Declaration of Conformity

EMDN CND Code :Y060406 Issue Date: Mar. 02, 2022

EMDN CND Description: Abdominal hernia supports Valid Date: Dec. 31, 2025

This declaration of conformity is issued under the sole responsibility of the manufacturer:
OPPO Medical Inc., address at: Seattle City Center, 1420 Fifth Ave., Ste. 220080, Seattle, WA98101, USA
Single registration number (SRN): US-MF-000005202

Authorised Representative: MT Promedt Consulting GmbH, address at: Altenhofstrasse 80,
D-66386 St. Ingbert Germany Single registration number (SRN): DE-AR-000000085

The basic UDI-DI: 8402706CL110KU

We herewith declare that the products as referred to in Attachment I are in conformity with the requirements set out in the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL.

Conformity Assessment Procedure: MDR (EU) 2017/745 Art 52. (7), ANNEX II (Technical documentation) & ANNEX III (Technical documentation on post-market surveillance)

Risk Classification of the Product: Class I Rule: MDR (EU) 2017/745, Annex VIII, Section 4, 4.1, Rule 1"All non-invasive devices are classified as class I."

Applied Standards: EN ISO13485:2016, EN1041:2008, EN ISO14971:2012, EN ISO 15223-1:2016, EN ISO 10993-1:2009, EN ISO10993-5:2009, EN ISO 22523:2006,EN 62366:2008, ISO 10993-10:2021, ISO 9001:2015

Intended purpose: The devices help resolve inguinal hernias. They target the specific location of a hernia and keep protruding tissue in place to relieve discomfort.

Mar. 02, 2022/ Seattle
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Date/Place

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Jackson Chiang
President

Attachment 1

EMDN CND Code :Y060406

EMDN CND Description: Abdominal hernia supports

Reference number	Product Trade name
2049	Hernia Truss Double-Sided
2149	Hernia Truss Single-Sided
2249	Hernia Truss with Removable Pad