

55714-1706-1
ALLERGIES
National Drug Code Directory

The Drug Listing Act of 1972 requires registered drug establishments to provide the Food and Drug Administration (FDA) with a current list of all drugs manufactured, prepared, propagated, compounded, or processed by it for commercial distribution. (See Section 510 of the Federal Food, Drug, and Cosmetic Act (Act) (21 U.S.C. § 360)). Drug products are identified and reported using a unique, three-segment number, called the National Drug Code (NDC), which serves as a universal product identifier for drugs. FDA publishes the listed NDC numbers and the information submitted as part of the listing information in the NDC Directory which is updated daily.

55714-1706-1 Allergies

<i>Labeler Name</i>	Newton Laboratories, Inc.	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	55714-1706-1	The labeler code, product code, and package code segments of the National Drug Code number, separated by hyphens. Asterisks are no longer used or included within the product and package code segments to indicate certain configurations of the NDC.
<i>Proprietary Name</i>	Allergies	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	55714-1706-01	It should be noted that many NDCs are displayed on drug packaging in a 10-digit format. Proper billing of an NDC requires an 11-digit number in a 5-4-2 format. Converting NDCs from a 10-digit to 11-digit format requires a strategically placed zero, dependent upon the 10-digit format.
<i>Product NDC</i>	55714-1706	The labeler code and product code segments of the National Drug Code number, separated by a hyphen. Asterisks are no longer used or included within the product code segment to indicate certain configurations of the NDC.
<i>Product Type Name</i>	HUMAN OTC DRUG	Indicates the type of product, such as Human Prescription Drug or Human OTC Drug. This data element corresponds to the "Document Type" of the SPL submission for the listing. The complete list of codes and translations can be found at www.fda.gov/edrls under Structured Product Labeling Resources.
<i>Non Proprietary Name</i>	Cow Milk, Skim Milk, Cow Milk Fat, Casein Lactococcus Lactis Cultured, Casein Lactococcus Lactis Penicilium Roqueforti Cultured, Casein Lactococcus Lactis Cultured Penicillium Camemberti Cultured, Egg, Felis Catus Skin, Canis Lupus Familiaris Skin, Oryctolagus Cuniculus Skin, Equus Caballus Skin, Serinus Canaria Feather, Gallus Gallus Feather, Anas Platyrhynchos Feather, Anser Anser Feather, Melopsittacus Undulatus Feather, Phasianus Colchicus Feather, Meleagris Gallopavo Feather, Felis Catus Hair	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	28 g in 1 BOTTLE, GLASS (55714-1706-1)	A description of the size and type of packaging in sentence form. Multilevel packages will have the descriptions concatenated together. For example: 4 BOTTLES in 1 CARTON/100 TABLETS in 1 BOTTLE.

<i>Marketing Category Name</i>	UNAPPROVED HOMEOPATHIC	Product types are broken down into several potential Marketing Categories, such as NDA/ANDA/BLA, OTC Monograph, or Unapproved Drug. One and only one Marketing Category may be chosen for a product, not all marketing categories are available to all product types. Currently, only final marketed product categories are included. The complete list of codes and translations can be found at www.fda.gov/edrls under Structured Product Labeling Resources.
<i>Product Marketing Start Date</i>	20110301	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	PELLET	The translation of the DosageForm Code submitted by the firm. The complete list of codes and translations can be found www.fda.gov/edrls under Structured Product Labeling Resources.
<i>Route Name</i>	ORAL	The translation of the Route Code submitted by the firm, indicating route of administration. The complete list of codes and translations can be found at www.fda.gov/edrls under Structured Product Labeling Resources.

