

I.A.Q.

Environment Conditioning Technology

Five Supporting Technologies

- Filtration
- Ionization
- Ozone
- Germicidal UV
- UV w/Target Plates

PASSIVE versus ACTIVE

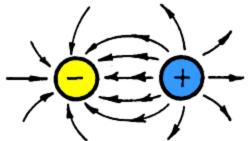
- Filtration—bring the pollution to the solution
 - Low–Resistant Airflow Technologies
 - HEPA Grade MERV Rated Technologies
- Purification—take the solution to the pollution
 - Heart of systems—UV W/Target Technology
 - Multi-strategic approach with Multiple Technologies

Multi-Strategic Approach

- 1) <u>lonization</u>—to precipitate particles
- 2) Advanced Oxidation—utilizing low levels of ozone, UV, and UV w/target technologies to break down VOCs and deactivate pathogens
- 3) Filtration—to capture particles

1) Ionization

<u>Ions</u> are charged particles which travel out into the environment, charging particulate in the air—oppositely charged particles attract one another, allowing them to gain sufficient mass to be filtered or fall to the ground. activTek technology may incorporates one or both of two forms of ionization:



- Needlepoint 2M ions/cu cm
- * "RF Ionizer Effect" produced by an electromagnetic pulse to excite particles inducing a charge and reducing particles.

2) Advanced Oxidation

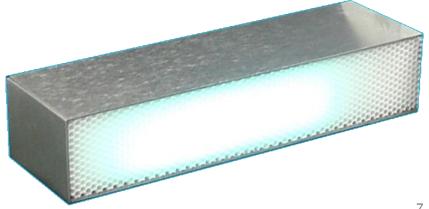
Advanced Oxidation—utilizing low levels of ozone, UV, and UV w/target plate technologies to break down VOCs and deactivate pathogens.

Introducing a unique technology...

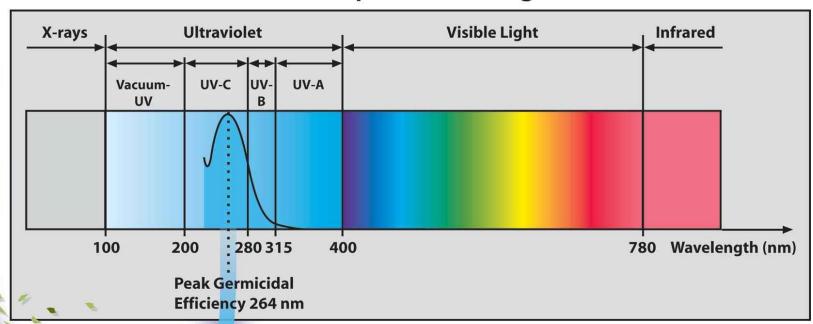
ActivePure™

A "Certified Space Technology"

Go to: www.spacefoundation.org



The Spectrum of Light

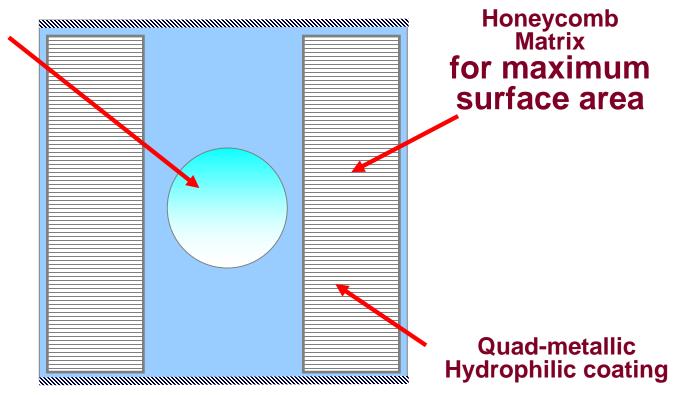


UV-C Lig

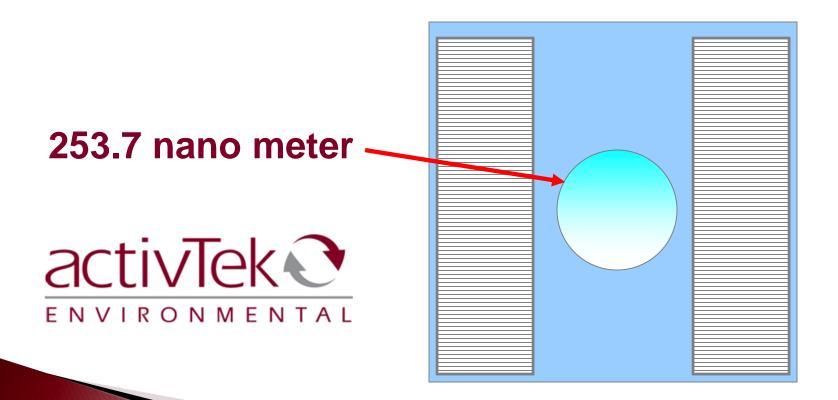
UV-C Radiation used for disinfection is most effective at a wavelength of 264 nm.

