

### 1. PRODUCT

**Standart Surgical Gown** 

#### 2. PRODUCT CONFIRMITIES

ISO 9001, ISO 13485, EN 13795, ISO 11135:2014, ISO 14644-3:2019, ISO 14791:2012, ISO 14937: 2009, ISO 10993-7:2008, EN 20811

#### 3. DESCRIPTION:



Products	Surgical gown, Full Protection, assure air permeability Adjustable, with velcro closure at the neck, it is provided with 4 ties. Ultrasonic seems, Raglan type. Including two towels 40*40 and CSR Wrapping 60*60 cm  Validity 5 years from the date of production
Fabric	SMS
Weight	42 gr/m2.
Color	Blue – 150 +-10 Cm



#### 4. PACKING

The product is individually packaged. On the packaging is the inscription with the name and address of the manufacturer, the product model, description, dimensions, date of manufacture, validity date, lot number, etc.

### **5. STORAGE**

0°C - + 30°C, Avoid Sunlight, Avoid Humidity

### 6. Term of validity

5 years from the date of production









# **QUALITY MANAGEMENT SYSTEM CERTIFICATE**

Universal GmbH

This certificate is granted to the organization,

### Zeyni Medikal Tekstil Ins. San. Ve Tic. Ltd. Sti.

Yunus Emre Mah. 12. Yildirim Sok. No 17 Yildirim-Bursa/Turkey

by review of RA1-5076 numbered report for the scope

Manufacture And Sales Of Disposable Surgical Sterile Gowns, Drapes And Custom Packs, Non-Sterile Medical Textile Products

> to certify that a management system in accordance with standard's clauses is established and being implemented

> > **DIN EN ISO 9001:2015**

Certificate No: QMS 0520 006861

Original Certification Date: 22.04.2020

**Revised Date:** 13.04.2022

**Expiry Date**: 21.04.2024

**Certification Period**: 3 years (1st year)









# MEDICAL QUALITY MANAGEMENT SYSTEM CERTIFICATE

Universal GmbH

This certificate is granted to the organization,

Zeyni Medikal Tekstil Ins. San. Ve Tic. Ltd. Sti.

Yunus Emre Mah. 12. Yildirim Sok. No 17 Yildirim-Bursa/Turkey

by review of RA1-5076 numbered report for the scope

Manufacture And Sales Of Disposable Surgical Sterile Gowns, Drapes And Custom Packs, Non-Sterile Medical Textile Products

to certify that a management system in accordance with standard's clauses is established and being implemented

**DIN EN ISO 13485:2016** 

Certificate No: MDMS 0423 009795

Original Certification Date: 22.04.2020

**Revised Date :** 13.04.2022

**Expiry Date**: 21.04.2024

**Certification Period**: 3 years (1st year)







Medical Devices Regulation (EU) 2017/745 Annex XI Part A (Class Is, Im and Ir Devices)



Certificate Number: M.2022.MDR.1007

Manufacturer Name : Zeyni Medikal Tekstil İnş. San. Ve Tic. Ltd. Şti.

Manufacturer Address : Yunusemre Mah. 12. Yıldırım Sk. No:17 Yıldırım BURSA / TURKEY

Single registration number-SRN : TR-MF-000017474

Authorised Representative Name

(If applicable)

Authorised Representative Address : NA

**Product Scope** : See the product list on the following page(s).

: NA

Based on the conformity assessment for the abovementioned manufacturer's quality assurance system in accordance with (EU) 2017/745 Medical Devices Regulation Annex XI Part A, UDEM Adriatic d.o.o hereby declares that the requirements of Annex XI Part A of the Regulation (EU) 2017/745 have been met for the listed products in this certificate.

The manufacturer has established, documented and implemented a quality management system, which is subject to periodic surveillance assessments by UDEM Adriatic d.o.o. according Annex XI Part A Section 7 of the aforementioned Regulation.

For the devices covered by this certificate, the involvement of UDEM Adriatic d.o.o. in the conformity assessment procedures is limited: in the case of devices placed on the market in sterile condition, to the aspects relating to establishing, securing and maintaining sterile conditions; in the case of devices with a measuring function, to the aspects relating to the conformity of the devices with the metrological requirements; in the case of reusable surgical instruments, to the aspects relating to the reuse of the device, in particular cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use.

The report referenced below summarizes the result of assessments/examinations and includes reference to relevant CS, harmonized standards and test reports.

 Report Number
 : MDR.1130

 Date of Issue
 : 22/11/2022

Recertification Date : Reissue Date/No : -

**Date of Expiry** : 21/11/2027

If any, Previous Certificate(s) No: NA

UDEM Adriatic d.o.o. General Manager



UDEM Adriatic d.o.o. is a Notified Body (identification no 2696) under (EU) 2017/745 Medical Devices Regulation.

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Medical Devices Regulation (EU) 2017/745 Annex XI Part A (Class Is, Im and Ir Devices)



Certificate Number: M.2022.MDR.1007

### PRODUCT LIST COVERED BY THE CERTIFICATE

PRODUCT NAME	BASIC UDI-DI	RISK CLASS	EMDN CODE	WODEL	TYPE	INTENDED PURPOSE
	868 234012 101001GZ 868 234012 101002H3 868 234012 101003H5 868 234012 101004H7 868 234012 101005H9 868 234012 101006HB 868 234012 101019HL 868 234012 101020H5 868 234012 101022H7 868 234012 101022H9 868 234012 101023HB 868 234012 101023HB	Class Is	T0299	SHIRT & PANTS SHORTS	S/M/L/ XL/ XXL/ XXXL	Examination Clothes are used to wear during the surgical operations and examination purposes as sterile (Helping prevention to infection risk).
	868 234012 102001H8 868 234012 102002HA 868 234012 102003HC	Class Is	T030502	BEDDING DUVET COVER PILLOW CASE COVER SHEET		The bedding sets are that patients have to use before / after surgical operations in the operating room. It is completely used for patients to minimize the risk of infection.
Sterile Disposable Surgical Drapes	868 234012 101007HD 868 234012 101008HF 868 234012 101009HH 868 234012 101010H2 868 234012 101011H4 868 234012 101012H6 868 234012 101013H8 868 234012 101014HA 868 234012 101015HC 868 234012 101016HE 868 234012 101017HG 868 234012 101017HG	Class Is	T04010103	COLONOSCOPY PANTS COLONOSCOPY SHORTS	S/M/L/ XL/ XXL/ XXXL	Examination Clothes are used to wear during the surgical operations and examination purposes as sterile.  Examination Clothes are sterile products which use conditions as minimizing the risk of microbial contamination and present the protection and safety. So sterile Examination Clothes are using conditions as minimizing the risk of microbial contamination.
	868 234012 103001BCQ 868 234012 103001TDU 868 234012 103001SDS 868 234012 103002BCT 868 234012 103002TDX 868 234012 103002SDV	Class Is	T020199	BIFLEX PLAIN DRAPE TRIPLEX PLAIN DRAPE SMS PLAIN DRAPE	Width: Min:20 Max:200 Length: Min:20 Max:250	Sterile Plain drapes are used in
	868 234012 103003BCW 868 234012 103003TE2 868 234012 103003SDY 868 234012 103004BCZ 868 234012 103004TE5 868 234012 103004SE3 868 234012 103005BD4 868 234012 103005TE8			BIFLEX DRAPE WITH SIDE ADHESIVE TAPE TRIMPLEX DRAPE WITH SIDE ADHESIVE TAPE SMS DRAPE WITH SIDE ADHESIVE TAPE	Width: Min:20 Max:200 Length: Min:20 Max:300	operating rooms to cover the needed area of the patients during surgical operations, to protect the patient and the surgical team around him from situations that threaten their health and prevent any infection from spreading to the patient, surgical team and the operating environment.
	868 234012 103005SE6 868 234012 103006BD7 868 234012 103006TEB 868 234012 103006SE9 868 234012 103007BDA 868 234012 103007TEE 868 234012 103007SEC			SMS FENESTRATED DRAPE BIFLEX FENESTRATED DRAPE	Width: Min:20 Max:200 Length: Min:20 Max:300	

