



# EU Declaration of Conformity

Regarding Medical Device Regulation (EU) 2017/74



Escape Mobility hereby declares that the product named below has been designed, manufactured, tested and found to comply with the relevant sections of the reference standards. The products comply with all essential requirements of the Directives. The product carries the CE Mark

**Manufacturer**

Name: Escape Mobility Company LTD

Address: 85 Great Portland Street, First Floor, London, United Kingdom, W1W 7LT

**EU Representative**

Name: Amstermed B.V.

Address: T.a.v. de heer R. Rajemahadik, Sapporodreef 20, 2152 KI Nieuw-Vennep

SRN: NL-AR-000001971

**Competent Authority**

Name: Ministerie van Volksgezondheid, Welzijn en Sport

Address: P.O. Box, 20350, The Hague

Country: Netherlands

SRN: NL-CA-058

**Conformity Assessment Procedure**

Annex VII

**Harmonised Standards referenced or applied**

EN ISO 14971:2019

EN ISO 15223-1:2016

ISO 10993-1:2018

EN ISO 10993-5:2009

EN ISO 10993-10:2013

ISO 20417:2021

IEC 62365-1:2015

ISO 7176-28:2012

EN 61000-6-2:2005

EN 61000-6-4:2007:A1+2011

ISO/TR 20416:2020

**Product Information**

Name: Emergency Evacuation Chair

UDI-ID

Models

05061075900041 Escape Mobility VOLT

05061075900294 Escape Mobility VOLT NARROW

05061075900089 Escape Mobility VOLT-XS

05061075900317 EM PRO-EMS-H

05061075900324 EM PRO-EMS-P

EMDN: V080502

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**Classification**                      Class I in accordance with Rule 1, Annex VIII Regulation (EU) 2017/745

**Signature**



Name: Mike Vermin  
 Position: Managing Director  
 Date: 08/10/2024

**Place of Signature**                      Heerlen, Netherlands

This declaration of conformity is issued under the sole responsibility of Escape Mobility. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. All supporting documentation is retained at the premises of the manufacturer.