






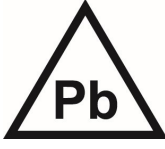





















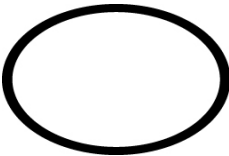
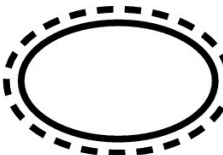
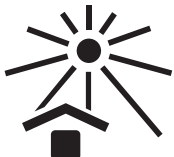
























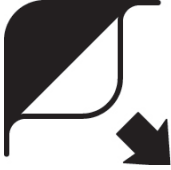



REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
NO STANDARD REFERENCE/NOT A UNIVERSALLY RECOGNIZED OR STANDARDIZED SYMBOL		
		United Kingdom Responsible Person
		Quantity
		Device does not Contain or Have Presence of Natural Latex Rubber
		Customer Feedback
		Read Accompanying Documents for Important Safety-Related Information
		Rated Type CF Patient Protection
		Contains the CMR Substance, Cobalt
		Contains the CMR Substance, Lead
		Contains Nickel
		Contains Diocetyl Bis(2-Ethylhexyl Mercaptoacetate)
ISO 15223-1 - MEDICAL DEVICES - SYMBOLS TO BE USED WITH MEDICAL DEVICE LABELS, LABELLING AND INFORMATION TO BE SUPPLIED		
5.1.1		Manufacturer

REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
5.1.2		Authorized Representative in the European Community/European Union
5.1.3		Date of Manufacture
5.1.4		Use-by-Date
5.1.5		Batch Code
5.1.6		Catalogue Number
5.1.7		Serial Number
5.1.8		Importer
5.1.9		Distributor
5.1.11		Country of Manufacture - Mexico
5.1.11		Country of Manufacture - United States of America
5.2.3		Sterilized Using Ethylene Oxide
5.2.3/5.2.11		Sterilized Using Ethylene Oxide/Single Sterile Barrier System

REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
5.2.3/5.2.14		Sterilized Using Ethylene Oxide/Single Sterile Barrier System with Protective Packaging Outside
5.2.4		Sterilized using Irradiation
5.2.4/5.2.11		Sterilized using Irradiation/Single Sterile Barrier System
5.2.6		Do Not Resterilize
5.2.7		Non-Sterile
5.2.8		Do Not Use if Package is Damaged and Consult Instructions for Use
5.2.11		Single Sterile Barrier System
5.2.14		Single Sterile Barrier System with Protective Packaging Outside
5.3.2		Keep Away from Sunlight
5.3.4		Keep Dry
5.3.6		Upper Limit of Temperature

REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
5.3.7		Temperature Limit
5.4.2		Do not Re-use
5.4.3		Consult Instructions for Use or Consult Electronic Instructions for Use
5.4.4		Caution
5.4.7		Contains a Medicinal Substance
5.4.8		Contains Biological Material of Animal Origin
5.4.10		Contains Hazardous Substances
5.6.3		Non-pyrogenic
5.7.3		Patient Identification
5.7.4		Patient Information Website
5.7.5		Healthcare Center or Doctor

REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
5.7.6		Date
5.7.7		Medical Device
5.7.10		Unique Device Identifier
BS EN 15986 - SYMBOL FOR USE IN THE LABELLING OF MEDICAL DEVICES. REQUIREMENTS FOR LABELLING OF MEDICAL DEVICES CONTAINING PHTHALATES		
A.2		Contains Phthalate: DEHP
FDA GUIDANCE USE OF SYMBOLS IN LABELING (CFR 801)		
	Rx Only	Caution: Federal law restricts this device to sale by or on the order of a (licensed healthcare practitioner).
ASTM F2503 - STANDARD PRACTICE FOR MARKING MEDICAL DEVICES AND OTHER ITEMS FOR SAFETY IN THE MAGNETIC RESONANCE ENVIRONMENT		
		MR Safe
		MR Conditional
		MR Unsafe
		MR Conditional - 3.0 Tesla
		MR Conditional - 1.5, 3.0 Tesla

REFERENCE NUMBER	SYMBOL	EXPLANATORY TEXT/TITLE OF SYMBOL
ISO 7000 - GRAPHICAL SYMBOLS FOR USE ON EQUIPMENT REGISTERED SYMBOLS		
3079		Open Here
MDD 93/42/EEC, MDR 2017/745, REGULATION (EC) 765/2008		
		CE Marking of Conformity
MEDICINES AND MEDICAL DEVICES ACT 2021		
		UK Conformity Assessed
SWISSMEDIC REGULATIONS		
		Authorized Representative in Switzerland