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868 234012 103008BDD			Width:	
868 234012 103008TEH			Min:22	
	1		Max:75	
868 234012 103008SEF	1 1			
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868 234012 103009TEL	1 1		Min:100	
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868 234012 103018BDJ	1 1			
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Medical Devices Regulation (EU) 2017/745 Annex XI Part A (Class Is, Im and Ir Devices)



Certificate Number: M.2022.MDR.1007

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868 234012 1030271EW			
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868 234012 104002HQ			
868 234012 104003HS			
868 234012 104004HU			
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868 234012 104007J2			
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868 234012 104009J6			<i>i</i>
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868 234012 104019J9			
868 234012 104020HS			
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868 234012 104022HW			
868 234012 104023HY			
868 234012 104024J2			
868 234012 104025J4			
868 234012 104026J6			
868 234012 104027J8			
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868 234012 104039JF			
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868 234012 104045JA			
868 234012 104046JC			- 3
868 234012 104047JE			
868 234012 104048JG			
868 234012 104049JJ			
868 234012 104050J3			
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	868 234012 104062JA					
	868 234012 104063JC					
	868 234012 104064JE					
	868 234012 104065JG					
	868 234012 104066JJ					
	868 234012 104067JL					
	868 234012 104068JN					
	868 234012 104069JQ					
	868 234012 104070J9					
	868 234012 104071JB					
	868 234012 104072JD					
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	868 234012 104074JH					
	868 234012 104075JK					
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1	868 234012 105017JC				1	
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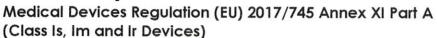
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868 234012 105035JE			
868 234012 105036JG			
868 234012 105037JJ			
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868 234012 105071JJ			
868 234012 105072JL	1	1	
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868 234012 105075JS			
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868 234012 105079K2			
868 234012 105080JK			
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868 234012 105082JP			
868 234012 1050833K	1		
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868 234012 105092JS			
868 234012 105093JU			
868 234012 105094JW			
868 234012 105095JY			
868 234012 105096K2			

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	868 234012 105097K4					
	868 234012 105098K6					
	868 234012 105099K8		l			
	868 234012 105100HY			1		
	868 234012 108048KC			1		
	868 234012 108049KE		1			
1	868 234012 108050JX					
	868 234012 108051JZ			1		
	868 234012 108052K3			1		
	868 234012 108053K5					
	868 234012 108054K7					
1	868 234012 106001J4					
	868 234012 106002J6			1	Width:	
	868 234012 106003J8	1			Min:75	Sterile Table Covers are used in
	868 234012 106004JA		1		Max:200	operating rooms to cover the
	868 234012 106005JC	Class Is	T030102	TABLE COVER		instrument table in order to keep safe sterile area to keep put the
	868 234012 106006JE				Length:	instruments on while surgical
	868 234012 106007JG				Min:90	operation.
	868 234012 106008JJ	İ			Max:230	
	868 234012 106009JL			4.		
					Width:	
		1			Min:35 Max:120	
			1		MUX.120	
				INSTRUMENT COVER	Length:	
			=	(FLOROSCOPY OR	Min:35	
	868 234012 107001JB			IMAGE INTENSIFIER COVER)	Max:230	
	868 234012 107002JD					
	868 234012 107003JF				Diamete	
	868 234012 107004JH				r: Min:Ø 60 –	
	868 234012 107005JK				Max:Ø 80	Sterile covers are used in operating rooms to cover the instruments for keeping the instruments safe and sterile
	868 234012 107006JM					
	868 234012 107007JP			CAMERA COVER	13x250 ir	
	868 234012 107008JR					
	868 234012 107009JT	Class Is	T030101			during Surgery operations and
	868 234012 107010JC		1000101		Width:	create environment that is the
	868 234012 107011JE			SCOPY COVER	Min:35 most sterile who max:150 prone to infecti	most sterile when the patients are
	868 234012 107012JG	i				prone to infection while their
	868 234012 107013JJ					wounds open at operation.
	868 234012 107014JL		-		Length:	
	868 234012 107015JN				Min:35	
	868 234012 107016JQ				Max:150	30
	868 234012 107017JS					.
				MICROSCOPE COVER	2250 x	
				HILDROGOFE COVER	1220	
				20	Ø100x	
	!			C-ARM COVER	225x	-
					Ø100	
	868 234012 108001JJ			ABDOMINAL SURGICAL	annoneconomico	
	868 234012 108002JL			DRAPE	200x300	
	868 234012 108003JN			LAPAROTOMI	2507200	Sterile Plain drapes are used in
	868 234012 108004JQ 868 234012 108005JS 868 234012 108007JW			SURGICAL DRAPE	350X300	operating rooms to cover the
				LAPAROSCOPY	250X300	needed area of the patients
				SURGICAL DRAPE	200/1000	during surgical operations, to
		Class Is	T020102	MINOR SURGERY DRAPE	75X75	protect the patient and the surgical team around him from
	868 234012 108009K2	-,		DRESSING DRAPE	60X60	situations that threaten their
	868 234012 108010JK			ARTHROSCOPY DRAPE		health and prevent any infection
	868 234012 108011JM			WITHOUT POUCH	200X300	from spreading to the patient,
	868 234012 108012JP			EXTREMITY DRAPE	200X300	surgical team and the operating
	868 234012 108013JR			LAIREIVIII I DRAFE	200A300	environment.
	868 234012 108014JT			HAND SURGERY DRAPE	200X300	

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Certificate Number: M.2022.MDR.1007

