daytot

Manufacturer's Name;	Daytot Ltd.
Manufacturer's Address;	29 Quarterlands Road Lisburn BT27 5TN Northern Ireland
SRN (Single Registration Number);	XI-MF-000025474
Basic UDI-DI;	5065012224207QJ
Name of Device(s);	Joey
Intended Use	Joey is an early intervention crawler for infants at risk of motor delay.
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-labelling of their products according to the above regulation. EC self conformity assessment for Class 1 devices in accordance with the Regulation.

CE

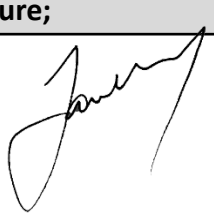
This declaration of conformity is issued under the sole responsibility of the manufacturer states above. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by a Quality Management System.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

Name & Title;	**Signature;**	**Date & Place;**
James Leckey CEO		24/05/2022 97 Saintfield Road, Belfast BT8 7HN

Manufacturer's Name;	Daytot Ltd.
Manufacturer's Address;	29 Quarterlands Road Lisburn BT27 5TN Northern Ireland
SRN (Single Registration Number);	XI-MF-000025474
Basic UDI-DI;	5065012224208QL
Name of Device(s);	Neo+
Intended Use	Neo+ is a multipurpose positioning support for babies aged 0-18 months showing atypical posturing due to surgery, disability or delay.
Product Code(s);	208-600 (UDI-DI 05065012224122)
Classification;	Class 1, as per Annex VIII of the regulation
Conformity Assessment Process;	The manufacturer uses the following procedures for the CE-labelling of their products according to the above regulation. EC self conformity assessment for Class 1 devices in accordance with the Regulation.

CE

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		24/05/2023 97 Saintfield Road, Belfast BT8 7HN