DIRECTA

EU/UK Declaration of Conformity

Manufacturer	**DIRECTA AB** Finvids väg 8, SE-194 47 Upplands Väsby, Sweden
SRN	SE-MF-000002500
Basic UDI-ID	731023999520AQ
Product/trade name	**DryDent® Sublingual** **DryDent® Parotid**
Product code/REF	416067, 416068, 416069, 416072, 416073
Intended purpose	To absorb saliva and moisture during dental treatments. Used by dentist.
Risk Class	Class I, rule 5
Common specification (CS)	N/A
Notified body	N/A
Conformity Assessment Procedure	Article 52, paragraph 7 Annex II and III
Statement	We, Directa AB, confirm that this EU declaration of conformity is issued under our sole responsibility and that the device covered by this declaration is in conformity with Regulation (EU) 2017/745 and UK MDR 2002 on medical devices.

Upplands Väsby, 2022-12-19

Henric Karsk
CEO

First version of EU Declaration of Conformity (MDR) issued year 2021.
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