



Product Service

# EC Certificate

## Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)  
(Devices in Class IIa, IIb or III)

No. G1 18 04 40575 004

**Manufacturer:** asia-med GmbH

Kirchplatz 1  
82049 Pullach  
GERMANY



**Facility(ies):**

asia-med GmbH  
Kirchplatz 1, 82049 Pullach, GERMANY

asia-med GmbH  
Schützenstrasse 22, 98527 Suhl, GERMANY

**Product Category(ies):** Acupuncture needles

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

**Report No.:** 713126616

**Valid from:** 2018-06-26

**Valid until:** 2023-06-25



**Date,** 2018-06-12

Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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