Dublin, 01 January 2024

DECLARATION OF CONFORMITY
Regulation (EU) 2017/745 Of the European Parliament and of The Council on Medical Devices (MDR)

We,

LM Global Design Ltd.

with registered place of business in:
Suite 123, The Capel Building,
Mary's Abbey, Dublin 7 D07 VY68,
Ireland
SRN: N/A

as a legal manufacturer hereby declare under our sole responsibility, that the medical device listed below with related accessories, meets the general safety and performance requirements of Annex I and that it is in conformity with the Regulation (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on medical devices (MDR).

The product specified below is a "technical aid for the disabled", classified as Class I, medical device. The classification is based on the requirements of annex VII & Rule I, of the MDR.
Conformity assessment was carried out according to Art. 52, pt. 7 and Annex II of the MDR.
The CE mark has been affixed on the product according to Annex V of the MDR.

TRADE NAME: **COMMODE CHAIR LM** - ITEM NUMBER: **KING-CMD-00**
TRADE NAME: **COMMODE CHAIR W/ARMRESTS LM** - ITEM NUMBER: **KING-CMDA-00**
TRADE NAME: **COMMODE CHAIR W/BACKREST LM** - ITEM NUMBER: **KING-CMDB-00**
TRADE NAME: **COMMODE CHAIR W/BACKREST & ARMRESTS LM** - ITEM NUMBER: **KING-CMDBA-00**

BASIC UDI-DI CODE: **539153269LMCMDFC**
ACCESSORY LIST: **Soft Cushions (KING-CCMD2/4-00/KING-CCMDH4-00, KING-CBRX-00), Bucket and Lid (KING-BC-00)**

Following harmonized norms and/or common specifications were used for conformity evaluation: EN ISO 14971:2019+A11:2021, PN-EN 1041:2010 + A1 2013, EN ISO 12182:2012, EN ISO 21856:2022, EN ISO 14001:2015, EN ISO 13485:2016+A11:2021.

Luca Mammi,
Managing Director
LM Global Design Limited

CE

Ref. Declaration of Conformity file Commode Chairs LM Rev 00 as of 01 2024