	868 234012 108015JV	T	T	IA IEE GUD CERVICE	T	T
	868 234012 108016JX			SHOULDER SURGERY	200X300	4
	868 234012 108017JZ			DRAPE SURGERY	200X300	
	868 234012 108018K3 868 234012 108020JN			HIP SURGERY DRAPE	200X250	
	868 234012 108021JQ 868 234012 108022JS			ANGIOGRAHY SURGERY DRAPE	150X240	
	868 234012 108023JU 868 234012 108024JW			BY PASS SURGERY DRAPE	240X350	
	868 234012 108025JY			VALVE REPLACEMENT SURGERY DRAPE	240X350	
	868 234012 108026K2 868 234012 108027K4			EMBRIO TRANSFER (E.T.) SURGERY DRAPE	50X90	
	868 234012 108028K6			I.V.F SURGERY DRAPE	150X225	
				DELIVERY SURGERY DRAPE	75X150	
				CASEREAN SURGERY DRAPE WITHOUT POUCH	200X300	
				OVUM PICK UP (O.P.U) SURGERY DRAPE	130X150	
				CRANIOTOMY SURGERY DRAPE	230X300	
				SHUNT SURGERY DRAPE	150X300	
				SPINAL SURGERY DRAPE	200X300	
				PERCUTAN (PCNL) SURGERY DRAPE	150X300	
				TUR SURGERY DRAPE	200X240	
				CYSTOSCOPY SURGERY DRAPE	75X150	
				E.N.T. SURGERY DRAPE	150X200	
				ARTHROSCOPY DRAPE WITH POUCH	200X300	
	868 234012 108006JU			CASEREAN SURGERY DRAPE WITH POUCH	200X300	
	868 234012 108019K5					
	868 234012 108029K8 868 234012 108030JR 868 234012 108031JT 868 234012 108032JV 868 234012 108033JX 868 234012 108034JZ 868 234012 108035K3 868 234012 108036K5 868 234012 108037K7 868 234012 108037K7 868 234012 108039KB 868 234012 108040JU 868 234012 108040JU 868 234012 108041JW 868 234012 108041JW 868 234012 108041JW 868 234012 108045K6 868 234012 108045K6 868 234012 108045K6 868 234012 108045K6		T02010101			
				SMS EYE SURGERY DRAPE (SINGLE POUCH)	Width: Min:100 Max:150	Sterile Plain drapes are used in operating rooms to cover the needed area of the patients during surgical operations, to protect the patient and the
		Class is			Length: Min:120 Max:240	surgical team around him from situations that threaten their health and prevent any infection from spreading to the patient, surgical team and the operating environment.
				SMS EYE SURGERY DRAPE (DOUBLE POUCH)	Width: Min:100 Max:150 Length: Min:120 Max:240	

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				BIFLEX EYE SURGERY DRAPE (SINGLE POUCH)	Width: Min:100 Max:150 Length: Min:130 Max:240	
				BIFLEX EYE SURGERY DRAPE (DOUBLE POUCH)	Width: 150 Length: Min:150 Max:240	
Sterile	868 234012 100001GS 868 234012 100002GU 868 234012 100003GW 868 234012 100004GY 868 234012 100005H2 868 234012 100005H2 868 234012 100007H6 868 234012 100009HA 868 234012 100009HA 868 234012 100010GT 868 234012 100011GV 868 234012 100013GZ 868 234012 100013GZ 868 234012 100015H5 868 234012 100015H5 868 234012 100015H5 868 234012 100015H5 868 234012 100015H6 868 234012 100055HH 868 234012 100059HR	Class Is	T020401	STANDARD SURGICAL GOWN FULL ULTRASONIC STANDARD SURGICAL GOWN HALF ULTRASONIC STANDARD SURGICAL GOWN STANDARD CHEMOTHERAPY GOWN	S/M/L/ XL/ XXL/ XXXL	Sterile gowns are used to wear /cover during the surgical operations and examination purposes as sterile (Helping prevention to infection risk). Gowns are being barrier between sterile and non-sterile areas and helping prevention to infection risk used as sterile products. So sterile gown use conditions as minimizing the risk of microbial contamination.
Disposable Surgical Gowns	868 234012 100019HD 868 234012 100020GW 868 234012 100021GY 868 234012 100023H2 868 234012 100023H4 868 234012 100024H6 868 234012 100024H6 868 234012 100025H8 868 234012 100027HC 868 234012 100027HC 868 234012 100029HG 868 234012 10003GZ 868 234012 10003GZ 868 234012 100033H3 868 234012 100033H5 868 234012 100033H7 868 234012 100033H7 868 234012 100035HB 868 234012 100035HB 868 234012 100035HB 868 234012 100037HF 868 234012 100037HF 868 234012 100039HK 868 234012 100049H4 868 234012 100044H6 868 234012 100044H6 868 234012 100044HC 868 234012 100044HC 868 234012 100044HC	Class Is	T020402	REINFORCED SURGICAL GOWN FULL ULTRASONIC REINFORCED SURGICAL GOWN HALF ULTRASONIC REINFORCED SURGICAL GOWN FULL REINFORCED SURGICAL GOWN FULL ULTRASONIC FULL REINFORCED SURGICAL GOWN HALF ULTRASONIC FULL REINFORCED SURGICAL GOWN REINFORCED CHEMOTHERAPY GOWN FULL REINFORCED CHEMOTHERAPY GOWN GOWN	S/M/L/ XL/ XXL/ XXXL	Reinforced gowns are the surgical gowns which operators and nurses has to wear in all kinds of surgical operations in operation area. Sterile gowns are used to wear/cover during the surgical operations and examination purposes as sterile (helping prevention to infection risk). They cover doctors, nurses and the patients totally in order to minimize the infection risk. Gowns are being barrier between sterile and non-sterile areas. Polyethylene is used for penetrating the water absorption.

