

## PROHLÁŠENÍ O SHODĚ A PRODUKTOVÝ LIST

### Nitrilové rukavice NITRIL IDEAL

#### ČÁST I: POPIS PRODUKTU

Typ:	jednorázové nesterilní ochranné rukavice
Materiál:	100% syntetický nitril
Barva:	černá
Provedení:	pravolevé, hladké, korálková manžeta
Pudr:	není přidán
Skladování:	rukavice neztrácejí své vlastnosti při skladování v suchu při teplotě od 10 do 30 °C
Životnost:	5 let od data výroby při dodržení podmínek skladování
Balení:	100 ks v krabičce, 10 krabiček v kartonu

#### ČÁST II: SPECIFIKACE PRODUKTU

Délka (mm):	min. 220 (XS, S), min. 230 (M, L, XL)
Šířka (mm):	XS – $70 \pm 10$ S – $80 \pm 10$ M – $95 \pm 10$ L – $110 \pm 10$ XL – $120 \pm 10$
Tloušťka (mm):	prsty: $0.09 \pm 0.02$ (typická hodnota 0.08 – 0.09) dlaň: $0.07 \pm 0.02$ (typická hodnota 0.06 – 0.07)
Prodloužení do přetržení (%):	min. 500
Pevnost v tahu (MPa):	min. 14
AQL:	1.5



### ČÁST III: NORMY A NAŘÍZENÍ

Tímto potvrzujeme, že výše uvedený výrobek je v souladu s:

Obecné: PPER (EU) 2016/425 Cat. I  
EN 420

Potravinářství: EC 1935/2004  
EC 2023/2006

### ČÁST IV: POLOŽKY

Pol. č.	Velikost	Hmotnost (g)	Rozměry (mm)	Kvalita (g)	EAN
100177	XS	310	200x110x60	2.9 ± 0.2	8594177201749
100083	S	340	200x110x60	3.2 ± 0.2	8594177200407
100084	M	371	200x110x60	3.5 ± 0.2	8594177200414
100085	L	402	200x110x60	3.8 ± 0.2	8594177200421
100086	XL	491	200x110x60	4.1 ± 0.2	8594177200438

### ČÁST V: NÁHLED PRODUKTU



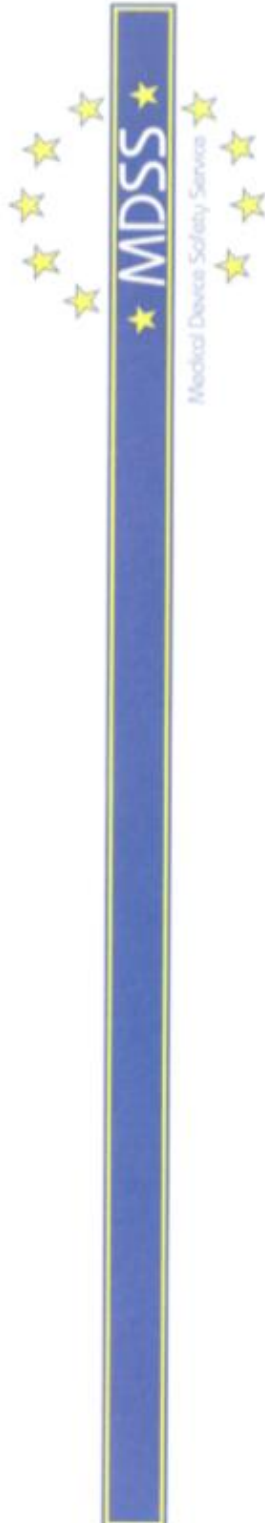
### ČÁST VI: PŮVOD PRODUKTU

Výrobce:

SRI TRANG GLOVES , 57 Wireless Road, Bangkok 10330, Thailand  
INTCO Medical Technology Co., Ltd, No. 29 Zhangliu Road, Zibo, Shandong, China

Distributor:

Espeon s.r.o., U větrolamu 1212/53, 184 00 Praha 8, [info@espeon.cz](mailto:info@espeon.cz) , [www.espeon.cz](http://www.espeon.cz)



## 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:

**Sri Trang Gloves (Thailand) Public Company Limited**  
10 Soi 10, Phetkasem Road, Hat Yai  
Songkhla 90110  
Thailand

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 dated June 22, 2020**

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.

