

**55714-4575-2**  
**DUST - MOLD - DANDER**  
**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.

**55714-4575-2 Dust - Mold - Dander**

<i>Labeler Name</i>	Newton Laboratories, Inc.	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	55714-4575-2	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>	Dust - Mold - Dander	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	55714-4575-02	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>	55714-4575	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 PRESCRIPTION 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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.
<i>Non Proprietary Name</i>	Felis Catus Skin, Canis Lupus Familiaris Skin, Ortcyolagus Cuniculus Skin, Equus Caballus Skin, Serinus Canaria Feather, Gallus Gallus Feather, Anas Platyrynchos 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	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	57 g in 1 BOTTLE, GLASS (55714-4575-2)	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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.
<i>Product Marketing Start Date</i>	20110601	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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> 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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.

<i>Substance Name</i>	<p>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;  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; WHEAT  GLUTEN; ASPERGILLUS  NIGER VAR. NIGER;  CLADOSPORIUM  HERBARUM; HOUSE  DUST; CANDIDA  ALBICANS; HISTAMINE  DIHYDROCHLORIDE;  SACCHAROMYCES  CEREVISIAE; USTILAGO  MAYDIS; ANTIMONY  TRISULFIDE; APIS  MELLIFERA; ARSENIC  TRIOXIDE; BAPTISIA  TINCTORIA ROOT;  BLATTA ORIENTALIS;  BROMINE; BRYONIA  ALBA ROOT;  CENTAUREA  BENEDICTA;  CHELIDONIUM MAJUS;  CINCHONA OFFICINALIS  BARK; IPECAC;  LYCOPODIUM  CLAVATUM SPORE;  STRYCHNOS NUX-  VOMICA SEED;  PODOPHYLLUM;  TOXICODENDRON  PUBESCENS LEAF;  SILICON DIOXIDE; IRIS  VERSICOLOR ROOT;  SOLIDAGO VIRGAUREA  FLOWERING TOP</p>	<p>This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.</p>
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<i>Pharmaceutical Classes</i>	<p>Standardized Animal Skin Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Increased IgG Production [PE], Animal Fur [CS], Allergens [CS], Non-Standardized Animal Skin Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Increased IgG Production [PE], Cells, Epidermal [EXT], Allergens [CS], Non-Standardized Animal Skin Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Increased IgG Production [PE], Cells, Epidermal [EXT], Allergens [CS], Non-Standardized Animal Skin Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Increased IgG Production [PE], Cells, Epidermal [EXT], Allergens [CS], Non-Standardized Feather Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Increased IgG Production [PE], Feathers [CS], Allergens [CS], Non-Standardized Feather Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Feathers [CS], Allergens [CS], Increased IgG Production [PE], Non-Standardized Feather Allergenic Extract [EPC], Increased Histamine Release [PE], Cell-mediated Immunity [PE], Feathers [CS], Allergens</p>	<p>These are the reported pharmaceutical class categories corresponding to the SubstanceNames listed above.</p>
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	<p>[CS],Increased IgG Production [PE],Non-Standardized Feather Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Feathers [CS],Allergens [CS],Non-Standardized Feather Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Feathers [CS],Allergens [CS],Non-Standardized Feather Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Feathers [CS],Allergens [CS],Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Dander [CS],Allergens [CS],Salivary Proteins and Peptides [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Animal Fur [CS],Allergens [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Cell-mediated Immunity [PE],Increased Histamine Release [PE],Increased IgG Production [PE],Animal Fur [CS],Allergens [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Animal Fur [CS],Allergens [CS],Salivary Proteins and Peptides [CS],Non-Standardized Animal Hair Allergenic Extract</p>	
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	<p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Allergens [CS],Animal Fur [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Animal Fur [CS],Allergens [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Allergens [CS],Animal Fur [CS],Non-Standardized Animal Hair Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Allergens [CS],Animal Fur [CS],Non-Standardized Food Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Allergens [CS],Dietary Proteins [CS],Grain Proteins [EXT],Non-Standardized Fungal Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Fungal Proteins [CS],Allergens [CS],Non-Standardized Fungal Allergenic Extract</p> <p>[EPC],Increased Histamine Release [PE],Cell-mediated Immunity [PE],Increased IgG Production [PE],Fungal Proteins [CS],Allergens [CS],Non-Standardized House Dust Allergenic Extract [EPC],Increased Histamine Release [PE]</p>	
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<i>Status</i>	Deprecated	<p>Possible status values:</p> <ul style="list-style-type: none"> <li>&lt;ul&gt; &lt;li&gt;&lt;strong&gt;Active&lt;/strong&gt; &lt;br/&gt;Active NDC Code&lt;/li&gt; &lt;li&gt;&lt;strong&gt;Deprecated&lt;/strong&gt; &lt;br/&gt;Deprecated NDC Code&lt;/li&gt; &lt;li&gt; &lt;strong&gt;Unfinished&lt;/strong&gt; (Unapproved) &lt;br/&gt;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. &lt;br/&gt; &lt;strong&gt; FDA does not review and approve unfinished products. Therefore, all products having "unfinished" status are considered unapproved. &lt;/strong&gt; &lt;/li&gt; &lt;/ul&gt;</li> </ul>
<i>Last Update Date</i>	2020-01-01	The date that a record was last updated or changed.

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