EC Declaration of Conformity

Manufacturers Name: Van der Bend B.V.
Manufacturers Address: Kloosterweg 34, 3232 LC Brielle (NL)
SRN (Single Registration Number): NL-MF-000001575
Basic UDI-DI: See product reference list
Name of the Device(s): Van der Bend Patch Test Chambers
Classification: I

Conformity assessment route: The patch test chambers has been classified as Class I according to Annex VIII rule 1, and is in conformity with the general safety and performance requirements and provisions of the Regulation MDR 2017/745 and (are)(is) in conformity with the relevant harmonized standards: ISO 14971:2019
ISO 15223-1:2014
ISO 10993-5:2009
ISO 10993-10:2010
and is subject to the procedure set out in Annex II & III of the Regulation MDR 2017/745

This declaration of conformity is issued under the sole responsibility of Van der Bend B.V.
We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

All supporting documentation is retained at the premises of the manufacturer.

(Product reference list)

<table>
<thead>
<tr>
<th>Name manufacturer</th>
<th>Productname</th>
<th>Article reference</th>
<th>BASIC-UDI (GTIN)</th>
<th>Classification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Van der Bend B.V.</td>
<td>Van der Bend Patch Test Chambers</td>
<td>4600</td>
<td>08717056720104</td>
<td>Class I (rule 1)</td>
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</tbody>
</table>

Signature: Annemarie van der Bend
Place and date of issue: Brielle, 18-05-2021