EMERGO EUROPE
Quality Assurance & Regulatory Affairs

CE Registration Certificate

This is to certify that, in accordance with the In Vitro Diagnostic Medical Device Directive 98/79/EC, Emergo Europe agrees to perform all duties and responsibilities as the Authorized Representative for

Diagnostic Biosystems
1020 Serpentine Lane, Unit 114
Pleasanton, CA 94566
U.S.A.

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

See Attached
Dated
26 September 2007

Emergo Europe Registration Number: NL/CA01/601529

The Manufacturer has provided Emergo Europe with the appropriate Declaration(s) of Conformity confirming that the In Vitro Diagnostic Medical Devices fulfill the applicable requirements of Directive 98/79/EC.

26 September 2007

Rene van de Zande
President & CEO
Emergo Europe