

58809-214-02

**TEXACLEAR TEXAS ALLERGY RELIEF PLUS CEDAR
FEVER**

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.

58809-214-02 Texaclear Texas Allergy Relief Plus Cedar Fever

<i>Labeler Name</i>	GM Pharmaceuticals, INC	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	58809-214-02	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.
<i>Proprietary Name</i>	Texaclear Texas Allergy Relief Plus Cedar Fever	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	58809-0214-02	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>	58809-214	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>	60 mL in 1 BOTTLE, DROPPER (58809-214- 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.
<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>		This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
	PLANTAGO LANCEOLATA POLLEN; PROSOPIS JULIFLORA POLLEN; QUERCUS VIRGINIANA POLLEN; ULMUS AMERICANA POLLEN; AMBROSIA BIDENTATA POLLEN; BASSIA SCOPARIA POLLEN; BETULA NIGRA POLLEN; DISTICHLIS SPICATA POLLEN; EUPATORIUM CAPILLIFOLIUM POLLEN; HOLCUS LANATUS POLLEN; MORUS RUBRA POLLEN; PINUS STROBUS POLLEN; NASTURTIUM OFFICINALE; AMARANTHUS SPINOSUS POLLEN; FRAXINUS AMERICANA POLLEN; JUGLANS NIGRA POLLEN; JUNIPERUS VIRGINIANA POLLEN; PLATANUS OCCIDENTALIS POLLEN; POPULUS DELTOIDES SUBSP. DELTOIDES POLLEN; POPULUS DELTOIDES SUBSP. MONILIFERA POLLEN; QUERCUS RUBRA POLLEN; SORGHUM HALEPENSE POLLEN; TRITICUM AESTIVUM POLLEN; URTICA DIOICA POLLEN; AMBROSIA PSILOSTACHYA POLLEN; RUMEX CRISPUS POLLEN; SECALE CEREALE POLLEN; TAXODIUM DISTICHUM POLLEN; XANTHIUM STRUMARIUM POLLEN; QUERCUS NIGRA POLLEN; QUERCUS STELLATA POLLEN; SOLIDAGO CANADENSIS POLLEN; AMARANTHUS RETROFLEXUS POLLEN; AMBROSIA CONFERTIFLORA POLLEN; EUCALYPTUS GLOBULUS POLLEN; MYRRH; SALSOLA KALI POLLEN; QUERCUS MACROCARPA POLLEN; SYAGRUS ROMANZOFFIANA POLLEN; ZEA MAYS POLLEN;	

	CHENOPODIUM ALBUM POLLEN; ARTEMISIA TRIDENTATA POLLEN; GOLDENSEAL; FAGUS GRANDIFOLIA POLLEN; FRAXINUS VELUTINA POLLEN; BROUSSONETIA PAPYRIFERA POLLEN; PINUS ECHINATA POLLEN; URTICA URENS WHOLE; AMARANTHUS PALMERI POLLEN; AMARANTHUS TUBERCULATUS POLLEN; CARYA ILLINOINENSIS POLLEN; LIQUIDAMBAR STYRACIFLUA POLLEN; MELALEUCA QUINQUENERVIA POLLEN; MORUS ALBA POLLEN; HESPEROCYPARIS ARIZONICA POLLEN; JUNIPERUS OCCIDENTALIS POLLEN; OLEA EUROPAEA POLLEN; ECHINACEA ANGUSTIFOLIA ROOT; ACER SACCHARUM POLLEN; ANTHOXANTHUM ODORATUM POLLEN; AVENA SATIVA POLLEN; PHYTOLACCA AMERICANA ROOT; BAPTISIA TINCTORIA ROOT; ACER NEGUNDO POLLEN; AMBROSIA ACANTHICARPA POLLEN; AMBROSIA ARTEMISIIFOLIA POLLEN; AMBROSIA TRIFIDA POLLEN; ARTEMISIA VULGARIS POLLEN; CARYA OVATA POLLEN; CELTIS OCCIDENTALIS POLLEN; CYCLACHAENA XANTHIFOLIA POLLEN; JUNIPERUS ASHEI POLLEN; LIGUSTRUM VULGARE POLLEN; MYRICA CERIFERA POLLEN; POA PRATENSIS POLLEN; POPULUS ALBA POLLEN; RUMEX ACETOSELLA POLLEN; SALIX NIGRA POLLEN; SCHINUS MOLLE POLLEN; ACACIA DEALBATA POLLEN;	
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[illegible]

<i>Status</i>	Active	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.
<i>Last Update Date</i>	2025-12-16	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/>.