National Drug Code Directory

Current through April 10, 2020

You have searched Finished drug products

Search Results: *'74651-0001-1'*

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CSV Excel

Display 50 **records per page**

Search for text in the table:

Proprietary Name	NDC Package Code	Strength	Dosage Form	Route	Appl. No.	Labeler Name	Product NDC	Nonproprietary Name	Substance Name	Product Type Name	Start Marketing Date	End Marketing Date	Market Category	Package Description	Pharm Class	DEA
+ GMW-Gold	74651-0001-1	9.2 g/100mL	LIQUID	TOPICAL		DONGBO BIO CO.,LTD	74651-0001	zeolite	ZEOLITE A	HUMAN OTC DRUG	04/04/2020	N/A	UNAPPROVED DRUG OTHER	1000 mL in 1 BOTTLE (74651-0001-1)	N/A	N/A

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Background Information (https://www.fda.gov/Drugs/InformationOnDrugs/ucm142438.htm)

Drug questions email: **DRUGINFO@FDA.HHS.GOV (mailto:DRUGINFO@FDA.HHS.Gov)**

See also: **Drug Registration and Listing Instructions (https://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/DrugRegistrationandListing/ucm078801.htm)**
National Drug Code Directory Data Files (https://www.fda.gov/Drugs/InformationOnDrugs/ucm142438.htm)

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Center for Drug Evaluation and Research
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