


L&R successfully completed MDR conformity! Delivery capability and security of supply ensured



 MDR
Successfully completed!

The MDR (Medical Device Regulation) is the EU Regulation concerning medical devices (EU 2017/745). It regulates the placing on the market of medical devices throughout Europe and defines the requirements for the conformity assessment of medical devices.

The MDR and the standards defined therein are intended to improve

- safety,
- traceability and
- transparency relating to medical devices

for patients and healthcare providers.

- As in the past, L&R ensures that only compliant products will reach the market.
- L&R has now also achieved compliance under the terms of the MDR.
- The company can continue to reliably supply its products to healthcare systems and to its customers.

L&R successfully completed MDR conformity! Delivery capability and security of supply ensured

You're in safe hands with L&R!

Safe, high-quality products

Dependable supply

Transparent communication
about product availability

- On 24 May 2024, L&R was awarded MDR certificates for its Class Is, IIa and IIb products and its procedure packs (e.g. Kitpack and CNP sets) by the notified body TÜV Süd.
- Our products are MDR-compliant – and as safe and effective as ever.
- Our internal quality management system and all of our regulatory processes meet the MDR requirements and have been MDR-certified since 2019.

- Existing L&R products will continue to be legally marketed within the statutory transition period until 2027-2028.



Successfully
completed!

What is the UDI?

The Unique Device Identifier (UDI) is a unique numeric or alphanumeric code for a medical device.

This enables clear and unambiguous identification and helps with its traceability.

The UDI is made up of the following components:

- A unique device identifier (UDI-DI)
- A production identifier (UDI-PI)

