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# EU DECLARATION OF CONFORMITY

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| **Manufacturer** | **Etac Immedia A/S**<br>Parallelvej 3<br>DK-8751 Gedved<br>Denmark |
| **SRN** | DK-MF-000019241 |
| **Statement** | This EU declaration of conformity is issued under the sole responsibility of the manufacturer. The device(s) covered by present declaration is/are in conformity with EU Regulation 2017/745 on medical devices. |
| **Basic UDI-DI** | **57080126001MM** |
| **Intended purpose** | For placing a sling easily. |
| **Product / device name** | Immedia SlingOn |
| **Brand** | Immedia |
| **Risk class of the device** | Class I |
| **Place** | Gedved, Denmark |
| **Date of issue** | 1 June 2022 |
| **Name and function** | Michael Bruun, Senior Vice President |

Signature, on behalf of Etac Immedia A/S