58809-215-90

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.

58809-215-90 Texaclear Texas Allergy Relief Plus Cedar Fever

Labeler Name	GM Pharmaceuticals, INC	Name of Company corresponding to the labeler code segment of the ProductNDC.
NDC Package Code	58809-215-90	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	Texaclear Texas Allergy Relief Plus Cedar Fever	Also known as the trade name. It is the name of the product chosen by the labeler.
11 Digit NDC Code	58809-0215-90	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	58809-215	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	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.
Package Description	90 min in 1 BOTTLE (58809-215-90)	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	20231110	This is the date that the labeler indicates was the start of its marketing of the drug product.

Dosage Form Name	TABLET, CHEWABLE	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	SOLIDAGO CANADENSIS POLLEN; PHYTOLACCA AMERICANA ROOT; AMBROSIA ARTEMISIIFOLIA POLLEN; POPULUS DELTOIDES SUBSP. MONILIFERA POLLEN; PINUS ECHINATA POLLEN; SORGHUM HALEPENSE POLLEN; QUERCUS MACROCARPA POLLEN; QUERCUS NIGRA POLLEN; OLEA EUROPAEA POLLEN; BROUSSONETIA PAPYRIFERA POLLEN; ARTEMISIA VULGARIS POLLEN; POPULUS NIGRA POLLEN; AMBROSIA PSILOSTACHYA POLLEN; QUERCUS RUBRA POLLEN; SECALE CEREALE POLLEN; CARYA OVATA POLLEN; RUMEX ACETOSELLA POLLEN; AMARANTHUS TUBERCULATUS POLLEN; RUMEX CRISPUS POLLEN; CELTIS OCCIDENTALIS POLLEN; PLANTAGO LANCEOLATA POLLEN; POPULUS DELTOIDES SUBSP. DELTOIDES POLLEN; AMBROSIA ACANTHICARPA POLLEN; CHENOPODIUM ALBUM POLLEN; BAPTISIA TINCTORIA ROOT; TAXODIUM DISTICHUM POLLEN; JUGLANS NIGRA POLLEN; ECHINACEA ANGUSTIFOLIA ROOT; MYRRH; ULMUS AMERICANA POLLEN; GOLDENSEAL; EUPATORIUM CAPILLIFOLIUM POLLEN; PROSOPIS JULIFLORA POLLEN; MORUS ALBA POLLEN; SYAGRUS ROMANZOFFIANA POLLEN; AMBROSIA CONFERTIFLORA POLLEN; FRAXINUS AMERICANA POLLEN; CYCLACHAENA XANTHIFOLIA POLLEN;	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.

	URTICA DIOICA POLLEN; CARYA ILLINOINENSIS POLLEN; PINUS STROBUS POLLEN; AMBROSIA BIDENTATA POLLEN; ARTEMISIA TRIDENTATA POLLEN; XANTHIUM STRUMARIUM POLLEN; JUNIPERUS VIRGINIANA POLLEN; BETULA NIGRA POLLEN; PLATANUS OCCIDENTALIS POLLEN; ACER SACCHARUM POLLEN; EUCALYPTUS GLOBULUS POLLEN; MYRICA CERIFERA POLLEN; SCHINUS MOLLE POLLEN; BASSIA SCOPARIA POLLEN; POA PRATENSIS POLLEN; QUERCUS VIRGINIANA POLLEN; JUNIPERUS ASHEI POLLEN; MELALEUCA QUINQUENERVIA POLLEN; QUERCUS GAMBELII POLLEN; QUERCUS STELLATA POLLEN; QUERCUS ALBA POLLEN; AMARANTHUS RETROFLEXUS POLLEN; LIGUSTRUM VULGARE POLLEN; AMBROSIA TRIFIDA POLLEN; MORUS RUBRA POLLEN; SALSOLA KALI POLLEN; ATRIPLEX WRIGHTII POLLEN; DISTICHLIS SPICATA POLLEN; AMARANTHUS SPINOSUS POLLEN; ANTHOXANTHUM ODORATUM POLLEN; LIQUIDAMBAR STYRACIFLUA POLLEN; HOLCUS LANATUS POLLEN; JUNIPERUS OCCIDENTALIS POLLEN; POPULUS ALBA POLLEN; SALIX NIGRA POLLEN; URTICA URENS WHOLE; HESPEROCYPARIS ARIZONICA POLLEN; PASPALUM NOTATUM POLLEN; FAGUS GRANDIFOLIA POLLEN; ACER NEGUNDO POLLEN; AMARANTHUS PALMERI POLLEN; AVENA SATIVA POLLEN; ACACIA DEALBATA	

	POLLEN; FRAXINUS VELUTINA POLLEN; ZEA MAYS POLLEN; TRITICUM AESTIVUM POLLEN; NASTURTIUM OFFICINALE	
Strength Number	6; 5; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 5; 6; 6; 5; 5; 6; 5; 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; 5; 6; 6; 6; 6; 6; 6; 6; 6; 6; 6; 5	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]/min; [hp_X]/min	These are the units to be used with the strength values above, listed in the same order as the SubstanceName and SubstanceNumber.

Pharmaceutical Classes	Allergens [CS], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity	These are the reported pharmaceutical class categories corresponding to the SubstanceNames listed above.

	[PE], Cell-mediated Immunity [PE], Cell-mediated Immunity	

	[PE], Cell-mediated Immunity [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE]	

Status	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.
Last Update Date	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/.