

EC CERTIFICATE Full Quality Assurance System

Certificate No.:

Project No.:

Valid Until:

9905-2017-CE-CZS-NA-PS rev.2.0

PRJC-89738-2008-PRC-SVK

27 May 2024

This is to certify that the quality system of:

EKOM spol. s.r.o.

Priemyselná 5031/18, 921 01 Piešťany, Slovak Republic

For design, production and final product inspection/testing of:

DENTAL ACTIVE DEVICES

Has been assessed with respect to:

The conformity assessment procedure described in Annex II excluding section 4 of Council Directive 93/42/EEC on Medical Devices, as amended

and found to comply

Further details of the product(s) and conditions for certification are given overleaf

Place and date: Høvik,08 April 2021 For the issuing office: Notified Body 2460 DNV Product Assurance AS



Sholeh Gheissar Principal Assessor



Certificate No.: 9905-2017-CE-CZS-NA-PS rev.2.0

Place and date: Høvik,08 April 2021

Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
0.0	Supersedes DNV GL (NB0434) certificate No. 106328-2011-CE-CZS-NA following transfer of notified body function to DNV Nemko Presafe AS (NB2460) at Recertification	2017-05-22
1.0	Extension in scope - new products added	2020-05-18
2.0	Re-certification	2021-04-08

Products covered by this Certificate:

Product Description	Product Name	Class
Dental suction equipment	ASPINA – model DO-M	lla
Dental compressors with suction unit	DUO, models as listed in the List of Models dated 20 January 2021	lla
Dental Compressors	DK50, models as listed in the List of Models dated 20 January 2021	lla

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address
EKOM spol. s.r.o.	Priemyselná 5031/18, 921 01 Piešťany, Slovak Republic



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Place and date: Høvik,08 April 2021

Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a
 defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning
 liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the Notified Body of any intended updating of the quality system and the Notified Body will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies
 the quality system. the Notified Body reserves the right, on a spot basis or based on
 suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number.

End of Certificate