

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
W3 Pro	Knee Massager

Subject	Directive/Regulation	Standard
CE	2014/35/EU	EN55014-1:2021
		EN55014-2:2021
		EN 60335-2-32:2003+A1:2008+A2:2015
		EN 60335-1:2012+AC:2014+A11:2014+A13:2017+A1:2019+A14:2019+A2:2019+A15:2021
		EN 62233:2008+AC: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*

