Product Licence

Product Number:	80079080	

Brand Name:

Senokot[®] Ginger Relief™

Issued to:

Name of licensee:

Purdue Pharma 575 Granite Court Pickering, Ontario, L1W 3W8 Canada

Authorized for the following:

Dosage form: Tablet

Recommended route of administration: Oral

Recommended Conditions of Use:

Recommended Use and Purpose:

Traditionally used in Herbal Medicine to help relieve digestive upset/disturbances including lack of appetite, nausea, digestive spasms, indigestion, dyspepsia and flatulent colic.

Helps prevent nausea and vomiting associated with motion sickness, and/or seasickness.

Recommended dose:

		Adults, adoleso (indigestions c	scents and children older than 6 years old claims)			
Dosage			Frequency			
Min: 1	Max: 2	Units: Tablets	Min: 1	Max: 2	times a day	
Additional Dosage Information: n/a						
Directions of Use: Maximum 5 tablets/day.						

Sub-population group: Adults, adolese (motion sickness)		cents and children older than 6 years old ess)					
Dosage		Frequency					
Min: 1	Max: 3	Units: Tablets	Min:	1	Max:	3	times a day
Additional Dosage Information: n/a							
Directions of Use: Take a single dose (1-3 tablets) 30 minutes before travel and every 4 hours as needed. Maximum 3 tablets/day.							

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Duration Statement: n/a

Risk Information

Cautions and Warnings:

Consult a health care practitioner if symptoms persist or worsen.

Contraindications: n/a

Known Adverse Reactions: n/a

Symptoms and Treatment of Overdosage: n/a

Medicinal Ingredients:

Proper Name	Common Name	Quantity per Dosage Unit	Extract	Potency	Source Material
Zingiber officinale	Ginger	150 mg	3.500 : 1	525 mg	Dried Ginger Root

Non-Medicinal Ingredients:

Croscarmellose sodium Lactose monohydrate Magnesium stearate Povidone K-30 Silicon dioxide Talc

This Licence is issued by the Minister of Health under the authority of section 7 of the Natural Health Products Regulations. Sale of the described natural health product, including any changes thereto pursuant to section 11 of the Regulations, is subject to the Food and Drugs Act and to the Natural Health Products Regulations.

Issued: 2017-07-14	Revised/Amended: n/a

Director General NHPD