

Laboratory Report

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Your date of order:	23.02.2016	Receipt of samples / sampling:	24.02.2016			
Your order no .:	PO: 4500771813	Period of analysis:	0106.03.2016			
BMA order no .:	AU160223-07	Date of report:	22.03.2016			
BMA sample no.:	160224-10/1	Report no.:	BE160223-07/1/K1 Ersatz für BE160223-07/1			
Report writing:	U. Stephan		Page 1 of 2			

Sample

Cloth: MicronQuick (The sample was sent by the customer)

Analyses

Microbiological examination of products Evaluation of the capability of the cloth MicronQuick to reduce bacteria on floor surfaces

1. Method(s) and material

The present study is based on EN 1174-2, DIN EN ISO 846, method C the customer's instructions and BMA-Laboratory reports of AU141212-07. The test was performed in a clean bench.

Bacteria test strain: Pseudomonas aeruginosa (DSM-Nr. 288)

Neutral cleaner: Tana Green care Neutral-Reiniger 04631 (TANA Chemie GmbH, Mainz, Germany)

Test surface: PVC floor covering, non structured (98 cm x 27 cm), disinfected; subdivided into 39 sample squares (9 cm x 7 cm)

Samples 1.1 to 1.3: Negative control after disinfection, samples 1.4 to 1.9: Positive control after bacteria application, samples 1.10 to 1.39: treated sample squares after bacteria application and cleaning with the test cloth.

Applied bacteria suspension

5 ml of P. aeruginosa suspension 7 x 10⁹ cfu (colony forming unit); calculated amount per test sample (9 cm x 7 cm): 1,9 x 10⁸.

Eluation and determination of bacteria from the sample squares

Samples were incubated in 15 ml 0,9% NaCl solution in a falcon tube and shaked 20 min end to end. The bacteria concentration of the suspension was analysed using the spread plate method (100 μ l plating volume of dilution series) and/or the filtration method for samples with expected high or low bacteria contamination respectively. The agar plates were cultivated 5 days at 30°C.

Cleaning procedure

The cloth was fixed in a double layer to lab a testing device (provided by the customer) consisting of a massive plastic block (9 cm x 6 cm) and ensuring a homogenous pressure. It was moistened by spraying 5 ml water with 1% neutral cleaner (without disinfectants). Then the cloth with the block was wiped once across the test surface by moving it in form of an 8 at a speed of approx. 5 cm/s based on instrutions of the manufacturer.

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2. Results

2.1 Amount of aerobic mesophilic bacteria after disinfection

The results of the measurements and analyses exclusively refer to the examined sample(s).

			Mean concentration of mesophilic bacteria	
Sample / Sample identification	Sample No.	Sample description	Sample square (63 cm ²) [CFU/15 ml]	[CFU/m²]
Control	1 1-1 3	Negative control after	1 (a)	1.6 x 10 ²
160224-10/1	1.0	disinfection		1,0 × 10

^(a) Detection limit filtration: 1 cfu/15 ml

^(b) Detection limit spread plate: 150 cfu/15 ml

2.2 Amount of Pseudomonas aeruginosa before and after cleaning

The results of the measurements and analyses exclusively refer to the examined sample(s).

			Mean concentration of P. aeruginosa	
Sample / Sample identification	Sample No.	Sample description	Sample square (63 cm ²) [CFU/15 ml]	[CFU/m ²]
	1.4-1.9	Positive control after bacteria application	2,2 x 10 ^{5 (b)}	3,5 x 10 ⁷
Cloth: MicronQuick 160224-10/1	1.10-1.39	Treated sample squares after bacteria application and cleaning with the test cloth	16 ^(a)	2,5 x 10 ³
		Reduction [%]	99,99	

^(a) Detection limit filtration: 1 cfu/15 ml

^(b) Detection limit spread plate: 150 cfu/15 ml

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