

Test Report

Bactericidal activity of the disinfectant

“TEGO 2000”

According to BS EN 1276 (1997) “TEGO 2000“ possesses bactericidal activity at a concentration of 0.125 % or 0.5 % for **obligatory** purposes at 20°C with a contact time of 5 min and under low or high level soiling (0.3 g/l or 3.0 g/l bovine albumin) conditions for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Enterococcus hirae* ATCC 10541, *Pseudomonas aeruginosa* ATCC 15442 and *Escherichia coli* ATCC 10536.

According to BS EN 1276 (1997) “TEGO 2000“ possesses bactericidal activity at a concentration of 0.5 % or 1.0 % for specific purposes at 10°C with a contact time of 5 min and under low or high level soiling (0.3 g/l or 3.0 g/l bovine albumin) conditions for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Enterococcus hirae* ATCC 10541, *Pseudomonas aeruginosa* ATCC 15442 and *Escherichia coli* ATCC 10536.

Customer

Evonik Goldschmidt GmbH
Goldschmidtstr. 100
45127 Essen

Test laboratory

Technische Mikrobiologie
Dr. Jutta Höffler GmbH
Ahrensburger Str. 162
22045 Hamburg

Hamburg, 2009-05-20



DAC - PL - 0217 - 03
Accreditation according to
DIN EN ISO/IEC 17025

Bankverbindung · Hamburger Sparkasse · Konto Nr. 1208/115 319 · BLZ 200 505 50

IBAN DE 12 2005 0550 1208 1153 19 · BIC: HASPDEHH XXX · Geschäftsführerin: Dr. rer. nat. Jutta Höffler

Amtsgericht Hamburg Handelsregister B Nr. 58 682 · St.Nr.: 08/850/00703 · UST.IdNr. DE 171622643



Test report

BS EN 1276, bactericidal activity

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Goldschmidtstr. 100
45127 Essen

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22045 Hamburg

Disinfectant sample

Name of the product **TEGO 2000**
ID.No **ES 68757650**
Manufacturer..... **Evonik Goldschmidt GmbH,
Goldschmidtstr. 100
45127 Essen**
Date of delivery.....**2009-04-30**
Storage conditions.....**room temperature**
Active substance(s).....**amphoteric surfactants**

Test method and its validation

Method.....**quantitative suspension test,
dilution-neutralization method**
Neutralizer**polysorbate 80 (30 g/l),
saponin (30 g/l), histidine (1 g/l),
lecithin (3 g/l)**



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Experimental conditions

Period of analysis.....	2009-05-06 bis 2009-05-20
Diluent used for product test solutions	sterile hard water (300 mg/kg CaCO ₃)
Product test concentrations	0.01; 0.125; 0.25; 0.5; 1.0 % (V/V)
Appearance of the product and its dilutions	yellowish to colourless clear solutions
Stability of the mixture	no precipitation during the test
Temperature of incubation.....	37°C ± 1°C
test organisms	Staphylococcus aureus ATCC 6538 Pseudomonas aeruginosa ATCC 15442 Escherichia coli ATCC 10536 Enterococcus hirae ATCC 10541

Obligatory test conditions

Contact time.....	5 min ± 10 sec
Test temperature	20°C ± 1°C
Interfering substance(s).....	0.3 g/ l bovine albumin 3.0 g/ l bovine albumin

Specific test conditions

Contact time.....	5 min ± 10 sec
Test temperature	10°C ± 1°C
Interfering substance(s).....	0.3 g/ l bovine albumin 3.0 g/ l bovine albumin

Test results

s. tables D.1.1; D.1.2; D.2.1; D.2.2

Conclusion

According to BS EN 1276 (1997) the batch ES 68757650 of the product **TEGO 2000**

possesses bactericidal activity
at a concentration of **0.125 %** in hard water

for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Pseudomonas aeruginosa* ATCC 15442, *Escherichia coli* ATCC 10536 and *Enterococcus hirae* ATCC 10541 for **obligatory** purposes at **20°C** with a **5 min** contact time and under low level soiling (0.3 g/l bovine albumin) conditions.

According to BS EN 1276 (1997) the batch ES 68757650 of the product **TEGO 2000**

possesses bactericidal activity
at a concentration of **0.5 %** in hard water

for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Pseudomonas aeruginosa* ATCC 15442, *Escherichia coli* ATCC 10536 and *Enterococcus hirae* ATCC 10541 for **obligatory** purposes at **20°C** with a **5 min** contact time and under high level soiling (3.0 g/l bovine albumin) conditions.