2020-06-22



Dr. Philipp Hohenbrink  
Senior Consultant  
MDSS GmbH



**Annex A dated June 22, 2020**  
**Manufacturer: Sri Trang Gloves (Thailand) Public Company Limited**

UMDNS Code Description Notified Medical Device Product Name & Catalogue Number	UMDNS Code	Risk Class	Cat. Code	Registration Number	NB No. / NB Certificate No.	NB Cert. valid until YYYY-MM-DD
Gloves, Examination/Treatment	11-882	I	10	DE/CA09/0170/S40/001-02	N.A	N.A
Latex Powdered Gloves, Non Sterile; Latex Powder Free Gloves, Non Sterile; Nitrile Powder Free Gloves, Non Sterile						
Latex Powdered Gloves, Non Sterile: LX01, NP02						
Latex Powder Free Gloves, Non Sterile (Offline Chlorination): LC01						
Latex Powder Free Gloves, Non Sterile (Online Chlorination): LO01						
Nitrile Powder Free Gloves, Non Sterile (Offline Chlorination): NC01, NCO2						
Nitrile Powder Free Gloves, Non Sterile (Online Chlorination): NO01, NO02, NO03, NO04						
Gloves, Examination/Treatment	11-882	Is	10	DE/CA09/0170/S40/002-02	0123/G2S0991880008	2022-12-12
Latex Powder Free Offline Chlorination Gloves, Sterile; Nitrile Powder Free Online Chlorination Gloves, Sterile; Latex Powdered Gloves, Sterile						
Latex Powder Free Offline Chlorination Gloves, Sterile						
Nitrile Powder Free Online Chlorination Gloves, Sterile						
Latex Powdered Gloves, Sterile						

*Handwritten signature*





# Certificate of Verification

MDSS GmbH hereby declares  
that an Authorized Representative's Mandate according to the  
EU Regulation 2017/745 (MDR) is in place and that the following tasks have been carried out  
in accordance with the requirements of the MDR on behalf of the Manufacturer:

**Sri Trang Gloves (Thailand) Public Company Limited**  
**10 Soi 10, Phetkasem Road**  
**90110 Hat Yai Songkhla**  
**THAILAND**

MDSS verified that the EU declaration of conformity and technical documentation have been  
drawn up and, where applicable, that an appropriate conformity assessment procedure has  
been carried out by the manufacturer;

MDSS keeps available a copy of the technical documentation, the EU declaration of conformity  
and, if applicable, a copy of the relevant certificate, including any amendments and  
supplements, issued in accordance with Article 56, at the disposal of competent authorities  
for the period referred to in Article 10(8);

MDSS complied with the registration obligations laid down in Article 123.3(d) and until Eudamed  
is fully functional, the corresponding provisions of Directives 90/385/EEC and/or 93/42/EEC  
have been applied.

Details of the device(s) covered by the Certificate are listed hereafter.

Issued: 2024-01-15  
(YYYY-MM-DD)

This Certificate is valid without signature. The document can be traced within MDSS' electronic system.

Certificate No.: 740659

This certificate is subject to the following terms and conditions:

It is only valid for the device(s) listed hereafter;

It is not a proof for compliance to CE marking;

The Manufacturer shall inform MDSS of any significant change(s) to the device(s) listed hereafter and MDSS will verify the change(s) and determine if a renewed  
certificate has to be issued;

As in accordance with the Directive 85/374/EEC Art. 1, the producer is liable for damages caused by a defect in his product(s). The Manufacturer in addition  
confirms that the requirements of Art. 10.16 of the MDR are fulfilled.

This Certificate of Verification is valid for 5 years or until expiry of the EU Declaration of Conformity or NB Certificate if applicable, whichever comes first.

TF-MD-NF-01-102 rev. 09	Nitrile Examination Gloves, Powder Free, Polymer Coated, Accelerator Free, Non-Sterile NC02	56286	I	EU Declaration of Conformity (NC02) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-103 rev. 10	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO01	56286	I	EU Declaration of Conformity (NO01) Signed 5 January 2024	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-104 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO02	56286	I	EU Declaration of Conformity (NO02) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-105 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Accelerator Free, Non-Sterile NO03	56286	I	EU Declaration of Conformity (NO03) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-106 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO04	56286	I	EU Declaration of Conformity (NO04) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945

\*The registration number has been issued by the German Competent Authority.

