APRAGAZ

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BELAC EN ISO/IEC 17021 Cert N° 084-QMS

NOTIFIED BODY EC CERTIFICATE OF CONFORMITY In accordance with Appendix II of the Medical Devices Directive 93/42/EC & the latest amendment 2007/47/EC, excluding section 4.		Certificate N° 13/BE/3103-0-REV 0
Manufacturer	Name	Dumont Instruments & Co S.A.
	Address	42, Rue des Anciens Etangs B – 1190 Brussels BELGIUM

Concerned Equipment:

Design, manufacturing and / or marketing of diamond burs and disks, steel and tungsten carbide burs, polishers / abrasives for dental use.

This is to certify that the Quality Management System of the above mentioned manufacturer has been assessed against the requirements of Annex II of the Medical Devices Directive 93/42/EC and conforms to the requirements for the equipment shown above. The approval is subject to the continued maintenance of the Quality System in accordance with the requirements of the above Directive, this shall be controlled by intermediate audits, inspections and surveys.

The manufacturer is allowed to affix the "CE" mark followed with our notified body identification number 0029 to the approved equipment in the conditions described in the Directive.

The approval is valid until 05/05/2018

Date : **06/05/2013**

Name : **Ch. Leplat**

Position **General Manager**

Signature

Notified body identification number : **0029**

Notified body stamp :

APRAGAZ
Belgium
Inspecting Authority

Notified body reference : **P15210/001**