auditdata
Your Partner in Audiology Solutions

CE

Declaration of Conformity

According to Medical Device Directive (MDD) 93/42/EEC

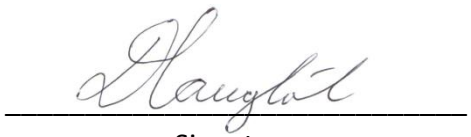
Manufacturer		
	Auditdata A/S Dalbergstroeget 5-7 2630 Taastrup Denmark	
Conformity Assessment Procedure		
	Annex II of Medical device Directive MDD 93/42/EEC	
Registration		
	Danish Health and Medicines Authority Axel Heides Gade 1 2300 Copenhagen S, Denmark	
Product Identification		
	Category:	Office Management System
	Brand:	AuditBase
	Model:	AuditBase
	Lot/Batches/Serial number:	From release 4.19.3.5
MDD Directive		
	Class I, Rule 11, MDD 93/42/EEC	

The medical device is in compliance with the essential requirements according Annex I of the Medical Device Directive 93/42/EEC.

We declare under our sole responsibility that the products, to which this declaration relates, are in conformity with the Essential Requirements Annex I of the above directive.

Taastrup, December 4th 2014

Dan Haugbøl, QA & Regulatory Manager



Signature