TANDEM™ Diabetes Care

Symbols Glossary

The following table provides a comprehensive list of symbols used on Tandem Diabetes Care™ packaging and labeling as well as the associated descriptions and indications.

Symbol	Description	Indication
	Authorized Representative in Switzerland	Indicates the authorized representative in Switzerland.
	Authorized Representative in the European Community	Indicates the authorized representative in the European Community
	Batch Code	Indicates the manufacturer's code so that the batch or lot can be identified.
	*Bluetooth*® Wireless or Enabled Technology	Indicates Bluetooth wireless or enabled technology
	Conformité Européene (CE) Mark	Indicates that the manufacturer or importer of that product affirms its compliance with the relevant EU legislation and the product may be sold anywhere in the European Economic Area (EEA).
	Catalog Number	Indicates the manufacturer's catalog number so that the medical device can be identified.
	Consult Instructions for Use or Consult Electronic Instructions for Use	Indicates the need for the user to consult the Instructions for Use.
	Country of Manufacture	Indicates the country of manufacture of products.

Symbol	Description	Indication
	Date of Manufacture	Indicates the date when the medical device was manufactured.
	Degree of Ingress Protection Provided by Enclosure	Indicates protection against the effects of temporary immersion in water.
	Distributor	Indicates the entity distributing the medical device into the locale.
	Do Not Re-Use	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.
	Do Not Use if Package is Damaged	Indicates that the device must not be used if the package holding the device is damaged, for example on packaging of medical devices.
	General Warning Sign	Indicates a general warning.
	Importer	Indicates the entity importing the medical device into the locale.
	Keep Dry	Indicates a medical device that needs to be protected from moisture.
	MN Number	Indicates configuration number created by Tandem.
	MR Unsafe	Indicates that the product should not enter the MRI scanner room.
	Manufacturer	Indicates that this symbol shall be accompanied by the name and address of the manufacturer (i.e., the person placing the medical device on the market), adjacent to the symbol.
	Medical Device	Indicates the item is a medical device.

Symbol	Description	Indication
	Model Number	Indicates the model number or type number of a product.
	Non-Ionizing Electromagnetic Radiation	Indicates generally elevated, potentially hazardous, levels of nonionizing radiation, or to indicate equipment or systems, e.g., in the medical electrical area that include RF transmitters or that intentionally apply RF electromagnetic energy for diagnosis or treatment.
	Non-Pyrogenic	Indicates that the product is non-pyrogenic.
	Prescription Only	Indicates requirement of a prescription in the United States.
	Product Subjected to a Sterilization Process	Indicates a medical device that has been subjected to a sterilization process.
	Recycle: Electronic Equipment	Indicates do not dispose of this product in unsorted municipal waste stream.
	Refer to Instruction Manual / Booklet	Indicates that the instruction manual/booklet must be read.
	Regulatory Compliance Mark (RCM)	Indicates compliance with ANZ radiocommunications requirements.
	Serial Number	Indicates the manufacturer's number so that a specific medical device can be identified.
	Single Sterile Barrier System	Indicates a single sterile barrier system.
	Sterilized by EO Treatment	Indicates a medical device that has been sterilized using ethylene oxide.
	Sterilized Using Irradiation	Indicates a medical device that has been sterilized using irradiation.

Symbol	Description	Indication
	Storage Humidity Range	Indicates the range of humidity to which the medical device can be safely exposed.
	Storage Temperature Range	Indicates the temperature limits to which the medical device can be safely exposed.
	Triman Logo	Indicates recyclable product and packaging sold in France.
	Type BF Applied Part	Indicates a type BF applied part complying with IEC 60601-1.
	U-100 Insulin	Indicates the insulin infusion pump is compatible with U-100 insulin.
	Unique Device Identifier	Indicates a carrier that contains unique device identifier information.
	Use-By Date	Indicates the date after which the medical device is not to be used.