EMERGO EUROPE

CE Registration Certificate

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:

6 July 2009
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.

July 2009

Rene van de Zande
President & CEO
Emergo Europe

EMERGO EUROPE

Annex A to the Emergo Europe CE Registration Certificate

dated July 2009

Medical Device	Class Per MDD 93/42/EEC	Registration Date
Calset	Class I	31 January 2006
Trimax	Class I	31 January 2006
Microlux DL	Class I	31 January 2006
Microlux Transilluminator	Class I	31 January 2006
Rite-Lite	Class I	31 January 2006
CoMax Composite Dispenser	Class I	6 July 2009