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	868 234012 100046HG	T	T		T	
	868 234012 100047HJ					
	868 234012 100048HL					
	868 234012 100049HN					
	868 234012 100050H7					
l	868 234012 100051H9					
	868 234012 100052HB					
	868 234012 100053HD					
	868 234012 100054HF					
l	868 234012 100061HC					
	868 234012 100062HE					
	868 234012 100063HG					
	868 234012 100064HJ	1				
İ	868 234012 100065HL					
	868 234012 100066HN					
	868 234012 100067HQ					
	868 234012 100068HS					
	868 234012 100069HU					
	868 234012 100070HD			2		
	868 234012 100071HF					
	868 234012 100072HH					
				SHORT SLEEVE PATIENT		
				EXAMINATION GOWN CUFFS PATIENT		Examination Clothes are used to
	868 234012 100073HK			EXAMINATION GOWN		wear during the surgical
l	868 234012 100074HM	011-	T000 400	LONG SLEEVE PATIENT		operations and examination
	868 234012 100075HP	Class Is	T020499	EXAMINATION GOWN		purposes as sterile
	868 234012 100076HR			LONG SLEEVE RUBBER		(Helping prevention to infection
	868 234012 100077HT			PATIENT EXAMINATION		risk.
				GOWN VISITOR GOWN		
	868 234012 200001H7					Charles Hair and Constant days
	868 234012 210001HJ			A. UNIVERSAL SURGICAL DRAPE SETS:		Sterile Universal Surgical drape sets are used in operating rooms
	868 234012 210002HL			1.UNIVERSAL SURGICAL		to cover the needed area of the
	868 234012 210003HN			DRAPE SET		patients during General Surgery
	868 234012 210004HQ					operations.
	868 234012 210005HS			B. GENERAL SURGICAL		
	868 234012 210006HU			DRAPE SETS:		Sterile General Surgical Drape sets
	868 234012 210007HW			1.GENERAL SURGICAL DRAPE SET		are used in operating rooms to cover the needed area of the
	868 234012 210008HY			2. ABDOMINAL		patients during General Surgery
	868 234012 210009J2			SURGICAL DRAPE SET		operations.
	868 234012 220001HV			3. LAPAROTOMY		•
	868 234012 220002HX			SURGICAL DRAPE SET		Sterile Abdominal Surgery drape
	868 234012 220003HZ			4. LAPAROSCOPY SURGICAL DRAPE SET		sets are used in operating rooms
	868 234012 220004J3			5. MINOR SURGICAL		to cover the needed area of the patients during Abdominal
	868 234012 220005J5			DRAPE SET		Surgery operations.
Sterile	868 234012 220006J7			6. DRESSING SURGICAL		
	868 234012 220007J9	Class is	T020102	DRAPE SET		Sterile Laparotomy Surgical Drape
Surgical	868 234012 220008JB	Olds is	1020102	7. DIALYSIS COVER		sets are used in operating rooms
Drape Sets	868 234012 220009JD			SURGICAL DRAPE SET 8. ABDOMINAL		to cover the needed area of the patients during laparotomy
	868 234012 230001J8			PERINEAL SURGICAL		Surgery operations.
	868 234012 230002JA			DRAPE SET		obligary operations.
	868 234012 230003JC			9.CIRCUMCISION		Sterile Laparoscopy Surgical
	868 234012 230004JE			COVER SURGICAL		Drape sets are used in operating
	868 234012 230005JG			DRAPE SET		rooms to cover the needed area
	868 234012 240001JK			C. ORTHOPEDY		of the patients during Laparoscopy Surgery operations.
	868 234012 240002JM			SURGICAL DRAPE SETS:		Laparoscopy solgery operations.
	868 234012 240003JP			1. ARTHROSCOPY		Sterile Minor Surgical drape sets
	868 234012 240004JR			DRAPE SET WITHOUT		are used in operating rooms to
	868 234012 240005JT			POUCH		cover the needed area of the
	868 234012 240006JV 868 234012 250001JW			2.ARTHROSCOPY		patients during Minor Surgery
	868 234012 250001JW			DRAPE SET WITH POUCH  3. LOWER EXTREMITY		operations.
	868 234012 250002J1 868 234012 250003K2			DRAPE SET		Sterile Dressing surgical drape sets
	868 234012 260001K9			4. UPPER EXTREMITY		are used after operations while
	000 2040 12 20000 TK7					,

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868 234012 270001KL
868 234012 280001KX
868 234012 280002KZ
868 234012 280003L3
868 234012 290001LA
868 234012 300001HL
868 234012 300002HN
868 234012 301001HT
868 234012 301002HV
868 234012 301003HX
868 234012 302001J2
868 234012 302002J4

DRAPE SET
5. HAND SURGERY
DRAPE SET
6. KNEE SURGERY
DRAPE SET
7. SHOULDER SURGERY
DRAPE SET
8. ARTHROPLASTY (HIP)
DRAPE SET
9. VERTEBRA DRAPE SET

D. CARDIOLOGY
SURGICAL DRAPE SETS;
1. ANGIOGRAPHY
DRAPE SET
2. ANGIOGRAPHY
CUSTEM SET
3.BY PASS DRAPE SET
4. VALVE REPLACEMENT
DRAPE SET
5. CARDIOVASCULAR
DRAPE SET

E. GYNECOLOGIC
SURGICAL DRAPE SETS:

1. EMBRIO TRANSFER
(E.T.) DRAPE SET
2. I.V.F DRAPE SET
3. DELIVERY DRAPE SET
4.CAESAREAN DRAPE
SET
5.O.P.U. (OVUM PICK
UP) DRAPE SET
6.PERINEAL SET

F. BRAIN SURGERY DRAPE SETS: 1. CRANIATOMY DRAPE SET 2. SHUNT DRAPE SET 3. SPINAL DRAPE SET

1.EYE DRAPE SET

H. ONCOLOGHY
SURGICAL DRAPE SETS:
1.CHEMOTHERAPHY

DRAPE SET

G. OPHTHALMOLOGY

SURGICAL-DRAPE SETS:

I. UROLOGICAL SURGICAL DRAPE SETS: 1.PERCUTANEUS DRAPE SET

2. TUR DRAPE SET
3. CYSTOSCOPY DRAPE
SET

J. RADIOLOGY SURGICAL DRAPE SETS: 1.BIOPSY DRAPE SET

K. EAR- NOSE-THROAT (E.N.T.) DRAPE SETS: 1.EAR-NOSE-THROAT (E.N.T.) DRAPE SET 2.THYROID DRAPE SET

L. ESTETIHCAL SURGERY DRAPE SETS: 1. RHYNOPLASTY DRAPE re-dressing the operated place and named as Dressing Set. The content has included all required items in order to use while cleaning and for re-dressing the operated area.

Sterile Dialysis Cover Surgical drape sets are used in operating rooms to cover the needed area of the patients during General Surgery operations.

Sterile Abdominal Perineal
Surgical drape sets are used in
operating rooms to cover the
needed area of the patients
during Gynecological Operations
for Ovum Pick Up operations.

Sterile Circumcision Surgical Cover drape sets are used in operating rooms to cover the needed area of the patients during General Surgery operations.

Sterile Arthroscopy Drape Sets Without Pouch are used in operating rooms to cover the needed area of the patients during Arthroscopic Surgery operations, draping in shoulder.

Sterile Arthroscopy Drape Sets With Pouch are used in operating rooms to cover the needed area of the patients during Arthroscopic Surgery operations.

Sterile Lower Extremity drape sets are used in operating rooms to cover the needed area of the patients during lower Extremity Surgery operations.

Sterile Upper Extremity drape sets are used in operating rooms to cover the needed area of the patients during Upper Extremity Surgery operations

Sterile Hand Surgery drape sets are used in operating rooms to cover the needed area of the patients during upper extremity surgery operations

Sterile Knee Surgery drape sets are used in operating rooms to cover the needed area of the patients during Knee Surgery operations

Sterile Shoulder Surgery drape sets are used in operating rooms to cover the needed area of the patients during Shoulder Surgery operations.

Sterile ARTHROPLASTY (HIP) drape sets are used in operating rooms

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SET
2. LIPOSCULPTURE
DRAPE SET
3. HAIR TRANSPLANT
DRAPE SET

M. DENTAL SURGERY DRAPE SETS:

1.DENTAL DRAPE SET 2.IMPLANT DRAPE SET

1 PAIR CHEMOTHERAPHY GLOVES 1 PAIR SURGERY **GLOVES LARGE** 1 PCS ABDOMINAL DRAPE 2 PCS ALCOHOLIC SWAB 3 PCS ANESTHESIA DRAPE 1 PCS ANESTHESIA DRAPE WITH ADHESIVE TAPE 1 PCS ANGIOGRAPHY DRAPE 1 PCS ARTHROSCOPY DRAPE WITH POUCH 1 PCS ARTHROSCOPY DRAPE WITHOUT POUCH 1 PCS BABY DRAPE 1 PCS BETADINE SPONGE 1 PCS BONNET 1 PCS CAESAREAN DRAPE 1 PCS CHEMOTHERAPHY DRAPE 1 PCS CRANIOTOMY DRAPE 1 PCS CYSTOSCOPY DRAPE 2 PCS DRAPE 2 PCS DRAPE WITH ADHESIVE

2 PCS DRAPE WITH

1 PCS EXTREMITY DRAPE 1 PCS EYE DRAPE WITH

I PCS FACE MASK

1 PCS FLUOROSCOPY

2 PCS HAND TOWEL

I PCS HOLE DRAPE

1 PCS I.V.F. DRAPE 2 PCS TOOLS TABLE

WITH ADHESIVE TAPE

1 PCS K.V.C. DRAPE

1 PCS LEG DRAPE

I PCS LAPAROSCOPY

ADHESIVE TAPE 4 PCS DRAPE WITH SIDE

ADHESIVE TAPE

1 PCS ELASTIC

BANDAGE

POUCH

COVER

DRAPE

DRAPE 2 PCS LEG COVER to cover the needed area of the patients during Hip replacement Operations.

