Declaration of Conformity

Manufacturers Name:	Gordon Ellis & Co.		-		
Manufacturers Address:	Trent Lane Castle Donington Derby DE74 2AT United Kingdom				Z
GMDN Code:	42800				GORDON
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:	5016181BATHSHORTENER97				
Name of the Device(s):	Ashby Bath Shortener	r			
Product Code(s):	6290				
Intended Purpose :	The device is intended to be used to reduce the internal length of a bath allowing a shorter person to rest their feet against a solid surface giving confidence with added safety and comfort 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:	01/12/2020 1st issue for MDR 2017/745 04/02/2022 The shape on the rear of the product has been adjusted.				
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 retaine	d at the premises of the	e manufacturer.			
Signature:	Place and Date (DD/MM/YYYY) of Issue:				
Œ	L	ocation:	Gordon Ellis & Co.	Date:	04/02/2022
Name:	F	unction:			
Ashley Kidd	Ashley Kidd 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.					