TF-MD-NF-01-102 rev. 09	Nitrile Examination Gloves, Powder Free, Polymer Coated, Accelerator Free, Non-Sterile NC02	56286	I	EU Declaration of Conformity (NC02) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-103 rev. 10	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO01	56286	I	EU Declaration of Conformity (NO01) Signed 5 January 2024	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-104 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO02	56286	I	EU Declaration of Conformity (NO02) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-105 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Accelerator Free, Non-Sterile NO03	56286	I	EU Declaration of Conformity (NO03) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945
TF-MD-NF-01-106 rev. 09	Nitrile Examination Gloves, Powder Free, Chlorinated, Non-Sterile NO04	56286	I	EU Declaration of Conformity (NO04) Signed 12 August 2022	N.A.	N.A.	DE/CA09/00180945

\*The registration number has been issued by the German Competent Authority.



## MODULE D CERTIFICATE

Issued To

Certificate 1 of 1

Sri Trang Gloves (Thailand) Public Company Limited  
110, 109/2, 352 and 110/3 Kanjanavanit Road, Pahtong, Hat Yai, Songkhla, 90230, Thailand  
207/1, Padang Besa Road, Sadao, Sadao, Songkhla, 90120, Thailand  
88/8, Moo 3, Samnak Kham, Sadao, Songkhla, 90320, Thailand  
85 Moo 6, KhuanThani, Kantang, Trang, 92110, Thailand  
189 Moo 7, Phlai Wat, Kanchanadit, Surat Thani, 84160, Thailand  
110/19 Ban Khao Mai Deang, Moo 7, Phlai Wat, Kanchanadit, Surat Thani, 84160, Thailand  
88/8, Moo 11, Khao Chai Rat, Pathio, Chumphon, 86210, Thailand

Has been found to conform to the requirements of Annex VIII (Module D) of the PPE Regulation (EU) 2016/425 – Conformity to type based on quality assurance of the production process, for the manufacture and distribution of Latex and Nitrile Disposable Gloves.

Standards	Module B Certificates*		
EN ISO 374-1: 2016 +A1: 2018	2777/10466-XX/E00-00	2777/10474-XX/E00-00	2777/16141-XX/E00-00
EN ISO 374-2: 2019	2777/10467-XX/E00-00	2777/14071-XX/E00-00	2777/16142-XX/E00-00
EN ISO 374-4: 2019	2777/10468-XX/E00-00	2777/14362-XX/E00-00	2777/24437-XX/E00-00
EN ISO 374-5: 2016	2777/10469-XX/E00-00	2777/14364-XX/E00-00	
EN ISO 21420: 2020	2777/10470-XX/E00-00	2777/15155-XX/E00-00	

\* Where 'XX' denotes the Module B certificate issue number

Date Issued 22/09/2023

Valid Until 29/09/2024  
SATRA Reference STE0349789/R4



Signed on behalf of SATRA Technology Europe Ltd – Notified Body Number: 2777



The issuance of this certificate is subject to the company maintaining its manufacturing and quality system to the required standard which will be monitored by: SATRA Technology Europe Limited, Braetown Business Park Clonee Dublin 15 D15 YN2P Ireland - Tel: +353 (0) 1 437 2484 Web: [www.satra.com](http://www.satra.com)



Product Service

## Certificate

No. Q5 099188 0012 Rev. 01

**Holder of Certificate:** **Sri Trang Gloves (Thailand)  
Public Company Limited**  
10 Soi 10, Phetkasem Road  
Hat Yai, Songkhla 90110  
THAILAND

**Certification Mark:**



**Scope of Certificate:** **Design and Development, Production and  
Distribution of Sterile and Non-Sterile  
Examination and Sterile and Non-Sterile  
Surgical Gloves**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:Q5 099188 0012 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:Q5 099188 0012 Rev. 01)

**Report No.:** 5675207Rev1-721430483

**Valid from:** 2023-11-01  
**Valid until:** 2026-10-31

**Date,** 2023-10-06



Christoph Dicks  
Head of Certification/Notified Body



	<b>山东英科医疗制品有限公司</b> Shandong Intco Medical Products Co., Ltd.	文件编号 Doc No. INTCO-PPE-NBR
		版 本 Ver. A3

### EU Declaration of Conformity

We declare that the following product is carried out in accordance with Regulation (EU) 2016/425.

