



Declaration

Company:

Sunrise Medical S.L.
Polígono Bakiola, 41
48498 Arrankudiaga
Vizcaya / ESPAÑA

Product:

(May include accessories)

Q100R
Q200R

We, Sunrise Medical declare under our sole responsibility that the product(s) to which this declaration relates, is a class 1 device, does not contain latex.
This was verified with conformity evaluation procedures according to Medical Device Directive Annex VII.

D.S.Davies – Director Project Management and R&D
Europe

A

29.10.19

Approval Name and Function

Revision

Approval Date


Signature (Sunrise Medical Approval representative)

GMS Form Number:

Revision: **B**

Effective Date: **01.02.2010**

Form Owner: Heads of Engineering

Form Approver: Global Head of Engineering

GMS Change Number: