

# Declaration of Conformity EU

Acc. To the EU-guideline regarding Medical Devices  
93/42/EWG, annex II dated July 12th 1993



Lichttherapie-technik  
Entwicklung  
Herstellung  
Vertrieb  
Service

We declare herewith, that the device described below corresponds to the fundamental requirements for security and health of EU-guideline regarding Medical Devices (93/42/EWG).

This declaration becomes invalid, if the device is not operated as agreed or if the device has been changed without consulting the manufacturer.

Name of the manufacturer: **A.L.T. Lichttherapie-technik GmbH  
Thura Mark 10, D – 06780 Zörbig**

Name of the device: **Idromed® 5 GS, Idromed® 5 PS**

Art.-No.: **100000 – 100050  
110000 – 110050**

From year of construction: **2008**

Accordance with the following guideline: **Medical Devices Act  
(93/42/EWG)**

Title, name and first name of the signatory: **Schnatwinkel, Mathias**

Function: **Managing Director**

Place and date of issue: **Zörbig, 17.03.2008**

Stamp:



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Signature: