

62713-961-01
ALLERGENA
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.

62713-961-01 Allergena

<i>Labeler Name</i>	Meditrend, Inc. DBA Progena Professional Formulations	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	62713-961-01	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>	Allergena	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	62713-0961-01	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>	62713-961	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>Proprietary Name Suffix</i>	Seasonal Allergy Tabs	A suffix to the proprietary name, a value here should be appended to the ProprietaryName field to obtain the complete name of the product. This suffix is often used to distinguish characteristics of a product such as extended release ("XR") or sleep aid ("PM"). Although many companies follow certain naming conventions for suffices, there is no recognized standard.
<i>Non Proprietary Name</i>	Allergy Relief	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	90 TABLET, ORALLY DISINTEGRATING in 1 BOTTLE (62713-961-01)	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>	20200511	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	TABLET, ORALLY DISINTEGRATING	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>	ACACIA POLLEN; ACER NEGUNDO POLLEN; ACER RUBRUM POLLEN; ACER SACCHARUM POLLEN; AGROSTIS GIGANTEA POLLEN; ALNUS INCANA SUBSP. RUGOSA POLLEN; ALNUS RUBRA POLLEN; AMARANTHUS PALMERI POLLEN; AMARANTHUS RETROFLEXUS POLLEN; AMBROSIA ARTEMISIIFOLIA POLLEN; AMBROSIA TRIFIDA POLLEN; ARTEMISIA VULGARIS POLLEN; BACCHARIS HALIMIFOLIA POLLEN; BAPTISIA TINCTORIA ROOT; BETULA LENTA POLLEN; BETULA NIGRA POLLEN; BETULA PAPYRIFERA POLLEN; CARYA ALBA POLLEN; CARYA GLABRA POLLEN; CARYA ILLINOINENSIS POLLEN; CARYA LACINIOSA POLLEN; CARYA OVATA POLLEN; CELTIS OCCIDENTALIS POLLEN; CHENOPODIUM ALBUM POLLEN; CUPRESSUS ARIZONICA POLLEN; CYNODON DACTYLON POLLEN; DACTYLIS GLOMERATA POLLEN; ECHINACEA, UNSPECIFIED; EUCALYPTUS GLOBULUS POLLEN; FAGUS GRANDIFOLIA POLLEN; FENUGREEK LEAF; FESTUCA PRATENSIS POLLEN; FRAXINUS AMERICANA POLLEN; FRAXINUS PENNSYLVANICA POLLEN; FRAXINUS VELUTINA POLLEN; GOLDENSEAL; IVA ANNUA POLLEN; JUGLANS NIGRA POLLEN; JUNIPERUS ASHEI POLLEN; JUNIPERUS PINCHOTII POLLEN; JUNIPERUS VIRGINIANA POLLEN; LIGUSTRUM VULGARE POLLEN; LIQUIDAMBAR STYRACIFLUA POLLEN; LOLIUM PERENNE POLLEN; MORELLA	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
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	<p>CERIFERA POLLEN; MORUS RUBRA POLLEN; MYRRH; NASTURTIUM OFFICINALE; PASPALUM NOTATUM POLLEN; PHOENIX DACTYLIFERA POLLEN; PHYTOLACCA AMERICANA ROOT; PINUS STROBUS POLLEN; PLANTAGO LANCEOLATA POLLEN; PLATANUS OCCIDENTALIS POLLEN; POA PRATENSIS POLLEN; POPULUS ALBA POLLEN; POPULUS DELTOIDES SUBSP. DELTOIDES POLLEN; POPULUS DELTOIDES SUBSP. MONILIFERA POLLEN; POPULUS TREMULOIDES POLLEN; PSEUDOTSUGA MENZIESII POLLEN; QUERCUS ALBA POLLEN; QUERCUS RUBRA POLLEN; QUERCUS VIRGINIANA POLLEN; ROBINIA PSEUDOACACIA POLLEN; RUMEX CRISPUS POLLEN; SALIX NIGRA POLLEN; SORGHUM HALEPENSE POLLEN; TAXODIUM DISTICHUM POLLEN; ULMUS AMERICANA POLLEN; ULMUS CRASSIFOLIA POLLEN; ULMUS PUMILA POLLEN; ULMUS RUBRA POLLEN; URTICA DIOICA POLLEN; XANTHIUM STRUMARIUM POLLEN</p>	
<p><i>Strength Number</i></p>	<p>30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 3; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 3; 30; 30; 3; 30; 30; 30; 30; 3; 30; 30; 30; 30; 30; 30; 30; 30; 30; 30; 3; 3; 30; 30; 3; 30; 30; 30; 30; 30; 30; 30; 30; 30</p>	<p>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.</p>

<i>Status</i>	Active	<p>Possible status values:</p> <ul style="list-style-type: none"> 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.
<i>Last Update Date</i>	2025-01-02	The date that a record was last updated or changed.

Food and Drug Administration
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Office of Compliance, Immediate Office
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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/>.