HEALTH CARE
unicos Health-care Business Venture Company

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Declaration of Conformity

CE 0120

Description of Device: **AUTO REFRACTO-KERATOMETER**

Model Name: **URK-800F**

Classification: **Class IIa**
according to the Rule 10 of Classification Criteria,
annex IX, MDD 93/42/EEC amended by 2007/47/EC

Conformity Assessment Route : **Annex Ⅱ(excluding section4), MDD 93/42/EEC As amended by 2007/47/EC**

Applied Standards: **IEC 60601-1: 2005+CORR.1(2006) + CPRR.2(2007)**
IEC 60601-1-2:2007(Third Edition)

Manufacturers Name: **UNICOS Co., Ltd.**
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We hereby declare that the complies with the Medical Devices Directive 93/42/EEC As amended by 2007/47/EC (and its relevant transposition into the national laws of the Member States in which the device is intended to be placed on the market) using Annex Ⅱ as the conformity assessment procedure via SGS(NB 0120) as the Notified Body.

Place, Date of issue : Jung-Gu, Daejeon, Korea Jan. 24, 2017

UNICOS CO.,LTD
K.C.KIM
K.C KIM PRESIDENT

Ki-chang, Kim/ President
On behalf of UNICOS.Co.,Ltd