EU Declaration of Conformity

Legal Manufacturer: MVS In Motion bv,
with registered office at: Westdijk 150, 2830 Tisselt (Willebroek), Belgium,
with SRN: BE-MF-000000664
declares under its sole responsibility that products listed below meet the provisions of Medical Device Regulation 2017/745
as a Class I Medical Device,
according to Rule 1 of Annex VIII.

Intended purpose:

- Hammers (542006300J3H): intended to test deep tendon reflexes with the aim to detect abnormalities in the peripheral or central nervous system.
- Wartenberg Pinwheel (542006300K3K): intended for sensory evaluation as it is rolled over the skin.

Conformity Assessment Route: Annex II & III of the Medical Device Regulation 2017/745.

List of Products:

REF	Description	BUDI-DI	UDI-DI
08-050101	MoVeS Buck Hammer	542006300J3H	05420063004527
08-050102	MoVeS Babinski Hammer	542006300J3H	05420063004534
08-050103	MoVeS Taylor Hammer	542006300J3H	05420063004541
08-050104	MoVeS Dejerine Hammer	542006300J3H	05420063004862
08-050105	MoVeS Troemner Hammer	542006300J3H	05420063004879
08-050201	MoVeS Wartenberg Pinwheel	542006300K3K	05420063004558

Date: 06 May 2021
Place: Tisselt (Willebroek), Belgium

Alexandra Vrancaert,
General Manager