Certificate of Conformity

Directive 93/42/EEC on Medical devices, Annex IV

Certificate No.: EU1407406 Date: 2014.07.25

Order No.: 266314

We hereby certify that a EC verification has been carried out on the series or batch of device(s) listed hereafter following the requirements of the national legislation: Regulation no1690 of 15th December 2005 relating to medical devices pursuant to act no. 6 of 12th January 1995 relating to medical devices, transposing directive 93/42/EEC into Norwegian law to which the undersigned is subjected. Confer EEA agreement, proposition no. 100 (1991-92) special appendix no. 2, volume 2A/3 A, goods, chapter XXX. We certify that the device(s) hereafter referenced conform to the relevant provisions of **Annex IV** of the aforementioned directive.

Manufacturer:	ALLIDEX 12, Rue de la Turbie 98000 Monaco
Device category:	Dental casting alloy
GMDN code:	35857
Models:	Gammarc CC4 VI
Batch / Serial number:	20732
Ref. of Test Report:	05023/01 0073
Date/period of testing:	2014.06.03
Nemko EC notification No.:	0470

Remarks:

On this basis, presupposed that the provisions in Annex VII are fulfilled, the manufacturer or his European authorized representative may draw up an EC Declaration of Conformity and affix the CE-marking as indicated below followed by the Nemko EC notification number on the conforming product. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC.

Arild Hansgård
For Nemko AS

Nemko Norway
Nemko AS, Gaustadalléen 30, P.O. Box 73 Blindern, 0314 Oslo, Norway
TEL +47 22 96 03 30 FAX +47 22 96 05 50 EMAIL info@nemko.com
ENTERPRISE NUMBER NO974404532