EMERGO*EUROPE*
QUALITY ASSURANCE & REGULATORY AFFAIRS

CE Registration Notification

This is to certify that, in accordance with the Medical Device Directive 93/42/EEC, Emergo Europe agrees to perform all duties and responsibilities as the Authorized Representative for

AdDent, Inc.
43 Miry Brook Road
Danbury, CT 06810
USA

as stipulated and demanded by the aforementioned Directive. The Dutch Competent Authorities have received Medical Device Registrations on the following date:

January 31, 2006
See attached product listing

Emergo Europe Registration Number: NL/CA01/601529

The Manufacturer has provided Emergo Europe with the appropriate Declaration(s) of Conformity confirming that the Medical Devices fulfill the applicable requirements of Directive 93/42/EEC.

May 14, 2007

Rene van de Zande
President
Emergo Europe

Molenstraat 15
2513 BH The Hague
The Netherlands

Tel: (+31) 70 345 8570
Fax: (+31) 70 346 7299

Email: europe@emergogroup.com
Website: www.emergogroup.com

EMERGOEUROPE
QUALITY ASSURANCE & REGULATORY AFFAIRS

Annex A to the Emergo Europe CE Registration Certificate

dated April 19, 2004

Medical Device	Class Per MDD 93/42/EEC	Registration Date
Calset	Class I	January 31, 2006
Trimax	Class I	January 31, 2006
Microlux DL	Class I	January 31, 2006
Microlux Transilluminator	Class I	January 31, 2006
Rite-Lite	Class I	January 31, 2006

Molenstraat 15
2513 BH The Hague
The Netherlands

Tel: (+31) 70 345 8570
Fax: (+31) 70 346 7299

Email: europe@emergogroup.com
Website: www.emergogroup.com