



EC DECLARATION OF CONFORMITY

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

We declare under our sole responsibility that

the medical device: **Disposable Hemoperfusion Cartridge**
model:
HA80, HA130, HA180, HA230, HA280, HA330,
HA330-II, HA380, HA430, HA480
**Disposable Plasma Bilirubin Perfusion Adsorption
Column**
model:
BS80, BS330, BS380

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)**
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Zhuhai 2019-08-07
Place, date /

Janhua Cao Management Representative
Name and function

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