# **Lindor®** lady Pads

**General Product Description/Intended Purpose**

Lindor® lady Pads are non-sterile anatomically shaped pads for the management of very light to moderate bladder weakness as well as fecal incontinence. The products can be used in home care environment and professional healthcare facilities and the application can be performed by lay person and/or health care professional. The pads are for single use only and should be regularly changed as needed. Lindor® lady Pad products account for Class I medical devices.

**Application/Indication**

Lindor® lady Pad products are for absorption of urine and fecals* due to incontinence.
**only Lindor® lady Pads 5 – 6 drops.*

**Catalogue Numbers**

| Article | Reference Number* | Pack Count |
|---|---|---|
| Lindor® lady Pad mini 2 drops | 167 050 | 12 pcs |
| Lindor® lady Pad normal 3 drops | 167 051 | 12 pcs |
| Lindor® lady Pad extra 4 drops | 167 052 | 12 pcs |
| Lindor® lady Pad maxi 5 drops | 167 053 | 14 pcs |
| Lindor® lady Pad maxi night 6 drops | 167 054 | 14 pcs |
| | | |
| Lindor® lady Pad mini 2 drops | 167 060 | 1 pc |
| Lindor® lady Pad normal 3 drops | 167 061 | 1 pc |
| Lindor® lady Pad extra 4 drops | 167 062 | 1 pc |
| Lindor® lady Pad maxi 5 drops | 167 063 | 1 pc |
| Lindor® lady Pad maxi night 6 drops | 167 048 | 1 pc |

** may not be available in all countries*

**Single-use device**

Reusing a single-use medical device is dangerous. Reprocessing devices, in order to reuse them, may seriously damage their integrity and their performance. Information available on request. Reuse of the product could lead to microbiological contamination and/or mild infections of the skin.

**Product disposal**

In order to minimise the risk of potential infection hazards, or environmental pollution, disposable components of Lindor® lady Pads should follow disposal procedures according to applicable and local laws, rules, regulations and infection prevention standards.

**Incident reporting**

For a patient/user/third party in the European Union and in countries with identical regulatory regime (Regulation (EU) 2017/745 on Medical Devices); if, during the use of this device or as a result of its use, a serious incident has

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occurred, please report it to the manufacturer and/or its authorised representative and to your national authority.

### Product Performance Characteristics

| Characteristics | mini 2 drops | normal 3 drops | extra 4 drops | maxi 5 drops | maxi night 6 drops |
|---|---|---|---|---|---|
| Total weight [g] | 9.4 | 14.5 | 20.9 | 33.7 | 38.1 |
| Absorbent core weight [g] | 7.3 | 11.0 | 15.8 | 28.4 | 32.8 |

≤ ± 10% tolerance

### Material Characteristics

| Component | Material |
|---|---|
| Topsheet | Polypropylene and Polyethylene – nonwoven, hydrophilic, white |
| Absorbent core: | • Kraft pulp*<br>• Superabsorbent polymer<br>• pH buffer agent<br>• Nonwoven, hydrophilic, white |
| Backsheet | Breathable textile backsheet, white |
| Elastics | Elastic film for leg cuff (except 2 Drops) |
| Adhesives | Hotmelts |
| Backing | 2D, 3D & 4D: Polyethylene film for single wrapping with release tape & silicone paper<br>5D & 6D: silicone paper |

*bleached elementary chlorine free

### Product Characteristics

| Dimensions in mm | mini 2 drops | normal 3 drops | extra 4 drops | maxi 5 drops | maxi night 6 drops |
|---|---|---|---|---|---|
| Total length | 220 | 265 | 335 | 400 | 400 |
| Total width | 98 | 112 | 132 | 162 | 162 |
| Length absorbent core | 200 | 233 | 305 | 365 | 365 |
| Width absorbent core | 78 | 90 | 112 | 138 | 138 |

≤ ± 10% tolerance

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**Labelling**

Lot-No. with 9-Digit Code
e.g.: 500102822

| | | | | | |
|---|---|---|---|---|---|
| LOT | 5 | 001 | 02 | 82 | 2 |
| | year 202X | internal key | internal key | internal key | cumulated check sum |

Date of manufacture

| | | | | |
|---|---|---|---|---|
| e.g.: | 2025 | 06 | 02 | |
| | year | month | day | |

Use-by Date

| | | | | |
|---|---|---|---|---|
| e.g.: | 2030 | 06 | 02 | |
| | year | month | day | |

Shelf Life: 5 years (Except for P1: 4 years)

Medical Device

MD

Unique Device Identification (UDI)

UDI

Keep dry

Do not reuse

CE-marking

CE

**Latest Date of Revision:** 2025-06-13