EU Declaration of Conformity
D-P005-50-56-04

Manufacturer: mo-Vis bvba

Address: Biebuyckstraat 15 D
B-9850 Deinze
Belgium

Hereby we declare that mo-Vis bvba, as a legal manufacturer, has the exclusive responsibility to draw up this Declaration of Conformity.

This declaration concerns the product group **Twister**, intended as small switches with light sensitivity, and consisting of the following products:

Twister Button (P005-75/76/77/78/79 and P-005-85/86/87/88/89), Twister Basic (P005-70/71/72/73/74 and P005-15/16/17/18/19), Twister on Bended Tube (P005-50/51/52/53/54 and P005-60/61/62/63/64), Satellite Twister (P005-40/41/42/43/44) and Twister Gooseneck (P018-20/21/22/23/24)

which are **Medical Devices Class I**, with reference to rule 1 of the Annex VIII of the Medical Device Regulation 2017/745.

mo-Vis declares that these products are in conformity with General Safety and Performance Requirements of the Medical Device Regulation 2017/745, Annex I and with the Medical Device Regulation 2017/745, Annex IV.

UDI: 540700832TWISTERSB

This Declaration of Conformity is valid for all devices described here above and until the issue of a revised Declaration of Conformity after change of the product.

Deinze, 22/10/2021

Rebecca Van Craeymeersch
mo-Vis bv
Quality Manager

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