

13537-669-28

ESIKA HYDRACOLOR 2 IN 1 SPF 25

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.

13537-669-28 Esika Hydracolor 2 In 1 Spf 25

<i>Labeler Name</i>	Ventura Corporation LTD	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	13537-669-28	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>	Esika Hydracolor 2 In 1 Spf 25	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	13537-0669-28	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>	13537-669	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>	Octinoxate And Oxybenzone	Sometimes called the generic name, this is usually the active ingredient(s) of the product.

[illegible]

<i>Status</i>	Deprecated	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>	2019-09-21	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/>.