Radiant Catalytic Ionization (RCI) A unique and proprietary System

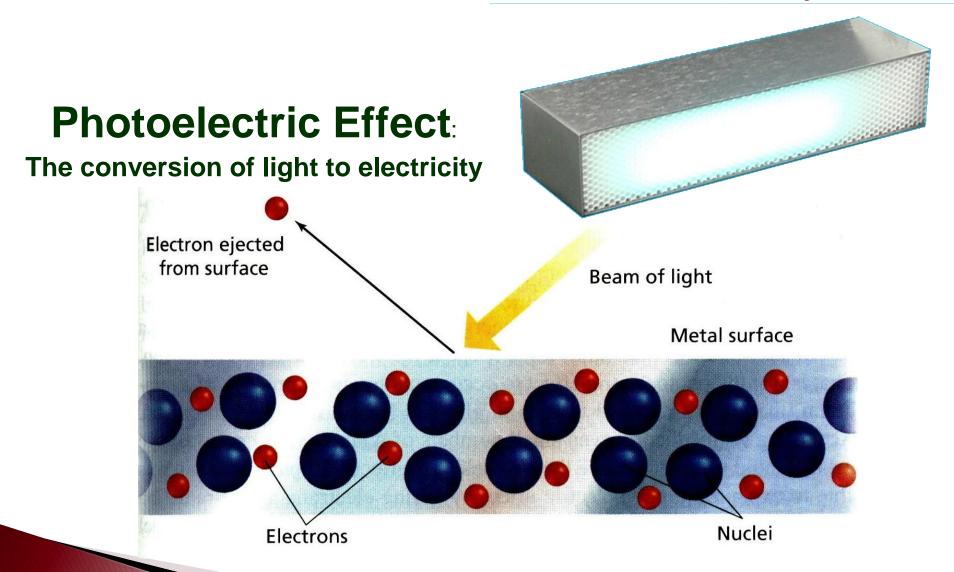
Broad spectrum UV lamp



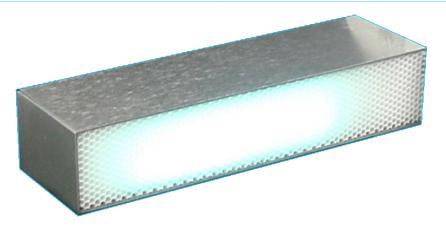
Germicidal UV (254 nm)



254nm UV Activates the Catalyst



The Catalyst Hydrophilic Coating

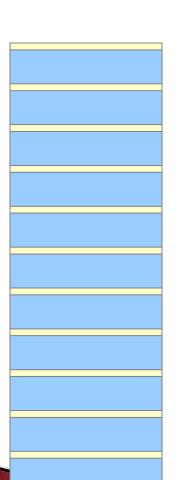


"Hydrophilic" – means the coating absorbs water vapor (H2O) from the air, creating an abundance of Hydrogen and Oxygen for creation of friendly oxidizers

Optimum Humidity is 40-60%

The Catalyst

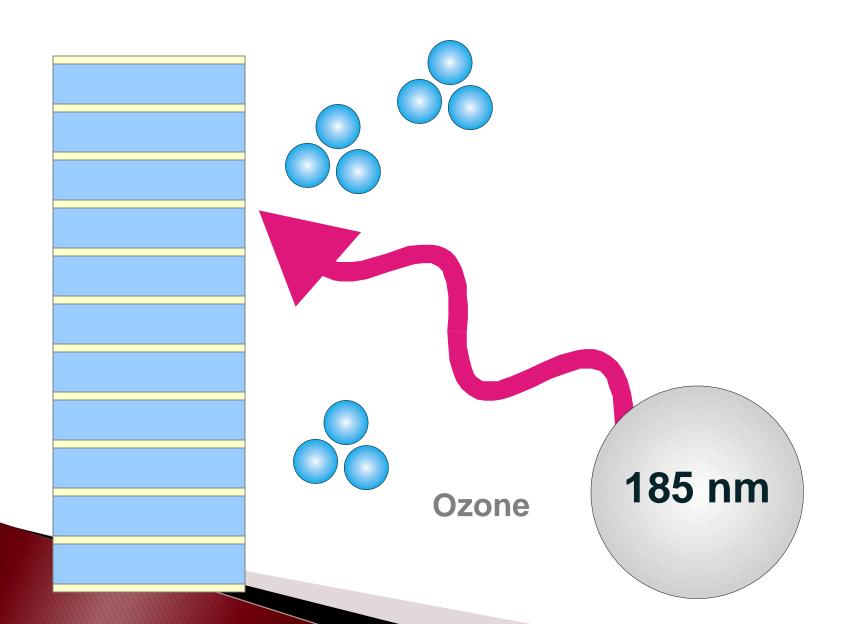
Quad-metallic Hydrophilic Catalytic Coating Titanium Dioxide, X, Y, Z



Each metal contributes to process:

- Titanium Dioxide forms two highly reactive substances Hydroxyl Radicals (OH) & Superoxide lons (O2-1) both very strong oxidizers
- X converts Nitric Oxide to Nitrogen and Oxygen
- Y speeds up the Titanium reactions by up to 3X. Proven to kill bacteria.
- Z has unique abilities to work with Hydrogen and can improve the functions of Titanium. Also, helps eliminates algae.

