MEDlight
natürlich gesund

Declaration of Conformity

within the meaning of
Council Directive 93/42/EEC concerning medical devices
including Directive Amendment 2007/47/EC

We

MEDlight GmbH
Füllenbruchstr. 201
32051 Herford
Germany

Declare under our sole responsibility that the device

Device name: **OCTAderm**

Class of medical device: **IIa**

REF/PZN: **1004 (UVA), 1005 (UVB narrow band)**

is designed and manufactured in compliance with Directive 93/42/EEC
and meets the essential requirements (Annex I).

The technical documentation is kept at the above address.

The conformity assessment was done in compliance with Annex V of Council Directive 93/42/EEC.
Applicable harmonised standards were applied.

CE 2409

This declaration is valid only in context of the Notified Body

CE Certiso Kft. (NB 2409), Office address: H-2092 Budakeszi, Erdő utca 101.
Postal address: H-2092 Budakeszi, Pf. 70

Issued certificate is valid for all deliveries after date of issue.

Herford, 2021-05-26
(place and date of issue)

(safety representative)