



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

Declaration of Conformity

We

Hubei Qianjiang Kingphar Medical Material Co., Ltd.
Yuanguang Road, 433100, Qianjiang, P. R. China

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

hereby declare in our own responsibility
that the products

Askina® Mullkompressen

Non-sterile gauze compresses 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: 694878811001PN
(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 2023-06-15

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



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Attachment I

Art. No.	Product name	Class
9031308	Askina® Mullkompressen	I
9031316	Askina® Mullkompressen	I
9031324	Askina® Mullkompressen	I
9033017	Askina® Mullkompressen	I
9033025	Askina® Mullkompressen	I
9033041	Askina® Mullkompressen	I