GUDID Data Fields

Labeler	Regulatory	Production	Characteristics
Labeler DUNS Number^	Publish Date	Lot/Batch Number (Y/N)	Single Use (Y/N)
Company Name*	Distribution End Date	Manufacturing Date (Y/N)	Combination Product#
Company Physical Address*^	Distribution Status*	Serial Number(Y/N)	HTC/P#
Customer Contact Phone	Premarket Exempt#	Expiration Date (Y/N)	Contains Rubber (Y/N)
Customer Contact E-Mail	Premarket Submission No.	Donation ID Number (Y/N)	Labeled 'Not made with Rubber'#
Device Identification (DI)	Supplement Number	**Packaging**	MRI Safety
Issuing Agency	FDA Listing Number^	Device Count	Size Type
Primary DI Number	FDA Product Code	Unit of Use DI Number	Size Value
Brand Name	FDA Product Code Name*	Kit#	Size Unit of Measure
Version/Model Number	GMDN Code^	Package DI Number	Size Type Text
Catalog Number	GMDN Name*	Quantity per Package	Storage & Handling Type
Device Description	GMDN Definition*	Package Contains DI Number	S&H Low Value
Second DI Issuing Agency	Prescription#	Package Type^	S&H High Value
Secondary DI Number	Over-the-Counter#	Package Discontinue Date	Storage & Handling Unit
Subject to DM, but Exempt#		Package Status*	Special Storage Conditions
DM DI Different (Y/N)#			Sterile Package (Y/N)
DM DI Number			Sterile Required (Y/N)
			Sterile Method

^Data elements not released to public *FDA GUDID System completes these fields #Checkbox