

66096-712-27

JOETTE CALABRESE SURVIVALIST KIT

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.

66096-712-27 Joette Calabrese Survivalist Kit

<i>Labeler Name</i>	OHM PHARMA INC.	Name of Company corresponding to the labeler code segment of the ProductNDC.
<i>NDC Package Code</i>	66096-712-27	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>	Joette Calabrese Survivalist Kit	Also known as the trade name. It is the name of the product chosen by the labeler.
<i>11 Digit NDC Code</i>	66096-0712-27	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>	66096-712	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>	Abrot Acon Agn Ail Ambr Ambro Am-c Ant-c Apis Arn Ars Aur-a Aur Bapt Bar-c Bell Bb Both Bov Brom Cadm-s Calad Calc Camph Carb-ac Card-m Cean Chel Chinin-a Cina Chin Cic Cob Colch Con Crot-h Crot-t Cupr Dulc Elaps Eup-per Fl-ac Galph Graph Gunp Ham Hekla Hell Hep Hydr-ac Hyos Hyper Kali-br Kali-c Kali-i Lath Laur Led Lob Merc-cy Merc Mez Mosch Mur-ac Naja Nat-c Nit-ac Nux-m Pyrog Ph-ac Pic-ac Plb Rhus-t Ruta Samb Sang Sarr Sep Stram Sul-ac Tab Terent Ter Teucr Verat Verat-v Vip	Sometimes called the generic name, this is usually the active ingredient(s) of the product.
<i>Package Description</i>	100 PELLET in 1 KIT (66096-712-27)	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 www.fda.gov/edrls under Structured Product Labeling Resources.
<i>Product Marketing Start Date</i>	20151026	This is the date that the labeler indicates was the start of its marketing of the drug product.
<i>Dosage Form Name</i>	PELLET	The translation of the DosageForm Code submitted by the firm. The complete list of codes and translations can be found www.fda.gov/edrls 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 www.fda.gov/edrls under Structured Product Labeling Resources.

<i>Substance Name</i>	ACONITUM NAPELLUS; AETHUSA CYNAPIUM; AILANTHUS ALTISSIMA FLOWERING TWIG; AMBERGRIS; AMBROSIA ARTEMISIIFOLIA POLLEN; AMMONIUM CARBONATE; ANTIMONY TRISULFIDE; APIS MELLIFERA; ARNICA MONTANA; ARSENIC TRIOXIDE; ARTEMISIA ABROTANUM FLOWERING TOP; ARTEMISIA CINA PRE- FLOWERING TOP; ATROPA BELLADONNA; BAPTISIA TINCTORIA ROOT; BARIUM CARBONATE; BORRELIA BURGDORFERI; BOTHROPS ATROX VENOM; BROMINE; CADMIUM SULFATE; CALCIUM SULFIDE; CAMPHOR (NATURAL); CEANOTHUS AMERICANUS LEAF; CHASTE TREE; CHELIDONIUM MAJUS; CICUTA VIROSA ROOT; CINCHONA OFFICINALIS BARK; CLAVICEPS PURPUREA SCLEROTIUM; COBALT; COLCHICUM AUTUMNALE BULB; CONIUM MACULATUM FLOWERING TOP; COPPER; CROTALUS HORRIDUS HORRIDUS VENOM; CROTON TIGLIUM SEED; DAPHNE MEZEREUM BARK; DATURA STRAMONIUM; DIEFFENBACHIA SEGUINE; EUPATORIUM PERfoliatum FLOWERING TOP; GALPHIMIA GLAUCA FLOWERING TOP; GOLD; GRAPHITE; HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK; HEKLA LAVA; HELLEBORUS NIGER ROOT; HYDROCHLORIC ACID; HYDROFLUORIC ACID; HYDROGEN CYANIDE; HYOSCYAMUS NIGER; HYPERICUM PERFORATUM; LATHYRUS SATIVAS	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.
-----------------------	---	--

	SEED; LEAD; LEDUM PALUSTRE TWIG; LOBELIA INFLATA; LYCOPERDON UTRIFORME FRUITING BODY; LYCOSA TARANTULA; MERCURIC CYANIDE; MERCURIUS SOLUBILIS; MICRURUS CORALLINUS VENOM; MILK THISTLE; MOSCHUS MOSCHIFERUS WHOLE; NAJA NAJA VENOM; NITRIC ACID; NUTMEG; OYSTER SHELL CALCIUM CARBONATE, CRUDE; PHENOL; PHOSPHORIC ACID; PICRIC ACID; POTASSIUM BROMIDE; POTASSIUM CARBONATE; POTASSIUM IODIDE; PRUNUS LAUROCERASUS LEAF; QUININE SULFATE; RANCID BEEF; RUTA GRAVEOLENS FLOWERING TOP; SAMBUCUS NIGRA FLOWERING TOP; SANGUINARIA CANADENSIS ROOT; SARRACENIA PURPUREA; SEPIA OFFICINALIS JUICE; SODIUM CARBONATE; SOLANUM DULCAMARA TOP; SULFURIC ACID; TEUCRIUM MARUM; TOBACCO LEAF; TOXICODENDRON PUBESCENS LEAF; TURPENTINE OIL; VERATRUM ALBUM ROOT; VERATRUM VIRIDE ROOT; VIPERA BERUS VENOM	
<i>Strength Number</i>	1; 200; 30; 30; 30; 30; 6; 1; 3; 3; 6; 3; 3; 200; 30; 200; 30; 30; 30; 6; 200; 30; 200; 6; 30; 3; 6; 30; 30; 3; 6; 6; 30; 200; 30; 30; 200; 30; 200; 200; 200; 30; 30; 30; 30; 30; 6; 1; 30; 30; 1; 30; 30; 30; 200; 6; 30; 30; 30; 30; 30; 30; 1; 6; 6; 200; 30; 200; 6; 30; 3; 200; 6; 30; 200; 30; 6; 30; 30; 30; 200; 30; 1; 6; 200; 30; 30	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>Pharmaceutical Classes</i>	Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Antimalarial [EPC], Bee Venoms [CS], Cell-mediated Immunity [PE], Copper [CS], Copper-containing Intrauterine Device [EPC], Decreased Embryonic Implantation [PE], Decreased Sperm Motility [PE], Dietary Proteins [CS], Food Additives [CS], Increased Histamine Release [PE], Increased IgG Production [PE], Increased IgG Production [PE], Increased IgG Production [PE], Inhibit Ovum Fertilization [PE], Non-Standardized Food Allergenic Extract [EPC], Non-Standardized Plant Allergenic Extract [EPC], Non-Standardized Plant Allergenic Extract [EPC], Plant Proteins [CS], Plant Proteins [CS], Pollen [CS], Seed Storage Proteins [CS], Standardized Chemical Allergen [EPC], Standardized Chemical Allergen [EPC], Standardized Insect Venom Allergenic Extract [EPC], Standardized Pollen 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">Active
Active NDC CodeDeprecated
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>	2021-12-31	The date that a record was last updated or changed.

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/>.