ULTRASONIX

Smart Ultrasound for better patient care

Ultrasonix Medical Corporation

Declaration of Conformity

Manufacturer's Address:	*130-4311 Viking Way, Richmond, B.C., V6V 2K9, Canada* *T (604) 437-9500* *F (604) 437-9502* www.ultrasonix.com
European Representative:	*MEDNET GmbH* *Borkstraße 10* *48163 Münster – Germany* *T 49-0-251-32266-0* *F 49-0-251-32266-22* *www.medneteurope.com*
Name of Device(s):	*SonixMDP*
Device Catalog Numbers:	*20.000.000*
MDD Annex IX Classification:	*Class II a*
Conformity Assessment Route:	*Annex V, section 3.2 – Production quality assurance, combined with Annex VII*

We hereby declare that the medical device(s) specified above comply with the European Medical Device Directive 93/42/EEC and its relevant transpositions into all national laws of the member states into which we place the device(s).

Notified Body:	*Notified Body #0543* *DGM* *Kollegievej 6* *2920 Charlottenlund* *Denmark*
Certificate:	*EC Certificate DGM-473*

(Authorized Signature)

Chas Yu

Quality Assurance Manager

21st December, 2009

(Date)