EU DECLARATION OF CONFORMITY

HEINE Optotechnik GmbH & Co. KG
Dornierstr. 6, 82205 Gilching, Germany
www.heine.com
Single Registration Number: DE-MF-000006269

STETHOSCOPES

We hereby declare, under sole responsibility, that the devices covered by the present EU declaration of conformity are in accordance with the MDR 2017/745.

The stethoscopes

<table>
<thead>
<tr>
<th>Product name</th>
<th>Basic UDI-DI</th>
<th>GMDN</th>
<th>UMDNS</th>
</tr>
</thead>
<tbody>
<tr>
<td>GAMMA 3.1 Pulse Stethoscope</td>
<td>4053755_S_01_37</td>
<td>13755</td>
<td>13-750</td>
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<tr>
<td>GAMMA 3.2 Acoustic Stethoscope</td>
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<tr>
<td>GAMMA 3.3 Acoustic Stethoscope</td>
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<tr>
<td>GAMMA C3 Cardio Stethoscope</td>
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</table>

are class I according to the risk classification of Annex VIII.

References to any common specifications: N/A

This declaration of conformity is valid until a revised declaration of conformity is issued.

Gilching, 20 January 2022
(Place and date of issue)

Thomas Sauerer / PRRC
(Name/function and signature)

HEINE OPTOTECHNIK
GmbH & Co. KG
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82205 Gilching