Section	Field Name	Туре	Description
	brand_name	string	The Proprietary/Trade/Brand name of the medical device as used in device labeling or in the catalog. This information may 1) be on a label attached to a durable device, 2) be on a package of a disposable device, or 3) appear in labeling materials of an implantable device. The brand
Ì		1	name is the name that is typically registered with USPTO and have the * and/or TM symbol.
			The state of the s
	catalog_number	string	This is an .exact field. It has been indexed both as its exact string content, and also tokenized. The catalog, reference, or product number found on the device label or accompanying packaging to identify a particular product.
	Catalog_named	30.1116	The catalog, reserved, or product number round on the device label of accompanying packaging to definity a particular product.
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
	commercial_distribution_end_date	date	Indicates the date the device is no longer held or offered for sale. See 21 CFR 807.3(b) for exceptions. The device may or may not still be available for purchase in the marketplace.
	commercial_distribution_status	string	Indicates whether the device is in commercial distribution as defined under 21 CFR 807.3(b).
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
			Value is one of the following
			In Commercial Distribution = In Commercial Distribution
	company_name	string	Not in Commercial Distribution = Not in Commercial Distribution Company name associated with the labeler DUNS Number entered in the DI Record.
	de Service Se have reduce		This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
	device_count_in_base_package device description	integer	Number of medical devices in the base package. Additional relevant information about the device that is not already captured as a distinct GUDID data attribute.
	has_donation_id_number	boolean	The Donation Identification Number is applicable to devices that are also regulated as HCT/Ps and is a number that is assigned to each
			donation. This number/code is required to be part of the UDI when included on the label in order to provide the means to track the device back to
			its manufacturing source or otherwise allow the history of the device manufacturing, packaging, labeling, distribution and use to be determined.
			Value is one of the following
			true = true false = false
			Talse = Talse
	has_expiration_date	boolean	The date by which the label of a device states the device must or should be used. This date is required to be part of the UDI when included on the
			label in order to provide the means to track the device back to its manufacturing source or otherwise allow the history of the device
			manufacturing, packaging, labeling, distribution and use to be determined.
			Value is one of the following
			true = true
	has_lot_or_batch_number	boolean	false = false The number assigned to one or more device(s) that consist of a single type, model, class, size, composition, or software version that are
		_ concult	manufactured under essentially the same conditions and that are intended to have uniform characteristics and quality within specified limits. This
		1	number is required to be part of the UDI when included on the label in order to provide the means to track the device back to its manufacturing
			source or otherwise allow the history of the device manufacturing, packaging, labeling, distribution and use to be determined.
			Value is one of the following
			true = true
			false = false
	has_manufacturing_date	boolean	The date on which a device is manufactured. This date is required to be part of the UDI when included on the label in order to provide the means
			to track the device back to its manufacturing source or otherwise allow the history of the device manufacturing, packaging, labeling, distribution
			and use to be determined.
			Value is one of the following
			true = true
	La contra de la contra del la contra del la contra del la contra del la contra de la contra del la contra	baston	false = false The number that allows for the identification of a device, indicating its position within a series. This number is required to be part of the UDI when
	has_serial_number	boolean	included on the label in order to provide the means to track the device back to its manufacturing source or otherwise allow the history of the
			device manufacturing, packaging, labeling, distribution and use to be determined.
			Value is one of the following
			true = true
			true = true false = false
	is_combination_product	boolean	false = false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and
	is_combination_product	boolean	false = false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and produced as a single entity, packaged together as a single package; or packaged separately for the intended use together as defined under 21 CFR
	is_combination_product	boolean	false = false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and produced as a single entity; packaged together as a single package; or packaged separately for the intended use together as defined under 21 CFR 3.2(e). At least one of the products in the combination product must be a device in this case.
	is_combination_product	boolean	false a false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and produced as a single entity, packaged together as a single package; or packaged separately for the intended use together as defined under 21 CFR 3.2(e). At least one of the products in the combination product must be a device in this case. Value is one of the following
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	is_combination_product is_cdirect_marking_exempt	boolean	false a false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and produced as a single entity; packaged together as a single package; or packaged separately for the intended use together as defined under 21 CFR 3.2(e). At least one of the products in the combination product must be a device in this case. Value is one of the following
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	is_direct_marking_exempt	boolean	false = false Indicates that the product is comprised of two or more regulated products that are physically, chemically, or otherwise combined or mixed and produced as a single entity, packaged together as a single package, or packaged separately for the intended use together as defined under 21 CFR 3.2(e). At least one of the products in the combination product must be a device in this case. Value is one of the following true a true Isable = false The device is exempt from Direct Marking requirements under 21 CFR 801.45. Value is one of the following true a true Isable = false Isable = false
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AND				this data element.
AND		public version number	string	Auto assigned version number, assigned just before file generation: All existing records will have version 1 assigned.
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Service Control Contro		publish_date	date	Indicates the date the DI Record gets published and is available via Public Search.
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Storage and Handling		string	Official correspondent middle initial.
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
Storage and Handling	storage.low.value	string	Official correspondent phone number.
1			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.eprocessed and reused.
Storage and Handling	storage.low.unit	string	Official correspondent company name (if different from owner operator company name).
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
Storage and Handling	storage.special conditions	string	First line of address for owner operator.
Storage and Handling	storage.type	string	Second line of address for owner operator.
OpenFDA fields	device_class	string	A risk based classification system for all medical devices ((Federal Food, Drug, and Cosmetic Act, section 513)
			Value is one of the following
			1 = Class I (low to moderate risk): general controls
			2 = Class II (moderate to high risk): general controls and special controls
			3 = Class III (high risk): general controls and Premarket Approval (PMA)
			U = Unclassified
			N = Not classified
			F = HDE
OpenFDA fields	device name	string	This is the proprietary name, or trade name, of the cleared device.
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
OpenFDA fields	fei number	array of strings	Facility identifier assigned to facility by the FDA Office of Regulatory Affairs.
OpenFDA fields	medical_specialty_description	string	Regulation Medical Specialty is assigned based on the regulation (e.g. 21 CFR Part 888 is Orthopedic Devices) which is why Class 3 devices lack the
			"Regulation Medical Specialty" field.
			This is an .exact field. It has been indexed both as its exact string content, and also tokenized.
OpenFDA fields	regulation_number	array of strings	The classification regulation in the Code of Federal Regulations (CFR) under which the device is identified, described, and formally classified (Code
		' -	of Federal regulations Title 21, 862.00 through 892.00). The classification regulation covers various aspects of design, clinical evaluation,
			manufacturing, packaging, labeling, and postmarket surveillance of the specific medical device.