

Declaration of Conformity



Manufacturers Name: Gordon Ellis & Co.

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

GMDN Code: 65280

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: 5016181BAMBERRYCUBESR2

Name of the Device(s): Bamberry Cubes

Product Code(s): 55913, 55914

Intended Purpose : The device is intended to raise the height of furniture such as an Armchair or a bed, to assist those with limited mobility as it reduces the movement required when sitting or standing 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 12182:2012 - 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: 17/011/2020 1st Issue for MDR 2017/745
01/08/2023 The device name has changed from 'Panda' to 'Bamberry'.

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:

A handwritten signature in black ink, appearing to be 'Ashley Kidd', written over a horizontal line.

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

Location: Gordon Ellis & Co. Date: 01/08/2023

Name:

Ashley Kidd

Function:

QHSE Manager

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

