General user information
for single use products in sterile packaging
Please read this information carefully before use.

Warnings and cautions
- This product is to be used by qualified medical personnel only
- This product is for single use only. Do not re-sterilise and/or re-use
- Do not use if the package is open or damaged
- Re-tighten closure plugs before use
- Do not use if protection caps are displaced
- Stress cracks could appear when used for a longer time together with solutions based on
  alcohol and lipids
- For administration of blood and blood components, check suitability of infusion pump
  before application
- For disconnectable connections, check tightness before use
- If equipped with an orange protection cap (vented), replace it with a closure plug (non-vented)
- Observe the medication manufacturer's instructions for use regarding incompatibility
  and the present Summary of Product Characteristics
- When applied for infusion, use fluid filter <20 μm (according to EN ISO 8536)
- Accepted good hygiene measures and working practices should be followed at all times
- The used product has to be disposed of correctly. Specially marked containers should be used if
  necessary. Local regulations must be observed

Re-use warnings
The re-use of the product may cause unpredicted health risks! A used product is a potential
Carrier of pathogens, which means a health hazard to the patient and the spreading of infectious
germs within the health care facility. Reprocessing of used products compromises safety. The
manufacturer cannot be held responsible for the sterility assurance of re-used products within
the health care facility.
<table>
<thead>
<tr>
<th><strong>Pressure</strong></th>
<th><strong>Gravity</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>Compatible for use under pressure and gravity - according to EN ISO 11135 &amp; EN ISO 8536</td>
<td>Compatible for use under gravity only - according to EN ISO 11136 &amp; EN ISO 8536</td>
</tr>
</tbody>
</table>

- **Non-pyrogenic**
  - Indicates a medical device that is non-pyrogenic.
  - EN ISO 10993

- **Contains or presence of**
  - Indicates the presence of identified process or substance.
  - ISO 7560

- **Do not use if package is damaged**
  - Indicates a medical device that should not be used if the package has been damaged or opened.
  - EN ISO 10993

- **Does not contain or no presence of**
  - Indicates the absence of identified product or substance.
  - ISO 7560/EN 8461

- **Drops per milliliter**
  - Indicates the number of drops per milliliter.
  - EN ISO 10993

- **Liquid filter with pore size**
  - Indicates an in vitro or in vivo pre-condition of a medical device that selects a filter of a particular nominal pore size.
  - EN ISO 10993

- **Pump, liquid pump**
  - Indicates a medical device with a value that allows flow in only one direction.
  - ISO 7560

- **Non-sterile**
  - Indicates a medical device that has not been sterilized and is not sterilizable.
  - STERILE

- **Batch code**
  - Indicates the manufacturer’s batch code on the device so that the trace or history can be identified.
  - EN ISO 10993

- **Date of manufacture**
  - Indicates the date when the medical device was manufactured.
  - EN ISO 10993

- **Manufacturer**
  - Indicates the medical device manufacturer as defined in list according to ISO 10993.
  - EN ISO 10993

- **End-use date**
  - Indicates the date after which the medical device is not to be used.
  - EN ISO 10993

| **Humidity limitation**
- Indicates the range of humidity to which the medical device can be safely exposed. (e.g., between 10% and 90%)
| **Keep away from sunlight**
- Indicates disposal of a medical device that needs protection from light sources.

- **Do not use if package is damaged**
  - Indicates a medical device that should not be used if the package has been damaged or opened.
  - EN ISO 10993

- **Fragile, handle with care**
  - Indicates a medical device that can be broken or damaged if not handled carefully.
  - EN ISO 10993

- **Temperature limit**
  - Indicates the temperature limits to which the medical device can be safely exposed. (e.g., between 10°C and 30°C)
  - EN ISO 10993

- **Cautions**
  - Indicates the need for the user to consult the instructions for use for relevant cautions.
  - EN ISO 10993

- **Consult instructions for use**
  - Indicates the need for the user to consult the instructions for use.
  - EN ISO 10993

- **Language information**
- **EN** = English, **DA** = Danish, **DE** = German, **FR** = French, **IT** = Italian, **NL** = Dutch, **NO** = Norwegian, **ES** = Spanish

- **Authorized representative**
  - Indicates the authorized representative in the European Community.
  - EN ISO 10993

- **Filling/storage volume**
  - Indicates the volume to be filled or stored with the device.
  - EN ISO 10993

- **Airline**
  - Indicates the ability to use the medical device with a special airline ticket.