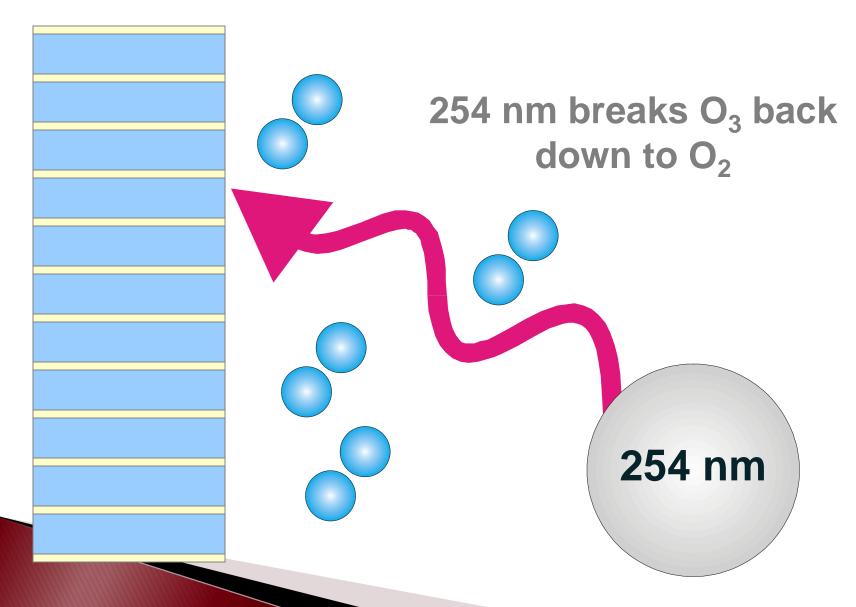
185 nm UV produces ozone



Oxidation Technology includes Low-Level Ozone

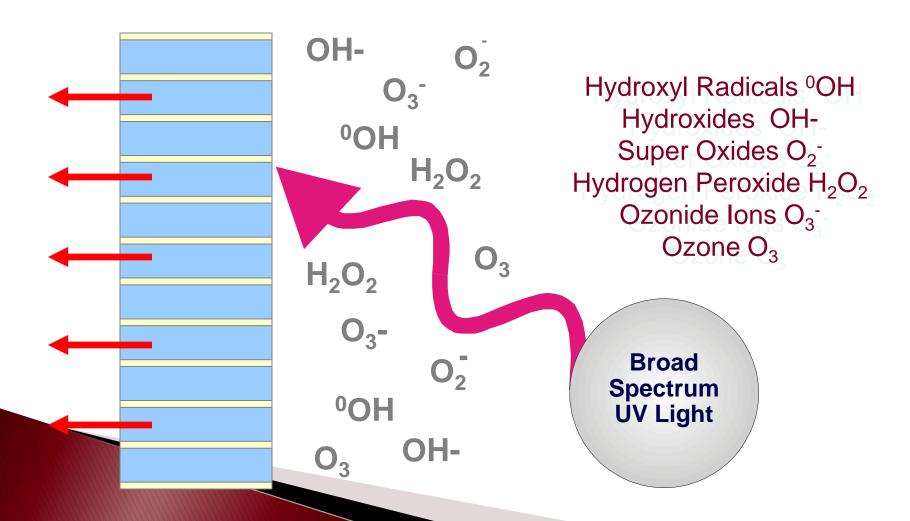
- Ozone reduces odors and VOC's including mold, bacteria & virus.
- Just like Oxygen & other naturally occurring elements Ozone is Dangerous at High Levels
- The last 40 years has shown that too low a level of ozone is also proving to be dangerous
- Ozone is proven to be effective even at low levels

