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Chemical and Microbiological Laboratory, Testing Laboratory No. 1273 certified by Czech Accreditation Institute according to ČSN EN ISO/IEC 17025.

Copy No.: 1
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Test report No. D90-1/2017

DETERMINATION OF BACTERICIDAL (EN 13727+A2, EN 14561), YEASTICIDAL (EN 13624, EN 14562) AND TUBERCULOCIDAL (EN 14348) ACTIVITY OF THE PRODUCT **GLOBACID SF MED 1.25%**

Sample ID: D90/2017
Sample name: **GLOBACID SF MED 1.25%**
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Producer: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404

Incoming date:
11.5.2017

Delivery date:
8.12.2017

Hodonín, 8.12.2017

Ing. Jana Šlitrová, Head of Laboratory

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

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Subject of testing:
Determination of bactericidal, yeasticidal and tuberculocidal activity of the product.

Identification of the sample:

Name of the product:	**GLOBACID SF MED 1.25%**
Batch number:	20042017
Date of manufacture:	20.04.2017
Expiry date:	19.04.2020
Manufacturer:	Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Incoming date:	11.5.2017
Storage conditions:	stated by the manufacturer
Active compounds in 100 g:	N-(3-aminopropyl)-N-dodecyl propane-1,3-diamine <1% DDAC <1%

Experimental conditions:

	Testing of disinfecting efficiency of chemical disinfecting and antiseptic agents by suspension method SOP-M-19-00 (EN 13727:2012+A2:2015)
Period of analysis:	12.9. – 13.9.2017
Test temperature:	20 °C ± 1 °C
Test method:	membrane filtration method
Filtration diluent:	rinsing liquid
Appearance of the product:	colourless liquid
Test concentration:	100% (concentrated)*
Contact time:	15, 30 and 60 min
Interfering substances:	3 g/l BSA and 3 ml/l sheep erythrocytes (dirty conditions)
Test organisms:	*Pseudomonas aeruginosa* ATCC 15442 *Staphylococcus aureus* ATCC 6538 *Enterococcus hirae* ATCC 10541
Incubation conditions:	37 °C ± 1 °C, 24 hours

Test procedure:

1. Preparation of the test suspension
2. Preparation of the product test solutions
3. Quantitative suspension test
4. Incubation and calculation
5. Expression and interpretation of the results

Note:
Bactericidal activity – the capability of a product to produce a reduction in the number of viable bacterial cells of relevant organisms under defined conditions by at least 5 orders (10^5).
$R = N_0 / N_a$ = the reduction in viability, or $\lg R = \lg N_0 - \lg N_a$
* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

The standard:
EN 13727:2012+A2:2015 Chemical disinfectants and antiseptics – Quantitative suspension test for the evaluation of bactericidal activity in the medical area – Test method and requirements (phase 2, step 1) October 2015

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

The Number of CFU in the tested product: 0 CFU/ml

1. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Pseudomonas aeruginosa* ATCC 15442

Tab No. 1.1 Verification of methodology, dirty conditions

Validation of suspension (N_{V0})			Validation of selected experimental conditions (A)			Membrane filtration control (B)			Method validation (C) Product conc.: 100%*		
V_{c1} 34 / V_{c2} 31	Φ_{Nvo}= 32.5		V_{c1} 24 / V_{c2} 31	Φ_A = 27.5		V_{c1} 30 / V_{c2} 27	Φ_B = 28.5		V_{c1} 19 / V_{c2} 30	Φ_C = 24.5	
$30 \le \Phi_{Nvo} \le 160$			$\Phi_A \ge 0.5\ \Phi_{Nvo}$			$\Phi_B \ge 0.5\ \Phi_{Nvo}$			$\Phi_C \ge 0.5\ \Phi_{Nvo}$		
x yes		no	x yes		no	x yes		no	x yes		no

Validation of suspension (N_{VB})	V_{c1}	29	V_{c2}	31	Φ_{NVB}	30	$30 \le \Phi_{NVB}\ (N_{VB}/1000) \le 160$		
							x yes		no

Tab No. 1.2 Test suspension

Test suspension N	N	V_{c1}	V_{c1}	Test suspension N_0 (time = 0)
$\Phi = 28 \times 10^8 = \lg 9.45$	10^{-7}	>165	>165	$\lg N_0 = \lg N/100 = \lg 7.45$
$9.17 \le \lg N \le 9.70$	10^{-8}	26	30	$7.17 \le \lg N_0 \le 7.70$
				x yes / no

Tab No. 1.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Pseudomonas aeruginosa* ATCC 15442

Test concentration (%)/contact time (min)/conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg(\Phi_a \times 10)$	**lg R ($\lg N_0 = \lg 7.45$)**
100*/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.30**
100*/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.30**
100*/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.30**

2. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Staphylococcus aureus* ATCC 6538

Tab No. 2.1 Verification of methodology, dirty conditions

Validation of suspension (N_{V0})			Validation of selected experimental conditions (A)			Membrane filtration control (B)			Method validation (C) Product conc.: 100%*		
V_{c1} 56 / V_{c2} 48	Φ_{Nvo}= 52		V_{c1} 53 / V_{c2} 40	Φ_A = 46.5		V_{c1} 42 / V_{c2} 49	Φ_B = 45.5		V_{c1} 48 / V_{c2} 43	Φ_C = 45.5	
$30 \le \Phi_{Nvo} \le 160$			$\Phi_A \ge 0.5\ \Phi_{Nvo}$			$\Phi_B \ge 0.5\ \Phi_{Nvo}$			$\Phi_C \ge 0.5\ \Phi_{Nvo}$		
x yes		no	x yes		no	x yes		no	x yes		no

Validation of suspension (N_{VB})	V_{c1}	47	V_{c2}	53	Φ_{NVB}	50	$30 \le \Phi_{NVB}\ (N_{VB}/1000) \le 160$		
							x yes		no

Tab No. 2.2 Test suspension

Test suspension N	N	V_{c1}	V_{c1}	Test suspension N_0 (time = 0)
$\Phi = 50 \times 10^8 = \lg 9.70$	10^{-7}	>165	>165	$\lg N_0 = \lg N/100 = \lg 7.70$
$9.17 \le \lg N \le 9.70$	10^{-8}	58	42	$7.17 \le \lg N_0 \le 7.70$
				x yes / no

Tab No. 2.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Staphylococcus aureus* ATCC 6538

Test concentration (%)/contact time (min)/conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg(\Phi_a \times 10)$	**lg R ($\lg N_0 = \lg 7.70$)**
100*/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.55**
100*/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.55**
100*/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.55**

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l
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3. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Enterococcus hirae* ATCC 10541

Tab No. 3.1 Verification of methodology, dirty conditions

Validation of suspension (N_{V0})			Validation of selected experimental conditions (A)			Membrane filtration control (B)			Method validation (C) Product conc.: 100%*		
V_{c1}	55	Φ_{Nv0}= 52.5	V_{c1}	45	Φ_A = 48	V_{c1}	47	Φ_B = 42.5	V_{c1}	52	Φ_C = 46.5
V_{c2}	50		V_{c2}	51		V_{c2}	38		V_{c2}	41	
$30 \leq \Phi_{Nv0} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nv0}$			$\Phi_B \geq 0.5\ \Phi_{Nv0}$			$\Phi_C \geq 0.5\ \Phi_{Nv0}$		
x	yes	no	x	yes	no	x	yes	no	x	yes	no

Validation of suspension (N_{VB})	V_{c1}	43	V_{c2}	49	Φ_{NVB}	46	$30 \leq \Phi_{NVB}$ (N_{VB}/1000) ≤ 160		
							x	yes	no

Tab No. 3.2 Test suspension

Test suspension N	N	V_{c1}	V_{c1}	Test suspension N_0 (time = 0)
$\Phi = 49.5 \times 10^8 = \lg 9.69$	10^{-7}	>165	>165	$\lg N_0 = \lg N/100 = \lg 7.69$
$9.17 \leq \lg N \leq 9.70$	10^{-8}	54	45	$7.17 \leq \lg N_0 \leq 7.70$
				x yes no

Tab No. 3.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Enterococcus hirae* ATCC 10541

Test concentration (%)/contact time (min)/conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_0 = \lg 7.69$)**
100*/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.54**
100*/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.54**
100*/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.54**

4. Evaluation of bactericidal activity of the product **GLOBACID SF MED 1.25%**

Tab No. 4.1 The efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on test strains – bactericidal activity

Bactericidal activity of the product (EN 13727:2012+A2:2015)						
Strain	Test temperature [°C]	Contact time [min]	Product test concentrations [%]	Interfering substances - conditions	lg R EN 13727:2012 +A2:2015	**lg R**
Pseudomonas aeruginosa ATCC 15442	20	15	100*	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	15	100*	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	15	100*	dirty	≥ 5	**> 5**
Pseudomonas aeruginosa ATCC 15442	20	30	100*	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	30	100*	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	30	100*	dirty	≥ 5	**> 5**
Pseudomonas aeruginosa ATCC 15442	20	60	100*	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	60	100*	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	60	100*	dirty	≥ 5	**> 5**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the test suspension, N_0 = the number of cfu/ml of the test suspension at the beginning of the contact time (time „0"), N_a = the number of survivors per ml in the test mixture at the end of the contact time and before the membrane filtration, N_v = the number of cfu/ml of the test suspension for validation, N_{V0} (A,C), N_{VB} (B) = the number of cfu/ml of the test suspensions for validation in the test mixture A, B, C at the beginning of the contact time „0", A,B,C = the number of survivors per ml in control tests (A – experimental conditions control, B – membrane filtration validation, C – method validation), $R = N_0 / N_a$ nebo $\lg R = \lg N_0 - \lg N_a$ the reduction in viability

* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

Prepared by: Ing. Eva Kremlová, Lab Technician

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

Experimental conditions:	**Testing of disinfecting efficiency of chemical disinfecting and antiseptic agents by suspension method** SOP-M-19-00 (EN 13624:2013)
Period of analysis:	11.9. – 13.9.2017
Test temperature:	20 °C ± 1 °C
Test method:	membrane filtration method
Filtration diluent:	rinsing liquid
Appearance of the product:	colourless liquid
Test concentration:	100% (concentrated)*
Contact time:	15, 30 and 60 min
Interfering substances:	3 g/l BSA and 3 ml/l sheep erythrocytes (dirty conditions)
Test organisms:	*Candida albicans* ATCC 10231
Incubation conditions:	30 °C ± 1 °C, 48 hours and additional period of 24 or 48 hours

Test procedure:

1. Preparation of the test suspension
2. Preparation of the product test solutions
3. Quantitative suspension test
4. Incubation and calculation
5. Expression and interpretation of the results

Note:
Fungicidal activity – the capability of a product to produce a reduction in the number of viable fungi belonging to reference strains under defined conditions by at least 4 orders (10^4).
Yeasticidal activity – the capability of a product to produce a reduction in the number of viable yeast cells of relevant test organisms under defined conditions by at least 4 orders (10^4).
$R = N_0 / N_a$ = the reduction in viability, or $\lg R = \lg N_0 - \lg N_a$
* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

The standard:
EN 13624:2013 Chemical disinfectants and antiseptics – Quantitative suspension test for the evaluation of fungicidal or yeasticidal activity in the medical area - Test method and requirements (phase 2, step 1) September 2013

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

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Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

5. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Candida albicans* ATCC 10231

Tab No. 5.1 Verification of methodology, dirty conditions

Validation of suspension (N_{V0})			Validation of selected experimental conditions (A)			Membrane filtration control (B)			Method validation (C) Product conc. 100%*		
V_{c1} 42	V_{c2} 44	Φ_{Nvo}= 43	V_{c1} 27	V_{c2} 21	Φ_A = 24	V_{c1} 39	V_{c2} 25	Φ_B = 32	V_{c1} 20	V_{c2} 37	Φ_C = 28.5
$30 \leq \Phi_{Nvo} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nvo}$			$\Phi_B \geq 0.5\ \Phi_{Nvo}$			$\Phi_C \geq 0.5\ \Phi_{Nvo}$		
x yes		no	x yes		no	x yes		no	x yes		no

Validation of suspension (N_{VB})	V_{c1} 31	V_{c2} 37	Φ_{NVB} 34	$30 \leq \Phi_{NVB}\ (N_{VB}/1000) \leq 160$
				x yes / no

Tab No. 5.2 Test suspension

Test suspension N	N	V_{c1}	V_{c1}	Test suspension N_0 (time = 0)
$\Phi = 43 \times 10^7 = \lg 8.63$	10^{-6}	>165	>165	$\lg N_0 = \lg N/100 = \lg 6.63$
$8.17 \leq \lg N \leq 8.70$	10^{-7}	37	49	$6.17 \leq \lg N_0 \leq 6.70$
				x yes / no

Tab No. 5.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Candida albicans* ATCC 10231

Test concentration (%)/contact time (min)/conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_0 = \lg 6.63$)**
100*/15/dirty	10^0	<14	<14	< 2.15	**≥ 4.48**
100*/30/dirty	10^0	<14	<14	< 2.15	**≥ 4.48**
100*/60/dirty	10^0	<14	<14	< 2.15	**≥ 4.48**

6. Evaluation of yeasticidal activity of the product **GLOBACID SF MED 1.25%**

Tab No. 6.1 The efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on test strains – yeasticidal activity

Fungicidal activity of the product (EN 13624:2013)						
Strain	Test temperature [°C]	Contact time [min]	Product test concentrations [%]	Interfering substances - conditions	lg R EN 13624:2013	**lg R**
Candida albicans ATCC 10231	20	15	100*	dirty	≥ 4	**> 4**
Candida albicans ATCC 10231	20	30	100*	dirty	≥ 4	**> 4**
Candida albicans ATCC 10231	20	60	100*	dirty	≥ 4	**> 4**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the test suspension, N_0 = the number of cfu/ml of the test suspension at the beginning of the contact time (time „0“), N_a = the number of survivors per ml in the test mixture at the end of the contact time and before the membrane filtration, N_v = the number of cfu/ml of the test suspension for validation, N_{V0} (A,C), N_{VB} (B) = the number of cfu/ml of the test suspensions for validation in the test mixture A, B, C at the beginning of the contact time „0“, A,B,C = the number of survivors per ml in control tests (A – experimental conditions control, B – membrane filtration validation, C – method validation), $R = N_0 / N_a$ nebo $\lg R = \lg N_0 - \lg N_a$ the reduction in viability

* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

Prepared by: Ing. Eva Kremlová, Lab Technician

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
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Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

Experimental conditions: **Testing of disinfecting efficiency of chemical disinfecting and antiseptic agents on carriers** SOP-M-22-12 (EN 14561:2006)

Period of analysis:	7.9. – 8.9.2017
Test temperature:	20 °C ± 1 °C
Test method:	dilution neutralization method
Neutralization medium:	Dey-Engley Neutralizing Broth M 1062
Appearance of the product:	colourless liquid
Test concentration:	100% (concentrated)
Contact time:	15, 30 and 60 min
Interfering substances:	3 g/l BSA and 3 ml/l sheep erythrocytes (dirty conditions)
Test organisms:	*Pseudomonas aeruginosa* ATCC 15442
	Staphylococcus aureus ATCC 6538
	Enterococcus hirae ATCC 10541
Incubation conditions:	37 °C ± 1 °C, 24 hours

Test procedure:

1. Preparation of the test suspension
2. Preparation of the product test solutions
3. Quantitative carrier test
4. Incubation and calculation
5. Expression and interpretation of the results

Note:
Bactericidal activity – the capability of a product to produce a reduction in the number of viable bacterial cells of relevant organisms on carriers under defined conditions by at least 5 orders (10^5). The drying time: 30-35 min
$R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

The standard:
EN 14561:2006 Chemical disinfectants and antiseptics – Quantitative carrier test for the evaluation of bactericidal activity for instruments used in the medical area - Test method and requirements (phase 2, step 2) May 2006

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

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Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
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Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

7. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Pseudomonas aeruginosa* ATCC 15442 on carriers

Tab No. 7.1 Verification of methodology, dirty conditions

Validation of suspension (N_{v0})			Validation of selected experimental conditions (A)			Neutralizer toxicity control (B)			Method validation (C) Product conc.: 100%		
V_{c1}	133	$\Phi_{Nv0} = 129$	V_{c1}	127	$\Phi_A = 123.5$	V_{c1}	127	$\Phi_B = 112.5$	V_{c1}	140	$\Phi_C = 126$
V_{c2}	125		V_{c2}	120		V_{c2}	98		V_{c2}	112	
$30 \leq \Phi_{Nv0} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nv0}$			$\Phi_B \geq 0.5\ \Phi_{Nv0}$			$\Phi_C \geq 0.5\ \Phi_{Nv0}$		
x	yes	no	x	yes	no	x	yes	no	x	yes	no

Tab No. 7.2 Test suspension

Test suspension (N)	N	V_{c1}	V_{c1}	$\Phi = 211 \times 10^7 = \lg 9.32$ $9.17 \leq \lg N \leq 9.70$
	10^{-7}	198	221	
	10^{-8}	20	25	
				x yes no

Tab No. 7.2.1 The control test suspension, dirty conditions

Test suspension (N_W)	N_W	V_{c1}	V_{c2}	$\Phi \times 10 = 445 \times 10^5 = \lg 7.65$ $\lg N_W = \lg 7.65$ $7.15 \leq \lg N_w \leq (\lg N - 1.3)\ 8.02$
	10^{-5}	59	30	
				x yes no

Tab No. 7.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Pseudomonas aeruginosa* ATCC 15442 on carriers

Test concentration (%) / contact time (min) / conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_W = \lg 7.65$)**
100/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.50**
100/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.50**
100/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.50**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the bacterial test suspension, N_W = the number of cfu/ml of the control bacterial test suspension, N_a = the number of survivors per ml in the test mixture at the end of the contact time, N_{v0}= the number of cfu/ml of the bacterial test suspension in the mixture A,B,C at the beginning of the contact time (time „0"), A,B,C = the number of survivors per ml in control tests (A – experimental conditions validation, B – neutralizer toxicity validation, C – method validation), $R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
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Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

8. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Staphylococcus aureus* ATCC 6538 on carriers

Tab No. 8.1 Verification of methodology, dirty conditions

Validation of suspension (N_{v0})			Validation of selected experimental conditions (A)			Neutralizer toxicity control (B)			Method validation (C) Product conc.: 100%		
V_{c1}	168	$\Phi_{Nvo} = 155.5$	V_{c1}	171	$\Phi_A = 155.5$	V_{c1}	165	$\Phi_B = 158$	V_{c1}	149	$\Phi_C = 149.5$
V_{c2}	143		V_{c2}	140		V_{c2}	151		V_{c2}	150	
$30 \leq \Phi_{Nvo} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nvo}$			$\Phi_B \geq 0.5\ \Phi_{Nvo}$			$\Phi_C \geq 0.5\ \Phi_{Nvo}$		
x	yes	no	x	yes	no	x	yes	no	x	yes	no

