EXODEN CURE POWDER- aluminum chlorohydroxy allantoinate powder
LIFEON Corp.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

70602-004_Exoden Cure Powder

Aluminum Chlorohydroxy Allantoinate 0.16%

Oral Health Care

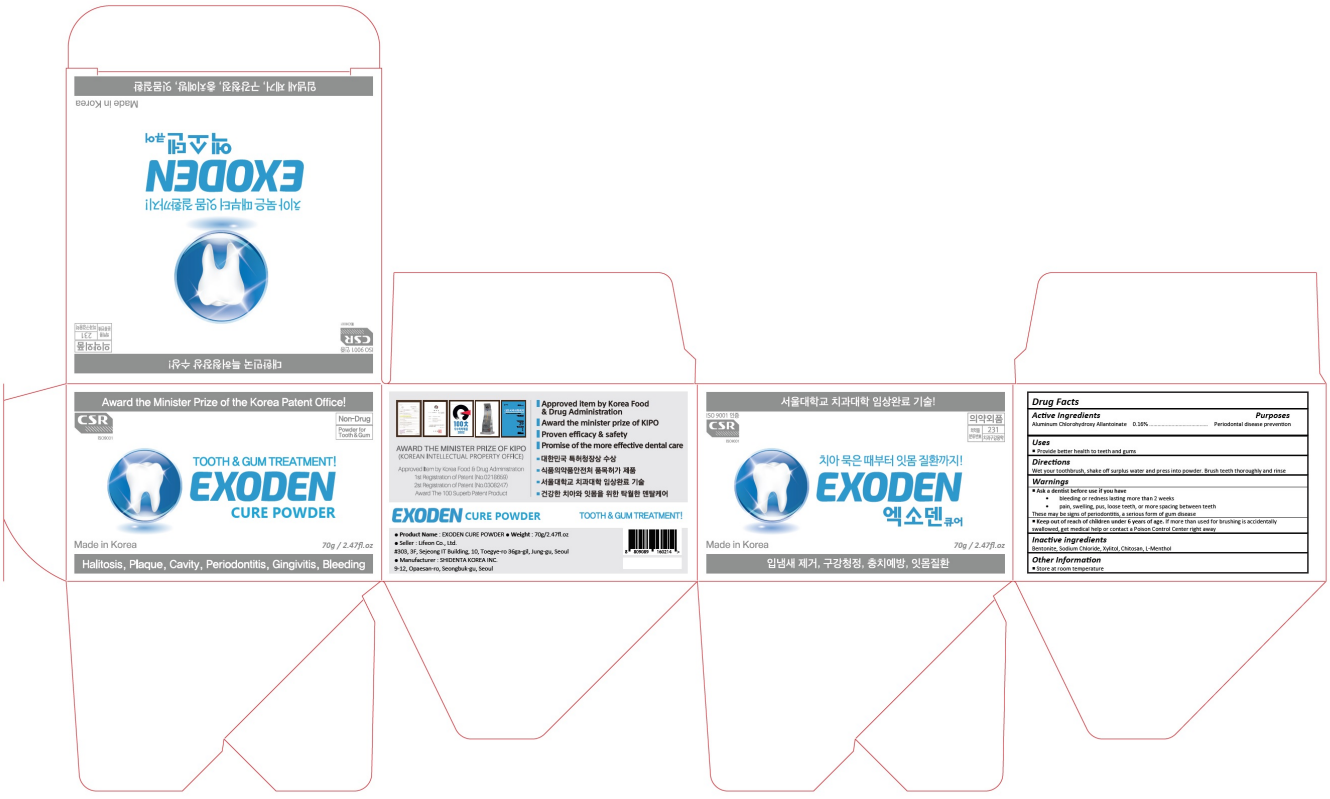
Provide better health to teeth and gums

Wet your toothbrush, shake off surplus water and press into powder. Brush teeth thoroughly and rinse

Ask a dentist before use if you have
• bleeding or redness lasting more than 2 weeks
• pain, swelling, pus, loose teeth, or more spacing between teeth
These may be signs of periodontitis, a serious form of gum disease

Keep out of reach of children under 6 years of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away

Bentonite, Sodium Chloride, Xylitol, Chitosan, L-Menthol



EXODEN CURE POWDER

aluminum chlorohydroxy allantoinate powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70602-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCLOXA (UNII: 18B8O9DQA2) (ALLANTOIN - UNII:344S277G0Z)	ALCLOXA	0.16 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
BENTONITE (UNII: A3N5ZCN45C)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
XYLITOL (UNII: VCQ006KQ1E)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70602-004-02	1 in 1 CARTON	03/09/2021	
1	NDC:70602-004-01	10 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:70602-004-04	1 in 1 CARTON	03/09/2021	
2	NDC:70602-004-03	30 g in 1 CONTAINER; Type 0: Not a Combination Product		
3	NDC:70602-004-06	1 in 1 CARTON	03/09/2021	
3	NDC:70602-004-05	70 g in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/09/2021	

Labeler - LIFEON Corp. (688528872)

Registrant - LIFEON Corp. (688528872)

Establishment

Name	Address	ID/FEI	Business Operations
Korea Shidenta		688171734	manufacture(70602-004)

Revised: 3/2022

LIFEON Corp.