

51672-1296-2
DIFLORASONE DIACETATE
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.

<i>Product Marketing Start Date</i>	20000424	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	CREAM	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>	TOPICAL	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>	DIFLORASONE DIACETATE	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
<i>Strength Number</i>	.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.
<i>Strength Unit</i>	mg/g	These are the units to be used with the strength values above, listed in the same order as the SubstanceName and SubstanceNumber.
<i>Pharmaceutical Classes</i>	Corticosteroid Hormone Receptor Agonists [MoA], Corticosteroid [EPC]	These are the reported pharmaceutical class categories corresponding to the SubstanceNames listed above.
<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>	2025-09-24	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
10903 New Hampshire Ave
Silver Spring, MD 20993-0002
Email: eDRLS@fda.hhs.gov

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/>.