

81395-001-02

UBLIM ALCOHOL DISINFECTANT
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

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<i>Labeler Name</i>	Jiangsu UBL Life Technology
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<i>Product Marketing Start Date</i>	20210218	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	CLOTH	The translation of the DosageForm Code submitted by the firm. The complete list of codes and translations can be found

Food and Drug Administration
Center for Drug Evaluation and Research
Office of Compliance, Immediate Office
Drug Registration and Listing Team
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Silver Spring, MD 20993-0002
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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/>.