# DECLARATION OF CONFORMITY

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|---|---|
| Name and address of the firm | **Hocoma AG**<br>Industriestrasse 4<br>8604 Volketswil<br>Switzerland<br>SRN: CH-MF-000014672 |
| EU Authorized Representative | Emergo Europe B.V.<br>Prinsessegracht 20<br>2514 AP The Hague<br>The Netherlands<br>Tel: (+31) (0)70 345-8570<br>Email: EmergoEurope@ul.com<br>SRN: NL-AR-000000116 |

We declare under our sole responsibility that

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| the medical device | **ArmeoSenso**<br>Basic UDI-DI: 764017466AXV9 |
| Intended purpose | ArmeoSenso is a medical device intended for patients with impaired function of the upper extremity. This device allows the patient to train specific exercises to increase, for example, muscle strength and range of motion in different joints, with the overall goal of improving functional abilities. |
| of class | **I**<br>according to rule 13 of annex VIII of regulation (EU) 2017/745 |

meets all the provisions of regulation (EU) 2017/745, which apply to it.

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| Conformity assessment procedure | Annex II and Annex III of regulation (EU) 2017/745 |
| Common Specifications | N/A |

Volketswil, 13.01.2022

Place, date

p.p.

**Petrus Broeksteeg**
**Head of Regulatory Affairs / PRRC**
Name and function
according to Article 15 [3] of regulation (EU) 2017/745

**Bernd Henningsen**
**Chief Specialist Regulatory Affairs / PRRC**
Name and function
according to Article 15 [3] of regulation (EU) 2017/745