



**Swift Silliker (Pty) Ltd t/a Mérieux NutriSciences**  
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**CERTIFICATE OF ANALYSIS**

**COA No.:** CT 57187/21  
**COA Date:** 21 September 2021  
**Page:** 1 of 6

**Customer:** IQ Green Solutions (Pty) Ltd.  
**Order No.:** N/A  
**Client Reference:** N/A

**Analysed by:** Swift Silliker Pty. Ltd. t/a Mérieux NutriSciences  
 7 Warrington Road,  
 Claremont,  
 Western Cape, South Africa.

**Received from:** IQ Green Solutions (Pty) Ltd.  
 15 Jan van Riebeeck Drive,  
 Paarl,

**TO** IQ GREEN SOLUTIONS  
 (PTY) LTD  
 15 JAN van RIEBEECK DRIVE  
 PAARL  
 7620  
 MOHAMED MOOSA  
[mohamed@iq-greensolutions.com](mailto:mohamed@iq-greensolutions.com)

**\*\*Information provided by customer**

**DATE RECEIVED:** 20/08/2021  
**TEST TYPE:** FUNGICIDAL ACTIVITY  
**METHOD NO.:** SWM.MIC.017

**a) Sample Identification**

<input type="checkbox"/> Product Name**:	IQ DISINFECTANT SOLUTION
<input type="checkbox"/> Batch Number**:	21R0034
<input type="checkbox"/> Manufacturer/ Supplier**:	IQ Green Solutions (Pty) Ltd.
<input type="checkbox"/> Manufacturing Date**:	16 July 2021
<input type="checkbox"/> Exp. Date**:	N/A
<input type="checkbox"/> Laboratory Ref No.:	CT 57187/21
<input type="checkbox"/> Storage Conditions:	Ambient
<input type="checkbox"/> Active Substances and their concentrations**:	Ano-Lyte® (HOCl +/- 0.05%)
<input type="checkbox"/> Appearance of the Product:	Liquid, clear, colourless
<input type="checkbox"/> Product diluent recommended by manufacturer for use:	Standard water

**b) Methods Used:**

SANS 51650:2011 – Evaluation of Fungicidal Activity of Chemical disinfectants & Antiseptics (Neutralization by dilution Method)

Directors: V. Stewart (Managing), A. Lambrechts, P. Sans (Finance), S. Schneider (Finance), J.F. Billiet (Finance) / Reg. No 2009/025067/07

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### c) Experimental Conditions - SANS 51650 Edition 2

<input type="checkbox"/> Test Strains:	<i>Aspergillus brasiliensis</i> ATCC16404 <i>Candida albicans</i> ATCC10231
<input type="checkbox"/> Product Test Concentrations**:	0.1%, 50% & 80% <b>(Please note:</b> a NEAT concentration is equal to 80% of sample. Products can only be tested at a concentration of 80 % or less, as some dilution is always produced by adding the test organisms and interfering substance.)
<input type="checkbox"/> Appearance of Diluted Product:	0.1% - Liquid, clear, colourless  50% - Liquid, clear, colourless  80% - Liquid, clear, colourless
<input type="checkbox"/> Interfering Substance:	0.3g/l Bovine Albumin, Clean conditions
<input type="checkbox"/> Contact Time:	5 minutes
<input type="checkbox"/> Test Temperature:	20°C
<input type="checkbox"/> Rinsing Solution:	Sodium thiosulphate (20g/l) + Tween 80 (30g/l) + Lecithin (3g/l)
<input type="checkbox"/> Incubation Conditions:	Aerobic incubation: 30°C ± 1°C
<input type="checkbox"/> Incubation Media:	Malt Extract Agar
<input type="checkbox"/> Testing Period:	31/08-17/09/2021

