

**DECLARATION OF CONFORMITY
TO COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993
CONCERNING MEDICAL DEVICES**



MANUFACTURER: NAME: Guangdong Transtek Medical Electronics Co.,Ltd.
ADDRESS: Zone A, No.105 ,Dongli Road ,Torch Development District,
Zhongshan,Guangdong,China

MEDICAL DEVICE: BLOOD PRESSURE MONITORS: TMB-1490/1490-A/1490-C/1491/
1491-A/1491-D

CLASSIFICATION - ANNEX IX: CLASS IIA, RULE 10

CONFORMITY ASSESSMENT ROUTE: MDD ANNEX II

WE, THE MANUFACTURER, HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES
MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE
93/42/EEC OF 14 JUNE 1993 CONCERNING MEDICAL DEVICES;
INCLUDING, AT 21 MARCH 2010, THE AMENDMENTS BY COUNCIL DIRECTIVE 2007/47/EEC.
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED: SEE ATTACHED

NOTIFIED BODY: TÜV SÜD PRODUCT SERVICE GMBH
RIDLERSTR 65, D-80339 M NCHEN, GERMANY

IDENTIFICATION NUMBER

CE 0123

(EC) CERTIFICATE(S): NO.G1 16 11 82800 026




EUROPEAN REPRESENTATIVE: MDSS-MEDICAL DEVICE SAFETY SERVICE GMBH
SCHIFFGRABEN, 41, 30175, HANNOVER, GEMANY

START OF CE-MARKING: 2014-9-16

PLACE, DATE OF DECLARATION: ZHONGSHAN, 2017-3-2

SIGNATURE:

NAME: 
POSITION: Vice President

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Standards applied:

Risk management	EN ISO 14971:2012
Labeling	EN 980: 2008
User manual	EN 1041: 2008
General requirements for safety	EN 60601-1: 2006/IEC 60601-1:2005+A1: 2012 EN 60601-1-11:2011
Non-invasive sphygmomanometers General requirements	EN 1060-1:1995+A2:2009 EN 1060-3:1997+A2:2009 IEC/EN 80601-2-30:2009
Electromagnetic compatibility	EN 60601-1-2:2007
Usability	EN 60601-1-6:2010 & EN 62366:2007
Software life-cycle	EN 62304:2006+AC: 2008
Biological evaluation	EN ISO 10993-1:2009 EN ISO 10993-5:2009 EN ISO 10993-10:2010
Clinical Investigation	MEDDEV.2.7.1: 2009 EN 1060-4: 2004