EU Declaration of Conformity

according to the Medical Devices Directive 93/42/EEC

(We)

Zhejiang Shuyou Surgical Instrument Co.,Ltd

3689#, East Yangguang Avenue, Dipu street Anji 313300

Huzhou city, zhejiang province China

(European Representative): Luxus Lebenswelt GmbH

Kochstr. 1, 47877, Willich, Germany

Declare under the product

Disposable Electrodes

(Class IIb by Annex IX,MDD93/42/EEC)

Caution:

The undersigned hereby declares that the medical device as Specified above conforms With the essential requirements listed in the Annex II excluding section 4 of the European Medical Device Directive 93/42EEC(MDD) ;

This declaration of conformity is based on the European Medical Device Directive 93/42EEC,Annex <II> excluding section 4 and is supported by a Notified body Tüv ;

Rheinland Product Safety GmbH Annex <II> excluding section 4 Certificate, with reference to articles 1 and 3 of the MDD ;

Notified Body: TUV Rheinland LGA Products GmbH, ID Number 0197

Conformity Assessment Certificate: 1.EC Certificate HD 60140608 0001

2.EN ISO13485 QMS Certificate SX 60152607 0001

General Manager

Of **Zhejiang Shuyou Surgical Instrument Co.Ltd**

2020.11.02 in Anji

(place and date of issue)

Suying

(name and signature or equivalent marking of authorized person)

This document covers the following product models

No.	Product model
1	MN01-FG001401
2	WN01-FG003001
3	WN02-FG002202
4	WN01-FG003201
5	MN01-FG001501
6	WN01-FG000602