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:

Aerolight-Xtra: AX6, AX9, AX12, AX15, AX18, AX21 & AX24
Aerolight-Classic: AC6, AC9, AC12 & AC15
Aerolight-Lifestyle: AL18, AL21, AL24, AL27 & AL28
Aerolight-Broadfold: AB15, AB18, AB21 & AB24
Aerolight-Travel: AXT
Aerolight-Max: AXM06, AXM09, AXM12, AXM15, AXM18, AXM21 & AXM24

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