



# EU Declaration of Conformity

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|---|---|
| <b>Manufacturer</b>                     | R82 A/S<br>Parallelvej 3<br>DK-8751 Gedved<br>Denmark<br>Phone: +45 7965 5888<br><a href="http://www.etac.com">www.etac.com</a>   |
| <b>SRN</b>                              | DK-MF-000000772   |
| <b>Statement</b>                        | This declaration of conformity is issued under the sole responsibility of the manufacturer.<br>The device(s) covered by this declaration of conformity is/are in conformity with EU Regulation 2017/745 on medical devices. |
| <b>Basic UDI-DI</b>                     | 57072920536033A   |
| <b>EMDN</b>                             | Y050202   |
| <b>Intended Purpose</b>                 | Standing frame for persons with disabilities  |
| <b>Device name</b>                      | Rabbit Up   |
| <b>Risk class of the device</b>         | Class I, Rule 1   |
| <b>Harmonized/Established Standards</b> | Separate list available upon request  |
| <b>Place</b>                            | Gedved, Denmark   |
| <b>Date of issue</b>                    | 2024-12-17  |
| <b>Name and function</b>                | Claus Riis – QA & RA Manager  |

A handwritten signature in blue ink, appearing to read 'C. Riis', is written above a horizontal line.

Signature, on behalf of manufacturer