


Reference No. KR/15/00007/CoCR1

Internal ref. 616475/01



Place and date Brussels, February 12, 2015

CERTIFICATE OF CONFORMITY for RoHS Directive (2011/65/EU)

Product	Ultrasound Bone Densitometer
Tested on request of	OSTEOSYS CO., LTD.
Name and address of the manufacturer	OSTEOSYS CO., LTD. 901~914, 9F, JNK Digitaltower, 111 Digital-ro 26, Guro-gu, Seoul, 152-848, Republic of Korea
Name and address of the factory	Same as above
Rating and principal characteristics	AC 100-240V, 50/60Hz
Trade mark (if any)	
Type/identification	SONOST 3000
Conclusion	The reviewed sample and technical information received comply with the hazardous substance limits of directive 2011/65/EU and the conformity assessment procedure as outlined in Decision 768/2008/EC, Annex II, Module A, Point 2 as well as RoHS harmonized standard EN 50581
Sufficient samples of the product/component/material have been analysed according to the following scope:	Pb, Hg, Cd, Cr (VI), PBB, PBDE according to standard IEC 62321(ed.1) - 2008-12-11
As shown in the report (reference No.)	F690101/LF-CTSAYGU14-04154R1

This Certificate of Conformity is based on analysis of the product and verifying the documentation being in accordance with the provisions of the Directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment. It does not imply an assessment of the production site. No amounts above the limits of these substances have been detected or if so specific exemptions are applicable. This result relates only to the item(s) tested. Other CE marking directives may apply but have not been considered during the RoHS assessment.

This Certificate of Conformity does not give the right to use any SGS mark on products, components or materials, their packaging or any marketing material or activities.

<p>On behalf of SGS Belgium NV-Division SGS CEBEC</p> <p>YURI HONG </p>	 <p>The form of the CE marking</p>
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