Declaration of Conformity

According to the Medical Devices Directive 93/42/EEC

Manufacturer's Name :	**HK Greatmade Tech Limited**
Manufacturer's Address :	**3th, B Building ,Baifuli Industiral Zone,shanghenglang village , LongHua town .**
	Shenzhen City , Guang Dong Province , China
Product :	**Non-Invasive Blood Pressure Cuff and Air Hose**
Type Designation/Trademark :	**N.I.B.P cuff , Class I**

Product Part No. Of Manufacturer: CF001A, CF002A, CF003A, CF004A,CF004LA,CF004TA , CF001B,CF002B,CF003B, CF004B, CF004LB, CF004TB, CF001C,CF002C,CF003C,CF004C, CF004LC,CF004TC,CF005,CF006.CF007A,CF007B,CF008,CF009,CF010,CF011.CF012,CF013,CF014,CF015,CF016 ,CF1601A/B,CF1602A/B ,CF1701A,CF1702B,CF006-H ,CF017,CF018,CF019 ,CF020. CF1501A/B,CF1502A/B,CF1503A/B, CF1504A/B, CF1505A/B, CF1506A/B, CF1507A/B, CF1508A/B,CF1509A/B,CF1510A/B,CF1512A/B,CF1101A,CF1102A,CF1103A,CF1104A,CF1105A,CF1106A/B,CF1107A/B,CF108A/B,CF1109A/B,CF1110A/B,CF1111A/B,CF1112A/B.CF1601A/B,CF1602A/B,CF1603A/B,CF1701-500A/A1,CF1701-1000A/A1,CF1701-3000A/A1,CF1701-500A1-BL,CF1701-1000A1-BL,CF1701-3000A1-BL,CF1801,CF021,CF022,CF023,CF025,CF026

Authorized representative established within the EU (if applicable):

Company Name :	**Prolinx GmbH**
Company Address :	**Brehmstr. 56,40239, Duesseldorf ,Germany**

Person responsible for making this declaration

Name, Surname :	**ZhangHanzhi**
Position/Title :	**Director , Owner**

Hereby we declares that the Medical device as indicated above conforms with the essential requirements listed in the Annex I of the European Medical Device Directive 93/42/EEC

Shenzhen China

Zhang Hanzhi

__Mar 19th,2018__

(Place) **(Date)**

(Company stamp and legal signature)