1. PPE: Disposable Nitrile Gloves

Glove sizes available: XS, S, M, L, XL, XXL (6, 7, 8, 9, 10, 11)

Product code :Blue:NGV/B/H/PEM(10013~10017), Violet:NGV/B/H/PEM(10033~10037),

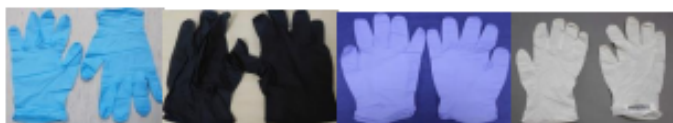
White: NGV/B/H/PEM(10023~10027),Black: NGV/B/H/PEM(10043~10047).

2. Manufacturer Name: Shandong Intco Medical Products Co., Ltd.

Address: No.9888,Qiawang Road, Naoshan Industrial Park. Qingzhou, Shandong, China

3. This declaration of conformity is issued under the sole responsibility of the manufacturer: Shandong Intco Medical Products Co., Ltd.

4. Object of the declaration: Disposable Nitrile Gloves



5. The object of the declaration described in point 4 is in conformity with the relevant Union harmonisation legislation: Regulation (EU)2016/425.

6. References to the relevant harmonised standards used, including the date of the standard, or references to the other technical specifications, including the date of the specification, in relation to which conformity is declared: EN ISO 21420:2020, EN ISO 374-1:2016 +A1:2018, EN374-4:2019, EN ISO 374-5:2016.

7. The notified body SATRA Technology Europe Ltd (Number: 2777) performed the EU type-examination (Module B) and issued the EU type-examination certificate.

8. The PPE is subject to the conformity assessment procedure Module C2 under surveillance of the notified body : SATRA Technology (Number: 2777),

Adress: Bracetown Business Park Clonee, D15 YN2P, Ireland.

Signed for and on behalf of: Shandong Intco Medical Products Co., Ltd.

签名 Singnature 程建建

职位 Position Quality Manager

日期 Date 11-Sep-2020



<b>SUBJECT</b>	<b>Chemical Test</b>																				
<b>TEST LOCATION</b>	<b>TÜV SÜD China</b>  TÜV SÜD Products Testing (Shanghai) Co., Ltd. B-3/4, No.1999 Du Hui Road, Minhang District Shanghai 201108, P.R. China																				
<b>CLIENT NAME</b>	<b>Shandong Intco Medical Products Co., Ltd</b>																				
<b>CLIENT ADDRESS</b>	No.9888 Qiwang Road, Naoshan Industry Park, Qingzhou, Shandong, China																				
<b>TEST PERIOD</b>	31-May-2021-07-Jun-2021																				
<b>RESULT SUMMARY</b>	<table border="0"> <tr> <td>1. The tested items <b>complied with</b> German Food &amp; Feed Acts of September 1, 2005 (LFGB), Section 30 and 31.</td> <td></td> </tr> <tr> <td>- Overall migration test</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>- Color release</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>- Extractable formaldehyde</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>- Total lead and zinc content</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>- Sensory test</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>2. The tested items <b>complied with</b> AFPS GS 2019: 01 PAK</td> <td></td> </tr> <tr> <td>- Polycyclic Aromatic Hydrocarbons (PAHs) content(Category 1)</td> <td style="text-align: right;"><b>PASS</b></td> </tr> <tr> <td>3. As per client's request</td> <td></td> </tr> <tr> <td>- Total Cadmium content</td> <td style="text-align: right;"><b>PASS</b></td> </tr> </table>	1. The tested items <b>complied with</b> German Food & Feed Acts of September 1, 2005 (LFGB), Section 30 and 31.		- Overall migration test	<b>PASS</b>	- Color release	<b>PASS</b>	- Extractable formaldehyde	<b>PASS</b>	- Total lead and zinc content	<b>PASS</b>	- Sensory test	<b>PASS</b>	2. The tested items <b>complied with</b> AFPS GS 2019: 01 PAK		- Polycyclic Aromatic Hydrocarbons (PAHs) content(Category 1)	<b>PASS</b>	3. As per client's request		- Total Cadmium content	<b>PASS</b>
1. The tested items <b>complied with</b> German Food & Feed Acts of September 1, 2005 (LFGB), Section 30 and 31.																					
- Overall migration test	<b>PASS</b>																				
- Color release	<b>PASS</b>																				
- Extractable formaldehyde	<b>PASS</b>																				
- Total lead and zinc content	<b>PASS</b>																				
- Sensory test	<b>PASS</b>																				
2. The tested items <b>complied with</b> AFPS GS 2019: 01 PAK																					
- Polycyclic Aromatic Hydrocarbons (PAHs) content(Category 1)	<b>PASS</b>																				
3. As per client's request																					
- Total Cadmium content	<b>PASS</b>																				

