Abbott
Diabetes Care

UK Declaration of Conformity	
Product Name	FreeStyle Libre Pro iQ Continuous Glucose Monitoring System (Reader Kit)
Manufacturer	Abbott Diabetes Care Ltd. Range Road, Witney, Oxfordshire, OX29 0YL, UK
First Date of UKCA Marking	See date of signing
GMDN	44611

The Medical Device Regulations 2002 (No.618)	
Classification	IIa
Rule	Rule 10
Conformity Route per Part II (MDR 2002)	Annex II, Section 3
UK Approved Body	BSI Assurance UK Ltd.
UK Approved Body Identification No.	0086
UK Approved Body Certificate(s) No.	UKCA 776995

The Radio Equipment Regulations 2017 (No 1206)	
Designated Standards	EN 300 330 V2.1.1
Conformity Route	Module A

The Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations 2012 (No 3032)	
Designated Standards	EN IEC 63000:2018

Abbott Diabetes Care hereby declare that the above-mentioned product is in conformity with the following legislative acts:

- The Medical Device Regulation 2002 (No.618)
- The Radio Equipment Regulations 2017 (No 1206)
- The Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations 2012 (No 3032)

All supporting documentation is retained under the control of Abbott Diabetes Care Ltd. and made available for review upon request. This declaration is issued under the sole responsibility of, and signed on behalf of the legal manufacturer, Abbott Diabetes Care Ltd.

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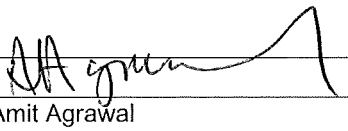
Signature	
Name	Amit Agrawal
Position	Director, Regulatory Affairs, EMEAP, ADC Ltd
Place	Witney, UK
Date	02-OCT-2023

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