

# SEDANA MEDICAL

## EC Declaration of Conformity

We:

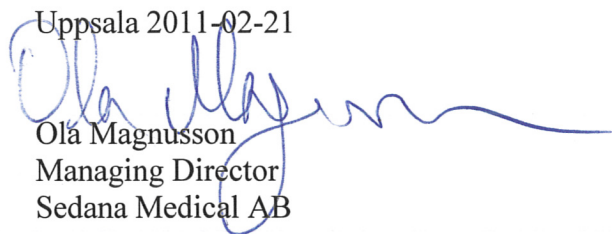
Sedana Medical AB  
Kungsgatan 62  
SE-753 18 Uppsala  
Sweden

declare under our sole responsibility that the products:

| Ref. No. | Product Description             | Classification |
|----------|---------------------------------|----------------|
| 26000    | AnaConDa™ System with syringe   | IIa            |
| 26022    | AnaConDa™ Syringe               | IIa            |
| 26042    | Filling Adaptor Sevoflurane     | I              |
| 26064    | Filling Adaptor Isoflurane      | I              |
| 26072    | Accessories for AnaConDa set-up | IIa            |

meet, (where applicable), the Essential Requirements (Annex I) and Annex VII coupled with Annex V for class IIa, Annex VII for class I products of Medical Devices Directive 93/42/EEC with the amendments in EU Directive 2007/47/EC

Uppsala 2011-02-21



Ola Magnusson  
Managing Director  
Sedana Medical AB