<i>Substance Name</i>		This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
	COW MILK; SKIM MILK; COW MILK FAT; CASEIN, LACTOCOCCUS LACTIS CULTURED, AGED; CASEIN, LACTOCOCCUS LACTIS CULTURED, PENICILLIUM ROQUEFORTI CULTURED, AGED; CASEIN, LACTOCOCCUS LACTIS CULTURED, PENICILLIUM CAMEMBERTI CULTURED, AGED; EGG; FELIS CATUS SKIN; ORYCTOLAGUS CUNICULUS SKIN; EQUUS CABALLUS SKIN; SERINUS CANARIA FEATHER; GALLUS GALLUS FEATHER; ANAS PLATYRHYNCHOS FEATHER; ANSER ANSER FEATHER; MELOPSITTACUS UNDULATUS FEATHER; PHASIANUS COLCHICUS FEATHER; MELEAGRIS GALLOPAVO FEATHER; FELIS CATUS HAIR; HUMAN DANDER; CANIS LUPUS FAMILIARIS HAIR; EQUUS CABALLUS HAIR; ORYCTOLAGUS CUNICULUS HAIR; HUMAN HAIR; HUMAN SKIN; SHEEP WOOL; BOS TAURUS HAIR; CAPRA HIRCUS HAIR; CAVIA PORCELLUS HAIR; MESOCRICETUS AURATUS HAIR; FD&C YELLOW NO. 5; D&C YELLOW NO. 10; FD&C YELLOW NO. 6; CARMOISINE; AMARANTH; PONCEAU 4R; FD&C RED NO. 3; ACID RED 1; FD&C BLUE NO. 2; SODIUM COPPER CHLOROPHYLLIN; ACID GREEN 50; SODIUM BENZOATE; POTASSIUM BENZOATE; PROPYLPARABEN; SODIUM DITHIONATE; 2- PHENYLPHENOL; SODIUM NITRITE; ASCORBIC ACID; SODIUM TRIPOLYPHOSPHATE; ASPARTAME; MONOSODIUM GLUTAMATE; BARLEY;	

	CORN; WHEAT GLUTEN; OAT; WHITE RICE; RYE; WHEAT; BEEF; PORK; LAMB; CHICKEN; TURKEY; QUAHOG, UNSPECIFIED; CRAB LEG, UNSPECIFIED; OYSTER, UNSPECIFIED; SHRIMP, UNSPECIFIED; SCALLOP, UNSPECIFIED; XEROPHYLLUM ASPHODELOIDES; AETHUSA CYNAPIUM; ALFALFA; ALETRIS FARINOSA ROOT; ARTEMISIA VULGARIS ROOT; BELLIS PERENNIS; DYSPHANIA AMBROSIOIDES; LYCOPUS VIRGINICUS; PHYTOLACCA AMERICANA ROOT; POPULUS TREMULOIDES LEAF; POPULUS TREMULOIDES BARK; PTELEA TRIFOLIATA BARK; SAMBUCUS NIGRA FLOWERING TOP; TRILLIUM ERECTUM ROOT; ELYMUS REPENS ROOT; URTICA URENS; WYETHIA HELENIOIDES ROOT; YUCCA FILAMENTOSA; ACONITUM NAPELLUS; ARUNDO PLINIANA ROOT; HELIANTHEMUM CANADENSE; DROSER ROTUNDIFOLIA; SOLANUM DULCAMARA TOP; LEDUM PALUSTRE TWIG; ONOSMODIUM VIRGINIANUM WHOLE; PULSATILLA VULGARIS; SANGUINARIA CANADENSIS ROOT; DATURA STRAMONIUM; ATROPA BELLADONNA; CAPSICUM; HYOSCYAMUS NIGER; SOLANUM NIGRUM WHOLE; TOBACCO LEAF; AMANITA MUSCARIA FRUITING BODY; ONION; GARLIC; ANACARDIUM OCCIDENTALE FRUIT; LYCOPERDON UTRIFORME FRUITING BODY; CAFFEINE; CINNAMON; CYNARA	
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	SCOLYMUS LEAF; ALPINE STRAWBERRY; ABRUS PRECATORIUS SEED; GLYCYRRHIZA GLABRA; MENTHA PIPERITA; NUTMEG; LACTOSE; BLACK MUSTARD SEED; ALLYLTHIOUREA; USTILAGO MAYDIS; GINGER; ASPERGILLUS NIGER VAR. NIGER; CLADOSPORIUM HERBARUM; BRAZIL NUT; HAZELNUT, UNSPECIFIED; PEANUT; ENGLISH WALNUT; TROUT, UNSPECIFIED; COD, UNSPECIFIED; HADDOCK; TUNA, UNSPECIFIED; SALMON, UNSPECIFIED; HERRING, UNSPECIFIED; AVENA SATIVA POLLEN; CYNOSURUS CRISTATUS POLLEN; ECHINOCHLOA CRUS- GALLI POLLEN; POA PRATENSIS POLLEN; ANTHOXANTHUM ODORATUM POLLEN; AGROSTIS GIGANTEA POLLEN; BROMUS SECALINUS POLLEN; PHLEUM PRATENSE POLLEN; FESTUCA PRATENSIS POLLEN; HOLCUS LANATUS POLLEN; LOLIUM PERENNE POLLEN; FAGUS SYLVATICA POLLEN; BETULA PENDULA POLLEN; QUERCUS ALBA POLLEN; FRAXINUS EXCELSIOR POLLEN; CORYLUS AMERICANA POLLEN; POPULUS NIGRA POLLEN; PLATANUS ORIENTALIS POLLEN; ULMUS GLABRA POLLEN; SALIX ALBA POLLEN; NARCISSUS PSEUDONARCISSUS; ROSA CANINA FLOWER; LILIUM CANDIDUM FLOWER; DIANTHUS CARYOPHYLLUS FLOWER; ULEX EUROPAEUS FLOWER; CYTISUS SCOPARIUS POLLEN; CALLUNA	
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	VULGARIS POLLEN; CRATAEGUS MONOGYNA POLLEN; HOUSE DUST; AMBROSIA ARTEMISIIFOLIA; ARSENIC TRIOXIDE; BROMINE; BRYONIA ALBA ROOT; OYSTER SHELL CALCIUM CARBONATE, CRUDE; MATRICARIA RECUTITA; CHELIDONIUM MAJUS; EUPHRASIA STRICTA; GELSEMIUM SEMPERVIRENS ROOT; HISTAMINE DIHYDROCHLORIDE; JUNIPERUS VIRGINIANA TWIG; POTASSIUM IODIDE; SOLANUM LYCOPERSICUM; LYCOPODIUM CLAVATUM SPORE; SODIUM CHLORIDE; TOXICODENDRON PUBESCENS LEAF; SCHOENOCAULON OFFICINALE SEED; SACCHARIN; SOLANUM TUBEROSUM; SULFUR; SACCHAROMYCES CEREVISIAE; APIS MELLIFERA; ECHINACEA, UNSPECIFIED; GOLDENSEAL; TARAXACUM OFFICINALE; ALOPECURUS PRATENSIS POLLEN; PRIMULA VULGARIS	
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Status	Deprecated	Possible status values: Active
Active NDC Code Deprecated
Deprecated NDC Code Unfinished (Unapproved)
The following status describes submitted unfinished drugs, including the marketing categories of Active Pharmaceutical Ingredient (API), Drug for Further Processing, Bulk for Human Drug Compounding, and Bulk for Animal Drug Compounding.
 FDA does not review and approve unfinished products. Therefore, all products having "unfinished" status are considered unapproved.
Last Update Date	2020-01-01	The date that a record was last updated or changed.

Food and Drug Administration
Center for Drug Evaluation and Research
Office of Compliance, Immediate Office
Drug Registration and Listing Team
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For all questions regarding this bundle please contact Support@DataLabs.Health. Also feel free to let us know about any suggestions or concerns. All additional information as well as customer support is available at <https://www.datalabs.health/>.