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| EMERGO EUROPE Authorized | le EO symbols as identified above) |
| EMERGO EUROPE | representative in the European |
| | representative in the European |
| EC NEF 6827 AT Arnhem (Indicators the | authorized representative in the |
| | nmunity/European Union) |
| · · · · · | representative in Switzerland |
| | authorized representative in |
| 6302 Zug Switzerland) | |
| Do not re-u | se |
| (∞) | edical device that is intended for one |
| single use) | |
| Catalogue i | number |
| - | manufacturer's catalogue number so |
| | cal device can be identified) |
| LOT Batch code | |
| | manufacturer's batch code so that the in be identified) |
| Use-by date | 1 |
| | date after which the medical device is |
| not to be used | 1) |
| | if pack is damaged |
| | edical device that should not be used if |
| | as been damaged or opened) |
| Manufactu | |
| manufacturer | ed by the name and address of the) |
| Date of ma | |
| | date when the medical device was |
| manufactured | , |
| | tructions for use or consult |
| | nstructions for use |
| | need for the user to consult the |
| instructions fo | • |
| | edical device that has not been |
| STERILE | sterilization process) |
| Caution | · · |
| | aution is necessary when operating the |
| | ds operator awareness in order to avoid |
| undesirable co | onsequences) |
| Lock | |
| Unlock | |