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 - 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

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:

Hannah Ker – Compliance Manager

Enable Access Ltd. Hatfield
13th May 2021

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