CERTIFICATE

This is to certify that the company

Vepro AG Germany

Max-Planck-Str. 1-3
64319 Pfungstadt
Germany

has implemented and maintains a **Quality Management System**.

Scope:
Design, production, distribution, installation and support of Software- and Hardware products for digital image management in the medical field.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

EN ISO 13485 : 2012 + AC : 2012

Certificate registration no.	527544 MP2012
Certificate unique ID	170649426
Effective date	2016-10-11
Expiry date	2019-10-10
Frankfurt am Main	2016-10-11

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-16021-01-01

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Dr. Thomas Feldmann
Head of Certification Body

Accredited Body: DQS Medizinprodukte GmbH, August-Schanz-Str. 21, 60433 Frankfurt a. M., Germany

EC-CERTIFICATE

(Full quality assurance system)

This is to certify that the company

Vepro AG Germany

Max-Planck-Str. 1-3
64319 Pfungstadt
Germany

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Software for digital image processing/-archiving and communication in the medical field (PACS) EMR Manager V9	Class IIa

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no.	527544 MR2
Certificate unique ID	170649427
Effective date	2016-10-31
Expiry date	2021-10-30
Frankfurt am Main	2016-10-31

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Dr. Thomas Feldmann
Head of Certification Body

August-Schanz-Straße 21, 60433 Frankfurt am Main,
Tel. +49 (0) 69 95427-300, medical.devices@dqs-med.de

DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.