

AF-QD-29	Revision	004
EC Declaration of Conformity	Effective From	17/10/22



Manufacturer's Name;	Daytot Ltd.
Manufacturer's Address;	97 Saintfield Road Belfast BT8 7HN Northern Ireland
SRN (Single Registration Number);	XI-MF-000025474
Basic UDI-DI;	5065012224207QJ
Name of Device(s);	Joey
Product Code(s);	207-600 (UDI-DI 05065012224054) 207-601 (UDI-DI 05065012224009) 207-602 (UDI-DI 05065012224016) 207-604 (UDI-DI 05065012224030) 207-605 (UDI-DI 05065012224047) 207-606 (UDI-DI 05065012224061)
Classification;	Class 1, as per Annex VIII of the regulation
Conformity Assessment Process;	The manufacturer uses the following procedures for the CE-labeling of their products according to the above regulation. EC self conformity assessment for Class 1 devices in accordance with the Regulation.



This declaration of conformity is issued under the sole responsibility of the manufacturer states above. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by a Quality Management System. All supporting documentation is retained at the premises of the manufacturer.

Name & Title;	Signature;	Date & Place;
James Leckey CEO		17/10/2022 97 Saintfield Road, Belfast BT8 7HN