma ■ a breathing problem such as emphysema or chronic bronchitis
■ difficulty in urination due to enlargement of the prostate gland

Ask a physician or pharmacist before use if you are
■ taking sedatives or tranquilizers

When using this product
■ you may get drowsy ■ avoid alcoholic beverages
■ be careful when driving a motor vehicle or operating machinery
■ alcohol, sedatives, and tranquilizers may increase the drowsiness effect

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions
to prevent motion sickness take it at least one hour before traveling
adults and children 12 years of age and over: take 1 to 2 (25 to 50 mg) tablets once daily, or as directed by a physician.

Other information
■ store at 20-30°C (68-86°F)

Inactive ingredients
acesulfame potassium, erythritol, hydroxypropyl cellulose, colloidal silicon dioxide, mannitol, menthol, sodium stearyl fumarate, yellow ferric oxide.

Carton Image -01

WAL-DRAM 2 QUICK-DISSOLVING

meclizine hydrochloride tablet, orally disintegrating

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-1407
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
ERYTHRITOL (UNII: RA96B954X6)	
MANNITOL (UNII: 3OWL53L36A)	
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	ROUND	**Size**	13mm
Flavor		**Imprint Code**	SJ
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-1407-01	2 in 1 CARTON	03/01/2016	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0363-1407-02	3 in 1 CARTON	10/23/2018	
2		6 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
OTC Monograph Drug	M009	03/01/2016	

Labeler - Walgreen Co. (008965063)

Registrant - Sato Pharmaceutical Co., Ltd. (690575642)