

**44911-0746-1**  
**BIO IMMUNE**  
**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.

**44911-0746-1 Bio Immune**

<i>Labeler Name</i>	Energique, Inc.	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	44911-0746-1	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>	Bio Immune	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	44911-0746-01	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>	44911-0746	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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.
<i>Non Proprietary Name</i>	Chelidonium Majus, Lobelia Inflata, Rumex Crispus, Sambucus Nigra, Senega Officinalis, Spongia Tosta, Taraxacum Officinale, Myristica Sebifera, Sticta Pulmonaria, Sulphur, Ustilago Maidis, Polyporus Officinalis, Candida Albicans, Carduus Marianus, Crotalus Horridus, Scrophularia Nodosa, Agaricus Muscarius, Bovista, Secale Cornutum, Anas Barbariae, Hepatis Et Cordis Extractum, Anthracinum, Arsenicum Album, Baptisia Tinctoria, Berberis Vulgaris, Botulinum, Bryonia (alba), Camphora, Cinchona Officinalis,	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	30 mL in 1 BOTTLE, DROPPER (44911-0746- 1)	A description of the size and type of packaging in sentence form. Multilevel packages will have the descriptions concatenated together. For example: 4 BOTTLES in 1 CARTON/100 TABLETS in 1 BOTTLE.

<i>Marketing Category Name</i>	UNAPPROVED HOMEOPATHIC	Product types are broken down into several potential Marketing Categories, such as NDA/ANDA/BLA, OTC Monograph, or Unapproved Drug. One and only one Marketing Category may be chosen for a product, not all marketing categories are available to all product types. Currently, only final marketed product categories are included. The complete list of codes and translations can be found at <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.
<i>Product Marketing Start Date</i>	20260108	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	LIQUID	The translation of the DosageForm Code submitted by the firm. The complete list of codes and translations can be found <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.
<i>Route Name</i>	ORAL	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 <a href="http://www.fda.gov/edrls">www.fda.gov/edrls</a> under Structured Product Labeling Resources.

<i>Substance Name</i>		This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
	<p>           AMANITA MUSCARIA            FRUITING BODY;            ARSENIC TRIOXIDE;            BACILLUS ANTHRACIS            IMMUNOSERUM RABBIT;            BAPTISIA TINCTORIA            ROOT; BERBERIS            VULGARIS ROOT BARK;            BRYONIA ALBA ROOT;            CAIRINA MOSCHATA            HEART/LIVER            AUTOLYSATE;            CAMPHOR (NATURAL);            CANDIDA ALBICANS;            CHELIDONIUM MAJUS            WHOLE; CINCHONA            OFFICINALIS BARK;            CLAVICEPS PURPUREA            SCLEROTIUM;            CLOSTRIDIUM            BOTULINUM; CROTALUS            HORRIDUS HORRIDUS            VENOM; GELSEMIUM            SEMPERVIRENS ROOT;            HUMAN SPUTUM,            BORDETELLA            PERTUSSIS INFECTED;            INFLUENZA A VIRUS            A/THAILAND/8/2022 IVR-            237 (H3N2) ANTIGEN            (FORMALDEHYDE            INACTIVATED);            INFLUENZA A VIRUS            A/VICTORIA/4897/2022            IVR-238 (H1N1) ANTIGEN            (FORMALDEHYDE            INACTIVATED);            INFLUENZA B VIRUS            B/AUSTRIA/1359417/2021            BVR-26 WHOLE;            INFLUENZA B VIRUS            B/PHUKET/3073/2013            BVR-1B WHOLE;            LARICIFOMES            OFFICINALIS FRUITING            BODY; LOBARIA            PULMONARIA; LOBELIA            INFLATA WHOLE;            LYCOPERDON            UTRIFORME FRUITING            BODY; MILK THISTLE;            POLYGALA SENEGA            ROOT; RANCID BEEF;            RUMEX CRISPUS ROOT;            SALMONELLA ENTERICA            ENTERICA SEROVAR            ENTERITIDIS;            SAMBUCUS NIGRA            FLOWERING TOP;            SCROPHULARIA            NODOSA WHOLE;            SHIGELLA            DYSENTERIAE;            SPONGIA OFFICINALIS         </p>	

	SKELETON, ROASTED; STREPTOCOCCUS PNEUMONIAE; SULFUR; TARAXACUM OFFICINALE; USTILAGO MAYDIS; VACCINIA VIRUS; VARIOLA VIRUS; VIROLA SEBIFERA RESIN	
<i>Strength Number</i>	10; 30; 30; 30; 30; 30; 30; 30; 12; 3; 30; 10; 30; 12; 30; 30; 30; 30; 30; 30; 8; 6; 3; 10; 12; 3; 30; 3; 30; 3; 12; 30; 3; 30; 6; 3; 6; 30; 30; 6	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>	[hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_X]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_C]/mL; [hp_X]/mL; [hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_X]/mL; [hp_C]/mL; [hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_X]/mL; [hp_C]/mL; [hp_X]/mL; [hp_X]/mL; [hp_X]/mL; [hp_C]/mL; [hp_C]/mL; [hp_X]/mL	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>	Allergens [CS], Allergens [CS], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Fungal Proteins [CS], Fungal Proteins [CS], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased IgG Production [PE], Increased IgG Production [PE], Non-Standardized Fungal Allergenic Extract [EPC], Non-Standardized Fungal Allergenic Extract [EPC]	These are the reported pharmaceutical class categories corresponding to the SubstanceNames listed above.

<i>Status</i>	Active	<p>Possible status values:</p> <ul style="list-style-type: none"> <li>&lt;li&gt;&lt;strong&gt;Active&lt;/strong&gt; &lt;br/&gt;Active NDC Code&lt;/li&gt;</li> <li>&lt;li&gt;&lt;strong&gt;Deprecated&lt;/strong&gt; &lt;br/&gt;Deprecated NDC Code&lt;/li&gt;</li> <li>&lt;li&gt;&lt;strong&gt;Unfinished&lt;/strong&gt; (Unapproved) &lt;br/&gt;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. &lt;br/&gt;&lt;strong&gt;FDA does not review and approve unfinished products. Therefore, all products having "unfinished" status are considered unapproved.&lt;/strong&gt;&lt;/li&gt;</li> </ul>
<i>Last Update Date</i>	2026-01-13	The date that a record was last updated or changed.

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For all questions regarding this bundle please contact [Support@DataLabs.Health](mailto: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/>.