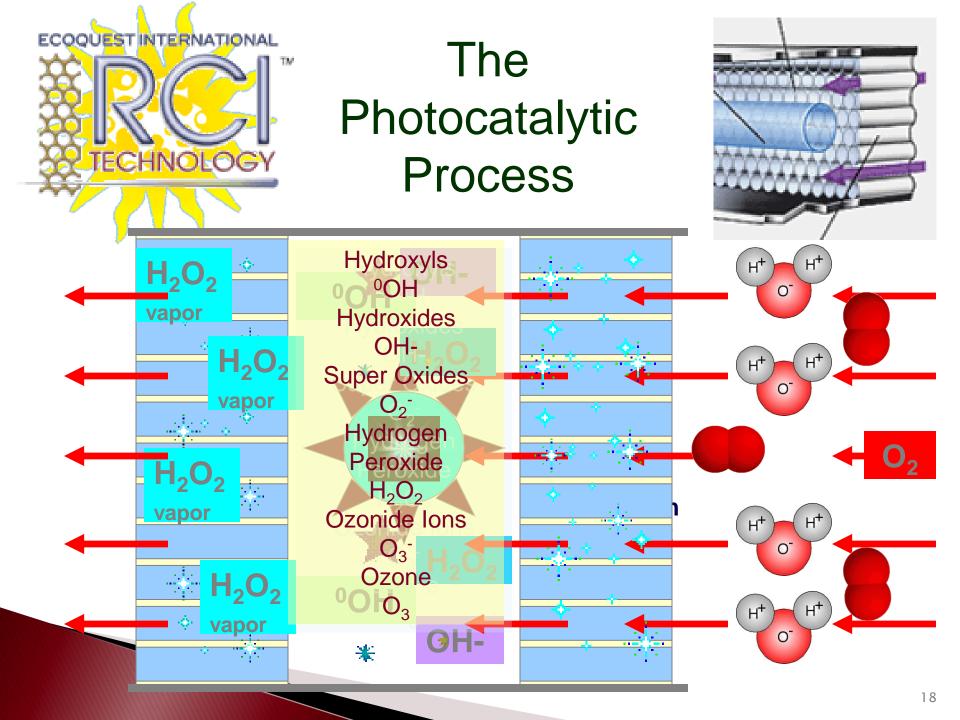
254 nm decomposition of ozone



AOP formation of "Friendly Oxidizers"

These oxidizers are then picked up in the air stream and dispersed throughout the building ...ACTIVELY purifying contaminants!





3) Filtration

- Filtration can be extremely effective in collecting particulates,
- Recommend MERV 7 (Minimum Efficiency Reporting Value) unless recommended by the manufacturer of the equipment,
- Harsh Environment pads protect the equipment from excessive accumulation of particulate,
- A combination of ACTIVE and PASSIVE technologies IS the synergistic answer!

How about some proof!



ActivePure Technology





University Testing on RCI Technology

Kansas State University:

Dr. James Marsden, Regents Distinguished Professor Microbial reduction on Surfaces (Mold, Bacteria & Virus)

University of Cincinnati:

Dr. Sergey Grinshpun, Department Head, Center for Health-Related Aerosol Studies, Department of Environmental Health

Reduction of the Aerosol Particle Concentration Airborne Microbial reduction (Virus)

University of Cincinnati Testing

Environ. Sci. Technol. 2007, 41, 606-612

Control of Aerosol Contaminants in Indoor Air: Combining the Particle Concentration Reduction with Microbial Inactivation

SERGEY A. GRINSHPUN,*
ATIN ADHIKARI, TAKESHI HONDA,†
KI YOUN KIM,‡ MIKA TOIVOLA,6
K. S. RAMCHANDER RAO, LAND
TIINA REPONEN

Center for Health-Related Aerosol Studies, Department of Environmental Health, University of Cincinnati, 3223 Eden Avenue, PO Box 670056, Cincinnati, Ohio 45267-0056

An indoor air purification technique, which combines unipolar ion emission and photocatalytic oxidation (promoted by a specially designed RCI cell), was investigated in two test chambers, 2.75 m³ and 24.3 m³, using nonbiological and biological challenge aerosols. The reduction in particle concentration was measured size selectively in real-time, and the Air Cleaning Factor and the Clean Air Delivery Rate (CADR) were determined. While testing with virions and bacteria, bioaerosol samples were collected and analyzed, and the microorganism survival rate was determined as a function of exposure time. We observed that the aerosol concentration decreased ~10

Another challenge is to decrease the indoor concentration of specific airborne contaminants, e.g., viable biological particles. While some indoor air purification techniques aim solely at the aerosol concentration reduction, others are designed to inactivate viable bioaerosols (e.g., viruses, bacteria, and fungi).

Some commercial air cleaners generate excessive ozone (either as a primary biocidal agent or as a bi-product); these devices have raised public health concerns (1). Among various guidelines for ozone exposures, the following thresholds have been specified for occupational environments: 0.2 ppm for 2 h (2), 0.05–0.10 for 8 h (2), 0.1 ppm for 8 h (3), and 0.05 ppm for instantaneous (no time limit specified) exposure (4). For comparison, the outdoor air standard is 0.08 ppm for 8 h (5). Ozone generators can inactivate viable microorganisms; however, the inactivation occurs at concentrations significantly exceeding health standards (6, 7).

