ENEMEEZ PLUS- docusate sodium and benzocaine liquid Enemeez LLC DBA Summit Pharmaceuticals

Enemeez® Plus

Drug Facts

Active ingredient (per mini enema)

Docusate Sodium USP 283 mg Benzocaine 20mg

Docusate Sodium USP 283 mg Stool Softener Laxative Benzocaine 20mg Anesthetic

Uses

For relief of occasional constipation (irregularity). This product generally produces a stimulus and bowel movement in 2 to 15 minutes.

Warnings

For rectal use only. This is not a suppository. Do not take orally. Individuals with sensitivity to Benzocaine should not use this product.

Ask a doctor before use if you have:

- abdominal pain, nausea or vomiting.
- a sudden change in bowel habits that persists over a period of 2 weeks.

Ask a doctor or pharmacist before use if you are presently taking mineral oil.

Stop use and ask a doctor if you:

- have rectal bleeding.
- failed to have a bowel movement after use. This may indicate a serious condition.
- encounter resistance. Forcing the tube may result in injury or damage to the rectum. If product is not
 administered properly and occasional rectal examinations are not performed, serious complications can
 arise. Occasional rectal exams are imperative for patients with impaired rectal function, especially loss of
 sensation.
- require a bowel program for longer than 1 week.
- experience rectal irritation or a rash around the anus appears.

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

Keep out of reach of children.

In case of accidental ingestion or overdose, get medical help or contact a Poison Control Center immediately.

Directions

Adults and children 12 years of age and older (with adult supervision) one to three units daily. Children 2 years and over but not more than 12 years should consult a doctor prior to use and not exceed one unit daily (with adult supervision).

Twist off and remove tip; Lubricate tip prior to insertion: Place a few drops of water or product on the shaft prior to insertion. Also apply one of these lubricants to the anus before inserting the enema. With steady

pressure, gently insert the tube into the rectum with care to prevent damage to the rectal wall. Squeeze to empty the contents. Continue squeezing the tube until it is removed from the rectum. Remove the disposable tube and discard. A small amount of liquid may remain in the tube after use.

Positioning:

For best results, lay on the left side with knees bent.

Alternate Positions:

- Administer while seated on the toilet.
- Kneel with the left arm folded comfortably, then lower head and chest forward until side of face is resting on the surface.

Other Information

Other Information Store at room temperature 15°-30° C (59°-86° F)

Inactive Ingredients

Inactive Ingredients Polyethylene Glycol and Glycerin USP

Questions or Comments: 1-888-273-9734 or www.enemeez.com

Manufactured by:

Enemeez, Inc., Phoenix, AZ 85050

Distributed by: Alliance Labs, L.L.C 2515 E. Rose Garden Lane, Suite #1 Phoenix, AZ 85050

ENEMEEZ PLUS

docusate sodium and benzocaine liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:17433-9877

Route of Administration RECTAL

Active Ingredient/Active Moiety

l	Ingredient Name	Basis of Strength	Strength
	DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	283 mg in 5 mL
l	BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 mg in 5 mL

Inactive Ingredients

	ggg.			
	Ingredient Name	Strength		
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)				
	GLYCERIN (UNII: PDC6A3C0OX)			

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:17433-9877-3	30 in 1 CONTAINER	06/15/2012				
1	NDC:17433-9877-0	5 mL in 1 TUBE; Type 0: Not a Combination Product					
2	NDC:17433-9877-6	35 in 1 CONTAINER	10/08/2021				
2	NDC:17433-9877-0	5 mL in 1 TUBE; Type 0: Not a Combination Product					
3	NDC:17433-9877-5	5 in 1 CONTAINER	01/14/2022				
3	NDC:17433-9877-0	5 mL in 1 TUBE; Type 0: Not a Combination Product					
4	NDC:17433-9877-7	6 in 1 CONTAINER	09/14/2023	02/23/2024			
4	NDC:17433-9877-0	5 mL in 1 TUBE; Type 0: Not a Combination Product					
5	NDC:17433-9877-2	2 in 1 CONTAINER	01/26/2024				
5	NDC:17433-9877-0	5 mL in 1 TUBE; Type 0: Not a Combination Product					

Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date OTC Monograph Drug M007 06/15/2012

Labeler - Enemeez LLC DBA Summit Pharmaceuticals (010717819)

Registrant - Enemeez LLC DBA Summit Pharmaceuticals (010717819)

Establishment							
Name	Address	ID/FEI	Business Operations				
Enemeez LLC DBA Summit Pharmaceuticals		010717819	manufacture(17433-9877)				

Revised: 2/2024 Enemeez LLC DBA Summit Pharmaceuticals