### d) Test Results: see tables 1-2.

### e) Summary of results

#### Fungicidal Efficacy

Organism	Experimental conditions	Product Conc.	Contact time	CFU/ML: Start	CFU/ML: End	Log Reduction (Log R= $\geq 5$ )	Evaluation
<i>Aspergillus brasiliensis</i> ATCC16404	Obligatory, Clean conditions	0.1%	15 minutes	6.64	>3.22	<3.42	FAIL
		50%		6.64	<2.15	>4.49	PASS
		80%		6.64	<2.15	>4.49	PASS
<i>Candida albicans</i> ATCC10231	Obligatory, Clean conditions	0.1%	15 minutes	6.40	>3.52	<2.88	FAIL
		50%		6.40	<2.15	>4.25	PASS
		80%		6.40	<2.15	>4.25	PASS

Directors: V. Stewart (Managing), A. Lamb rechts, P. Sans (France), S. Schneider (France), J-F. Billet (France) / Reg. No 2000/ 025067/07

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#### f) Conclusions

According to SANS 51650, the test product, **IQ DISINFECTANT SOLUTION**, when used at concentrations 50% & 80% has fungicidal activity ( $\log R \geq 4$ ) under the following test conditions:

- Contact time:** 15 minutes
- Temperature:** 20°C
- Interfering substance:** 0.3g/ L Bovine albumin – Clean conditions
- Test strains:** *Aspergillus brasiliensis* ATCC16404, *Candida albicans* ATCC10231

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Ref. Report section (d)

**ORGANISM: *Aspergillus brasiliensis* ATCC16404**

Obligatory Experimental Conditions

Table 1a: Validation test

Validation suspension ( $N_{V_0}$ )		Experimental Conditions Control (A)		Neutralizer Control (B)		Method Validation (C) (80% Product Concentration)	
	Ave		Ave		Ave		Ave
Vc1	38	Vc1	30	Vc1	24	Vc1	38
Vc2	44	Vc2	38	Vc2	28	Vc2	28
	41		34		26		33
Acceptance limits	$N_{V_0} = 30 - 160$	Acceptance limits	$\geq 0.5x N_{V_0}$	Acceptance limits	$\geq 0.5x N_{V_0}$	Acceptance limits	$\geq 0.5x N_{V_0}$
Complies	Yes	Complies	Yes	Complies	Yes	Complies	Yes

$0.5 \times N_{V_0} = 20.5$

Table 1b: Test suspensions

Dilution (Test suspension)	Vc1	Vc2	Average N (wm)	Log N	$N_0$	Log $N_0$
$10^{-5}$	480	410	$4.4 \times 10^7$	7.64	$4.4 \times 10^6$	6.64
$10^{-6}$	30	40				
Acceptance limits:	Log N is between 7.17 and 7.70		Complies	Yes		
Acceptance limits:	Log $N_0$ is between 6.17 and 6.70		Complies	Yes		
Acceptance limits:	Control of weighted mean counts: 12.7		Complies	Yes		

Table 1c: Log Reduction values

Product Concentration	Vc1	Vc2	$N_a$ (Ave Vc1 & Vc2 x 10)	Log $N_a$	Log Reduction ( $N_0$ : 6.64)	Contact time
0.1%	>165	>165	>1650	<3.22	<3.42	15 minutes
50%	<14	<14	<140	<2.15	>4.49	15 minutes
80%	<14	<14	<140	<2.15	>4.49	15 minutes

Where:

- VC = Viable Count
- N = Test suspension
- $N_0$  = Test suspension at beginning of contact time (t=0)
- $N_a$  = Test suspension (survivors) before neutralization
- $N_v$  = Validation suspension
- $N_{v_0}$  = Validation suspension at beginning of contact time
- A = number of cfu/mL of the experimental conditions control
- B = number of cfu/mL of the neutralization control
- C = number of cfu/mL of the method validation

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**ORGANISM: *Candida albicans* ATCC10231**

Obligatory Experimental Conditions

**Table 2a: Validation test**

Validation suspension ( $N_{V_0}$ )		Experimental Conditions Control (A)		Neutralizer Control (B)		Method Validation (C) (80% Product Concentration)	
	<b>Ave</b>		<b>Ave</b>		<b>Ave</b>		<b>Ave</b>
Vc1	34	Vc1	48	Vc1	35	Vc1	28
Vc2	39	Vc2	40	Vc2	45	Vc2	30
<b>Acceptance limits</b>	$N_{V_0} = 30 - 160$	<b>Acceptance limits</b>	$\geq 0.5x N_{V_0}$	<b>Acceptance limits</b>	$\geq 0.5x N_{V_0}$	<b>Acceptance limits</b>	$\geq 0.5x N_{V_0}$
<b>Complies</b>	Yes	<b>Complies</b>	Yes	<b>Complies</b>	Yes	<b>Complies</b>	Yes
0.5 x $N_{V_0} = 18.25$							