Photooxidation involving UV radiation and TiO₂ as a photocatalyst has been applied for gas-phase detoxification of organic contaminants (8, 9) and for inactivating microorganisms in water (10–12). Some effort has been made to explore its application for air cleaning inside a closed-loop system (13, 14). The investigators reported significant photocatalytic inactivation of stress-resistant Serratia marcesens that occurred when aerosolized bacteria circulated in a closed-loop duct equipped with a TiO₂ filter for a relatively long period of time. Pal et al. (15) found similar effect for Escherichia coli, Microbacterium sp., and Bacillus subtilis; Keller et al. (16) reported considerable inactivation of airborne E. coli passing through a photoreactor coated with TiO₂ film. The biocidal effect of the photocatalytic oxidation can be attributed to photogenerated valence-band holes, hydroxyl

University of Cincinnati Testing Findings Overview

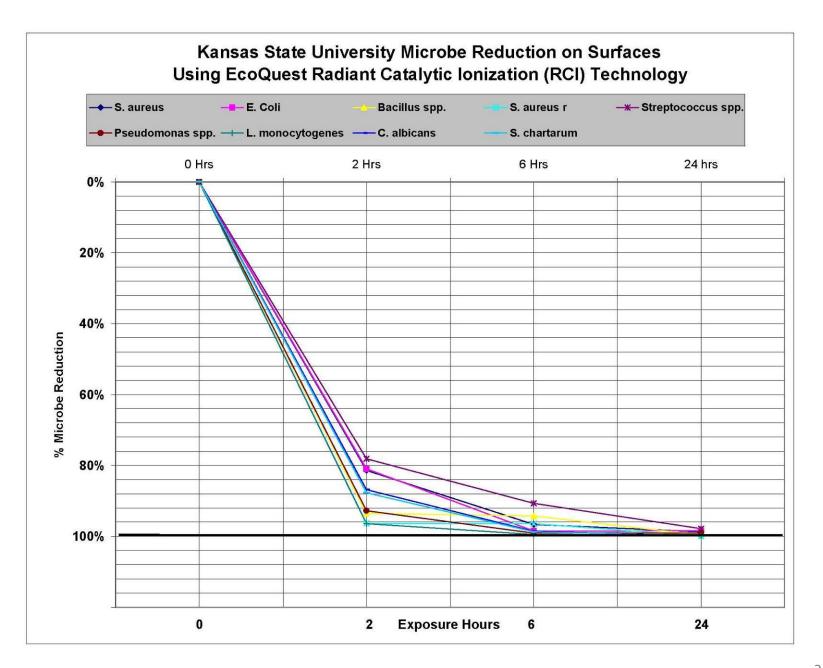
Testing included two technologies; Negative Ionization and Photo catalysis (an innovative proprietary Photo catalytic Reactor called Radiant Catalytic Ionization - RCI). Each technology was evaluated independently:

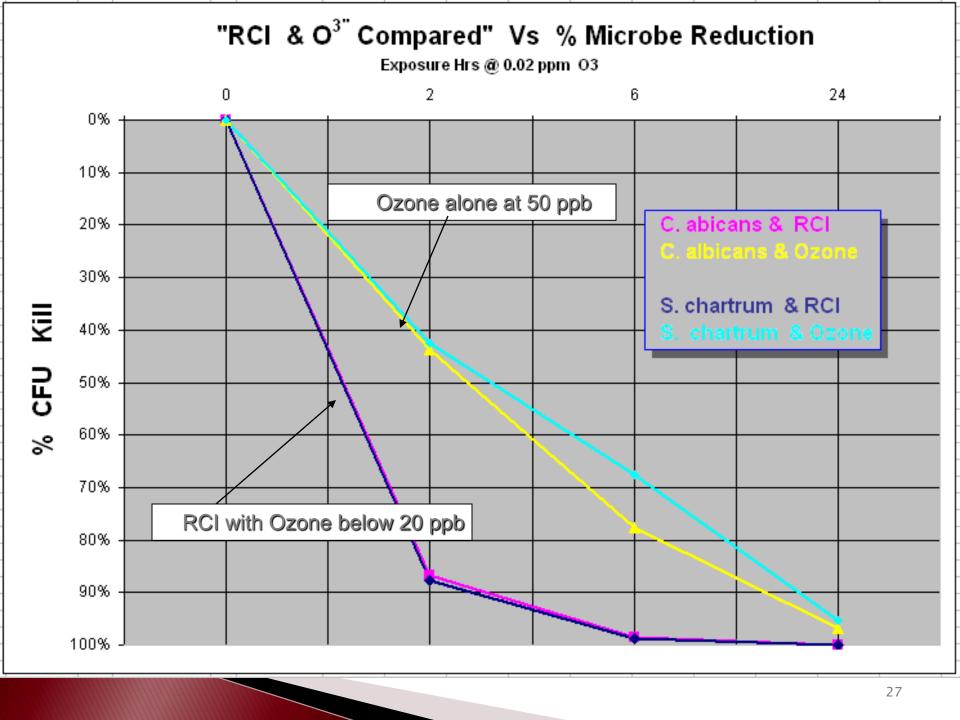
- Unipolar Ionization was able to reduce airborne particles from indoor air by <u>up to 250 times over natural decay</u>
- RCI was able to <u>inactivate approximately 90%</u> of airborne microorganisms in less than 60 minutes. The microorganisms tested were MS2 Virus and B. Subtilis (used as a surrogate for Anthrax)
- Dr. Grinshpun concluded the combination of the two technologies provided a more significant reduction of airborne bio contaminants than either of the two technologies working independently. This conclusion validates the synergistic effect of a multiple technology strategy.

