Designation Letter of the Person Responsible for Regulatory Compliance (PRRC)

…..according to art. 15(1) of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices (hereby ‘’MDR’’), in order to fulfil the obligations set forth by …:

If only one PRRC is designated according to art. 15 (1):

…

If several PRRC

…

I hereby confirm that the designated PRRC

….

I hereby acknowledge

….

<Signature>

Place and date(DD/MM/YY)<name>

<position)
<Stamp>

<Signature>

Place and date(DD/MM/YY)<name>

<position)
<Stamp>