Tab No. 8.2 Test suspension

Test suspension (N)	N	V_{c1}	V_{c1}	$\Phi = 170 \times 10^7 = \lg 9.23$ $9.17 \leq \lg N \leq 9.70$
	10^{-7}	156	183	
	10^{-8}	15	20	
				x yes no

Tab No. 8.2.1 The control test suspension, dirty conditions

Test suspension (N_W)	N_W	V_{c1}	V_{c2}	$\Phi \times 10 = 520 \times 10^5 = \lg 7.72$ $\lg N_W = \lg 7.72$ $7.15 \leq \lg N_w \leq (\lg N - 1.3)\ 7.93$
	10^{-5}	50	54	
				x yes no

Tab No. 8.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Staphylococcus aureus* ATCC 6538 on carriers

Test concentration (%) / contact time (min) / conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_W = \lg 7.72$)**
100/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.57**
100/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.57**
100/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.57**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the bacterial test suspension, N_W = the number of cfu/ml of the control bacterial test suspension, N_a = the number of survivors per ml in the test mixture at the end of the contact time, N_{v0}= the number of cfu/ml of the bacterial test suspension in the mixture A,B,C at the beginning of the contact time (time „0"), A,B,C = the number of survivors per ml in control tests (A – experimental conditions validation, B – neutralizer toxicity validation, C – method validation), $R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

9. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Enterococcus hirae* ATCC 10541 on carriers

Tab No. 9.1 Verification of methodology, dirty conditions

Validation of suspension (N_{v0})			Validation of selected experimental conditions (A)			Neutralizer toxicity control (B)			Method validation (C) Product conc.: 100%		
V_{c1}	155	$\Phi_{Nv0} = 157$	V_{c1}	148	$\Phi_A = 150.5$	V_{c1}	138	$\Phi_B = 142.5$	V_{c1}	122	$\Phi_C = 131$
V_{c2}	159		V_{c2}	153		V_{c2}	147		V_{c2}	140	
$30 \leq \Phi_{Nv0} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nv0}$			$\Phi_B \geq 0.5\ \Phi_{Nv0}$			$\Phi_C \geq 0.5\ \Phi_{Nv0}$		
x	yes	no	x	yes	no	x	yes	no	x	yes	no

Tab No. 9.2 Test suspension

Test suspension (N)	N	V_{c1}	V_{c1}	$\Phi = 171 \times 10^7 = \lg 9.23$ $9.17 \leq \lg N \leq 9.70$			
	10^{-7}	162	177				
	10^{-8}	16	21				
				x	yes		no

Tab No. 9.2.1 The control test suspension, dirty conditions

Test suspension (N_W)	N_W	V_{c1}	V_{c2}	$\Phi \times 10 = 1780 \times 10^4 = \lg 7.25$ $\lg N_W = \lg 7.25$ $7.15 \leq \lg N_w \leq (\lg N - 1.3)\ 7.93$			
	10^{-4}	166	190				
				x	yes		no

Tab No. 9.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Enterococcus hirae* ATCC 10541 on carriers

Test concentration (%) / contact time (min) / conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_W = \lg 7.25$)**
100/15/dirty	10^0	<14	<14	< 2.15	**≥ 5.10**
100/30/dirty	10^0	<14	<14	< 2.15	**≥ 5.10**
100/60/dirty	10^0	<14	<14	< 2.15	**≥ 5.10**

10. Evaluation of bactericidal activity of the product **GLOBACID SF MED 1.25%** on carriers

Tab No. 10.1 The efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on test strains – bactericidal activity on carriers

Bactericidal activity of the product on carriers (EN 14561:2006)						
Strain	Test temperature [°C]	Contact time [min]	Product test concentrations [%]	Interfering substances - conditions	lg R EN 14561:2006	**lg R**
Pseudomonas aeruginosa ATCC 15442	20	15	100	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	15	100	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	15	100	dirty	≥ 5	**> 5**
Pseudomonas aeruginosa ATCC 15442	20	30	100	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	30	100	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	30	100	dirty	≥ 5	**> 5**
Pseudomonas aeruginosa ATCC 15442	20	60	100	dirty	≥ 5	**> 5**
Staphylococcus aureus ATCC 6538	20	60	100	dirty	≥ 5	**> 5**
Enterococcus hirae ATCC 10541	20	60	100	dirty	≥ 5	**> 5**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the bacterial test suspension, N_W = the number of cfu/ml of the control bacterial test suspension, N_a = the number of survivors per ml in the test mixture at the end of the contact time, N_{v0}= the number of cfu/ml of the bacterial test suspension in the mixture A,B,C at the beginning of the contact time (time „0"), A,B,C = the number of survivors per ml in control tests (A – experimental conditions validation, B – neutralizer toxicity validation, C – method validation), $R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

