

EC Certificate

The LGA InterCert GmbH – **Notified Body 1275** according to the EC Directive concerning medical devices – herewith confirms that the following company



Q.Products AG
Säntisstraße 11
9401 Rorschach
Switzerland

applies a

Production Quality Assurance System

for the manufacture of the medical devices
as specified in the enclosure within the scopes: Lighttherapy / Phototherapy

according to

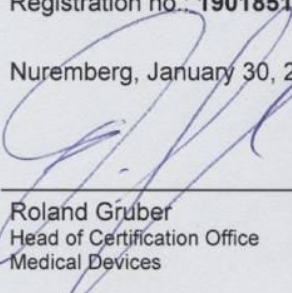
Directive 93/42/EEC, Annex V.

In a certification audit carried out by LGA InterCert GmbH, evidence was furnished proving that the quality system of the company fulfills the requirement of the above-mentioned directive.
The surveillance through the LGA InterCert GmbH is effected according to Annex V (4).

This certificate is valid until January 29, 2016
Registration no.: **1901851-014-000**

Audit report no.: 1901851
of: November 11, 2010

Nuremberg, January 30, 2011


Roland Gruber
Head of Certification Office
Medical Devices



It is allowed for the company to sign the products specified in the enclosure with the adjoining sign.

CE 1275

**Enclosure of the Certificate Registration no. 1901851-014-000 der
Q.Products AG, Säntisstraße 11, 9401 Rorschach, Switzerland**

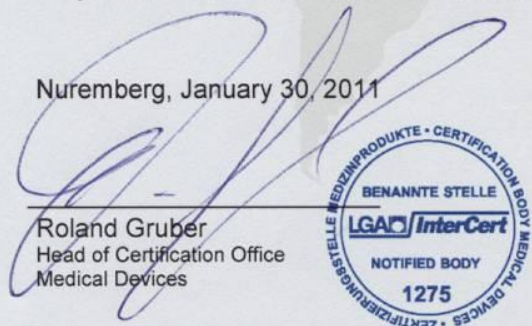
Period of Validity Januaray 30, 2011 until January 29, 2016

The above-mentioned certificate refers to the following medical devices:

No.	Product	UMDNS-No.	Class
1.	<p>light therapy- / phototherapy device Q.Light for the following fields of use and indications, Wound Healing, Acne, PDT, Burns, General Pain Treatment, Depressions, Sleep disorder</p> <ul style="list-style-type: none"> • Q.Light® 70 • Q.Light® 70 NT • Q.Light® 70 NT IR • Q.Light® 150 • Q.Light® 150 NT • Q.Light® 150 NT IR • Q.Light® 200 • Q.Light® 200 NT • Q.Light® 200 NT IR <p>Variants appropriate „article list“</p>	*	IIa

*At present no UMDNS number is available.

Nuremberg, January 30, 2011


Roland Gruber
Head of Certification Office
Medical Devices

