CVS MINERAL OIL- mineral oil liquid CVS

CVS Mineral Oi

Drug Facts

Active ingredient

Mineral Oil Heavy

Purpose

Laxative

Use

For relief of occasional constipation or irregularit. Generally produces bowel movement in 6 to 8 hrs.

Warnings

Do not take this product if you have abdominal pain, nausea, vomiting, change in bowel habits persisting more than 2 weeks, rectal bleeding or kidney failure are present, or use for a period of longer than 1 week unless directed by a doctor. Frequent or prolonged use may result in the dependence on laxatives.

Do not administer to infants or young children, bedridden or aged patients unless directed by a doctor.

Ask a doctor before use if

you are currently taking a stool softener laxative.

If pregnant or brest feeding ask a health professional before use.

When using this Product discotinue use

and consult a doctor if there is a failure to have a bowel movement after use.

Keep out of reach of children

In case of accidental overdose, seek professional assistance or contact Poison Control Center immediately. In case of eye contact wash gently with water for 15 minutes.

Directions

Adults and children 12 yrs and older: 1 to 2 tablespoonfulls at bedtime.

Children 6 to 12 yrs. of age: 1 to 3 teaspoonfuls at bedtime.

Children under 6 yrs. of age: Consult a doctor before use.

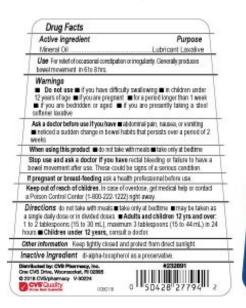
Inactive ingredient

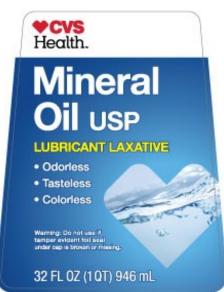
dl-alpha-tocopherol as a preservative.

Other information

Keep tightly closed and protect from direct light.

Principal display panel





CVS MINERAL OIL

mineral oil liquid

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-238

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	1000 mg in 1 mL

Inactive Ingredients

	Ingredient Name	Strength
l	.ALPHATOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:69842- 238-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/31/2020		
NDC:69842- 238-32	950 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/31/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	01/31/2020	

Labeler - CVS (062312574)

Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(69842-238), analysis(69842-238), pack(69842-238), label(69842-238)

Revised: 12/2023 CVS