FAZ LUXE

Medical Gauze Bandage (Sterile & None-sterile)

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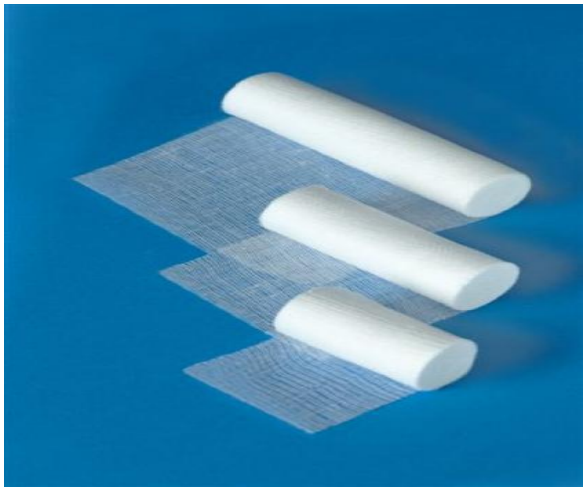

- **Specification:** Gauze bandages are made of medical gauze, in accordance with the National State Standard GOST. Sterile bandages are intended to protect wound from contamination, infection and mechanical impacts, while the non-sterile bandages are intended for fixing of dressings. For use in hospitals the bandages are produced in multiple packaging. For packaging of the bandages a three-layer material is used.
- **Material:** 100% cotton cloth.
- **Yarn:** Yarn cotton card straight lonely from pneumomechanical stretching machines tex. 20; Nm 50/1; Ne 29,5.
- **Density:** 17 g/cm², 23 g/cm², 24 g/cm², 27 g/cm², 30 g/cm², 32 g/cm², 34 g/cm², 36 g/cm².
- **Color:** Bleached white
- **Edge:** Unprocessed edge
- **Basic packing:** 1 roll/pack, 5 roll/pack, 6 roll/pack, 10 roll/pack, 12 roll/pack, 20 roll/pack,

25 roll/pack, 30 roll/pack. (individually packed in PE pocket)

- **Full size:** Length: 1 m, 2 m, 3 m, 4 m, 5 m, 7 m, 10 m.
- **Width:** 5 cm, 7 cm, 7,5 cm, 8,5 cm, 10 cm, 12 cm, 14 cm, 16 cm.
- **Certificate:** ISO 9001:2015; ISO 13485:2016; IQ Net 9001; IQ Net 13485; CE.
- **Note:** Personalized specifications are possibly at customer`s request.

Digitally signed by Ermicev Alexandr
Date: 2022.09.21 16:42:58 MSK
Reason: MoldSign Signature
Location: Moldova

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FAZO LUXE

СИСТЕМА СЕРТИФИКАЦИИ РУССКОГО РЕГИСТРА
RUSSIAN REGISTER CERTIFICATION SYSTEM

CERTIFICATE

This is to certify that the Quality Management System of

"FAZO-LUXE" LLC
21, Okhangrabo Str., Zangiota district, 111103, Gulistan community,
Tashkent region, Republic of Uzbekistan

has been assessed and found to be in accordance
with the requirements of

ISO 13485:2016

in respect of production and sales of polyethylene single-use shoe covers, sterile and non-sterile medical gauze bandages, non-sterile medical woven bandages, non-sterile medical absorbent cotton wool and bleached medical gauze

No: 17.0680.026
of 5th May, 2017

This certificate is valid until 5th May, 2020

Director General of Certification
Association "Russian Register"

Specification of the certification scope is provided in Annex. This certificate becomes invalid if conditions of certification are not fulfilled (http://www.rusregister.ru/doc/004.00-105.pdf). This Certificate is the property of Certification Association "Russian Register".

Certification Association "Russian Register": 101 Rimskogo-Korsakova Ave., 190121, Saint Petersburg, Russia

10-000066

RUSSIAN REGISTER РУССКИЙ РЕГИСТР

IQNet®

THE INTERNATIONAL CERTIFICATION NETWORK

CERTIFICATE

IQNet and
Certification Association "Russian Register"
hereby certify that the organization

"FAZO-LUXE" LLC
21, Okhangrabo Str., Zangiota district, 111103, Gulistan community,
Tashkent region, Republic of Uzbekistan

for the following field of activities

production and sales of polyethylene single-use shoe covers, sterile and non-sterile medical gauze bandages, non-sterile medical woven bandages, non-sterile medical absorbent cotton wool and bleached medical gauze

has implemented and maintains a

Management System

which fulfils the requirements of the following standard

ISO 13485:2016

Issued on : 5th May, 2017
Validity date : 5th May, 2020

Registration Number : **RU-17.0680.026**

Michael Drechsel
President of IQNet

Arkady Vladimirtsev,
Director General of
Russian Register

IQNet Partners*:
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Quality Austria *Austria* RR *Russia* SIGE *Mexico* SII *Israel* SIQ *Slovenia* SIRIM QAS International *Malaysia*
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* The list of IQNet partners is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com

FAZO LUXE

MDSS
Medical Device Safety Service

Certificate of CE-Registration

This is to certify that, in accordance with the Medical Device Directive 93/42/EEC, Medical Device Safety Service GmbH (MDSS) agrees to perform all duties and responsibilities as the Authorized Representative for:

Fazo-Luxe LLC
Mr. Nodir Yunusov
Ohangrabo Str 21, Tashkent district

Tashkent region
UZBEKISTAN

as stipulated and demanded by the aforementioned Directive. The German Competent Authority has allocated the medical devices of the Manufacturer registration numbers as foreseen in:

Annex A July 24 2018

The Manufacturer has provided MDSS with the appropriate Declaration(s) of Conformity confirming that the medical devices fulfill the applicable requirements of Directive 93/42/EEC. In compliance with German law, a safety officer has been appointed for Germany.

2018-07-24

Ludger Möller
President
MDSS GmbH