<table>
<thead>
<tr>
<th>Symbol</th>
<th>Title of Symbol</th>
<th>Description of Symbol</th>
</tr>
</thead>
<tbody>
<tr>
<td>![Manufacturer](ISO 15223-1, 5.1.1)</td>
<td>Manufacturer</td>
<td>Indicates the medical device manufacturer.</td>
</tr>
<tr>
<td>![Date of manufacture](ISO 15223-1, 5.1.3)</td>
<td>Date of manufacture</td>
<td>Indicates the date when the medical device was manufactured.</td>
</tr>
<tr>
<td>![Use-by date](ISO 15223-1, 5.1.4)</td>
<td>Use-by date</td>
<td>Indicates the date after which the medical device is not to be used.</td>
</tr>
<tr>
<td>![Batch code](ISO 15223-1, 5.1.5)</td>
<td>Batch code</td>
<td>Indicates the manufacturer’s batch code so that the batch or lot can be identified.</td>
</tr>
<tr>
<td>![Catalog number](ISO 15223-1, 5.1.6)</td>
<td>Catalog number</td>
<td>Indicates the manufacturer’s catalogue number so that the medical device can be identified.</td>
</tr>
<tr>
<td>![Do not use if package is damaged](ISO 15223-1, 5.2.8)</td>
<td>Do not use if package is damaged</td>
<td>Indicates a medical device that should not be used if the package has been damaged or opened.</td>
</tr>
<tr>
<td>![Keep away from sunlight](ISO 15223-1, 5.3.2)</td>
<td>Keep away from sunlight</td>
<td>Indicates a medical device that needs protection from light sources.</td>
</tr>
<tr>
<td>![Do not reuse](ISO 15223-1, 5.4.2)</td>
<td>Do not reuse</td>
<td>Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.</td>
</tr>
<tr>
<td>![Consult instructions for use](ISO 15223-1, 5.4.3)</td>
<td>Consult instructions for use</td>
<td>Indicates the need for the user to consult the instructions for use.</td>
</tr>
<tr>
<td>![Not made with natural rubber latex](ISO 15223-1, 5.4.5 and Annex B)</td>
<td>Not made with natural rubber latex</td>
<td>Indicates that natural rubber latex was not used in the manufacturing of the product, its container, or its packaging.</td>
</tr>
<tr>
<td>![Quantity](IEC 60878, 2794)</td>
<td>Quantity</td>
<td>To indicate the number of pieces in the package.</td>
</tr>
</tbody>
</table>
**INTENDED USE**
Protective cover or sheath placed over equipment. The cover helps prevent transfer of microorganisms, body fluids and particulate material to the patient and healthcare worker. The cover also provides a means for maintenance of a sterile field (sterile covers only). CIVCO Poly Covers are furnished sterile and non-sterile; single use patient / procedure, disposable.

**PATIENT POPULATION**
Cord Covers and System Drapes are not intended to be patient-contacting and do not carry any patient population considerations.

**PERFORMANCE CHARACTERISTICS**
Cord Covers and System Drapes protect integrity of equipment and reduce risk of cross-contamination.

**NOTE:** For a summary of clinical benefits for this product, visit www.CIVCO.com.

**CAUTION**
Federal (United States) law restricts this device to sale by or on the order of a physician.

**WARNING**
- Before use, you should be trained in ultrasonography. For instructions on the use of your transducer, see your system’s user guide.
- Always use a drape to protect equipment and clinicians from cross-contamination.
- Users of this product have an obligation and responsibility to provide the highest degree of infection control to patients, co-workers and themselves. To avoid cross-contamination, follow infection control policies established by your facility.
- Product is intended for single-use only.
- Do not use if integrity of packaging is violated.
- Do not use if expiration date has passed.
- Do not reuse, reprocess or resterilize single-use device. Reuse, reprocessing or resterilization may create a risk of contamination of the device, cause patient infection or cross-infection.
- If the product malfunctions during use or is no longer able to achieve its intended use, stop using the product and call CIVCO.
- Report serious incidents related to the product to CIVCO or appropriate regulatory authorities.

**NOTE:** Product is not made with natural rubber latex.

**COVERING THE CORD**
1. Hold cord cover so entire length does not unfold, possibly causing contamination.
2. Insert distal end of cord into cover and proceed toward proximal connection, sliding cover to desired position.
4. Inspect cover to ensure there are no holes or tears.

**COVERING THE SYSTEM**
1. Hold system drape so entire length does not unfold, possibly causing contamination.
2. Remove backing from adhesive strips and secure drape to control panel.
3. Inspect drape to ensure there are no holes or tears.

**STORAGE CONDITIONS**
- Avoid storing product in areas of temperature extremes or in direct sunlight.
- Store in a cool, dry place.

**DISPOSAL**
**WARNING**
- Dispose of single-use components as infectious waste.

**NOTE:** For questions or to order additional CIVCO products, please call +1 319-248-6757 or 1-800-445-6741 or visit www.CIVCO.com.