**Table 2b: Test suspensions**

Dilution (Test suspension)	Vc1	Vc2	Average N (wm)	Log N	$N_0$	Log $N_0$
$10^{-5}$	208	300	$2.5 \times 10^7$	7.40	$2.5 \times 10^6$	6.40
$10^{-6}$	18	28				
<b>Acceptance limits:</b>	Log N is between 7.17 and 7.70		<b>Complies</b>	Yes		
<b>Acceptance limits:</b>	Log $N_0$ is between 6.17 and 6.70		<b>Complies</b>	Yes		
<b>Acceptance limits:</b>	Control of weighted mean counts: 11.0		<b>Complies</b>	Yes		

**Table 2c: Log Reduction values**

Product Concentration	Vc1	Vc2	$N_a$ (Ave Vc1 & Vc2 x 10)	Log $N_a$	Log Reduction ( $N_0$ : 6.40)	Contact time
0.1%	>330	>330	>3300	>3.52	<2.88	15 minutes
50%	<14	<14	<140	<2.15	>4.25	15 minutes
80%	<14	<14	<140	<2.15	>4.25	15 minutes

**Where:**

- VC = Viable Count
- N = Test suspension
- $N_0$  = Test suspension at beginning of contact time (t=0)
- $N_a$  = Test suspension (survivors) before neutralization
- $N_v$  = Validation suspension
- $N_{v_0}$  = Validation suspension at beginning of contact time
- A = number of cfu/mL of the experimental conditions control
- B = number of cfu/mL of the neutralization control
- C = number of cfu/mL of the method validation

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### Test Validity

The test is valid when, for each test organism:

- $N$  (Test suspension) is between  $1,5 \times 10^7$  and  $5,0 \times 10^7$  ( $7,17 \leq \lg N \leq 7,70$ )
- $N_0$  (Test suspension) is between  $1,5 \times 10^6$  and  $5,0 \times 10^6$  ( $6,17 \leq \lg N_0 \leq 6,70$ )
- $N_{v0}$  is between 30 and 160 ( $3,0 \times 10^1$  and  $1,6 \times 10^2$ )
- $N_v$  is between  $3,0 \times 10^2$  and  $1,6 \times 10^3$
- $A, B, C$  are equal to or greater than  $0,5 \times N_{v0}$ .
- Control of weighted mean counts: quotient is not lower than 5 and not higher than 15.
- At least one of the test concentrations will demonstrate a log reduction of less than 4 log.

### Pass Requirements

#### Fungicidal activity for general purposes

The product shall be deemed to have passed the SANS 51650 standard if it demonstrates in a valid test, at least a 4 log reduction within 15 min at 20 °C with the chosen interfering substance simulating clean or dirty conditions as defined by SANS 51650, when the test organisms are *Aspergillus brasiliensis* and *Candida albicans*.

The fungicidal concentration for general purposes is the concentration active on the limiting strain.

#### Fungicidal activity for specific purposes

The fungicidal concentration for a specific purpose is the concentration of the tested product for which at least a 4 log reduction is demonstrated in a valid test, under the additional chosen test conditions. The product shall have passed the SANS 51650 standard under the obligatory test conditions. The fungicidal concentration for specific purposes may be lower than the one determined for general purposes.

### Special remarks regarding results:

- All controls and validation were within the basic limits.
- At least one concentration of the product demonstrated an lg reduction of less than 5 lg.
- No precipitate during the test procedure (test mixtures were homogeneous).

### Notes:

**Note 1:** Products can only be tested at a concentration of 80 % or less, as some dilution is always produced by adding the test organisms and interfering substance. A concentration indicated as NEAT therefore must be interpreted as a 80% solution.

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## Technical Signatory

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