SENNOSIDES- sennosides tablet
InvaTech Pharma Solutions LLC

Sennosides Tablets USP 8.6 mg

FOR FURTHER MANUFACTURING, PROCESSING OR REPACKING

KEEP OUT OF THE REACH OF CHILDREN

Laxative

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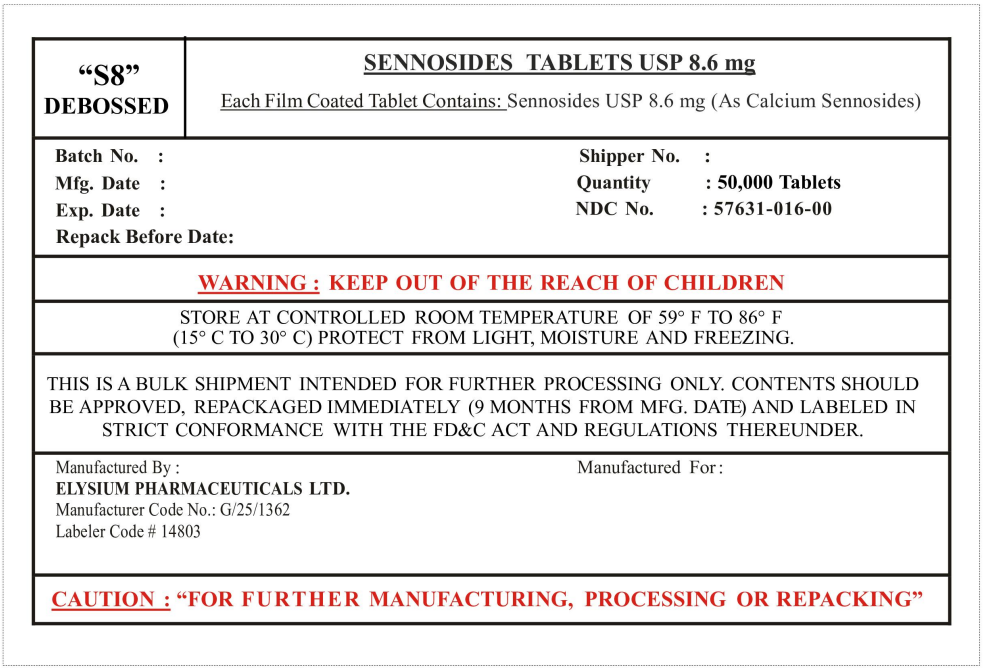
Microcrystalline Cellulose, Dicalcium Phosphate, Croscarmellose Sodium, Stearic acid Powder, Colloidal Silicon Dioxide, Magnesium Stearate, Hypromellose, Polyethylene glycol, , Titanium dioxide, FD&C red



"S8" DEBOSSED	SENNOSIDES TABLETS USP 8.6 mg Each Film Coated Tablet Contains: Sennosides USP 8.6 mg (As Calcium Sennosides)

Batch No. :
Mfg. Date :
Exp. Date :
Repack Before Date:

Shipper No. :
Quantity : 50,000 Tablets
NDC No. : 57631-016-00

WARNING : KEEP OUT OF THE REACH OF CHILDREN

STORE AT CONTROLLED ROOM TEMPERATURE OF 59° F TO 86° F
(15° C TO 30° C) PROTECT FROM LIGHT, MOISTURE AND FREEZING.

THIS IS A BULK SHIPMENT INTENDED FOR FURTHER PROCESSING ONLY. CONTENTS SHOULD BE APPROVED, REPACKAGED IMMEDIATELY (9 MONTHS FROM MFG. DATE) AND LABELED IN STRICT CONFORMANCE WITH THE FD&C ACT AND REGULATIONS THEREUNDER.

Manufactured By :
ELYSIUM PHARMACEUTICALS LTD.
Manufacturer Code No.: G/25/1362
Labeler Code # 14803

Manufactured For :

CAUTION : "FOR FURTHER MANUFACTURING, PROCESSING OR REPACKING"

SENNOSIDES

sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:57631-016
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNA ALATA LEAF (UNII: 4BXR6YZN92) (SENNA ALATA LEAF - UNII:4BXR6YZN92)	SENNA ALATA LEAF	8.6 mg

Inactive Ingredients

Ingredient Name	Strength
PARAFFINUM LIQUIDUM (UNII: T5L8T28FGP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
DICALCIUM PHOSPHATE (UNII: L11K75P92J)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	brown	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	S8
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57631-016-00	50000 in 1 BOX; Type 0: Not a Combination Product	01/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	01/13/2021	

Labeler - InvaTech Pharma Solutions LLC (078602180)

Establishment

Name	Address	ID/FEI	Business Operations
Elysium Pharmaceuticals Ltd		915664486	manufacture(57631-016) , analysis(57631-016)

Revised: 10/2025 InvaTech Pharma Solutions LLC