



Ultrasonix Medical Corporation

Declaration of Conformity

Manufacturer's Address: 130-4311 Viking Way, Richmond, B.C., V6V 2K9,
Canada
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European Representative: MEDNET GmbH
Borkstraße 10
48163 Münster – Germany
T 49-0-251-32266-0
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www.medneteuropa.com

Name of Device(s): Sonix OP
Device Catalog Numbers: 10.000.000
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)

Alan Remfry

Director Quality Assurance & Regulatory Affairs


(Date)