

# SIEMENS

## Compatibility certificate

### 1. Compatibility declaration, equipment designation and manufacturer

It is herewith certified that the equipment Pulse Oximeter 8600 FO of the manufacturer Nonin Medical, Inc. USA with the serial number 104502081 does not impair in its safety and functionality the magnetic resonance (MR) systems and products of the Siemens AG, Medical Solutions, designated in item 2 and specified individually. (Compatibility with regard to the effect of the above designated equipment on the named MR product). Possible functional restrictions of the MR products are stated under item 5. Possible unfavorable effects of the MR system on the above designated equipment are here expressly not object of the examination forming the basis of this certificate. The certificate therefore does not mean that the above equipment is unrestrictedly compatible with regard to effects caused by the MR systems.

### 2. MR systems affected

Serial no.	MR system or option (type designation)	Part number	System modification status (modification code)	Software versions (version numbers)	Configuration, included/ excluded accessory
1	Symphony	n.a.	n.a.	n.a.	n.a.
2	Sonata	n.a.	n.a.	n.a.	n.a.

### 3. Area of application

The equipment Pulse Oximeter 8600 FO is intended for the following purpose in combination with the magnetic resonance application; the compatibility certificate refers to this:

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| 1. Monitoring a patient's blood's pulse saturation during MRI scanning |
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### 4. Restrictions

During the use of this equipment, the following restrictions in the function and/or possibilities of use of the MR system (of the systems, options, accessories) result:

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| 1. n.a. |
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### 5. Warning notes

The following precautionary measures must be observed when using this equipment in combination with the MR products named under item 2:

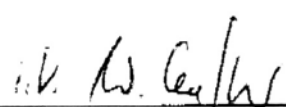
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| 1. Do not place the Pulse Oximeter 8600 FO in a distance less than 2 meter to the magnet. Warning notes must be included in the operating manual. |
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### 6. Validity:

The compatibility certificate is valid until it is revoked by the Siemens AG, Medical Solutions. The validity starts after attaching corresponding warning notes at the product and in the operating instructions.

Erlangen, 2003-08-06

Siemens Aktiengesellschaft

  
 Dr. Loeffler

  
 Lehne