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:

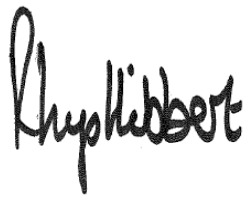
- BS6109: Part 2:1989, Appendix A.3

Product / code:

Permaramp - Original: PO9, PO12, PO15, PO18, PO21, PO24
Adjust: PA9, PA12, PA15, PA18, PA21, PA24
Superwide: All codes included
Quickrails: All codes included
High Rise: All codes included

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