# FIBER CAPLETS- calcium polycarbophil tablet, film coated SAFEWAY

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## Fiber Caplets Calcium Polycarbophil

## **Drug Fats**

## **Active Ingredient (in each caplet)**

Calcium polycarbophil 625 mg equivalent to 500 mg polycarbophil

### **Purpose**

**Bulk-forming laxative** 

#### Uses

- relieves occasional constipation to help restore and maintain regularity
- this product generally produces bowel movement in 12 to 72 hours

**Choking Warning:** Taking this product without adequate fluid may cause it to swell and block your throat or esophagus and may cause choking. Do not take this product if you have difficulty in swallowing. If you experience chest pain, vomiting, or difficulty in swallowing or breathing after taking this product, seek immediate medical attention.

# 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 your doctor or pharmacist before use if you are

taking any other drug. Take this product 2 or more hours before or after other drugs. All laxatives may affect how other drugs work.

# When using this product

- do not use for more than 7 days unless directed by a doctor
- do not take more than 8 caplets in a 24 hour period unless directed by a doctor

# Stop use and ask a doctor if

rectal bleeding occurs or if you fail to have a bowel movement after use of this or any other laxative. These could be signs of a serious condition.

# Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222.

#### **Directions**

- take each dose of this product with at least 8 ounces (a full glass) of water or other fluid. Taking this product without enough liquid may cause choking. (See choking warning).
- this product works naturally so continued use for one to three days is normally required to provide full benefit. Dosage may vary according to diet, exercise, previous laxative use or severity of constipation.

Age	Recommended dose	Daily maximum
adults & children 12 years of	2 caplets once a day	up to 4 times a day
age and over		
children under 12 years	consult a physician	

#### Other information

- each caplet contains: calcium 125 mg
- store at room temperature
- protect contents from moisture

#### DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING

calcium carbonate, caramel color, crospovidone, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, silicon dioxide, sodium lauryl sulfate.

#### **Questions or Comments?**

1-866-467-2748

#### PRINCIPAL DISPLAY PANEL

NDC# 21130-021-40

\*Compare to the active ingredients of Fibercon® Caplets\*

### Fiber Caplets

Calcium Polycarbophil

### **Bulk-Forming**

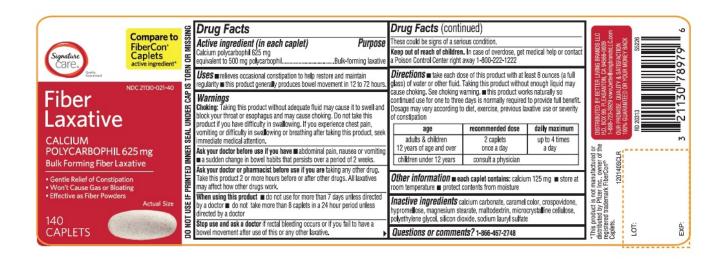
#### Fiber Laxative

- Gentle Relief of Constipation
- Won't Cause Gas or Bloating
- Effective as Fiber Powders

### 140 Caplets

# Distributed by:

\*This product is not manufactured or distributed by Pfizer Inc., owner of the registered trademark Fibercon® Caplets.



### **FIBER CAPLETS**

calcium polycarbophil tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-021
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
CALCIUM POLYCARBOPHIL (UNII: 8F049NKY49) (POLYCARBOPHIL - UNII: W25LM17A4W)	CALCIUM POLYCARBOPHIL	625 mg	

Inactive Ingredients	
Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARAMEL (UNII: T9D99G2B1R)	
CROSPOVIDONE (120 .MU.M) (UNII: 68401960MK)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics			
Color	BROWN (light)	Score	2 pieces
Shape	OVAL	Size	19mm
Flavor		Imprint Code	RP120
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:21130-021- 40	140 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/02/2020	

# **Labeler -** SAFEWAY (009137209)

Revised: 11/2024 SAFEWAY