Prepared By

*Nance Gao*

( Nance Gao )  
Report Drafter

Authorized By

*Leo Li*

( Leo Li )  
Authorized Signatory



**Note:** (1) General Terms & Conditions as mentioned overleaf. (2) The results relate only to the items tested.(3) The test report shall not be reproduced except in full without the written approval of the laboratory.(4) Without the agreement of the laboratory, the client is not authorized to use the test results for unapproved propaganda.

Chemical/Microbiology Laboratory:  
TÜV SÜD Products Testing (Shanghai) Co., Ltd  
B-3/4, No.1999 Du Hui Road, Minhang District  
Shanghai  
201108  
P.R. China

Phone : +86 (21) 6037 6375  
Fax : +86 (21) 6037 6345  
Email: food.chem@tuv-sud.cn  
Webpage: www.tuv-sud.cn

Regional Head Office:  
TÜV SÜD Certification and Testing  
(China) Co., Ltd.  
No.151 Heng Tong Road Shanghai  
200 070 P.R.China

**TUV**<sup>®</sup>

**RECEIPT DATE / TEST DATE**

31-May-2021/ 31-May-2021

**THE FOLLOWING SAMPLE(S) WAS/WERE SUBMITTED**

**BY/ ON BEHALF OF THE CLIENTS AS**

Sample Name: Disposable Nitrile Gloves  
Sample Specification: /  
Batch No./Date: /  
Manufacture: Shandong Intco Medical Products Co., Ltd

SAMPLE NO.	DESCRIPTION	PHOTOGRAPH
721664284	Blue glove	

**TEST RESULT(S)**

Note: The migration results in this report were tested and expressed based on single use articles.

1. Overall Migration Test

- In accordance with BFR Recommendations XXI
- Test method: With reference to EN 1186: Part 4 (Test methods for overall migration into olive oil by cell), EN 1186: Part 5 (Test methods for overall migration into aqueous food stimulants by cell) and BFR Recommendations XXI
- Migration ratio (S/V): 10dm<sup>2</sup>l

Simulant(s) Used	Test Condition	Result(s) [mg/dm <sup>2</sup> ]	Maximum Permissible Limit [mg/dm <sup>2</sup> ]
3% Acetic acid	40°C for 0.5 hours	3.15	10
10% Ethanol	40°C for 2 hours	0.950	10
Olive oil	40°C for 2 hours	<0.500	10

2. Color release

- Test method: With reference to Kunststoffe im Lebensmittelverkehr Book II, Teil B II, IX

Simulant(s) Used	Test Condition	Result(s)	Permissible Limit
10% Ethanol	50 °C for 5 hours	No bleeding	No bleeding
2% Acetic acid	50 °C for 5 hours	No bleeding	No bleeding
Peanut oil	50 °C for 5 hours	No bleeding	No bleeding
Water	50 °C for 5 hours	No bleeding	No bleeding

- Note:
1. No bleeding denotes no difference was found between blank and sample
  2. Bleeding denotes staining was found from sample

3. Extractable formaldehyde

- Test method: For compliance with the Recommendation of the BfR "Kunststoffe im Lebensmittelverkehr" Part XXI. Commodities based on Natural and Synthetic Rubber
- With reference to Section 2.7.1 of methods for the "Testing of commodities made of rubber"
- Test condition: 3% Acetic acid, 40°C for 0.5 hours
- Migration ratio (S/V): 6dm ³L