Sterile Vertebra drape sets are used in operating rooms to cover the needed area of the patients during Vertebra Surgery operations.

Sterile Angiography drape sets are used in operating rooms to cover the needed area of the patients during vascular visualization / Angiography Surgery operations.

Sterile Angiography custem drape sets are used in operating rooms to cover the needed area of the patients during vascular visualization / angiography operations.

Sterile By-pass drape sets are used in Cardiology / By-pass Surgical Operations for making new veins by-passing the blood circulation system and examination.

Sterile Valve Replacement drape sets are used in Cardiology / Valve Replacement Surgical Operations for replacing the valves in order to improve blood circulation system

Sterile Cardiovascular drape sets are used during Cardiology / All kinds of Cardiovascular Surgical operations, for making new veins by-passing the blood circulation system.

Sterile Embrio Transfer (E.T.) drape sets are used in operating rooms for Gynecological Operations transferring the embrio.

Sterile I.V.F. drape sets are used in Gynecological I.V.F operations.

Sterile Delivery drape sets are used in Gynecological Surgery operations for delivering the baby.

Sterile Caesarean drape sets are used during Gynecological / Caesarean Operations as Caesarean type of delivering the baby or others.

Sterile O.P.U drape sets are used in Gynecological Operations for Ovum Pick Up.

Sterile Perineal drape sets are used for surgical operations and examinations in perineal area such as saucerization of perineal or ischiorectal abscress, evacuation of thrombotic piles, pile ligation, gynecological

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	ADHESIVE TAPE		dilation and curettage etc.
	2 PCS LEGGINGS STANDARD		Storila Craniatamy drana sats are
	1 PCS MARKER & RULER		Sterile Craniotomy drape sets are used in open Brain Surgery
	1 PCS MAYO TABLE		Operations.
	DRAPE		Operations.
	1 PCS MINOR HOLE		Sterile Shunt drape sets are used
	DRAPE		in open Brain Surgery / Shunt
	1 PCS LAPAROTOMY		placement Surgery Operations
	DRAPE		and examinations.
	1 PCS LAPAROSCOPIC		aria oxariiranoris.
	ABDOMINAL PERINEAL		Sterile Spinal drape sets are used
	DRAPE		in Spine / Open Brain Surgery
	1 PCS NITRILE GLOVES		Operations.
	1 PCS NONWOVEN OP-		10007
	TAPE		Sterile Eye drape sets are used in
	1 PCS O.P.U. DRAPE		all kinds of Ophthalmology
	1 PCS OPAQUE SET		Operations.
	8 PCS OP-TAPE		
	2 PCS PACK OF	l	Sterile Chemotherapy drape sets
	STERILIZATION		are used while injecting
	1 PCS PATIENT DRAPE		chemotherapy drugs to the
	1 PCS PATIENTS MOUTH		patients to cover both the patient
	HOLE DRAPE		and the nurse for protection.
	1 PCS PERCUTANEOUS DRAPE		Storilo Boroutano que drano este
	1 PCS PERINE U DRAPE		Sterile Percutaneous drape sets are used in Urological Operations
	1 PCS PLAIN DRAPE		such as Nephrology surgery and
	1 PCS PROTECTIVE		examinations.
	GLASSES		examinations.
	2 PCS REFLECTOR		Sterile Spinal drape sets are used
	DRAPE		in Urological Operations such as
	3 PCS REINFORCED		T.U.R Surgery.
	SURGERY GOWN LARGE		
	3 PCS REINFORCED		Sterile Spinal drape sets are used
	SURGICAL GOWN		in Urological Operations such as
	1 PCS RHINOPLASTY		Cystoscopy operation.
	DRAPE		
	1 PCS SCALPEL TIP		Sterile Biopsy drape sets are used
	NO:11		in Biopsy Surgery Operations.
	1 PCS SCOPY COVER		01. 11. 5.12. 1
ı	1 PCS SHOULDER DRAPE		Sterile E.N.T drape sets are used in
- 1	1 PCS SHUNT DRAPE 1 PCS SIDA DRAPE		E.N.T Surgery Operations.
	4 PCS SIDE DRAPE WITH		Sterile Thyroid drape sets are used
	ADHESIVE TAPE		in Thyroid Surgery Operations.
	1 PCS SIDE TAPED		in myloid sorgery operations.
	DRAPE		Sterile Rhynoplasty drape sets are
	1 PCS SOLUTION CUP		used in Rhinoplasty Surgery
	250CC		Operations.
	1 PCS SOLUTION CUP		8
	500CC		Sterile Liposculpture drape sets
	10 PCS SPANCH		are used in Liposuction Surgery
	1 PCS SPINAL DRAPE		Operations.
	1 PCS SPLIT E.N.T. DRAPE		
	1 PCS SPONGE		Sterile Hair Transplant drape sets
	2 PCS STANDARD		are used in Hair transplant Surgery
- 1	SURGERY GOWN LARGE		Operations.
	1 PCS STANDART SURGERY GOWN		Starila Dantal draps sets are used
- 1	1 PCS STERILIZATION		Sterile Dental drape sets are used in Dental Surgery Operations.
	WRAPE		The Defind Surgery Operations.
	1 PCS STOCKINITTE (FOR		Sterile Implant drape sets are
-	HAND)		used in Implant Surgery
	I PCS T.U.R. DRAPE		Operations.
	1 PCS TABLE COVER		
	I PCS TABLE COVER		
	DRAPE		
	1 PCS THYROID DRAPE		
	4 PCS TOWEL		
	2 PCS TRANSPARENT		
- 1	DRAPE		

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HOSE COVER 1 PCS TUBE HOLDER (VELCRO) 1 PCS UNDER ARM DRAPE 2 PCS UNIVERSAL BAG 1 PCS UPPER EXTREMITE DRAPE 1 PCS VIDEO CAMERA COVER 1 PCS VINYL GLOVES 1 PCS WASTE BAG	
2 PCS WOUND PAD	

Conditions for or limitations to the validity of this certificate

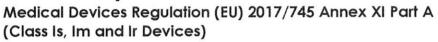
:NA

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	CERTIFICATE HIST	ORT
Rev. No.	Rev. Date	Description of Revision
00	22/11/2022	Initial Certification

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# **EC Declaration of Conformity**

issued in accordance with EC directive (EU) 2017/745 relating to Medical **Devices Regulation** 

Manufacturer:

Zeyni Medical Tekstil Ins. San. Ve Tic. Ltd. Sti.

