

58159-127-11

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

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<i>Labeler Name</i>	NURAY CHEMICALS PRIVATE LIMITED	Name of Company corresponding to the labeler code segment of the ProductNDC.
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<i>Strength Number</i>	35	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>	kg/35kg	These are the units to be used with the strength values above, listed in the same order as the SubstanceName and SubstanceNumber.

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/>