



UK DECLARATION OF CONFORMITY

In accordance with UK Government Guidance

Manufacturer	<p>Name Address Country Phone no. Web address</p>	UK Responsible Person	<p>Etac Ltd Unit 60, Hartlebury Trading Estate, Hartlebury, Kidderminster, DY10 4JD Email enquiries.uk@etac.com Web www.etac.com/uk Phone +44 (0)121 561 2222</p>
Statement	<p>This declaration of conformity is issued under the sole responsibility of the manufacturer. The device(s) covered by this declaration of conformity is/are in conformity with UK MDR 2002 (Medical Device Regulations 2002)</p>		
Basic UDI-DI	08401174SEATINGMWVCV		
Device description	.Attendant propelled wheelchair		
GMDN	41630		
Intended purpose	<p>The Convaid Coaster is an attendant-propelled device. Its intended function and use is to assist a disabled person in performing an activity that the person would otherwise find difficult to do or be unable to do such as said person from one place to another while sitting. The chair is intended to always be under the control and supervision of an attendant, and the occupant should never be left unattended.</p>		
Device name(s)	Coaster		
Parts covered in this declaration	See Annex A.		
Risk class of the device	Class I, rule 1		
Harmonized/Established Standards	See Annex B		
Place	Torrance, California		
Date of issue	February 3, 2026		
Name and function	Mark Murphy, QA/RA Manager, PRRC		

Mark Murphy

Signature, on behalf of Convaid Products, LLC.

Appendix A

Catalogue Number	Device Name	Variant	Device Description	UDI-DI
800127	Coaster	12	Attendant propelled wheelchair UK Preconfigured	00840117408586
800147	Coaster	14	Attendant propelled wheelchair UK Preconfigured	00840117408593
800167	Coaster	16	Attendant propelled wheelchair UK Preconfigured	00840117408609
800187	Coaster	18	Attendant propelled wheelchair UK Preconfigured	00840117408616

Appendix B

EN ISO 10993-1:2009 Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process

EN 12183:2022 Manual wheelchairs – Requirements and test methods

EN ISO 13485:2016 Medical devices – Quality management systems – Requirements for regulatory purposes

EN ISO 14971:2019/A11:2021 Medical devices – Application of risk management to medical devices