

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Ningbo MFLAB Medical Instruments Co., Ltd.
No.508, Yindong Road(N), Yinzhou Economic
Development Zone, Ningbo315145, China**

We declare under our sole responsibility that
the medical device:

Disposable Vacuum Blood Collection Systems

of class:

Article 9.1(for all devices other than those covered by
Annex II and devices for performance evaluation,
according to directive 98/79/EC

meets the provisions of the directive 98/79/EC Annex III 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: 98/79/EC Annex III

Registration No.: **SX601503630001**

Notified Body:

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

宁波蓝柏医疗器械有限公司
NINGBO MFLAB MEDICAL INSTRUMENTS CO.,LTD

Ningbo 2020-10-30

Place, date

李雅增

Name and function