Test Item(s)	Result(s) [µg/ml]	Maximum Permissible Limit [µg/ml]
Extractable Formaldehyde	<0.5	3

4. Total lead and zinc content

- Test method: Acid digestion, then followed by ICP-OES

Test Item(s)	Result(s) [%]	Maximum Permissible Limit [%]
Lead content	<0.001	0.003
Zinc content	0.250	3.0

5. Sensory test

- Test method: With reference to DIN 10955.
- The submitted sample was simulated in distilled water at 40°C for 2 hours. After this treatment treated water was examined by panels with regard to any divergence in smell and taste.

Sample(s)	Testing Parameter	Grading result(s)	Recommended level
721664284	Transfer of taste	0	<3
	Transfer of smell	0	<3

Note: 1. Available grading are listed as follow:

- Grading 0: No perceptible taste/smell deviation
- 1: Just perceptible taste/smell deviation
- 2: Weak taste/smell deviation
- 3: Clear taste/smell deviation
- 4: Strong taste/smell deviation

6. Polycyclic Aromatic Hydrocarbons (PAHs) content

- Test method: In accordance with AfPS GS 2019: 01 PAK

Compounds	Results [mg/kg]	Detection Limit [mg/kg]
Chrysene	ND	0.01
Benzo[a]anthracene	ND	0.01
Benzo[b]fluoranthene	ND	0.01
Benzo[j]fluoranthene	ND	0.01
Benzo[k]fluoranthene	ND	0.01
Benzo[e]pyrene	ND	0.01
Benzo[a]pyrene	ND	0.01
Indeno[1,2,3-cd]pyrene	ND	0.01
Dibenzo[ah]anthracene	ND	0.01
Benzo[ghi]perylene	ND	0.01
Naphthalene	0.0585	0.01
Phenanthrene	0.0619	0.01

Anthracene	ND	0.01
Fluoranthene	0.0109	0.01
Pyrene	0.0321	0.01
Sum of Phenanthrene, Pyrene, Anthracene and Fluoranthene	0.105	--
Group PAH	0.163	--
Category as in AfPS GS 2019: 01 PAK	<b>Category 1</b>	--

Note: 1. ND denotes not detected, less than Reporting Limit  
Limits and Categories for PAH in product according to AfPS GS 2019: 01 PAK

### Limits and Categories

Parameter [mg/kg]	Category 1	Category 2		Category 3	
		Materials not covered by category 1, with foreseeable skin contact for longer than 30 s (long-term skin contact) or repeated short term skin contact*	Products for use by children aged < 14 years (active or passive direct contact)	All other products acc to ProdSG	Products for use by children aged < 14 years (active or passive direct contact)
Benzo[a]pyrene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[e]pyrene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[a]anthracene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[b]fluoranthene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[k]fluoranthene	<0.2	<0.2	<0.5	<0.5	<1
Chrysene	<0.2	<0.2	<0.5	<0.5	<1
Dibenzo[a,h]anthracene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[ghi]perylene	<0.2	<0.2	<0.5	<0.5	<1
Indeno[1,2,3-cd]pyrene	<0.2	<0.2	<0.5	<0.5	<1
Phenanthrene, Anthracene, Fluoranthene and Pyrene	<1 sum	<5 sum	<10 sum	<20 sum	<50 Sum
Naphthalene	<1	<2		<10	
Sum 15 PAH	<1	<5	<10	<20	<50

\* Definition "short-term repetitive contact with the human skin" from REACH Annex XVII No. 50 amendment (COMMISSION REGULATION (EU) No 1272/2013)

### 7. Total Cadmium content

- Test method: Sample digested, analysed by ICP-MS

Test Item(s)	Result(s) [mg/kg]	Maximum Permissible Limit* [mg/kg]
Total Cadmium Content	<2	2

(SP) 01/01/2021  
 01/01/2021  
 01/01/2021

Test Report No.: 721664284  
Report Date: 9 June 2021



Note: 1. \* denotes Maximum Permissible Limit was provided by client

Note: This report is for internal use only such as internal scientific research ,education, quality control, product R&D

-END OF THE TEST REPORT-

