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Client Information							
Company Name:	Advanced Vapor Technologies	Sponsor:	Rick Hoverson				
Sponsor's Phone:	1-800-997-6584	E-mail:	rick@advap.com				
Test Information							
Test(s) Performed:	Quantitative Custom Surface Time-Kill Rangefinding Test (Study ID NG2478-A2)						
Protocol Followed:	Quantitative Time-Kill	Performed by:	B. Tanner, A. Gregg				
Sample Information							
Test Substance ID(s):	VaporJet PC 2400 with TANCS™	Number of Devices:	1				
Parameters							
Microorganism(s):	K. pneumoniae (NDM-1 Strain)	Exposure Temp.	Ambient (~22 - 24°C)				
Subculture Number:	1	Type of Carrier:	Clay Test Surfaces				
Growth Medium:	Tryptic Soy Broth	# of Replicates:	Duplicate, With Duplicate Plating				
Contact Time(s):	Various, See Table Below	Incubation Temp.:	36.0 ± 1°C				
Product Dilution(s):	Device; Ready to Use	Incubation Time:	18 - 24 Hours				
Neutralizer Used:	D/E Broth (10 mL)						
Controls							
Neutralized:	Passed (All)	Growth Control:	Passed (All)				
Broth Sterility:	Passed, Control D/E Tube Neg.	Agar Sterility:	Passed (All)				
Test Results							
Controls Performance:	Normal	Test(s) Valid?:	Yes				
Notes: After trea	tment of the surfaces by the VaporJet PC	2400 with TANCS™ (low	v setting) for the contact time, surfaces				
were left for ~20 second	s at room temperature, then harvested.	Additionally, after each m	ultiple-contact time series of				
	s sectioned and analyzed for target micro						
	nicroorganisms (likely <i>Bacillus</i> endospor		·				
	. ,	•					
Tests Completed:	3-Feb-2011	Report Sent:	9-Feb-2011				

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## Summary of Study Procedure

## <u>Preparation and Inoculation of Carriers</u>

- Small clay test surfaces were obtained for testing.
- Each brick selected was checked for fit into machined aluminum carrier holders.
- Test surfaces were sterilized prior to testing.
- Test surfaces were inoculated, then set aside until visibly dry (about 20 minutes).

#### Preparation of Test Device

- The Water reservoir for the steamer was filled with ordinary tap water.
- The test device was turned on and allowed to equilibrate for 10 minutes.
- The test device was set to the lowest setting for all testing.

#### **Test Execution**

- Inoculated, dried test surfaces were treated with the device for differing amounts of time, (0.5, 1, 2, and 5 seconds), and light to moderate pressure was applied during each test.
- The treated coupons were allowed an approximately 20 second rest time prior to harvesting.
- The treated coupons were harvested and enumerated relative to "time-zero" controls through elution with D/E broth and plating on Tryptic Soy Agar.
- Microbial reductions were calculated and reported.

#### **Additional Notes**

- A clean towel was used for set A, and then another clean towel was used for set B.
- A zone of inhibition test was performed using the test organism and the antibiotic Ertapenem (10  $\mu$ g) to verify the strength and strain of the test organism.

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• The test organism was shown to be Ertapenem (carbapenem) resistant.

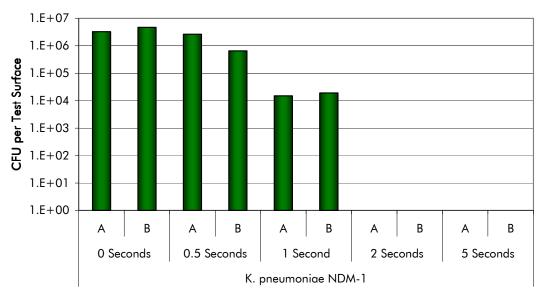
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## Summary Table and Chart

Microorganism	Exposure Time (seconds)	Replicate	CFU/Surface	Average CFU/Surface	Average Percent Reduction
K. pneumoniae NDM-1	Initial Dry Inoculum (No Exposure)	Α	3.30E+06	3.98E+06	n/a
		В	4.65E+06		n/α
	0.5	Α	2.65E+06	1.65E+06 58.490	58.49057%
		В	6.50E+05		30.47037 //
	1	Α	1.50E+04	1.70E+04	99.57233%
		В	1.90E+04		
	2	Α	<5	<5	99.99990%
		В	<5		77.77770/6
	5	Α	<5	<5	99.99990%
		В	<5		

## VaporJet PC 2400 with TANCS

Note: Towels analyzed post-treatment - all were free from contamination.



Note: Values Below Limit of Detection (5 CFU/Surface) Expressed as Zero on this Chart

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#### Additional Information

#### Method of Calculation for Percent Reduction:

% Reduction = 1- (C/B)\*100, where:

B = Average number of viable cells on the control pieces after 24 hours.

C = Average number of viable cells on the test pieces after 24 hours.

#### Photograph from the Study:



Test carriers inoculated with the test organism prior to treatment with cleaning instrument.

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# Photograph from the Study:



Towel fabric being removed from the head of the instrument, to be further analyzed. All towels were shown by lab analysis to be free from *K. pneumoniae* NDM-1 after use in the study.

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