REF

Article codes
Indicates the manufacturer's article number so that the medical device can be identified.

EC | REP

Authorized Representative in the EU

LOT

Batch code
Indicates the manufacturer's batch code so that the batch or lot can be identified.

CE

CE marking

Consult instructions for use
Consult electronic instructions for use.

Allergen IV

Contains potential Type IV chemical allergens

Date of manufacture

Do not use if package is damaged
Indicates that a medical device should not be used if the package has been damaged or opened.

For single use
Indicates a medical device that is intended for one single use only.

Importer
Indicates the entity of importing the medical device into the locale.

Keep away from sunlight
Indicates a medical device that needs protection from light sources.

Keep dry
Indicates a medical device that needs to be protected from moisture.

Legal manufacturer
Indicates the medical device manufacturer.

MD

Medical device
Indicates the item is a medical device.

NON STERILE

Non-sterile
Indicates a medical device that has not been subjected to a sterilization process.

Single sterile barrier system with protective packaging inside

Single sterile barrier system with protective packaging outside

STERILE

Sterile
Indicates a medical device that has been subjected to a sterilization process.

STERILE | R

Sterilized using irradiation
Indicates a medical device that has been sterilized using irradiation.

Temperature limit
Indicates the temperature limits to which the medical device can be safely exposed.

LATEX

Contains or presence of natural rubber latex
This product contains natural latex which may cause allergic reactions, including anaphylactic responses.

UKCA

UKCA marking

UDI

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.