Adress:

Yunus Emre Mah. 12. Yildirim Sk. No:17 Yildirim

Bursa/ TURKEY

**Commercial Name:** 

Zeyni

**Product Name:** 

Sterile Disposable Surgical Gowns

- **Standard Surgical Gowns**
- Full Ultrasonic Standard Surgical Gowns
- Half Ultrasonic Standard Surgical Gowns
- Standard Chemotherapy Gowns
- Reinforced Surgical Gown
- Full Ultrasonic Reinforced Surgical Gown
- Half Ultrasonic Reinforced Surgical Gown
- Full Reinforced Surgical Gown
- Full Ultrasonic Full Reinforced Surgical Gown
- Half Ultrasonic Full Reinforced Surgical Gown
- Reinforced Chemotherapy Gown
- Full Reinforced Chemotherapy Gown

ZEYNİ MEDİKAL TEKSTİL İNŞ. SANAYİ VE TİCARET LTD.ŞTİ. Yunusemre Mh. 12. Yıldınm Sk. No: 17 Yıldınm - BURSA / TÜRKİYE Tel.: 0224. 369 32 32 Faks: 0224. 369 32 36 Mobil: +90533 143 53 56 e-mail: info@zeynimedical.com.tr • www.zeynimedical.com.tr





Class I Sterile Medical Devices according to the Directive (EU) Classified as:

2017/745 Medical Devices Regulation

Applied directives: The Directive (EU) 2017/745 on medical devices, conformity

assesment according to Annex VII

Applied Harmonized Standards:EN ISO 13485:2016, EN ISO 15223-1:2016, EN 1041:2008+A1:2013

EN ISO 14155:2011 EN ISO 10993-1:2018, EN ISO 10993-5:2009 EN

ISO 10993-10:2014, EN 13795-1:2019

Notified Body: UDEM Adriatic d.o.o. Radnicki cesta 54/R3 Zagreb-Crotia

**Notified Body Number:** CE 2696

**EC Certificate:** M.2022.MDR.1007

**EC Certificate Validity:** 21.11.2027

The Company Zeyni Medical Tekstil Ins. San. Ve Tic. Ltd. Sti. Herewith declares that the above mentioned product meets all aplicable provisions of the Directive (EU) 2017/745. The product is safe under prescribed and reasonably foreseeable conditions of storage and use.

The company has implemented measures assuring that all the products of the mentioned type are safe and fulfil essential requirements of the (EU) 2017/745 Directive. If the device is modified without the agreement of the undersign, this declaration become invalid in relation to the modified product.

Date of issue: 22.11.2022

Baki Polat Canavar, CEO

On behalf of Company

ZEYNIMEDIKAL TEKSTIL INSAAT SAN VOTIC. LTD. STI. Yunusemre Mah 72 Xidyirin Sk. No:17 Yudiyiri BUTSSA Yildirin V.D.399 741 1400 Tic. Sic. No:102917 Mergis No:0998141140000001

ZEYNİ MEDİKAL TEKSTİL İNŞ. SANAYİ VE TİCARET LTD.ŞTİ.

Yunusemre Mh. 12. Yıldırım Sk. No: 17 Yıldırım - BURSA / TÜRKİYE Tel.: 0224, 369 32 32 Faks: 0224, 369 32 36 Mobil: +90533 143 53 56 e-mail: info@zeynimedical.com.tr • www.zeynimedical.com.tr





This Certificate has been awarded to

### ZEYNİ MEDİKAL TEKSTİL İNŞAAT SANAYİ VE TİCARET LİMİTED ŞİRKETİ

YUNUSEMRE MAHALLESİ 12. YILDIRIM SOKAK NO:17
YILDIRIM / BURSA / TÜRKİYE

In recognition of the organization's Management System which complies with

EN 13795-1:2019

The scope of activities covered by this certificate is defined below

MANUFACTURING, SALES OF DISPOSABLE, STERILE AND NON-STERILE SURGICAL GOWNS, INSULATED VISITOR GOWNS AND COVERS

TEK KULLANIMLIK, STERİL VE STERİL OLMAYAN CERRAHİ ÖNLÜK, İZOLASYONLU ZİYARETÇİ ÖNLÜĞÜ VE ÖRTÜLERİN İMALATI, SATIŞI

Certificate Number: SISTUREN082020161 Date of Issue of Original Certificate: 21.08.2020 Date of Issue of latest certificate: 23.09.2021

Expiry Date: 20.08.2022

Re-certification Due on: 21.07.2023

Managing Director

Note: This certificate is valid only if produced with the continuation letter after the surveillance is carried out successfully.

The Organization's documentation and Implementation has been reviewed and found to comply with the relevant standard rules. This certificate of Compliance is based on the evaluation of the mentioned scope given above. Organization is responsible for maintaining the responsibilities of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid

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Report No.

Page 1 of 6

### ZEYNİ MEDİKAL TEKSTİL İNŞ.SAN. VE TİC. LTD. ŞTİ.

111699085

YUNUSEMRE MAH. 12 YILDIRIM SOKAK NO:17 YILDIRIM/BURSA

Buyer	/
Test Item. :	Bilayered and Absorbent Drapes
Item No.:	/
Colour Name. :	/
Condition at delivery. :	Samples tested as received.
Test Scope. :	Parameters selected by customer
Test Specification:	Determination of hydrostatic pressure
Applicant's Provided - Care Instruction/Label:	

2021-10-18 (p.m)

Testing Period: 2021 10-21 to 2021-10-25

Sample Receiving date:

Test Result: Passed

For and on behalf of TÜV Rheinland Uluslararası Standartlar Sertifikasyon ve Denetim A.Ş

Tomris Hasançebi / Customer Relations Manager

Abdullah Akil / Physical Laboratory Manager

ashit

/ 2014-07-02 Version No / Date: 1.0

Kozyatagi Mah. Saniye Ermutlu Sok. No:12

Colakoglu Plaza B Blok 34742 Kadikoy Istanbul,

Tel. +902166653200, Fax +902166653299, e-mail: info@tr.tuv.com

TÜVRheinland®
Precisely Right.

Report No.: 111699085 Date: 10.25.2021

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Report No.: 111699085

Date: 10.25.2021

Material List: Page 3 of 6

Material No.	Material	Color	Location
M001	Textile	-	Textile
M002	Textile + coating	-	Textile w /coating

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Report No.: 111699085

Date: 10.25.2021

Conclusion: Page 4 of 6

**TEST PROPERTY** M001 M002 Hydrostatic Pressure Test # #

TÜVRheinland®
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Report No.: 111699085

Date: 10.25.2021

1.Hydrostatic Pressure Test Page 5 of 6

Test method : EN 20811:1993

Face Side: Original

Test Conditions: 65 cm2 Test Speed:65 cmH2O/dak

M001 <u>M002</u> <u>Requirement</u>

Pressure

Average 65 cm H2O 65 cm H2O

- END -

Report No.: 111699085

Date: 10.25.2021



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Esenyurt Firuzköy Bulvarı No:29 34325 Avcılar İstanbul/TÜRKİYE

TEST REPORT
DENEY RAPORU

20018576 -ing-Add

12-05

### **EKOTEKS**

Customer name: ZEYNİ MEDİKAL TEKTSİL İNŞ. SANAYİ VE TİC. LTD. ŞTİ.

