CE EC Declaration of Conformity

We, **Enable Access Ltd**., as manufacturer, declare under our sole responsibility that the products listed below meet the provisions of **Regulation (**EU**) 2017/745**. The listed products are **Class I medical devices** and conformity with the requirements of the Regulation have been assessed following the procedure outlined in **Article 52 (section 7)** of the Regulation. The products meet the General Safety & Performance Requirements listed in Annex I of the Regulation and, in addition, meet the requirements of the following standard:

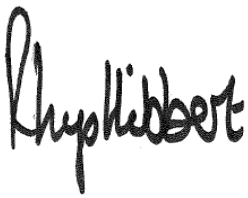
- BS6109: Part 2:1989, Appendix A.3

Product / code:

Rampcentre – Rollout-Trackway: RTP20, RT15, RT25, RT50

Technical documentation for the above products is kept by the manufacturer and can be made available by the Authorised Representative in the EU:
Enable Access Global Ltd., 27 Phibsboro Place, Dublin 7, D07 V20, Republic of Ireland.

Signed:



Rhys Hibbert – Compliance Officer

Enable Access Ltd. Leighton Buzzard
11th April 2025