LAXATIVE MAXIMUM STRENGTH- sennosides tablet, sugar coated
L.N.K. International, Inc.

Quality Plus 44-348

Active ingredient (in each tablet)

Sennosides USP, 25 mg

Purpose

Stimulant laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 6 to 12 hours

Warnings

Do not use

laxative products when abdominal pain, nausea, or vomiting are present unless directed by a doctor.

Ask a doctor before use if you have

noticed a sudden change in bowel habits that persists over a period of 2 weeks.

Ask a doctor or pharmacist before use if you are

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

When using this product,

do not use for a period longer than 1 week.

Stop use and ask a doctor if

rectal bleeding or failure to have a bowel movement occur after use of a laxative. These may be signs of a serious condition.

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

- swallow tablet(s) with a glass of water
- swallow tablet(s) whole; do not crush, break, or chew

adults and children 12 years and over	2 tablets once or twice daily
children 6 to under 12 years	1 tablet once or twice daily
children under 6 years	ask a doctor

Other information

- **each tablet contains:** calcium 40 mg, magnesium 5 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- see end flap for expiration date and lot number

Inactive ingredients

acacia, calcium carbonate, carnauba wax, corn starch, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, iron oxide black, magnesium stearate, methylparaben, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, propylene glycol, propylparaben, shellac glaze, silicon dioxide, sodium benzoate, sodium lauryl sulfate, sucrose, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

QUALITY
+PLUS

NDC 50844-348-08

*Compare to active ingredient
in ex•lax® Maximum Strength

MAXIMUM STRENGTH
LAXATIVE
Sennosides USP, 25 mg
Stimulant Laxative

Gentle, dependable

constipation relief

24
Coated Tablets

ACTUAL SIZE

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

Distributed by
LNK INTERNATIONAL, INC.
60 Arkay Drive
Hauppauge, NY 11788
USA

*This product is not manufactured or distributed by GSK Consumer Healthcare S.A., owner of the registered trademark ex•lax® Maximum Strength.
50844 ORG041934808

Quality Plus 44-348

LAXATIVE MAXIMUM STRENGTH

sennosides tablet, sugar coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50844-348
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	25 mg

Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SHELLAC (UNII: 46N107B71O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	blue	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	44;348
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:50844-348-02	1 in 1 CARTON	01/02/2003	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50844-348-08	2 in 1 CARTON	01/02/2003	
2		12 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	505G(a)(3)	01/02/2003	

Labeler - L.N.K. International, Inc. (038154464)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(50844-348)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(50844-348)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-348)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	pack(50844-348)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(50844-348)

Revised: 8/2023 L.N.K. International, Inc.