Address: Yunusemre Mah. 12. Yıldırım Sk. No:17 Yıldırım - BURSA

Buyer name:

Contact Person: Davut Daşdan

Order No:

Article No:

Name and identity of test item: Surgical Gowns-SMS

The date of receipt of test item: 12.05.2021

Re-submitted/re-confirmation

date:

**Date of test:** 12.05.2021-12.05.2021

Remarks:

Sampling: The results given in this report belong to the received sample by vendor.

End-Use:

Care Label: Not specified.

Number of pages of the report: 14

SealDateCustomer RepresentativeHead of Testing Laboratory12.05.2021Servin YURTSEVENSevim A. RAZAK<br/>12.05.2021

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05-26

REQUIRED TESTS	RESULT	COMMENTS
MICROPIOLOGICAL TECT		
MICROBIOLOGICAL TEST		
Microbial Cleanliness (Bioburden) (1)	P	
Wet-Bacterial Penetration <sup>(1)</sup>	P	
PHYSICAL PROPERTIES TESTS		
Tensile Stregth / Dry	P	
Tensile Stregth / Wet	P	
Bursting Strength / <b>Dry</b>	P	
Bursting Strength / Wet	P	

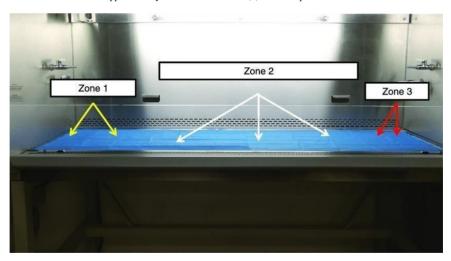
P: Pass

F: Fail

R: Refer to retailer technologist.

- (1) This report was reissued to add this test result.
- (2) Test results were evaluated according to EN 13795-1:2019 Standard Performance Properties Critical Sample Group limit values (Table 1)

REMARK: Original samples are kept for 3 months and all technical records are kept for 5 years unless otherwise specified. If requested, measurement uncertainty will be reported. But unless otherwise specified, measurement uncertainty is not considered while stating compliance with specification or limit values The reported uncertainty is based on a standard uncertainty multiplied by a coverage factor k=2, providing a level of confidence of approximately 95 %. Tests marked (\*) in this report are not included in the accreditation schedule.



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### **TEST RESULTS**

### **MICROBIAL CLEANLINESS (Bioburden)**

Test Metod: Ref: EN ISO 11737-1:2018 (\*)

The sample is put in extraciton liquid after shaking well, inoculated on the agar. After incubation at 30  $\pm$  1  $^{\circ}$  C for 72 hours, growth microorganisms are counted on the agar.

	<u>RESULTS</u>	REQUIREMENT
Microbial cleanliness (cfu/g)	206 cfu/100 cm <sup>2</sup>	≤300 cfu/100 cm <sup>2</sup>
1	208	
2	211	
3	206	
4	204	
5	198	
6	202	
7	203	
8	202	
9	201	
10	200	

<sup>\*</sup>cfu= Colony forming unit.

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### **TEST RESULTS**

**Test Method:** BS EN 22610: 2006 (Surgical gowns, garments and fresh air clothes used as medical devices for patients, hospital staff and equipment - Test method for determination of resistance to wet bacterial permeability) (\*)

A test sample is placed on the agar plate on a rotating disc. Bacteria carrier material and coating film are placed on the test sample and all parts are fixed on the disk. A finger is placed on the test sample to apply a certain force  $(3N \pm 0.02)$ . The finger moves on the test sample over the entire surface of the agar within 15 minutes. 5 studies are carried out for 15 minutes. 6. The study is repeated by inverting the sample.

Sample amount:	5 pieces 25x25cm2
Carrier Material:	30 μm thin, 25x25cm2 Polyurethane Film
Coating Material:	25x25cm2 HDPE Film
Microorganism:	Staphylococcus aureus ATCC 29213
Bacterial Concentration (kob / ml):	1-4x104 kob / ml
Incubation Conditions:	(36 ± 1) ° C 48 hours

RESULTS			
Number of Populating Bacteria (cfu)		Penetration Rate	
X1	55	RCUM1	0,01
X2	72	RCUM2	0,15
X3	154	RCUM3	0,34
X4	178	RCUM4	0,56
X5	156	RCUM5	0,74
Z	212		
T	827		

X1 ...... X5: Number of colonies growing in 5 parallel petri in the same sample

Z: number of colonies growing in the sixth petri dish

T: X1 + X2 + X3 + X4 + X5 + Z

RCUM1 = X1/T

RCUM2 = (X2 + X1)/T

RCUM3 = (X3 + X2 + X1)/T

RCUM4 = (X4 + X3 + X2 + X1)/T

RCUM5 = (X5 + X4 + X3 + X2 + X1)/T

### BARRIER INDEX (IB )

	Result	Expected value (*)
IB	4,14	≥2,8

IB = 6 - (CUM1 + CUM2 + CUM3 + CUM4 + CUM5)

\* EN 13795-1:2019 Surgical gowns - Requirements and test methods are evaluated according to Table-1.

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12-05

### **TEST RESULTS**

**TEST METHOD: EN 13795-1:2019** 

### SURGICAL GOWN -REQUIREMENTS AND TEST METHODS

### **ANNEX 1: SURGICAL GOWNS(\*);**

### TENSILE STRENGTH; EN 29073-3:1996 (\*)

Instron 5969 (Load: 50 kN), Strip Method. Speed: 100 mm/min±10, Gauge length 200 mm.

Pre-load was not applied. Without wetting samples.

The average results are given for weft and warp direction of five samples

Performed in the conditioned room (20±2°C-65%±4).

Dry;

Diy,	RESULT	REOUIREMENT
Weft	52.2 N	≥20N (Dry)
Warp	93.6 N	$\geq$ 20N (Dry)

### TENSILE STRENGTH; EN 29073-3:1996 (\*)

Instron 5969 (Load: 50 kN), Strip Method. Speed: 100 mm/min±10, Gauge length 200 mm.

Speed. 100 mm/mm±10, Gauge length 200 mm.

Pre-load was not applied. With wetting samples.

The average results are given for weft and warp direction of five samples

Performed in the conditioned room (20±2°C-65%±4).

Wet;

,	RESULT	REOUIREMENT
Weft	55.0 N	≥20N (Wet)
Warp	101.9 N	$\geq$ 20N (Wet)

#### BURSTING STRENGTH;; ISO 13938-1:1999

SDL ATLAS M229 tester. Test area: 30.5 mm diameter

Rate of increase in volume; 29 cm<sup>3</sup>/min. The average results are given of five samples.

Performed in the conditioned room (20±2°C-65%±4).