Kansas State University Testing:

ability of Unique Photocatalytic Device to Reduce Microbial Populations on Surfaces in 24 hours

- Staph (Staphylococcus aureus): 98.5% reduction
- MRSA (Methicillin Resistant Staphylococcus aureus): 99.8% reduction
- E-Coli (Escherichia coli): 98.1% reduction
- Anthrax family (Bacillus spp.): 99.38% reduction
- Strep (Streptococcus spp.): 96.4% reduction
- Pseudomonas aureuginosa: 99.0% reduction
- Listeria monocytogenes: 99.75% reduction
- Candida albicans: 99.92% reduction
- Black Mold (Stachybotrys chartarum): 99.93% reduction





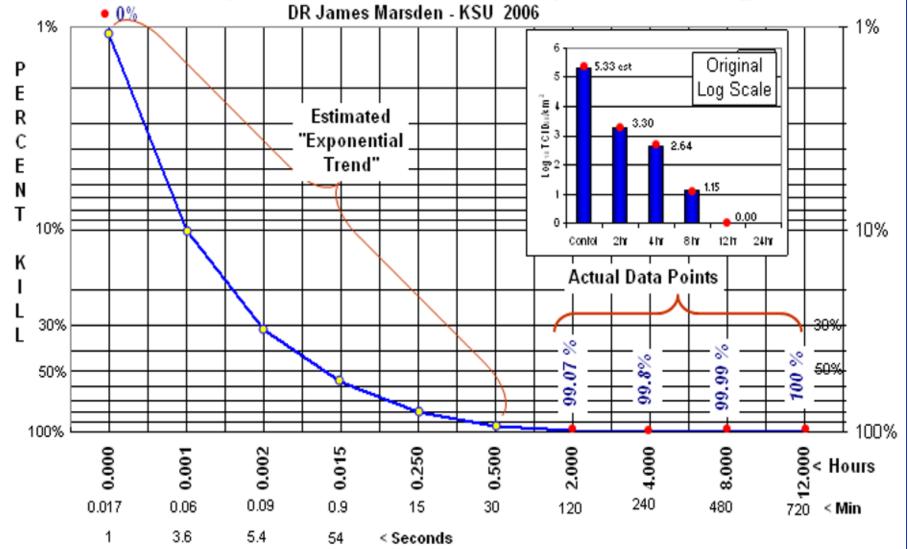
Avian Flu testing Dr. Jill Bieker Sandia National Laboratory

Spring 2006



KSU - Avian Influenza - H5N8 Summary

Stainless Steel Coupon - Percent Kill On Surfaces - After exposure to RCI technology



Field testing results





Ao Sr. Henrique Cury Atmosfera Pura Ltda / Ecoquest do Brasil

Prezado Sr. Henrique,

após a compra e utilização dos equipamentos purificadores de ar modelos EAGLE 5000 e FRESH AIR nos Biotérios de Suínos e Ratos respectivamente, Laboratório de Micro-cirurgia, Laboratório de Neurociências e Centro Cirúrgico de animais, gostaria de informar nossas avaliações.

- Quanto à eficiência no tratamento de odores e infecções.
- A utilização de dois equipamentos EAGLE 5000, no Biotério de suínos (pocilgas) com capacidade para 40 animais, foi suficiente para melhorar os odores em 90%.
 Isto se refletiu em melhores condições de trabalho para os técnicos; diminuição de custos no tratamento, uma vez que não foi mais necessário a utilização de máscaras; melhores condições de vida para os animais.
- 2. A utilização de 1 equipamento FRESH AIR no Biotério dos ratos foi suficiente para que reduzissemos as trocas de maravalha de 2 para 1 vez por semana, isto é 50%. Acreditamos que estas trocas possam vir a ser mais espaçadas. Também observamos que não houve infecções em nenhum animal até o momento (ainda não temos número estatisticamente significante).

Os níveis de amônia dentro do Biotério, que não tem um sistema de ar adequado, foram reduzidos em 70%.

 A utilização de 1 equipamento FRESH AIR no laboratório de micro-cirurgia durante e após o Curso de Microanatomia do Osso temporal, onde utilizamos brocas para perfurar ossos, o odor foi reduzido em 90%.

