



Banca Comercială „ENERGBANK” Societate pe Acțiuni
Republica Moldova, MD-2001, mun. Chișinău, str. Tighina 23/3
tel.: +(373 22) 544-377, 858-000; fax: +(373 22) 858-080
www.energbank.com; e-mail: office@energbank.com
IDNO 1003600008150, cod TVA 0202040
codul băncii: ENEGMD22
cod IBAN: MD10NB00000000035215845 la CD BNM
Capital social - 100 000 000 lei MD

Nr. 13101-08/180
din « 19 » 01 2017

“M-INTER-FARMA” S.A.
MD-2028, mun. Chișinău, str. Grenoble, 23

Prin prezenta B.C. “ENERGBANK” S.A. (codul băncii ENEGMD22), confirmă că “M-INTER-FARMA” S.A., cod fiscal 1003600005263 deține codul IBAN MD37EN00000022245246845 în Lei MD, Dolari SUA, EURO, Ruble Ruse, Hrivne Ucrainene, Lire Sterline și Franci Elvețieni.

Contabil șef adjuncț -
Șef Direcție Monitorizarea Tranzacțiilor



Vitalie Bulgaru



Fax: Gajinevschi Luminițe

Tel: (22) 858-020

REPUBLICA



MOLDOVA

CERTIFICAT DE ÎNREGISTRARE

PRIN PREZENTUL SE CERTIFICĂ, CĂ SOCIETATEA PE
ACIUNI "M-INTER-FARMA" ESTE ÎNREGISTRATĂ LA CAMERA
ÎNREGISTRĂRII DE STAT

Numărul de indentificare de stat - codul fiscal
1003600005263

Data înregistrării

12.05.1994

Data eliberării

17.12.2004

Iovu Galina, registrator de stat

*Functia, numele, prenumele persoanei
care a eliberat certificatul*

G. Iovu
semnatura

MD 0006726





I.P. "AGENȚIA SERVICII PUBLICE"

Departamentul înregistrare și licențiere a unităților de drept

EXTRAS

din Registrul de stat al persoanelor juridice

nr. 1638 din 28.01.2019

Denumirea completă: **SOCIETATEA PE ACȚIUNI «M-INTER-FARMA».**

Denumirea prescurtată: **«M-INTER-FARMA» S.A.**

Forma juridică de organizare: **Societate pe Acțiuni.**

Numărul de identificare de stat și codul fiscal: **1003600005263.**

Data înregistrării de stat: **12.05.1994.**

Sediul: **MD-2028, str. Grenoble, 23, mun. Chișinău, Republica Moldova.**

Modul de constituire: **nou creată.**

Obiectul principal de activitate:

- 1 **Practica medicală;**
- 2 **Comerțul cu ridicata al produselor agricole brute și animalelor vii;**
- 3 **Comerțul cu amănuntul al produselor farmaceutice;**
- 4 **Comerțul cu ridicata al produselor farmaceutice;**
- 5 **Activitatea farmaceutică;**
- 6 **Alte activități de asistență medicală;**
- 7 **Comerțul cu amănuntul în magazine nespecializate, cu vânzare predominantă de produse alimentare, băuturi și produse din tutun;**
- 8 **Practica stomatologică;**
- 9 **Comerțul cu ridicata al parfumurilor și produselor cosmetice;**
- 10 **Importul, fabricarea, comercializarea, asistența tehnică și (sau) reparația dispozitivelor medicale și (sau) a opticii;**
- 11 **Importul și (sau) fabricarea, depozitarea, comercializarea angro a substanțelor și materialelor chimice, toxice, articolelor și produselor chimice de menaj.**

Capitalul social: **5160000 lei.**

Administrator: MATEI VASILE, IDNP 0961110897556.

Prezentul extras este eliberat în temeiul art. 34 al Legii nr. 220-XVI din 19 octombrie 2007 privind înregistrarea de stat a persoanelor juridice și a întreprinzătorilor individuali și confirmă datele din Registrul de stat la data de: 28.01.2019.

Specialist coordonator
tel. 022-207-840



Lazar Aliona



CERTIFICAT

privind lipsa sau existența restanțelor față de bugetul public național

Nr.
№ A1919415

din
от 02.05.2019

1. Destinatar / Получатель

ACHIZIȚII PUBLICE

2. Date despre contribuabil / Информация о налогоплательщике

Denumirea Наименование	Codul fiscal / Numărul de identificare Фискальный код / Идентификационный номер
M-INTER-FARMA S.A.	1003600005263
Adresa sediului de bază (strada, numărul) Адрес основного месторасположения (улица, номер)	Codul - Denumirea localității Код - Наименование населенного пункта
Grenoble nr.23	0130-SEC.CENTRU

**3. Atestarea lipsei sau existenței restanțelor conform datelor Sistemului Informațional Automatizat /
Подтверждение отсутствия или наличия недоимки согласно данных Автоматизированной Информационной Системы**

La data emiterii prezentului certificat restanța la bugetul public național constituie/ На дату выдачи данной справки недоимка перед национальным публичным бюджетом составляет:
0,00 lei/лей.

