13 pozicija

Yafho
WOUND CARE

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YAFHO BIO-TECHNOLOGY CO., LTD.

Tel:86-20-85295649 Fax:86-20-85295797
http://www.yafho.com E-mail:info@yafho.com

Drape

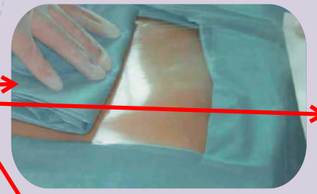
Incise Drape

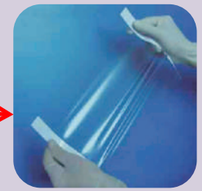
The surgical film is made of medical glue and polyurethane film, poliuretano plėvelė padengta akriliniais klijais
which is impermeable to bacteria, permeable to oxygen.
It is used for protecting the wound site from the bacteria and dirt in an operation.
apsauga nuo bakterijų pateikimo operacijos metu

Features:
1- Waterproof
2- Transparent
3- Elastic
4- Breathable

13 poz

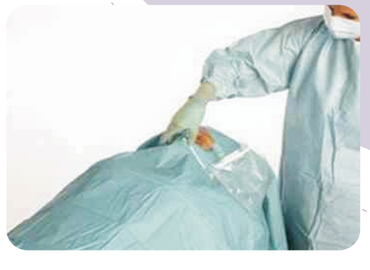
Ref. (iodine)	Ref.	Size(cm)	Package
632455	630455	45x45	20pcs/box,18boxes/ctn
632230	630230	20x30	30pcs/box,18boxes/ctn
632003	630003	28x45	20pcs/box,18boxes/ctn
632005	630005	45x55	10pcs/box,12boxes/ctn
632006	630006	28x30	20pcs/box,12boxes/ctn
632340	630340	30x40	20pcs/box,12boxes/ctn
632098	630098	32x45	20pcs/box,18boxes/ctn
632024	630024	96x60	10pcs/box,10boxes/ctn
632001	630001	10x14	100pcs/box,18boxes/ctn
632182	630182	14x12	100pcs/box,18boxes/ctn
632002	630002	15x28	30pcs/box,18boxes/ctn





Eye Drape

Ref.	Size(cm)	Package
630183	85x65	10pcsx18box,180pcs/ctn



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Yafho WOUND CARE

EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60148977 0001

Report No.: 17047222 012

Manufacturer:

Yafho Bio-Technology Co., Ltd.
Second Floor Room 202 and
Third Floor
No. 81, Junfeng Road, Huangpu District
Guangzhou
510760 Guangdong
P.R. China

Products:

Aspects of manufacture concerned with securing and maintaining sterile conditions of Wound Dressings, Surgical Drapes,Sterile Instrument Covers

Replaces Approval, Registration No.: DD 60144595 0001

Expiry Date: 2024-05-26

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2020-08-28

Date: 2020-08-28

Notified Body

Herbert Zhong

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.