

DECLARATION OF CONFORMITY

MANUFACTURER

Rexxam Co., Ltd. Kagawa Factory
958 Ikeuchi, Konan-cho,
Takamatsu-shi
Kagawa-ken 761-1494
Japan

AUTHORIZED EUROPEAN REPRESENTATIVE

Rexxam Czech s.r.o.
Prumyslova 937, 334 01 Prestice, Czech Republic

MEDICAL DEVICE

Model Name : Auto Ref-keratometer ACCUREF K-900
Classification : Class I (Devices with a measuring function)
Serial number : effective from Y9BN3992 (Valid until a product is changed.)

This declaration is only valid for the manufactured products with the document released by us of each product.

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, VII 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, 1.April.2020

Place and date of issue



Name, signature and
position of manufacture

Development Dept.1 Manager
Shunji Takashima