This Brazilian Hospital Laboratory was conducting tests with lab rats. The rats were operated on to insert probes in their brains. Prior to installation of Stand-alone RCI Air Purification System the laboratory was loosing 10% of the rats after each surgery. Since installation of the purifier, no rats have been lost.

Test Data from a home in Central Florida in USA before and after 212mm (9") in-duct system installation

Micro-5(™) Cassette Analysis of Fungal Spores & Other Airborne Particulates by Optical Microscopy (EMSL Method M001)

Client Sample Number Client Sample to 388233 25										
Volume Sample Location Spore Types Raw Count Spores/m² % of Total Associated Call Call Call Call Spores/m² % of Total Call	Lab Sample Number	34	40601072-00	01						
Sample Location	Client Sample ID		386233							
Spore Types										
AgrocyberCoprinus										
Alternaria		Raw Count	Spores/m³	% of Total	Raw Count	Spores/m³	% of Total	Raw Count	Spores/m³	% of Total
Arthrinium Arthrospores	Agrocybe/Coprinus	-	-	-						
Arthrospores		-	-	-						
Ascospores Aspergillusi/Penicillium Basidiospores Bipolaris Chaetomium Cladosporium Cladosporium Cladosporium Cladosporium Cladosporium Ganoderma Fusarium F	Arthrinium	-	-	-						
Aspergillus/Pericillum 333 13300 40.3 Basisiospores 12 480 1.5 Bigolaris - - - Chaedomium - - -	Arthrospores	>300	>12000	36.3						
Basidiospores 12 480 1.5 Bipolaris - - -	Ascospores	1	40	0.1						
Chaetomium	Aspergillus/Penicillium	333	13300	40.3		Б	C			
Chaetomium	Basidiospores	12	480	1.5		$ B\epsilon$	etore	e trea	arme	nt
Curvularia - - - -	Bipolaris	-	-	-			(
Curvularia - - - -	Chaetomium	-	-	-		to	ct d	ata o	بيادي	Stod
Epicocoum	Cladosporium	34	1360	4.1		נכ	St u	ata E	value	attu
Fusanum	Curvularia	-	-	-		B 4		20	2006	_
Fusanum	Epicoccum	-	-	-		IMI	arch	78.	ノ()() 6)
Memnoniela	Fusarium	-	-	-			ai ci i	20/	2000	,
Myxomycete	Ganoderma	-	-	-						
Paecilomyces	Memnoniella	4	160	0.5		N /		D	٠. ـ ـ ـ	
Paecilomyces	Myxomycete	4	160	0.5		_ ^	aste	r Bea	iroor	n
Rust		117	4680	14.2						
Rust	Pithomyces/Ulocladium	_	-	-						
Scopulariopsis 17		_	-	-						
Stachylotrys		17	680	2.1						
Torula Unidentifiable Spores 5 200 0.6 Zygomycetes Total Fungi >827 >33100 100 Fibrous Particulate 20 800 Hyphal Fragment 78 3120 Insect Fragment 10 400 Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2		_	-	-						
Unidentifiable Spores 5 200 0.6 Zygomycetes Total Fungi >827 >33100 100 Fibrous Particulate 20 800 Hyphal Fragment 78 3120 Insect Fragment 10 400 Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2		_	-							
Zygomycetes	Unidentifiable Spores	5	200	0.6						
Total Fungi >827 >33100 100 Fibrous Particulate 20 800 Hyphal Fragment 78 3120 Insect Fragment 10 400 Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2	· ·	_	-	-						
Fibrous Particulate 20 800 Hyphal Fragment 78 3120 Insect Fragment 10 400 Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2		>827	>33100	100						
Insect Fragment 10 400 Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2			800							
Pollen 2 80 Analytical Sensitivity 40 Skin Fragments (1-4) 2	Hyphal Fragment	78	3120							
Analytical Sensitivity 40 Skin Fragments (1-4) 2	Insect Fragment	10	400							
Skin Fragments (1-4) 2	Pollen	2	80							
	Analytical Sensitivity		40							
Background (1-5) 2	Skin Fragments (1-4)		2							
	Background (1-5)		2							

Micro-5(™) Cassette Analysis of Fungal Spores & Other Airborne Particulates by Optical Microscopy (EMSL Method M001)