4. Valabil pînă la / Действителен до 17.05.2019

5. Autentificarea organului fiscal / Подтверждение налогового органа

Sef DDF CENTRU

L.S./M.L.
Execuție



[Signature]
Semnătura/Подпись

VIORICA CĂUȘ

Numele și prenumele/Фамилия и имя

Este extras din Sistemul Informațional al SFS SIA „Contul curent al contribuabilului”// 02.05.2019 ora 14:18:05
cu aplicarea prevederilor pct. 82-83 Ordin IFPS nr.400 din 14.03.2014 (Monitorul Oficial 72-77/399, 28.03.2014)

NOTA (0,00)



TECHNICAL DATA SHEET

Latex surgical gloves powder-free

MANUFACTURER:	TG MEDICAL SDN BHD (Malaysia)
CATALOGUE NUMBER:	TG-0260, TG-0265, TG-0270, TG-0275, TG-0280, TG-0285, TG-0290
CLASSIFICATION:	- Class IIa Medical Device in accordance with MDD 93/42/EEC - Personal Protective Equipment Category III according to Directive 89/686/EEC
METHOD OF STERILIZATION:	Gamma radiation
CURRENT TECHNICAL STANDARDS ARE APPLICABLE TO THE DEVICE:	<ul style="list-style-type: none"> in compliance with requirements of ASTM D3577, EN 455, resistant to penetration of viruses in accordance with ASTM F1671, resistant to penetration of chemicals according to EN 374-3, in accordance with EN 420, tested in accordance with EN 388 Quality Assurance System: Manufacturing process in accordance with US FDA Quality System Regulation (QSR), BS EN ISO 9001 Quality System and EN ISO 13485

PERFORMANCE CHARACTERISTICS:

- Flexible, soft, with non-slip external surface, anatomically shaped, fit perfectly to the hand, providing the user with comfort of work and at the same time provide a protective barrier against microorganisms
- Powder free - avoid skin irritation and postoperative complications caused by the powdering agent
- Easy putting on thanks to the internal polymer layer
- Designed for wide use in surgery



TECHNICAL DATA

RAW MATERIAL:	natural rubber latex
COLOUR:	natural latex, white and cream
CUFF:	beaded
POWDER:	powder-free
PROTEIN LEVEL:	79 µg/g
COVERAGE:	inner layer covered with polymer
AQL:	1,0
RESIDUES FROM CHEMICAL AGENTS USED FOR PRODUCTION:	below the limit of detection, which reduces the risk of type IV allergy and skin irritation



Distributor:

ZARYS International Group sp. z o.o. sp.k.

ul. Pod Borem 18, 41-808 Zabrze, Poland, phone +48 32 376 07 00 – 02, 06 fax +48 32 370 38 08

Date: 25.10.2018

Page 1 of 2

TG MEDICAL SDN. BHD.
TEST REPORT

Type Of Glove
Glove Code
AQL Required
Reference Standard

Latex Surgical Powder Free Glove (Natural, Palm Textured)
ENW7B

1.0

The above consignment of goods have been inspected against EN 455:00 Part 1, EN 455:15 Part 2 & EN 455:15 Part 3 standard where samples selected at random using Single Sampling Plans for Normal Inspection of MIL-STD 105E / ISO 2859-1 and also EN 455:09 Part 4 requirement (maximum complied with 5 years).

Declared - Quantity
- Size

Size	Quantity (pairs)	Quantity (pcs)
6.0	24,000	48,000
6.5	24,000	48,000
7.0	24,000	48,000
7.5	24,000	48,000
8.0	24,000	48,000
8.5	24,000	48,000
9.0	24,000	48,000
Total	168,000	336,000