According to BS EN 1276 (1997) the batch ES 68757650 of the product **TEGO 2000**

possesses bactericidal activity
at a concentration of **0.5 %** in hard water

for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Pseudomonas aeruginosa* ATCC 15442, *Escherichia coli* ATCC 10536 and *Enterococcus hirae* ATCC 10541 for specific purposes at **10°C** with a **5 min** contact time and under low level soiling (0.3 g/l bovine albumin) conditions.

According to BS EN 1276 (1997) the batch ES 68757650 of the product **TEGO 2000**

possesses bactericidal activity
at a concentration of **1.0 %** in hard water

for the referenced strains of *Staphylococcus aureus* ATCC 6538, *Pseudomonas aeruginosa* ATCC 15442, *Escherichia coli* ATCC 10536 and *Enterococcus hirae* ATCC 10541 for specific purposes at **10°C** with a **5 min** contact time and under high level soiling (3.0 g/l bovine albumin) conditions.

Technische Mikrobiologie

Hamburg, 2009-05-20

Dr. Jutta Höffler GmbH



Dr. Jutta Höffler

Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas/ BS EN 1276

product: TEGO 2000, Lot ES 68757650

test conditions: 5 min, 20°C, bovine albumin 0.3 g/l

Table D.1.1 Test results

Test organisms	Validation test				Bacterial test suspension (5.4.1.4)	Test procedure at concentration (V/V) (see 5.5.2)				
	Bacterial suspension (see A.2)	Experimental conditions (see A.4.1.a)	Neutralizer toxicity control (see A.4.1.b)	Dilution-neutralization control (see A.4.1.c)			0.01 %	0.125 %	0.25 %	0.5 %
Staphylococcus aureus ATCC 6538	V _c : 113/105 N _v : 1.1 x 10 ³	V _c : 108/148 A: 1.3 x 10 ²	V _c : 176/156 B: 1.7 x 10 ²	V _c : 122/144 C: 1.3 x 10 ²	10 ⁻⁶ : 222/230 10 ⁻⁷ : 26/23 N: 2.3 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <7.7 x 10 ³	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵
Pseudomonas aeruginosa ATCC 15442	V _c : 288/262 N _v : 2.8 x 10 ³	V _c : 154/144 A: 1.5 x 10 ²	V _c : 186/146 B: 1.7 x 10 ²	V _c : 200/128 C: 1.6 x 10 ²	10 ⁻⁶ : >300/>300 10 ⁻⁷ : 50/50 N: 5.0 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	0/0 <1.5 x 10 ² >3.3 x 10 ⁵	0/0 <1.5 x 10 ² >3.3 x 10 ⁵	0/0 <1.5 x 10 ² >3.3 x 10 ⁵
Escherichia coli ATCC 10536	V _c : 88/112 N _v : 1.0 x 10 ³	V _c : 70/100 A: 8.5 x 10 ¹	V _c : 120/128 B: 1.2 x 10 ²	V _c : 58/52 C: 5.5 x 10 ¹	10 ⁻⁶ : 152/160 10 ⁻⁷ : 16/19 N: 1.6 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <5.3 x 10 ³	0/0 <1.5 x 10 ² >1.1 x 10 ⁵	0/0 <1.5 x 10 ² >1.1 x 10 ⁵	0/0 <1.5 x 10 ² >1.1 x 10 ⁵
Enterococcus hirae ATCC 10541	V _c : 111/155 N _v : 1.3 x 10 ³	V _c : 160/166 B: 1.6 x 10 ²	V _c : 120/184 B: 1.5 x 10 ²	V _c : 192/148 C: 1.7 x 10 ²	10 ⁻⁶ : 240/220 10 ⁻⁷ : 37/27 N: 2.4 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	0/0 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵
V _c	viable count									
N	Number of cfu/ml of the bacterial test suspension (see 5.4.1.4)									
N _v	Number of cfu/ml of the bacterial suspension (see A.2)									
R	reduction in viability (see 5.6.3)									
N _a	number of cfu/ml in the test mixture (see 5.5.2.2.3)									
A	number of cfu/ml of the experimental conditions validation (see A.4.1a)									
B	number of cfu/ml of the neutralizer toxicity validation (see A.4.1b)									
C	the number of cfu/ml of the dilution-neutralization validation (see A.4.1.c)									

Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas/ BS EN 1276

product: TEGO 2000, Lot ES 68757650

test conditions: 5 min, 20°C, bovine albumin 3.0 g/l

Table D.1.2 Test results										
Test organisms	Validation test				Bacterial test suspension (5.4.1.4)	Test procedure at concentration (V/V) (see 5.5.2)				
	Bacterial suspension (see A.2)	Experimental conditions (see A.4.1.a)	Neutralizer toxicity control (see A.4.1.b)	Dilution-neutralization control (see A.4.1.c)			0.125 %	0.25 %	0.5 %	1.0%
Staphylococcus aureus ATCC 6538	V _c : 113/105 N _v : 1.1 x 10 ³	V _c : 156/112 A: 1.3 x 10 ²	V _c : 176/156 B: 1.7 x 10 ²	V _c : 128/128 C: 1.3 x 10 ²	10 ⁻⁶ : 222/230 10 ⁻⁷ : 26/23 N: 2.3 x 10 ⁸	V _c N _a R	48/70 5.9 x 10 ² 3.9 x 10 ⁴	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵
Pseudomonas aeruginosa ATCC 15442	V _c : 288/262 N _v : 2.8 x 10 ³	V _c : 160/128 A: 1.4 x 10 ²	V _c : 186/146 B: 1.7 x 10 ²	V _c : 178/172 C: 1.8 x 10 ²	10 ⁻⁶ : >300/>300 10 ⁻⁷ : 50/50 N: 5.0 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	12/8 <1.5 x 10 ² >3.3 x 10 ⁵	0/0 <1.5 x 10 ² >3.3 x 10 ⁵	0/0 <1.5 x 10 ² >3.3 x 10 ⁵
Escherichia coli ATCC 10536	V _c : 88/112 N _v : 1.0 x 10 ³	V _c : 104/74 A: 8.9 x 10 ¹	V _c : 120/128 B: 1.2 x 10 ²	V _c : 60/62 C: 6.1 x 10 ¹	10 ⁻⁶ : 152/160 10 ⁻⁷ : 16/19 N: 1.6 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <5.3 x 10 ³	>300/>300 >3.0 x 10 ³ <5.3 x 10 ³	0/0 <1.5 x 10 ² >1.1 x 10 ⁵	0/0 <1.5 x 10 ² >1.1 x 10 ⁵
Enterococcus hirae ATCC 10541	V _c : 111/155 N _v : 1.3 x 10 ³	V _c : 188/212 B: 2.0 x 10 ²	V _c : 120/184 B: 1.5 x 10 ²	V _c : 112/190 C: 1.5 x 10 ²	10 ⁻⁶ : 240/220 10 ⁻⁷ : 37/27 N: 2.4 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	0/0 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵
V _c	viable count									
N	Number of cfu/ml of the bacterial test suspension (see 5.4.1.4)									
N _v	Number of cfu/ml of the bacterial suspension (see A.2)									
R	reduction in viability (see 5.6.3)									
N _a	number of cfu/ml in the test mixture (see 5.5.2.2.3)									
A	number of cfu/ml of the experimental conditions validation (see A.4.1a)									
B	number of cfu/ml of the neutralizer toxicity validation (see A.4.1b)									
C	the number of cfu/ml of the dilution-neutralization validation (see A.4.1.c)									

Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas/ BS EN 1276

product: TEGO 2000, Lot ES 68757650

test conditions: 5 min, 10°C, bovine albumin 0.3 g/l

Table D.2.1 Test results

Test organisms	Validation test				Bacterial test suspension (5.4.1.4)	Test procedure at concentration (V/V) (see 5.5.2)				
	Bacterial suspension (see A.2)	Experimental conditions (see A.4.1.a)	Neutralizer toxicity control (see A.4.1.b)	Dilution-neutralization control (see A.4.1.c)			0.01 %	0.125 %	0.25 %	0.5 %
Staphylococcus aureus ATCC 6538	V _c : 113/105 N _v : 1.1 x 10 ³	V _c : 138/118 A : 1.3 x 10 ²	V _c : 98/96 B : 9.7 x 10 ¹	V _c : 116/104 C : 1.1 x 10 ²	10 ⁻⁶ : 222/230 10 ⁻⁷ : 26/23 N : 2.3 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <7.7 x 10 ³	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵
Pseudomonas aeruginosa ATCC 15442	V _c : 288/262 N _v : 2.8 x 10 ³	V _c : 214/198 A : 2.1 x 10 ²	V _c : 200/220 B : 2.1 x 10 ²	V _c : 168/192 C : 1.8 x 10 ²	10 ⁻⁶ : >300/>300 10 ⁻⁷ : 50/50 N : 5.0 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	2/0 <1.5 x 10 ² >3.3 x 10 ⁵
Escherichia coli ATCC 10536	V _c : 88/112 N _v : 1.0 x 10 ³	V _c : 68/70 A : 6.9 x 10 ¹	V _c : 68/92 B : 8.0 x 10 ¹	V _c : 60/66 C : 6.3 x 10 ¹	10 ⁻⁶ : 152/160 10 ⁻⁷ : 16/19 N : 1.6 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ >5.3 x 10 ³	>300/>300 >3.0 x 10 ³ >5.3 x 10 ³	0/0 <1.5 x 10 ² >1.1 x 10 ⁵	0/0 <1.5 x 10 ² >1.1 x 10 ⁵
Enterococcus hirae ATCC 10541	V _c : 111/155 N _v : 1.3 x 10 ³	V _c : 106/100 B : 1.0 x 10 ²	V _c : 112/80 B : 9.6 x 10 ¹	V _c : 100/106 C : 1.0 x 10 ²	10 ⁻⁶ : 240/220 10 ⁻⁷ : 37/27 N : 2.4 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	10/18 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵
V _c	viable count									
N	Number of cfu/ml of the bacterial test suspension (see 5.4.1.4)									
N _v	Number of cfu/ml of the bacterial suspension (see A.2)									
R	reduction in viability (see 5.6.3)									
N _a	number of cfu/ml in the test mixture (see 5.5.2.2.3)									
A	number of cfu/ml of the experimental conditions validation (see A.4.1a)									
B	number of cfu/ml of the neutralizer toxicity validation (see A.4.1b)									
C	the number of cfu/ml of the dilution-neutralization validation (see A.4.1.c)									

Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas/ BS EN 1276

product: TEGO 2000, Lot ES 68757650

test conditions: 5 min, 10°C, bovine albumin 3.0 g/l

Table D.2.2 Test results

Test organisms	Validation test				Bacterial test suspension (5.4.1.4)	Test procedure at concentration (V/V) (see 5.5.2)				
	Bacterial suspension (see A.2)	Experimental conditions (see A.4.1.a)	Neutralizer toxicity control (see A.4.1.b)	Dilution-neutralization control (see A.4.1.c)			0.125 %	0.25 %	0.5 %	1.0%
Staphylococcus aureus ATCC 6538	V _c : 113/105 N _v : 1.1 x 10 ³	V _c : 106/76 A : 9.1 x 10 ¹	V _c : 98/96 B : 9.7 x 10 ¹	V _c : 144/108 C : 1.3 x 10 ²	10 ⁻⁶ : 222/230 10 ⁻⁷ : 26/23 N : 2.3 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <7.6 x 10 ³	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵	0/0 <1.5 x 10 ² >1.5 x 10 ⁵
Pseudomonas aeruginosa ATCC 15442	V _c : 288/262 N _v : 2.8 x 10 ³	V _c : 188/168 A : 1.8 x 10 ²	V _c : 200/220 B : 2.1 x 10 ²	V _c : 176/208 C : 1.9 x 10 ²	10 ⁻⁶ : >300/>300 10 ⁻⁷ : 50/50 N : 5.0 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	>300/>300 >3.0 x 10 ³ <1.7 x 10 ⁴	96/90 9.3 x 10 ² 5.4 x 10 ⁴	0/0 <1.5 x 10 ² >3.3 x 10 ⁵
Escherichia coli ATCC 10536	V _c : 88/112 N _v : 1.0 x 10 ³	V _c : 104/122 A : 1.1 x 10 ²	V _c : 68/92 B : 8.0 x 10 ¹	V _c : 60/50 C : 5.5 x 10 ¹	10 ⁻⁶ : 152/160 10 ⁻⁷ : 16/19 N : 1.6 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <5.3 x 10 ³	>300/>300 >3.0 x 10 ³ <5.3 x 10 ³	0/0 <1.5 x 10 ² >1.1 x 10 ⁵	0/0 <1.5 x 10 ² >1.1 x 10 ⁵
Enterococcus hirae ATCC 10541	V _c : 111/155 N _v : 1.3 x 10 ³	V _c : 112/66 B : 8.9 x 10 ¹	V _c : 112/80 B : 9.6 x 10 ¹	V _c : 102/110 C : 1.1 x 10 ²	10 ⁻⁶ : 240/220 10 ⁻⁷ : 37/27 N : 2.4 x 10 ⁸	V _c N _a R	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	>300/>300 >3.0 x 10 ³ <8.0 x 10 ³	0/0 <1.5 x 10 ² >1.6 x 10 ⁵	0/0 <1.5 x 10 ² >1.6 x 10 ⁵
V _c	viable count									
N	Number of cfu/ml of the bacterial test suspension (see 5.4.1.4)									
N _v	Number of cfu/ml of the bacterial suspension (see A.2)									
R	reduction in viability (see 5.6.3)									
N _a	number of cfu/ml in the test mixture (see 5.5.2.2.3)									
A	number of cfu/ml of the experimental conditions validation (see A.4.1a)									
B	number of cfu/ml of the neutralizer toxicity validation (see A.4.1b)									
C	the number of cfu/ml of the dilution-neutralization validation (see A.4.1.c)									