Prepared by: Mgr. Mirka Horáková, Ph.D., Lab Technician

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

Experimental conditions: **Testing of disinfecting efficiency of chemical disinfecting and antiseptic agents on carriers** SOP-M-22-12 (EN 14562:2006)

Period of analysis:	4.9. – 7.9.2017
Test temperature:	20 °C ± 1 °C
Test method:	dilution neutralization method
Neutralization medium:	Dey-Engley Neutralizing Broth M 1062
Appearance of the product:	colourless liquid
Test concentration:	100% (concentrated)
Contact time:	15, 30 and 60 min
Interfering substances:	3 g/l BSA and 3 ml/l sheep erythrocytes (dirty conditions)
Test organisms:	*Candida albicans* ATCC 10231
Incubation conditions:	30 °C ± 1 °C, 48 hours and additional period of 24 or 48 hours

Test procedure:

1. Preparation of the test suspension
2. Preparation of the product test solutions
3. Quantitative carrier test
4. Incubation and calculation
5. Expression and interpretation of the results

Note:
Fungicidal activity – the capability of a product to produce a reduction in the number of relevant organisms on carriers under defined conditions by at least 4 orders (10^4).
Yeasticidal activity – the capability of a product to produce a reduction in the number of viable fungi belonging to reference strain *Candida albicans* on carriers under defined conditions by at least 4 orders (10^4).
The drying time: 30-35 min
$R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

The standard:
EN 14562:2006 Chemical disinfectants and antiseptics – Quantitative carrier test for the evaluation of fungicidal or yeasticidal activity for instruments used in the medical area - Test method and requirements (phase 2, step 2) May 2006

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

11. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Candida albicans* ATCC 10231 on carriers

Tab No. 11.1 Verification of methodology, dirty conditions

Validation of suspension (N_{vo})			Validation of selected experimental conditions (A)			Neutralizer toxicity control (B)			Method validation (C) Product conc. 100%		
V_{c1}	43	$\Phi_{Nvo} = 36.5$	V_{c1}	39	$\Phi_A = 37.5$	V_{c1}	33	$\Phi_B = 35.5$	V_{c1}	39	$\Phi_C = 35$
V_{c2}	30		V_{c2}	36		V_{c2}	38		V_{c2}	31	
$30 \leq \Phi_{Nvo} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nvo}$			$\Phi_B \geq 0.5\ \Phi_{Nvo}$			$\Phi_C \geq 0.5\ \Phi_{Nvo}$		
x yes		no	x yes		no	x yes		no	x yes		no

Tab No. 11.2 Test suspension

Test suspension (N)	N	V_{c1}	V_{c1}	$\Phi = 47.5 \times 10^7 = \lg 8.68$ $8.17 \leq \lg N \leq 8.70$
	10^{-6}	> 330	> 330	
	10^{-7}	45	50	
				x yes / no

Tab No. 11.2.1 The control test suspension, dirty conditions

Test suspension (N_W)	N_W	V_{c1}	V_{c2}	$\Phi \times 10 = 1615 \times 10^3 = \lg 6.21$ $\lg N_W = \lg 6.21$ $6.15 \leq \lg N_w \leq (\lg N - 1.3)\ 7.38$
	10^{-3}	155	168	
				x yes / no

Tab No. 11.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Candida albicans* ATCC 10231 on carriers

Test concentration (%) / contact time (min) / conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_W = \lg 6.21$)**
100/15/dirty	10^0	<14	<14	< 2.15	**≥ 4.06**
100/30/dirty	10^0	<14	<14	< 2.15	**≥ 4.06**
100/60/dirty	10^0	<14	<14	< 2.15	**≥ 4.06**

12. Evaluation of yeasticidal activity of the product **GLOBACID SF MED 1.25%**

Tab No. 12.1 The efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on test strains – yeasticidal activity

Yeasticidal activity of the product (EN 14562:2006)						
Strain	Test temperature [°C]	Contact time [min]	Product test concentrations [%]	Interfering substances - conditions	lg R EN 14562:2006	**lg R**
Candida albicans ATCC 10231	20	15	100	dirty	≥ 4	**> 4**
Candida albicans ATCC 10231	20	30	100	dirty	≥ 4	**> 4**
Candida albicans ATCC 10231	20	60	100	dirty	≥ 4	**> 4**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the fungal test suspension, N_W = the number of cfu/ml of the control fungal test suspension, N_a = the number of survivors per ml in the test mixture at the end of the contact time, N_{v0}= the number of cfu/ml of the fungal test suspension in the mixture A,B,C at the beginning of the contact time (time „0"), A,B,C = the number of survivors per ml in control tests (A – experimental conditions validation, B – neutralizer toxicity validation, C – method validation), $R = N_W / N_a$ nebo $\lg R = \lg N_W - \lg N_a$ = the reduction in viability

