Supplier EU Declaration of Conformity to Directive 2011/65/EU (RoHS)

Dongguan Actherm Medical Corp.
3F, No.21, Hai-Bin Road, Wusha Community, Chang-An Town, 523858 Dongguan City, Guangdong Province, China

DECLARATION OF CONFORMITY according to Annex VI of Directive 2011/65/EU (RoHS)

1. Number (unique identification of the Electrical and Electronic Equipment, EEE)	Product name/model: **Thermoval rapid / kids** **(Actherm model name: ACT5030 Express)**
2. Name and address of the manufacturer or his authorised representative:	**Dongguan Actherm Medical Corp.** 3F, No.21, Hai-Bin Road, Wusha Community, Chang-An Town, 523858, Dongguan City, Guangdong Province, China European Representative: MedPath GmbH Mies-van-der-Rohe-Strasse 8, 80807 Munich,Germany
3. This declaration of conformity is issued under the sole responsibility of the manufacturer (or installer):	Dongguan Actherm Medical Corp.
4. Object of the declaration (identification of EEE allowing traceability. It may include a photograph, where appropriate):	Digital Clinical Thermometer (pen-type)
5. The object of the declaration, described above, is in conformity with Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (*):	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 %) Diisobutyl phthalate (DIBP) (0,1 %)
6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:	EN IEC 63000
7. Additional information:	
Exemptions: the following exemptions according to the Annexes of the above mentioned directive and their scope and dates of applicability are relevant:	*Apply 6(a), 6(b), 6(c), 7(a) and 7(c)-I as listed in Annex III of Directive 2011/65/EU.*

The technical documentation forming the basis of this declaration will be kept for 10 years after the EEE has been placed on the market (or 10 years after the relevant contracts have expired).

Signed for and on behalf of:	Dongguan Actherm Medical Corp.
Place and date of issue:	Dongguan, China, December 01, 2023
Name, Function and signature:	YiChuan Chou (Director) YiChuan Chou .....

(*) OJ L 174, 1.7.2011, p. 88-110.

Note: Text in *italic* is a HARTMANN requirement.