

83202-6841-3

**DR. NO I (WOUND RECOVERY, BURN WOUND
RECOVERY, ACNE TREATMENT, FERMENTED
LETTUCE EXTRACT CONCENTRATE)**

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

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<i>Labeler Name</i>	HumanEnos LLC	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>		

<i>Product Marketing End Date</i>	20250212	This is the date the product will no longer be available on the market. If a product is no longer being manufactured, in most cases, the FDA recommends firms use the expiration date of the last lot produced as the EndMarketingDate, to reflect the potential for drug product to remain available after manufacturing has ceased. Products that are the subject of ongoing manufacturing will not ordinarily have any EndMarketingDate. Products with a value in the EndMarketingDate will be removed from the NDC Directory
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Food and Drug Administration
Center for Drug Evaluation and Research
Office of Compliance, Immediate Office
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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/>.