

## DECLARATION OF CONFORMITY

Medical Device Directive 93/42 EU

DENLUX AS  
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DK-2750 Ballerup  
DENMARK

Phone: +45 4465 0661

CVR No. DK28305494

TO WHOM IT MAY CONCERN

< Company name >

< Address >

< Postal code & City >

< Country >

It is hereby declared that DENLUX A/S (Dental Electronic A/S), Denmark holds the sole responsibility for conformity and that below mentioned Medical Devices manufactured by DENLUX A/S (Dental Electronic A/S), Denmark fulfil requirements in the Medical Device Directive 93/42/EU, RoHS Directive 2011/65/EU and REACH Regulation 1907/2006/EU.

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Type / product: Pulp vitality Tester / Pulp tester

Brand name: Pulppen B1000 Analogue (DENLUX part No. 11090)  
Pulppen DP2000 Digital (DENLUX part No. 11290)  
Pulppen DP3000 Digital (DENLUX part No. 11390)

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Classification: According to MDD 93/42/EU, Annex IX  
Rule: 10  
Class: IIa

Certification: According to MDD 93/42/EU, Annex VI

Certification body: SLG Prüf- und Zertifizierungs GmbH (NB No. 0494)  
Burgstädterstrasse 20  
09232 Hartmannsdorf  
DEUTSCHLAND

Date: xx/xx 20xx

Signature: \_\_\_\_\_

Mr. Jesper Lund, CEO  
DENLUX A/S, Dental Electronic A/S