Prepared by: Mgr. Mirka Horáková, Ph.D., Lab Technician

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

Experimental conditions: **Testing of disinfecting efficiency of chemical disinfecting and antiseptic agents by suspension method**
SOP-M-19-00 (EN 14348:2005)

Period of analysis:	11.9. – 2.10.2017
Test temperature:	20 °C ± 1 °C
Test method:	membrane filtration method
Filtration diluent:	rinsing liquid
Appearance of the product:	colourless liquid
Test concentration:	100% (concentrated)*
Contact time:	15, 30 and 60 min
Interfering substances:	3 g/l BSA and 3 ml/l sheep erythrocytes (dirty conditions)
Test organisms:	*Mycobacterium terrae* ATCC 15755
Incubation conditions:	37 °C ± 1 °C, 21 days

Test procedure:

1. Preparation of test suspension
2. Preparation of product test solutions
3. Quantitative suspension test
4. Incubation and calculation
5. Expression and interpretation of results

Note:
Mycobactericidal activity – the capability of a product to produce a reduction in the number of viable cells of *Mycobacterium terrae* and *Mycobacterium avium* under defined conditions by at least 4 orders (10^4).
Tuberculocidal activity - the capability of a product to produce a reduction in the number of viable cells of *Mycobacterium terrae* under defined conditions by at least 4 orders (10^4).
$R = N_0 / N_a$ nebo $\lg R = \lg N_0 - \lg N_a$ the reduction in viability
* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

The standard:
EN 14348:2005 Chemical disinfectants and antiseptics – Quantitative suspension test for the evaluation of mycobactericidal activity of chemical disinfectants in the medical area including instrument disinfectants - Test method and requirements (phase 2, step 1) January 2005

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

Sample ID: D90/2017
Rep No: 131
Sample name: **GLOBACID SF MED 1.25%**
Sampled: by client
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017

Sampling date: 5.5.2017
Sample delivered: 11.5.2017
Testing date: 4.9. – 2.10.2017
Delivered amount: 1 l

13. Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Mycobacterium terrae* ATCC 15755

Tab No. 13.1 Verification of methodology, dirty conditions

Validation of suspension (N_{vo})			Validation of selected experimental conditions (A)			Membrane filtration control (B)			Method validation (C) Product conc.: 100%*		
V_{c1}	31	$\Phi_{Nvo} = 31$	V_{c1}	25	$\Phi_A = 27.5$	V_{c1}	18	$\Phi_B = 29.5$	V_{c1}	27	$\Phi_C = 24.5$
V_{c2}	31		V_{c2}	30		V_{c2}	41		V_{c2}	22	
$30 \leq \Phi_{Nvo} \leq 160$			$\Phi_A \geq 0.5\ \Phi_{Nvo}$			$\Phi_B \geq 0.5\ \Phi_{Nvo}$			$\Phi_C \geq 0.5\ \Phi_{Nvo}$		
x	yes	no	x	yes	no	x	yes	no	x	yes	no

Tab No. 13.2 Test suspensions

Test suspension N	N	V_{c1}	V_{c1}	Test suspension N_0 (time = 0)
$\Phi = 32 \times 10^8 = \lg 9.51$	10^{-7}	> 165	> 165	$\lg N_0 = \lg N/100 = \lg 7.51$
$9.17 \leq \lg N \leq 9.70$	10^{-8}	28	36	$7.17 \leq \lg N_0 \leq 7.70$
				x yes no

Tab No. 13.3 Testing the efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on *Mycobacterium terrae* ATCC 15755

Test concentration (%)/contact time (min)/conditions	Dilution after test procedure	V_{c1}	V_{c2}	$\lg N_a = \lg (\Phi_a \times 10)$	**lg R ($\lg N_0 = \lg 7.51$)**
100*/15/dirty	10^{-1}	18	19	3.27	**4.24**
100*/30/dirty	10^{-1}	<14	<14	< 3.15	**≥ 4.36**
100*/60/dirty	10^{-1}	<14	<14	< 3.15	**≥ 4.36**

14. Evaluation of mycobactericidal and tuberculocidal activity of the product **GLOBACID SF MED 1.25%**

Tab No. 14.1 The efficacy of chemical disinfectant **GLOBACID SF MED 1.25%** on test strain – mycobactericidal and tuberculocidal activity