**RESULT Dry**; 141.2 kPa  $\geq$  40 kPa (Dry)

Height at Burst\* 12.9 mm

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12-05

### TEST RESULTS

**TEST METHOD: EN 13795-1:2019** 

SURGICAL GOWN -REQUIREMENTS AND TESTMETHODSANNEX

1: SURGICAL GOWN (\*);

BURSTING STRENGTH;; ISO 13938-1:1999

SDL ATLAS M229 tester. Test area: 30.5 mm diameter

Rate of increase in volume; 45.2 cm³/min. The average results are given of five samples. Performed in the conditioned room (20±2°C-65%±4).

RESULT

REOUIREMENT

Wet; 139.4 kPa  $\geq 40 \text{ kPa} \text{ (Wet)}$ 

Height at Burst\* 14.3 mm

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12-05

The resistance to dry microbial penetration [Material]Test Method: EN ISO 22612:2005

#### **Test principle:**

The test is carried out on test pieces each fixed in a container. In every container except one a portion of talc contaminated with Bacillus subtilis is poured on the test piece. One container is left uncontaminated as a control. A sedimentation plate is inserted at the base of each container at a short distance below the test piece. The apparatus supporting the containers is then vibrated by a pneumatic ball vibrator. The talc that penetrates is captured on the sedimentation plate, the sedimentation plates are removed and incubated. The numbers of colonies produced are counted.

### **Test equipment:**

Resistance to dry microbial penetration test Incubator Electronic balance Autoclave

#### The environmental conditions of the laboratory and test condition:

Test environment temperature: 24.5°C, Relative humidity: 56.0%

Test environment monitoring: total bacteria: 0 CFU/plate, total fungi: 0 CFU/plate, blank experiment: aseptic growth

Culture medium: TGE agar medium; Other materials: talc and ethyl alcohol.

Dimensions of the test specimens: 200mm×200mm

Sample: 12 pieces

Vibration frequency: 20800 times/min; Vibration time: 30 min.

Test bacteria: The fourth generation of spores of bacillus subtilis ATCC 9372

Concentration of bacterium: 1.8×10<sup>8</sup> CFU/g

20018576 ing-Add

### **Results:**

Sample	Measured value (CFU)	Requirement (CFU)	Conclusion
1	19		
2	17	<200	
3	31	≤300	
4	15		
5	10	(Surgical Gown performance	D
6	25	less critical productarea)	Pass
7	16	-	
8	28	EN 13795-1:2019	
9	17		
10	8		

20018576 – ing-Add

12-05

The resistance to wet bacterial penetration[Material]Test Method: EN ISO 22610:2006

#### **Test principle:**

A test specimen is placed on an agar plate. A sheet of donor material, of corresponding size and carrying the bacteria, is placed on the test specimen with the contaminated side face down and covered by a sheet of approximately 10 µm high density polyethylene (HDPE) film. Two tithing conical steel rings hold the three sheets together, applying a tensile force. An abrasion-resistant finger is placed on top of the materials with a specified force to bring the test specimen in contact with the agar. The finger is moved over the entire surface of the plate in less than 15 min by means of a pivoted lever moved by an exocentric cam. The assemblage of materials, stretched by the weight of the steel rings, ensures that only a small area of the test specimen is brought into contact with the agar surface at any one time. Due to the combined effect of rubbing and liquid migration, bacteria may pass from the donor material through the test specimen down to the agar surface. After being tested for 15 min, the agar plate is replaced by a fresh one, and the test is repeated with the same donor and test specimen. Allowing 15 min for each test, five tests are performed with the same pair of donor and test specimen. In this way, the test allows for an estimation of the penetration over time. Finally, the bacterial contamination on the top side of the test specimen is estimated using the same technique. The agar plates are incubated in order to observe the bacterial colonies, which are then enumerated.

#### **Test equipment:**

The resistance to wet bacterial penetration test Incubator Electronic balance Autoclave

### The environmental conditions of the laboratory and test condition:

Test environment temperature: 24.5 °C, Relative humidity: 56.0%

Test environment monitoring: total bacteria: 0 CFU/plate, total fungi: 0 CFU/plate, blank experiment: aseptic growth

Culture medium: Tryptone Soya Agar, Tryptic Soy Broth, peptone water and nutrient agar medium.

Dimensions of the test specimens: 25cm×25cm

The carrier material: solvent-cast polyurethane (PU) film of 30  $\mu m$  thickness

Nutrient agar to from the brim: 3 mm

Test bacteria: The fifth generation of staphylococcus aureus ATCC 29213

Concentration of bacterium: 2.3×10<sup>4</sup> CFU/ml

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12-05

### **Results:**

Sample	Barrier index	Requirement Barrier index	Conclusion	
1	6.7	≥6		
2	6.6	(Surgical Gown: performance	Pass	
3	6.6			
4	6.6	critical product area)		
5	6.6	EN 13795-1:2019		

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12-05

### Lint and other particles generation in the dry state[Material]

**Test Method:** EN ISO 9073-10:2004

### **Test principle:**

This procedure describes a modified Flex method in which the sample is subjected to a combined twisting and compression action in a test chamber. During the flexing, air is withdrawn from chamber and particulates in the air stream are counted and classified in a particle counter. Depending on the choice of counter, the size ranges can fall within the limits of  $0.3 \, \mu m$  or  $0.5 \, \mu m$  to  $25 \, \mu m$ .

### **Test equipment:**

Flex tester with particle counter

### The environmental conditions of the laboratory:

Test environment temperature: 20.2°C, Relative humidity: 64.7%

20018576 ing-Add

### **Results:**

Size of particles counted (µm)	Sample		Measured value Coefficient of linting log <sub>10</sub>	Requirement Coefficient of linting log <sub>10</sub>	Conclusion
		1	2.1		
		2	2.1	<u>-</u> ≤4.0	
	A: Face	3	2.1	_ ≥4.0	
		4	2.1		
2 25		5	2.2	(Surgical Gown: performance	Pass
3~25		1	2.0	critical product area)	
		2	2.1	1	
	B: Face	3	2.0	1	
		4	2.2	EN 13795-1:2019	
		5	2.1		

20018576 – ing-Add

12-05

#### Static hydrostatic resistance[Material]Test

**Method:** EN ISO 811:2018

#### **Test principle:**

The hydrostatic head supported by a fabric is a measure of the opposition to the passage of water through the fabric. A specimen is subjected to a steadily increasing pressure of water on one side of the fabric, under standard conditions, until penetration occurs in three places. The pressure at which the water penetrates the fabric at the third place is noted.

### **Test equipment:**

Hydrostatic tester

Water, grade 3 water in accordance with ISO 3696.

#### The environmental conditions of the laboratory and test condition:

Pretreatment: Condition the test specimens at 20.1°C air at 65.2% RH for 24 h

Face side tested

Temperature of the water: 20.0℃

Rate of increasing water pressure: 10cmH<sub>2</sub> O/min

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12-05

### **Results:**

Sample	Measured value (cmH <sub>2</sub> O)	Requirement (cmH <sub>2</sub> O)	Conclusion
1	125 18	≥100 ≥10	Pass
		(Surgical Gown performance critical product area) EN 13795-1:2019	