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

Labeler Name	OHM PHARMA INC.	Name of Company corresponding to the labeler code segment of the ProductNDC.
NDC Package Code	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.
Proprietary Name	Joette Calabrese Survivalist Kit	Also known as the trade name. It is the name of the product chosen by the labeler.
11 Digit NDC Code	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.
Product NDC	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.
Product Type Name	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.
Non Proprietary Name	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.
Package Description	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.

Marketing Category Name	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.
Product Marketing Start Date	20151026	This is the date that the labeler indicates was the start of its marketing of the drug product.
Dosage Form Name	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.
Route Name	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.

Substance Name	ARTEMISIA ABROTANUM FLOWERING TOP; ACONITUM NAPELLUS; AETHUSA CYNAPIUM; CHASTE TREE; AILANTHUS ALTISSIMA FLOWERING TWIG; AMBERGRIS; AMBROSIA ARTEMISIIFOLIA POLLEN; AMMONIUM CARBONATE; ANTIMONY TRISULFIDE; APIS MELLIFERA; ARNICA MONTANA; ARSENIC TRIOXIDE; GOLD; BAPTISIA TINCTORIA ROOT; BARIUM CARBONATE; ATROPA BELLADONNA; BORRELIA BURGDORFERI; BOTHROPS ATROX VENOM; LYCOPERDON UTRIFORME FRUITING BODY; BROMINE; CADMIUM SULFATE; DIEFFENBACHIA SEGUINE; OYSTER SHELL CALCIUM CARBONATE, CRUDE; CAMPHOR (NATURAL); PHENOL; MILK THISTLE; CEANOTHUS AMERICANUS LEAF; CHELIDONIUM MAJUS; QUININE SULFATE; ARTEMISIA CINA PRE-FLOWERING TOP; CINCHONA OFFICINALIS BARK; CICUTA VIROSA ROOT; COBALT; COLCHICUM AUTUMNALE BULB; CONIUM MACULATUM FLOWERING TOP; CROTALUS HORRIDUS HORRIDUS VENOM; CROTON TIGLIUM SEED; COPPER; SOLANUM DULCAMARA TOP; MICRURUS CORALLINUS VENOM; EUPATORIUM PERFOLIATUM FLOWERING TOP; HYDROFLUORIC ACID; GALPHIMIA GLAUCA FLOWERING TOP; GRAPHITE; HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK; HEKLA LAVA; HELLEBORUS NIGER ROOT; CALCIUM SULFIDE; HYDROGEN	This is the active ingredient list. Each ingredient name is the preferred term of the UNII code submitted.

	CYANIDE; HYOSCYAMUS NIGER; HYPERICUM PERFORATUM; POTASSIUM BROMIDE; POTASSIUM CARBONATE; POTASSIUM IODIDE; LATHYRUS SATIVAS SEED; PRUNUS LAUROCERASUS LEAF; LEDUM PALUSTRE TWIG; LOBELIA INFLATA; MERCURIC CYANIDE; MERCURIUS SOLUBILIS; DAPHNE MEZEREUM BARK; MOSCHUS MOSCHIFERUS WHOLE; HYDROCHLORIC ACID; NAJA NAJA VENOM; SODIUM CARBONATE; NITRIC ACID; NUTMEG; RANCID BEEF; PHOSPHORIC ACID; PICRIC ACID; LEAD; TOXICODENDRON PUBESCENS LEAF; RUTA GRAVEOLENS FLOWERING TOP; SAMBUCUS NIGRA FLOWERING TOP; SANGUINARIA CANADENSIS ROOT; SARRACENIA PURPUREA; CLAVICEPS PURPUREA SCLEROTIUM; SEPIA OFFICINALIS JUICE; DATURA STRAMONIUM; SULFURIC ACID; TOBACCO LEAF; LYCOSA TARANTULA; TURPENTINE OIL; TEUCRIUM MARUM; VERATRUM ALBUM ROOT; VERATRUM VIRIDE ROOT; VIPERA BERUS VENOM	
Strength Number	6; 1; 200; 200; 30; 30; 30; 30; 6; 1; 3; 3; 200; 200; 30; 3; 200; 30; 30; 30; 30; 30; 1; 200; 6; 30; 30; 6; 3; 3; 3; 30; 30; 30; 3; 6; 30; 6; 30; 30; 200; 30; 30; 200; 200; 30; 30; 6; 30; 6; 1; 30; 200; 6; 30; 30; 1; 30; 200; 6; 200; 30; 30; 30; 30; 30; 30; 200; 6; 200; 30; 1; 6; 30; 200; 30; 6; 6; 30; 30; 30; 30; 6; 200; 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.

Strength Unit	[hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_X]/1; [hp_X]/1; [hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_M]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1; [hp_C]/1	These are the units to be used with the strength values above, listed in the same order as the SubstanceName and SubstanceNumber.

Pharmaceutical Classes	Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Allergens [CS], Antimalarial [EPC], Bee Venoms [CS], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], Cell-mediated Immunity [PE], 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 Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], Increased Histamine Release [PE], 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.

Status	Active	Possible status values: - Active Active NDC Code - Deprecated 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.
Last Update Date	2021-12-31	The date that a record was last updated or changed.

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