Mycobactericidal and tuberculocidal activity of the product (EN 14348:2005)						
Strain	Test temperature [°C]	Contact time [min]	Product test concentrations [%]	Interfering substances - conditions	lg R EN 14348:2005	**lg R**
Mycobacterium terrae ATCC 15755	20	15	100*	dirty	≥ 4	**> 4**
Mycobacterium terrae ATCC 15755	20	30	100*	dirty	≥ 4	**> 4**
Mycobacterium terrae ATCC 15755	20	60	100*	dirty	≥ 4	**> 4**

Note: V_c = value is the number of cfu per ml, Φ = average V_{c1} a V_{c2} (1. + 2. duplicate V_c values), N = the number of cfu/ml of the test suspension, N_0 = the number of cfu/ml of the test suspension at the beginning of the contact time (time „0“), N_a = the number of survivors per ml in the test mixture at the end of the contact time and before the membrane filtration, N_v = the number of cfu/ml of the test suspension for validation, N_{vo}= the number of cfu/ml of the test suspension in the mixture A,B,C at the beginning of the contact time (time „0“), A,B,C = the number of survivors per ml in control tests (A – experimental conditions control, B – membrane filtration validation, C – method validation), $R = N_0 / N_a$ nebo $\lg R = \lg N_0 - \lg N_a$ the reduction in viability
* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

Prepared by: Ing. Eva Kremlová, Lab Technician

Description: *Testing the efficacy of chemical disinfectants and antiseptics*

☐Sample ID: D90/2017 — Sampling date: 5.5.2017
Sample ID: D90/2017 — Sampling date: 5.5.2017
Rep No: 131 — Sample delivered: 11.5.2017
Sample name: **GLOBACID SF MED 1.25%** — Testing date: 4.9. – 2.10.2017
Sampled: by client — Delivered amount: 1 l
Sampling point: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Client: Goodpoint Chemicals Ltd, Urda tee 3, Jälgimäe, Saku vald, Harjumaa, Estonia 76404
Batch No: 20042017 — Page: 15

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Interpretation:

Results of tests are in Tabs.

According to EN 13727:2012+A2:2015 the tested concentrated* product **GLOBACID SF MED 1.25%,** batch No. 20042017, in contact times 15, 30 and 60 min under dirty conditions at temperature 20 °C ± 1 °C by the membrane filtration method **decreased** the number of alive microbes *Pseudomonas aeruginosa* ATCC 15442, *Staphylococcus aureus* ATCC 6538, *Enterococcus hirae* ATCC 10541 by at least 5 (lg) orders.

According to EN 13624:2013 the tested concentrated* product **GLOBACID SF MED 1.25%,** batch No. 20042017, in contact times 15, 30 and 60 min under dirty conditions at temperature 20 °C ± 1 °C by the membrane filtration method **decreased** the number of alive microbes *Candida albicans* ATCC 10231 by at least 4 (lg) orders.

According to EN 14561:2006 the tested concentrated product **GLOBACID SF MED 1.25%,** batch No. 20042017, in contact times 15, 30 and 60 min under dirty conditions at temperature 20 °C ± 1 °C by the dilution neutralization method **decreased** on carriers the number of alive microbes *Pseudomonas aeruginosa* ATCC 15442, *Staphylococcus aureus* ATCC 6538, *Enterococcus hirae* ATCC 10541 by at least 5 (lg) orders.

According to EN 14562:2006 the tested concentrated product **GLOBACID SF MED 1.25%,** batch No. 20042017, in contact times 15, 30 and 60 min under dirty conditions at temperature 20 °C ± 1 °C by the dilution neutralization method **decreased** on carriers the number of alive microbes *Candida albicans* ATCC 10231 by at least 4 (lg) orders.

The tested concentrated* product **GLOBACID SF MED 1.25%,** batch No. 20042017, in contact times 15, 30 and 60 min under dirty conditions at temperature 20 °C ± 1 °C by the membrane filtration method **decreased** the number of alive microbes *Mycobacterium terrae* ATCC 15755 by at least 4 (lg) orders (EN 14348:2005).

* Product can only be tested at a concentration of 97% (RTU) or less, as some dilution is always produced by adding the test organisms and interfering substance.

Conclusion:

The product **GLOBACID SF MED 1.25%** is capable of reducing the number of viable bacterial, mycobacterial and vegetative yeast cells of the relevant organisms in the suspension and on carriers under defined conditions to the declared values, and consequently, may be called bactericidal, tuberculocidal and yeasticidal.

8.12.2017, Hodonín

ACCREDITED BY CZECH ACCREDITATION INSTITUTE -2- Chemla, spol. s r.o. 695 01 Hodonín CZ č. 1273

Ing. Barbora Stoklásková, Leader of Study