

58809-213-01

TEXACLEAR ALL SEASON ALLERGY RELIEF
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.

58809-213-01 Texaclear All Season Allergy Relief

<i>Labeler Name</i>	GM Pharmaceuticals, INC	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	58809-213-01	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>	Texaclear All Season Allergy Relief	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	58809-0213-01	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>	58809-213	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>	Antigens All Zones Grasses Trees And Weeds. Baptisia Tinctoria, Echinacea, Hydrastis Canadensis, Myrrha, Nasturtium Aquaticum, Phytolacca Decandra, Urtica Urens	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	30 mL in 1 BOTTLE, DROPPER (58809-213- 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>	20231110	This is the date that the labeler indicates was the start of its marketing of the drug product.

<i>Dosage Form Name</i>	SOLUTION/ DROPS	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>	SUBLINGUAL	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 DEALBATA POLLEN; ACER NEGUNDO POLLEN; ACER SACCHARUM POLLEN; AMARANTHUS PALMERI POLLEN; AMARANTHUS RETROFLEXUS POLLEN; AMARANTHUS SPINOSUS POLLEN; AMARANTHUS TUBERCULATUS POLLEN; AMBROSIA ACANTHICARPA POLLEN; AMBROSIA ARTEMISIIFOLIA POLLEN; AMBROSIA BIDENTATA POLLEN; AMBROSIA CONFERTIFLORA POLLEN; AMBROSIA PSILOSTACHYA POLLEN; AMBROSIA TRIFIDA POLLEN; ANTHOXANTHUM ODORATUM POLLEN; ARTEMISIA TRIDENTATA POLLEN; ARTEMISIA VULGARIS POLLEN; ATRIPLEX WRIGHTII POLLEN; AVENA SATIVA POLLEN; BAPTISIA TINCTORIA ROOT; BASSIA SCOPARIA POLLEN; BETULA NIGRA POLLEN; BROUSSONETIA PAPYRIFERA POLLEN; CARYA ILLINOINENSIS POLLEN; CARYA OVATA POLLEN; CELTIS OCCIDENTALIS POLLEN; CHENOPODIUM ALBUM POLLEN; CYCLACHAENA XANTHIFOLIA POLLEN; DISTICHLIS SPICATA POLLEN; ECHINACEA ANGUSTIFOLIA ROOT; EUCALYPTUS GLOBULUS POLLEN; EUPATORIUM CAPILLIFOLIUM POLLEN; FAGUS GRANDIFOLIA POLLEN; FRAXINUS AMERICANA POLLEN; FRAXINUS VELUTINA POLLEN; GOLDENSEAL; HESPEROCYPARIS ARIZONICA POLLEN; HOLCUS LANATUS POLLEN; JUGLANS NIGRA POLLEN; JUNIPERUS ASHEI POLLEN; JUNIPERUS	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
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	<p> OCCIDENTALIS POLLEN; JUNIPERUS VIRGINIANA POLLEN; LIGUSTRUM VULGARE POLLEN; LIQUIDAMBAR STYRACIFLUA POLLEN; MELALEUCA QUINQUENERVIA POLLEN; MORUS ALBA POLLEN; MORUS RUBRA POLLEN; MYRICA CERIFERA POLLEN; MYRRH; NASTURTIIUM OFFICINALE; OLEA EUROPAEA POLLEN; PASPALUM NOTATUM POLLEN; PHYTOLACCA AMERICANA ROOT; PINUS ECHINATA POLLEN; 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 NIGRA POLLEN; PROSOPIS JULIFLORA POLLEN; QUERCUS ALBA POLLEN; QUERCUS GAMBELII POLLEN; QUERCUS MACROCARPA POLLEN; QUERCUS NIGRA POLLEN; QUERCUS RUBRA POLLEN; QUERCUS STELLATA POLLEN; QUERCUS VIRGINIANA POLLEN; RUMEX ACETOSELLA POLLEN; RUMEX CRISPUS POLLEN; SALIX NIGRA POLLEN; SALSOLA KALI POLLEN; SCHINUS MOLLE POLLEN; SECALE CEREALE POLLEN; SOLIDAGO CANADENSIS POLLEN; SORGHUM HALEPENSE POLLEN; SYAGRUS ROMANZOFFIANA POLLEN; TAXODIUM DISTICHUM POLLEN; TRITICUM AESTIVUM POLLEN; ULMUS </p>	
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<i>Status</i>	Active	<p>Possible status values:</p> <ul style="list-style-type: none">Active
Active NDC CodeDeprecated
Deprecated NDC CodeUnfinished (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>	2025-12-16	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/>.