

EU Declaration of Conformity



We, **SKG Health Technologies Co.,Ltd.** as **Manufacturer**,declare under our sole responsibility that the following product(s) with Brand **SKG**, produced in **China**, is in compliance with all relevant CE directives/regulations.

Model Number	Product Type
G7 Pro-E	Neck Massager

Subject	Directive/Regulation	Standard
RED	2014/53/EU	EN55014-1:2017+A11:2020
		EN55014-2:2015
		EN301489-1 V2.2.3
		EN301489-17 V3.2.4
		EN300328 V2.2.2
		EN62479:2010
		EN60335-1:2012+A11:2014+A13:2017+A1:2019+A14:2019+A2:2019
		EN60335-2-32:2003+A1:2008+A2:2015
		EN62233:2008
RoHS	2011/65/EU+(EC)2015/863	IEC62321

Importer: Name: **Rolux Leuchten GmbH**
 Address:Margarete-steiff-StraBe 18 28844 Weyhe

Signed by and on behalf of (manufacturer): SKG Health Technologies Co.,Ltd.
 Place: 23A Floor,Building 3,Zhongke R&D Park,No.009,Gaoxin South 1st Road, High-tech Zone
 Community Yuehai street, Nanshan District,Shenzhen City,China

Name: Chenzhe Function: Compliance Engineer Date:4 Dec. 2023

Signature:



Chenzhe