

SIEMENS

EC DECLARATION OF CONFORMITY

Manufacturer Siemens Shanghai Medical Equipment Ltd.(SSME)
Facility Siemens Shanghai Medical Equipment Ltd.XP Facility
278 Zhou Zhu Road, Shanghai 201318,China
Authorized Representative Siemens Healthcare GmbH
Henkestr. 127, 91052, Erlangen, Germany
Type of device X-Ray Radiography System
Medical device Multix Select DR
Product identification 10569624
Classification Class IIb (according to Annex IX to Council Directive 93 42/EEC)

We declare that the above medical device is in conformity with the following Directive(s):

Council Directive 93/42/EEC

The conformity of the full quality assurance system according to Annex II without Chapter II.4 is certified by:

TÜV SÜD Product Service GmbH
Ridlerstrasse 65
80339 Muenchen
Germany

SSME
Original
原件

The identification number of the notified body for implementation of the procedure set out in Annex II to the above Directive is 0123.

Directive 2011/65/EU of the European Parliament and of the Council.

This declaration of conformity is issued under the sole responsibility of SSME.

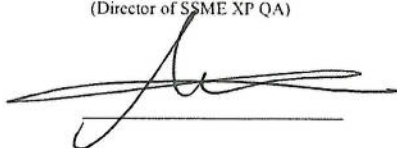
This declaration supersedes any declaration issued previously for the same product.

Place and date Shanghai, July 15, 2016

Name Mr. LU Hong Liang
(Head of SSME XP)

Mr. Liu Cong Zhi
(Director of SSME XP QA)

Signature



For conditions of guarantee and liability please refer to our General Conditions of Sale.

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