EU Certificate

for the assessment of the quality management system

according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter I+III

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the manufacturer

Hocoma AG

Single Registration Number (SRN): CH-MF-000014672
Industriestraße 4, 8604 Volketswil, Switzerland

Name, address of the authorized representative:
Emergo Europe B.V., Prinsessegracht 20, 2514 The Hague, Netherlands

applies a quality management system according to Annex IX Chapter I+III of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. This certificate is based on the assessments listed in CNo50682-00.

EU Certificate no.: 50682-60-00

Certificate valid from: 2023-01-28
Certificate valid to: 2027-09-24

DEKRA Certification GmbH • D-70565 Stuttgart, Handwerkstraße 15

Karin Leicht
DEKRA Certification GmbH, Stuttgart, 2023-01-28
Notified Body ID number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-092

Annex to the EU Certificate no. 50682-60-00

valid from 2023-01-28 to 2027-09-24

Revision status of the annex: 0 dated 2023-01-28

Following devices/device categories are included in this certificate:

Class IIa

- MDA 0313
  - **Energetically driven gait orthoses**
    - LokomatPro; Basic UDI-DI: 764017466L6U8
    - LokomatNanos; Basic UDI-DI: 764017466LNVN
  - **Energetically driven devices for exercise therapy**
    - ErigoPro, ErigoBasic; Basic UDI-DI: 764017466ERV9
    - ArmeoPower; Basic UDI-DI: 764017466APUR
    - Andago V2.0; Basic UDI-DI: 764017466ANUM

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