

## **EU Declaration of Conformity**

2625 Patton Road Roseville, MN 55113 651-294-2200 www.ablenetinc.com

This declaration of conformity is issued under the sole responsibility of the manufacturer. The device covered by the present declaration is in conformity with all regulations below, including compliance with related Essential Requirements.

### Object of the declaration:

| Product Name:  | Buddy Button - Green |
|--|----------------------|
| Product Model Designator:  | Buddy Button - Green |
| Product Part Number(s):  | 57200                |
| Basic UDI-DI   | 00186648000654       |
| <required for="" mdr=""></required>                                  |                      |
| Control Indicator:   | 2020W20 thru 2025W20 |
| Global Medical Device<br>Nomenclature Code (GMDN)<br>and Description |                      |
| Product  |                      |
| Options/Accessories:   | N/A                  |



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The object of the declaration described above is in conformity with the following regulations:

| EU Regulation                       | Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices  |
|-------------------------------------|--|
| Device Classification:              | Class I based on Annex VIII and Rule 1   |
| Conformity Assessment Path          | Not Applicable – Class I device  |
| Name/Address/ID of Notified Body:   | Not Applicable – Class I device  |
| Standards and Common Specifications | The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below.                 |
|                                     | BS EN ISO 14971:2012 – Medical Devices – Application of risk<br>management to medical devices<br>BS EN ISO 13485:2016 – Medical Devices – Quality management<br>systems – Requirements for regulatory purposes |

| EU Regulation                       | Restriction of Hazardous Substances in Electrical and Electronic Equipment Directive 2011/65/EU  |
|-------------------------------------|--|
| Standards and Common Specifications | The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below. |
|                                     | Supply and Component Controls - CPSIA testing of all components. Xref testing  |



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### Additional information:

| EU Authorized<br>Representative: | EUCEREP<br>Roald Dahllaan 33<br>5629MC – Eindhoven<br>The Netherlands |
|----------------------------------|---|
|                                  |   |

| Signature (signed for and on behalf of AbleNet): | Date of Issue: <dd month="" yyyy=""> 15 April 2021</dd>   |
|--|---|
| Soe Ny J.  |   |
| Printed Name:                                    | Place of Issue:<br>AbleNet Inc – Roseville, MN            |
| Joe Volp   | ADICIACT IIIO TAGGEVIIIC, IVIIA                           |
| Title: Director of Marketing                     | Document Number:<br>DoC_57200_Buddy Button - Green_041521 |