PLEO SANUVIS- lactic acid, l-
Sanum Kehlbeck GmbH & Co. KG

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Pleo™ Sanuvis
PORTABLE SIPS
4X, 6X, 12X, 30X, 200X

Oral Homeopathic Medicine

50 doses, each 2 mL
(.06 fl oz)

INDICATIONS

Digestive Aid.

DIRECTIONS FOR USE

Snap off top portion of sipping container. Insert glass sipping straw.

DOSAGE

1 SIP, 1–3 times weekly.

INGREDIENTS

2 mL contains L(+)-Lactic acid 4X, 6X, 12X, 30X, 200X in a base of purified saline solution.

WARNING

If symptoms persist more than a few days, contact a licensed practitioner. As with any drug, if you are pregnant or nursing a baby, seek the advice of a health care professional before using this product.

Keep this and all medications out of the reach of children. In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Protect from light and heat.

Tamper Evident

Do not use this product if the glass vial is broken or if imprinted security strip on carton is torn.

Made in Germany

Distributed by:
SANUM USA Corp.
1465 Slater Road
Ferndale, WA 98248

Manufactured By:
Sanum-Kehlbeck GmbH & Co. KG

Rev. 09/2000

PRINCIPAL DISPLAY PANEL - 2 mL Carton

6302-2

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convenient, disposable
single dose containers

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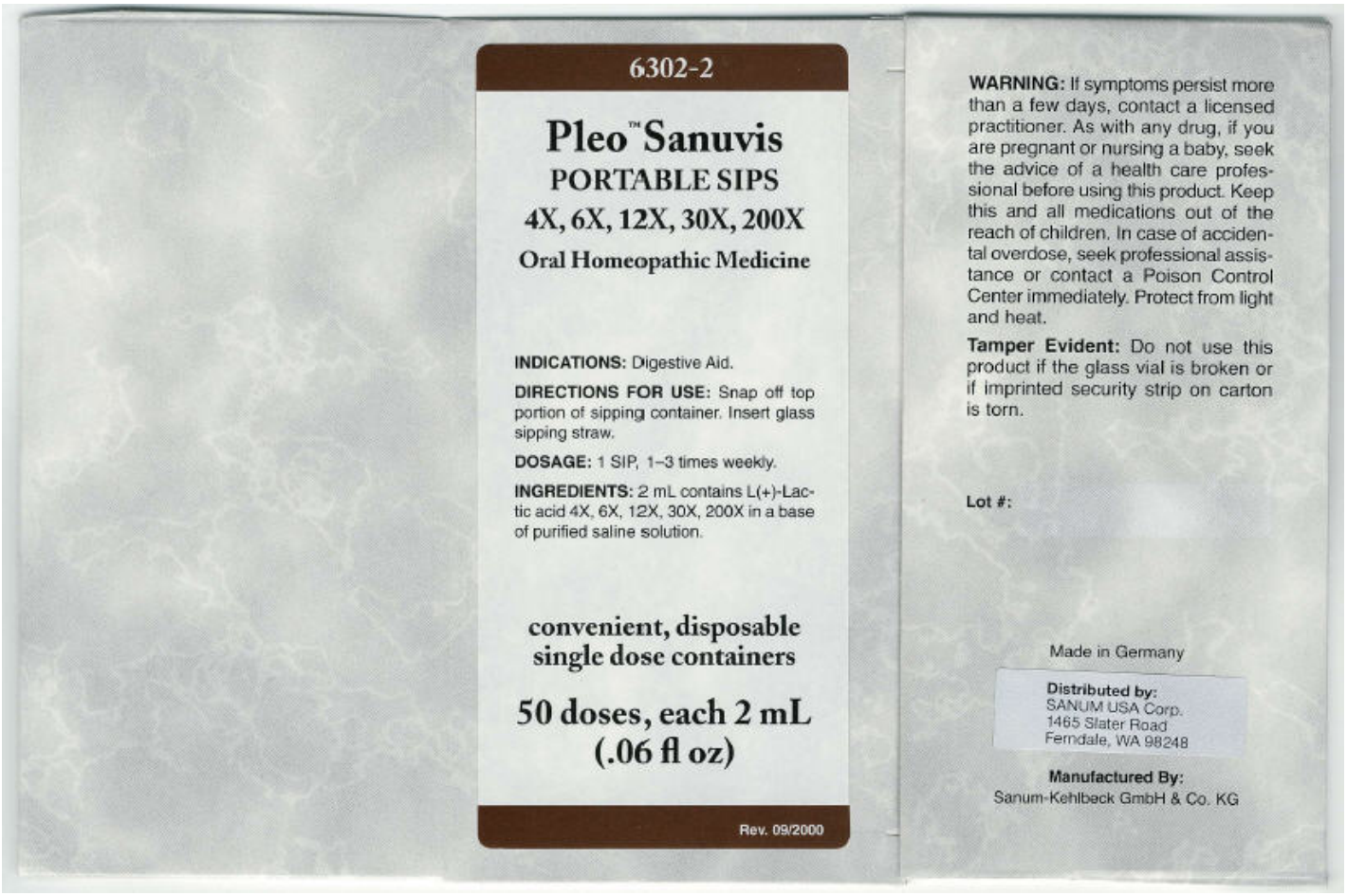
(.06 fl oz)

Rev. 09/2000



PLEO SANUVIS

lactic acid, l- kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60681-6302

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:60681-6302-1	10 in 1 CARTON		
1		1 in 1 VIAL, GLASS		
2	NDC:60681-6302-2	50 in 1 CARTON		
2		1 in 1 VIAL, GLASS		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1		2
Part 2		2
Part 3		2
Part 4		2
Part 5		2

Part 1 of 5

LACTIC ACID, L-

lactic acid, l- liquid

Product Information	
Item Code (Source)	NDC:60681-6302
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
lactic acid, l- (UNII: F9S9FFU82N) (lactic acid, l- - UNII:F9S9FFU82N)	lactic acid, l-	4 [hp_X] in 2 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

Part 2 of 5

LACTIC ACID, L-

lactic acid, l- liquid

Product Information	
Item Code (Source)	NDC:60681-6302
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
lactic acid, l- (UNII: F9S9FFU82N) (lactic acid, l- - UNII:F9S9FFU82N)	lactic acid, l-	6 [hp_X] in 2 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

Part 3 of 5

LACTIC ACID, L-

lactic acid, l- liquid

Product Information

Item Code (Source)	NDC:60681-6302
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
lactic acid, l- (UNII: F9S9FFU82N) (lactic acid, l- - UNII:F9S9FFU82N)	lactic acid, l-	12 [hp_X] in 2 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

Part 4 of 5

LACTIC ACID, L-

lactic acid, l- liquid

Product Information

Item Code (Source)	NDC:60681-6302

Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
lactic acid, l- (UNII: F9S9FFU82N) (lactic acid, l- - UNII:F9S9FFU82N)	lactic acid, l-	30 [hp_X] in 2 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

Part 5 of 5

LACTIC ACID, L-

lactic acid, l- liquid

Product Information	
Item Code (Source)	NDC:60681-6302
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
lactic acid, l- (UNII: F9S9FFU82N) (lactic acid, l- - UNII:F9S9FFU82N)	lactic acid, l-	200 [hp_X] in 2 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/24/2004	

Labeler - Sanum Kehlbeck GmbH & Co. KG (318386133)

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