
EC DECLARATION OF CONFORMITY

The following EC declaration of conformity exemplifies the required content according to Directive 93/42/EEC.

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Jafron Biomedical Co.,Ltd.**
No. 98, Technology Sixth Road, High-tech Zone,
519085, Zhuhai, Guangdong, China

We declare under our sole responsibility that

the medical device: **Disposable Hemoperfusion Cartridge**
model:
HA60, HA80, HA100, HA130, HA150, HA180,
HA230, HA280, HA330, HA330- II , HA380, HA430,
HA480

of class: /

II b

according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure: / **Directive 93/42/EEC Annex II, excluding Section 4**

Registration No.: **HD 60138860 0001**

Notified Body: **TÜV Rheinland LGA Products GmbH**
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197

European Authorized Representative : **Shanghai International Holding Corp. GmbH (Europe)**
Eiffestrasse 80, 20537 Hamburg, Germany
Tel: +49-40-2513175

Zhuhai 2022-08-19

Place, date /

Jianhua Cai

Management Representative

Name and function