59623-010-02
SEASONAL ALLERGY PET FORMULA
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.

59623-010-02 Seasonal Allergy Pet Formula

Labeler Name	Integrative Healing Institute, LLC	Name of Company corresponding to the labeler code segment of the ProductNDC.
NDC Package Code	59623-010-02	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.
Proprietary Name	Seasonal Allergy Pet Formula	Also known as the trade name. It is the name of the product chosen by the labeler.
11 Digit NDC Code	59623-0010-02	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.
Product NDC	59623-010	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.
Product Type Name	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.
Non Proprietary Name	Hydrastis Canadensis, Kali Muriaticum, Gelsemium Sempervirens, Kali Bichromicum, Sabadilla, Histaminum, House Dust, Grass Mix, Trees Mix, Weeds Mix	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
Package Description	59 mL in 1 BOTTLE, DROPPER (59623-010-02)	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.
Marketing Category Name	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.
Product Marketing Start Date	20131107	This is the date that the labeler indicates was the start of its marketing of the drug product.

Dosage Form Name	LIQUID	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.
Route Name	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.

Substance Name	GOLDENSEAL; POTASSIUM CHLORIDE; GELSEMIUM SEMPERVIRENS ROOT; POTASSIUM DICHROMATE; SCHOENOCAULON OFFICINALE SEED; HISTAMINE DIHYDROCHLORIDE; HOUSE DUST; ANTHOXANTHUM ODORATUM; BROMUS SECALINUS POLLEN; CYNODON DACTYLON POLLEN; BARLEY; SORGHUM HALEPENSE POLLEN; POA PRATENSIS POLLEN; MELILOTUS OFFICINALIS TOP; PHLEUM PRATENSE TOP; TRITICUM AESTIVUM POLLEN; ZEA MAYS POLLEN; AMARANTHUS RETROFLEXUS POLLEN; AMBROSIA ARTEMISIIFOLIA POLLEN; RUMEX ACETOSELLA POLLEN; SOLIDAGO VIRGAUREA POLLEN; ARTEMISIA VULGARIS POLLEN; ALNUS RUBRA POLLEN; FRAXINUS PENNSYLVANICA POLLEN; FRAXINUS AMERICANA POLLEN; BETULA PAPYRIFERA POLLEN; BETULA NIGRA POLLEN; BETULA LENTA POLLEN; BETULA PUBESCENS FLOWER BUD; ACER NEGUNDO POLLEN; ULMUS AMERICANA POLLEN; ULMUS PUMILA POLLEN; ULMUS RUBRA POLLEN; CARYA GLABRA POLLEN; CARYA OVATA POLLEN; CARYA LACINIOSA POLLEN; CARYA TOMENTOSA POLLEN; ACER RUBRUM POLLEN; ACER SACCHARINUM POLLEN; ACER SACCHARUM POLLEN; QUERCUS MARILANDICA POLLEN; QUERCUS MACROCARPA POLLEN; QUERCUS STELLATA POLLEN; QUERCUS	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.

	RUBRA POLLEN; QUERCUS ALBA POLLEN; PINUS NIGRA POLLEN; PINUS TAEDA POLLEN; PINUS SYLVESTRIS POLLEN; PINUS STROBUS POLLEN; MORUS RUBRA POLLEN; JUNIPERUS VIRGINIANA POLLEN; PLATANUS OCCIDENTALIS POLLEN; JUGLANS REGIA POLLEN; POPULUS ALBA POLLEN; PLANTAGO LANCEOLATA POLLEN	
Strength Number	4; 6; 12; 12; 12; 30; 12; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6	These are the strength values (to be used with units below) of each active ingredient, listed in the same order as the SubstanceName field above.
Strength Unit	[hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_X]/mL	These are the units to be used with the strength values above, listed in the same order as the SubstanceName and SubstanceNumber.

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	2018-02-05	The date that a record was last updated or changed.

Food and Drug Administration
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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/.