MEDIKRO

RoHS Declaration of Conformity

Issued by Medikro Oy

- 1. List of electrical and electronic equipment which under RoHS directive, 2011/65/EU amended by 2015/863 Compliant:
 - M911 Medikro® Ambi,
 - M914 Medikro® Primo,
 - M915 Medikro® Pro,
 - M920 Medikro® Pro and
 - their OEM variants.

OEM units are designed, manufactured and tested according to the same quality system requirements as the Medikro units.

- 2. We hereby confirm that items listed above is in conformity with the RoHS directive 2011/65/EU as amended by (EU) 2015/863 on the restriction of the use of certain hazardous substances in electrical and electronic equipment.
- 3. The items listed above will not exceed the limit for restricted substances and maximum concentration values tolerated by weight in homogeneous materials
 - Lead (0,1 %)
 - Mercury (0,1 %)
 - Cadmium (0,01 %)
 - Hexavalent chromium (0,1 %)
 - Polybrominated biphenyls (PBB) (0,1 %)
 - Polybrominated diphenyl ethers (PBDE) (0,1 %)
 - Bis(2-ethylhexyl) phthalate (DEHP) (0.1 %)
 - Butyl benzyl phthalate (BBP) (0.1 %)
 - Dibutyl phthalate (DBP) (0.1 %)
 - Di-isobutyl phthalate (DIBP) (0.1 %)

Place and Date Kuopio, 2020-06-22

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Tuukka Eloranta Medikro Oy, Managing Director