

Declaration of Conformity

Manufacturers Name: Gordon Ellis & Co.

Manufacturers Address: Trent Lane
Castle Donington
Derby
DE74 2AT
United Kingdom

GMDN Code: 37162

Authorised Representative Name: UNITED KINGDOM
Gordon Ellis & Co.

EUROPE
Advena Ltd

Authorised Representative Address: Trent Lane
Castle Donington
Derby
DE74 2AT
United Kingdom

Tower Business Centre, 2nd Floor
Tower Street
Swatar
BKR 4013
Malta

SRN (Single Registration Number): GB-MF-000012829

MT-AR-000000234

Basic UDI-DI: 5016181BAMBERRYTRANSFERV3

Name of the Device(s): Bamberry Transfer board

Product Code(s): 55932
55933
55935

Intended Purpose : The device is intended to help with assisted or unassisted transfer of a person with limited strength or mobility from one position to another i.e. from a wheel chair to a chair, bed, or car, and is used as an aid to daily living.

Classification: Class I

Conformity Assessment Route: Gordon Ellis & Co. uses the requirements for the CE-labelling of their products according the Regulations MDR 2017/745 - Article 19 Annex II and Annex III.

Gordon Ellis & Co. test their products in accordance with BS EN 21856:2022 - Assistive Products for Persons with Disability - General Requirements and Test Methods and Risk Assess to BS EN ISO 14971:2019 - Application of Risk Management to Medical Devices.

Update Information: 20/11/2020 1st Issue for MDR 2017/745
01/08/2023 The device name has changed from 'Panda' to 'Bamberry'.
16/11/2023 Packaging update, board construction method updated.

This declaration of conformity is issued under the sole responsibility of Gordon Ellis & Co. 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.

Signature:

Place and Date (DD/MM/YYYY) of Issue:



Location: Gordon Ellis & Co. Date: 28/11/2023

Name:

Function:

Lee Rice

Product Development Manager

Signing of this declaration of conformity authorises the manufacturer to affix the CE Mark to the product in accordance with the above directive.