Lab Sample Number	_	40000044 004	.,							
Client Sample ID	3	40602014-000 386105	J1							
Volume		25								
Sample Location	١,	20 naster bedroor	m							
Spore Types	Raw Count	Spores/m³	% of Total	Raw Cour	nt Spores/r	m³ % o	of Total	Raw Count	Spores/m³	% of Total
Agrocybe/Coprinus	-	-	-							
Alternaria	_	-	-							
Arthrinium	-	-	-							
Arthrospores	-	-	-							
Ascospores	2	80	25							
Aspergillus/Penicillium	2	80	25		_	_ .	_	_		
Basidiospores	-	-	-		— Д	tter	1 tr	eatm	nent	test
Bipolaris	1	40	12.5		, ,		Ci	Cati	Citc	
Chaetomium	-	-	-		٦,	a ta	\bigcirc	/alua	tod	
Cladosporium	1	40	12.5		u	ala	CV	/aiua	teu	
Curvularia	-	-	-		-			200		
Epicoccum	-	-	-		- 11	ine		, 200	6	
Fusarium	-	-	-		50			, 200		
Ganoderma	-	-	-							
Myxomycete	-	-	-		N /			D - 4.		_
Nigrospora	2	80	25		— V	ast	er	Bedi	roon	1
Paecilomyces	-	-	-							· .
Pithomyces/Ulocladium	-	-	-							
Rust	-	-	-							
Scopulariopsis	-	-	-							
Stachybotrys	-	-	-							
Torula	-	-	-							
Unidentifiable Spores	-	-	-							
Zygomycetes	-	-	-							
Total Fungi	8	320	101							
Fibrous Particulate	4	160								
Hyphal Fragment	2	80								
Insect Fragment	2	80								
Pollen	1	40								
Analytical Sensitivity		40								
Skin Fragments (1-4)		2								
Background (1-5)		1								

Summary of Results

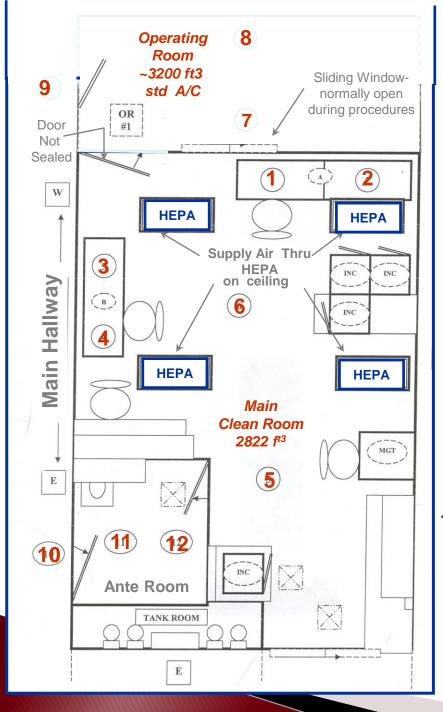
- Aspergillus/Penicillium
- Cladosporium
- ArthrosporesTotal Fungi
- Fibrous Particulate
- Hyphal Fragment
- Insect Fragment
- Pollen
 Total Particulate

Spores/cu meter

<u>Before</u>	After
13,300	80
1,360	80
>12,000	0
>33,100	320

800	160
3,120	80
400	80
80	40
4.420	360

Test Data from Invitro Fertilization Lab in a Large Midwestern Hospital in USA



Hospital - Clean Room

- Particulate count/m3 measured @ 0.5 um
 - Typical office 2 -3 million/m3 @ 0.5 um
- Before and After 2-9" PCO Units in A/C ducts

Loc	Class	Before/m3	After/m3	<u>lmp %</u>
01	5	1391.4	77.7	94%
02	5	692.2	21.2	97%
03	5	621.5	7.1	99%
04	5	374.3	7.1	98%
05	6	11,569.1	2,366.1	80%
06	6	22,236.5	332.0	99%
07	7	337,161.9	165,484.6	51%
80	7	249,121.4	153,887.2	38%
09	8	1,614,245.0	334,401.6	79%
10	8	1,304,842.0	373,078.3	71%
11	7	247,614.7	53,614.7	78%
12	7	203,518.4	50,302.2	75%

All particle counts were taken in accordance with ISO 14644-1 standards

Microbial Colonies - Before & After RCI

<u>Location</u>	Before	<u>After</u>
1-2 Horizontal Flow Clean Bench	0	0
3- 4 Horizontal Flow Clean Bench	0	0
5-6 Main Clean Room	14	0
7-8 Operating Room	9	0
9- 10 Hallway	27	8
11-12 Ante Room	23	15 ₃₇

3/

Three points to remember about activTek Environment Conditioning Technology......

- Nature based technologies: We clean air indoors like nature cleans the air outdoors using:
 - Two types of Ionization to reduce particles
 - UV and a Catalyst to create Oxygen & Hydrogen based friendly oxidizers to reduce microbial contaminants (Photo Catalytic Oxidation)
 - Low-level ozone to reduce odors
- 2. The <u>only</u> residential Air Purification Technology recognized by the Space Foundation and awarded Certified Space Technology status
- 3. The only Air and Surface Purification with validation testing from multiple universities proving the ability to reduce up to 99.99% of Mold, Fungi, Bacteria and Viruses, including MRSA and Avian Bird Flu on surfaces.