

DECLARATION OF CONFORMITY

MANUFACTURER

Rexxam Co.,Ltd. Kagawa factory
958 Ikeuchi,Konan-cho,Takamatsu-shi KAGAWA-KEN 761-1494 JAPAN

AUTHORIZED EUROPEAN REPRESENTATIVE

REXXAM ELECTRONICS IRELAND CO.LIMITED
Donore Road, Drogheda, County Louth, Ireland

MEDICAL DEVICE

Model Name : Auto Ref-keratometer ACCUREF K-900
Classification : Class I (Devices with a measuring function)
Serial number : effective from 51BN0011(Valid until a product is changed.)

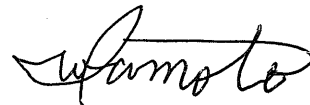
The undersigned hereby declares that the medical device as specified above conforms with the essential requirements listed in Annex I of EC Directive 93/42/EEC.

This declaration of conformity is based on the EC Directive 93/42/EEC Annex V and supported by the TUV Rheinland LGA Products GmbH(0197) (TILLYSTRASSE 2, 90431 NUREMBERG - GERMANY-) Annex V certificate, with reference to articles 3 of directive 93/42/EEC.

We declare that we are exclusively responsible for this declaration.

Kagawa Factory, 13.Jun.2011

Place and date of issue



Name, signature and
position of manufacture

Development Dept.1 Manager
Masakatsu Iwamoto