



湖北省潜江市江赫医用材料有限公司
HUBEI QIANJIANG KINGPHAR MEDICAL MATERIAL CO, LTD

Qianjiang, 2023-06-16

Declaration of Conformity

We

Hubei Qianjiang Kingphar Medical Material Co., Ltd.
Yuanguang Road, 433100, Qianjiang, P. R. China
SRN (Single Registration No): CN-MF-000009051

Manufacturer according to Medical Device Regulation (EU) 2017/745

hereby declare in our own responsibility
that the products

Askina® Vliesstoffkompressen

Non sterile non woven swabs are ideally used for primary care of acute wounds, for the absorption of blood and exudate and for cleansing the skin's surface. For single use.

Basic UDI-DI: 694878811002PQ

(article numbers see attachment I)

are in conformity with the requirements of the Medical Device Regulation (EU) 2017/745

Conformity Assessment Procedure

according to article 52 section 7
of the Regulation named above

Classification

according to annex VIII of the Regulation named above
Class I


Authorized Representative for the European Union (Article 11)

Shanghai International
Holding Corp. GmbH (Europe)
Eiffenstraße 8
20537 Hamburg
Germany

Valid until 2026-06-15

Name: Wang Zhongxiao

Function: President

Signature: 



湖北省潜江市江赫医用材料有限公司
HUBEI QIANJIANG KINGPHAR MEDICAL MATERIAL CO, LTD

Attachment I

Art. No.	Product name	Class
9034722	Askina® Vliesstoffkompressen	I
9034730	Askina® Vliesstoffkompressen	I
9034706	Askina® Vliesstoffkompressen	I
9034714	Askina® Vliesstoffkompressen	I
9034749	Askina® Vliesstoffkompressen	I