

CTI Declaration of Conformity

The submitted sample of the following equipment has been tested for CE marking according to the following European Directives: Medical Devices Directive 93/42/EEC.
The manufacturer's technical documentation, as required for Class I devices, has been reviewed and comply with the requirements in Annex VII, section 3 of the Medical Devices Directive 93/42/EEC.

Applicant name & address : Wuhan bms Medicaltech Co., Ltd
C/D Unit, 5th Floor, B9, Hi-tech Medical Devices Park, No.818
Biolake Gaoxin Avenue, Wuhan, China.

Manufacturer name & address : Wuhan bms Medicaltech Co., Ltd
C/D Unit, 5th Floor, B9, Hi-tech Medical Devices Park, No.818
Biolake Gaoxin Avenue, Wuhan, China.

Product Name : Tube Sealer

Trade Name : 
experts in blood technology

Model(s) : Mobile3724, Station3712, MSS3773

Serial No. : N/A

Technical Data : Input: 200-240V~, 50/60Hz, 200VA

Order No. / Report No. : EED33H000078/EED33H000078

Test Standards	
EN 60601-1: 2006+A1: 2013	Medical electrical equipment Part 1: General requirements for basic safety and essential performance
EN 60601-1-2: 2007	Medical electrical equipment —Part 1-2: General requirements for basic safety and essential performance — Collateral standard: Electromagnetic compatibility — Requirements and tests

This Declaration is for the exclusive use of CTI's Client and is provided pursuant to the agreement between CTI and its Client. The observations and test results referenced from this Declaration are relevant only to the sample tested. This Declaration by itself does not imply that the material, product, or service is or has ever been under a CTI certification program.
Note: This Declaration is part of the full test report(s) and should be read in conjunction with it.



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Approved Signatory
Date: Mar. 24, 2016

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