

DECLARATION OF CONFORMITY

iWALKFree, Inc. located at 130 N. Marina Drive, Long Beach, California, 90803, USA hereby declares that the product listed below:

PART NUMBER	DESCRIPTION	USA FDA CLASSIFICATION
HFC30303	IWALK3.0 Hands Free Crutch	CLASS I (non-sterile)


meets the standards listed below, and as a company has implemented and maintains a full quality assurance system which applies to the product at every stage from design, manufacture, to final controls, and meets the provisions of the MDR Regulation (EU) 2017/745, and the applied harmonized standards:

- EN ISO 13485: Quality Management System
- EN ISO 11334-1:2007: Assistive Products for Walking Manipulated by One Arm
- EN ISO 12182:2012: Assistive Products for Persons with Disability
- EN ISO 13287:2004 / ASTM TM144:1999: Safety, Protective & Occupational Footwear
- EN ISO 4649:2010 / ASTM D5963:2015: Rubber, Determination of Abrasion Resistance

MDR Conformity Assessment Route: Annex II & Annex III

Authorized Representative for EU:
 CEpartner4U (SRN: NL-AR-000.000.111)
 Esdoorniaan 13
 3951 DB Maarn
 The Netherlands

EU Registration (Farmatec | CIBG):
 iWALKFree, Inc. – iWALK Hands Free Crutch (NL-CA-002.2017.41653)

Location: Long Beach, California, USA	Name / Title: Brad Hunter, President
Date: May 17 th , 2021	Signature: 



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