Size	Water Tight Test			Major Defects, AQL 2.5			Minor Defects, AQL 4.0			Result
	Inspection level : G1, AQL 1.0			Sample size (pcs)	Acceptance (pcs)	Defects (pcs)	Sample size (pcs)	Acceptance (pcs)	Defects (pcs)	
	Sample size (pcs)	Acceptance (pcs)	Defects (pcs)							
6.0	200	5	3	200	10	2	200	14	6	Pass
6.5	200	5	1	200	10	6	200	14	9	Pass
7.0	200	5	1	200	10	3	200	14	4	Pass
7.5	200	5	0	200	10	1	200	14	5	Pass
8.0	200	5	2	200	10	5	200	14	5	Pass
8.5	200	5	3	200	10	1	200	14	4	Pass
9.0	200	5	1	200	10	3	200	14	2	Pass

2. Dimensions

Sampling : 13 test pieces

Acceptance : Median of test pieces shall comply with requirements

Defects found : Nil
Result : Pass

Sample No.	Size	Length (mm)	Width (mm)	Thickness (single wall) (mm)		
				Fingertip	Palm	Cuff
1	6.0	286	76	0.15	0.14	0.12
2		286	77	0.16	0.14	0.12
3		288	83	0.16	0.14	0.11
4	6.5	290	84	0.17	0.13	0.11
5		285	91	0.16	0.15	0.11
6		285	91	0.15	0.14	0.12
7	7.0	286	95	0.16	0.14	0.12
8		285	96	0.17	0.15	0.11
9		285	104	0.17	0.14	0.12
10	8.0	286	105	0.16	0.14	0.11
11		284	109	0.16	0.13	0.11
12		286	110	0.15	0.15	0.12
13	9.0	286	115	0.16	0.14	0.12

EN 455:15 Part 2 Requirement:

Size	Length	Width
5.0	Min 250	67 ± 4
5.5		72 ± 4
6.0		77 ± 5
6.5	Min 260	83 ± 5
7.0		89 ± 5
7.5		95 ± 5
8.0	Min 270	102 ± 6
8.5		108 ± 6
9.0		114 ± 6
9.5	Min 280	121 ± 6

3. Physical Properties

Sampling : 13 test pieces

Acceptance : Median of test pieces shall comply with requirements

Defects found : Nil
Result : Pass

Sample No.	Size	Before Aging	After Aging
		Force at Break (Newton)	Force at Break (Newton)
1	6.0	11.5	10.3
2		10.4	9.9
3		9.6	10.3
4	6.5	10.5	10.5
5		10.4	9.5
6		10.6	10.3
7	7.0	10.7	9.8
8		11.2	10.3
9		9.6	11.1
10	8.0	11.2	9.9
11		10.3	10.5
12		10.6	10.2
13	9.0	10.8	9.8

EN 455:15 Part 2 Requirement:

Median Value of Force at Break (N)	
Before Aging	After Aging
≥ 9	≥ 9

Note :

A test result is the median of three individual test measurement values.

4. Powder Residue

Sampling: N = 5

Requirement : Max 2 mg / glove

Size	mg / glove	Result
6.0	0.8	Pass
6.5	1.2	Pass
7.0	0.8	Pass
7.5	1.2	Pass
8.0	1.0	Pass
8.5	1.2	Pass
9.0	1.0	Pass



CONCLUSION : We hereby certify that the above consignment of goods were determined to meet the acceptable limit of the specifications as referring to the of above findings randomly selected samples.

Prepared By : Noorul Shamsidah Hamsudin
QA Manager

Verified By : Noor 'Akilah Saidin
QA Deputy General Manager

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to
Registration No.: DD 60034769 0001
Report No.: 15039020 002

Manufacturer: Shanghai Horin Protective
Products Co., Ltd.
No.3999 Huadong Road

201201 Shanghai China

Scope: Aspects of manufacture concerned with securing and
maintaining sterile conditions:

- Face Masks
- Surgical Gowns
- Non-woven Caps
- Non-woven Shoe Covers
- Plastic Shoe Covers
- Coveralls

Date 01.10.2015



Certification Body

X. Ren





APPROVAL

EC Directive 93/42/EEC Annex V, Article 3
Quality Assurance System Production

Registration No.: DD 60034769 0001

Report No.: 15039020 002

Manufacturer: Shanghai Horin Protective
Products Co., Ltd.
No. 3999 Huadong Road,

201201 Shanghai China

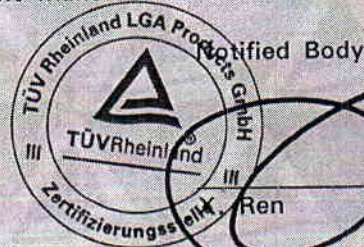
Scope: Manufacture of Medical Devices
(see attachment for products included)

Replaces Approval, Registration No.: DD 60034043 0001

Date of Expiry: 27.09.2020

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex V, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex V, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Date 01.10.2015



TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. **CE**