

EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

No. CE 591626
Issued To: **Natures Naturals Ltd**
5 Moorfield Farm
Warkton Village
Kettering
Northamptonshire
NN16 9XJ
United Kingdom

In respect of:

Manufacture of Hand-held deep-tissue pulsed electromagnetic stimulators

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):



Frank Lee, EMEA Compliance & Risk Director

First Issued: **21 January 2013**

Date: **13 September 2016**

Expiry Date: **13 September 2021**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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Certificate History

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Date	Reference Number	Action
21 January 2013	7905320	Initial Issue. Transfer from another notified body
11 December 2015	8451112	Change of address from Ringstead Business Centre, 1-3 Spencer Street, Ringstead, Kettering, NN14 4BX. To 5 Moorfield Farm, Warkton Village, Kettering, Northamptonshire, NN16 9XJ
13 September 2016	8497704	Renewal. Amend scope to remove Painsolv product name.