

Smart Ultrasound for better patient care

Ultrasonix Medical Corporation

Declaration of Conformity

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Name of Device(s): SonixMDP

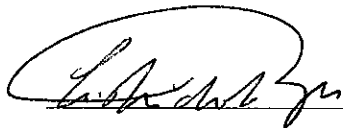
Device Catalog Numbers: 20.000.000

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)