HEMORRHOIDAL STARCH - starch suppository
H and P Industries, Inc. dba Triad Group

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

ACTIVE INGREDIENT

Topical starch, 51%

PURPOSE

Protectant

USE

- Provides temporary relief of the itching, burning and discomfort associated with hemorrhoids and other anorectal disorders
- Provides a coating to protect irritated tissue

WARNINGS

For rectal use only.

When using this product
do not exceed the recommended daily dosage unless directed by a doctor.

Stop use and ask a doctor

- in case of bleeding
- if condition worsens or does not improve within 7 days.

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

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS
Adults:

- When practical, cleanse the affected area with mild soap and warm water. Rinse thoroughly.
- Gently dry by patting or blotting with toilet tissue or soft cloth before application of this product.
- Detach one suppository from strip of suppositories.
- Remove film before inserting into the rectum by holding the suppository upright. Carefully separate

film by inserting tip of fingernail at film split.
- Carefully peel film slowly and evenly down both sides to expose suppository
- Avoid excessive handling of suppository, which is designed to melt at body temperature. If suppository seems soft, hold in film wrapper under cold water for 2 or 3 minutes.
- Insert one suppository rectally up to 6 times daily or after each bowel movement

Children under 12 years: ask a doctor.

OTHER INFORMATION
Store below 86° F (30° C).

INACTIVE INGREDIENTS
benzyl alcohol, hydrogenated vegetable oil, tocopheryl acetate

PACKAGE INFORMATION - REPRESENTATIVE LABEL
Compare to the active ingredient in TUCKS®*

HEMORRHOIDAL
SUPPOSITORIES
TOPICAL STARCH

- Relieves itching, burning and discomfort
- Protects and soothes irritated tissue

12 SUPPOSITORIES

*This product is not manufactured or distributed by Pfizer consumer Healthcare, Inc., owner of the registered trademark TUCKS®.

HEMORRHOIDAL STARCH

starch suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50730-1512
Route of Administration	RECTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
STARCH, CORN (UNII: O8232NY3SJ) (STARCH, CORN - UNII:O8232NY3SJ)	STARCH, CORN	0.51 g

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
FAT, HARD (UNII: 8334LX7S21)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50730-1512-1	12 in 1 BOX		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	08/01/2006	

Labeler - H and P Industries, Inc. dba Triad Group (050259597)

Revised: 12/2009 H and P Industries, Inc. dba Triad Group