Testing Results Booklet



Allergen study In Vitro Lab

Study conducted by Indoor Biotechnologies



Study objective

To prove that betterair's Enviro-Biotics® degrade allergens, by using them as a nutrient, samples were taken for bacterial concentration (using a classic microbiology method) and allergen concentration (using the INDOOR MARIA® assay, MARIA SOP-001 ver.1.3).





Figure 1: Enviro-Biotics[®] concentration over time after incubation with Dust mite (Der p2) Allergen

Figure 2: Dust mite (Der p2) Allergen concentration over time after incubation with Enviro-Biotics®





Figure 3: Enviro-Biotics[®] concentration over time after incubation with Dog (Can f1) Allergen

Figure 4: Dog (Can f1) Allergen concentration over time after incubation with Enviro-Biotics®

Conclusion

Our Enviro-Biotics® can grow and prosper by using allergens as a nutrient source, hence dramatically decrease the concentration of allergens and reduce the side effects of ubiquitous indoor allergens on the residents.

University of Oregon study

The BioBE (Biology and Built Environment) center at the University of Oregon, conducted fully controlled trials on the Better Air Enviro-Biotics® technology inside a controlled chamber simulating real life conditions. The U of O performed the testing that generated the data presented below.



In these trials, BioBE has focused on the microbial viability of Enviro-Biotic™ treatment on surfaces as well as the antagonistic effect against indoor pathogenic microbes.

Oregon 2018

Plates were preloaded with either live or sterilized human-associated microbial communities (HAMC), which was obtained from homogenized and filtered vacuum cleaner dust sourced from three distinct occupied Buildings.

The course of treatment was two weeks using betterair's Biotica 800 device with standard settings in the ESBL Climate Chamber (25.75 m3) with humidity, air temperature, and air exchange rates held constant throughout the duration of the experiment and within the range typical of residential/commercial buildings. Air exchange was held at the minimum acceptable via ASHRAE standards for occupied office buildings. All incoming air was filtered to MERV 15.

Proven efficacy against pathogens

The probiotic treatment exhibited a clear antagonistic effect against three typical indoor microbial pathogens:

Escherichia coli – a pathogenic bacteria causing a wide range of human related infections.

Staphylococcus epidermidis - a Gram-positive bacterium that infects patients with compromised immune systems, people with catheters and is a frequent contaminant of specimens sent to the diagnostic laboratory.

University of Oregon study continued



Fig. 1: Counts of live E. coli and S. epidermidis over time. Enviro-Biotics® treated plates received treatment from day 1 to day 14. The treatment period is defined by the dashed vertical lines.



Fig. 2: Counts of live Cladosporium sp. over time. Enviro-Biotics® treated plates received treatment from day 1 to day 14. The treatment period is defined by the dashed vertical lines. Oregon 2018 Cladosporium sp. - an indoor pathogenic mold. Its airborne spores are significant allergens that severely affect asthmatics.

The inhibition effect over E. Coli and S. epidermidis is depicted in Fig. 1. Continuous application for 14 days suppressed the pathogenic bacteria counts in comparison to the control.

Therefore, it can be concluded that from the 8th day onwards, a significant antagonistic effect was demonstrated over the Cladosporium sp. pathogenic fungi.

Following the 14 days treatment regime, the E. coli and S. epidermidis counts in the untreated-control increased dramatically, while their counts in the treated samples were kept low due to the probiotic inhibition effect.

The antagonistic effect against Cladosporium sp. was even more impressive as can be seen in Fig. 2: Cladosporium sp. counts increased significantly more over time in control versus treated samples (Fig. 2; χ 2 = 12.3, P < 0.001).

University of Oregon study continued



Conclusion

In conclusion the Enviro-Biotics® treatment exhibited a clear antagonistic effect against three typical indoor microbial pathogens: Escherichia coli, Staphylococcus epidermidis and Cladosporium sp. It is noted that these species typical indoor microbial pathogens with a potential to be pathogenic and harmful to people, able to cause a wide range of human related infections or indoor acquired illnesses.

Oregon 2018

Field household study to evaluate Enviro-Biotics[™] on atopic dogs



Australia 2018

The objective was to evaluate owner's perception of their allergic dog's pruritus (atopic skin condition) and malodour following frequent use of Enviro-Biotics[™] over a 21-day period. This study will also evaluate the population pattern of S. pseudintermedius and Enviro-Biotics[™] in a subgroup of the study animals.

Dogs with atopic dermatitis have been shown to have less species of bacteria on their skin compared to healthy dogs, therefore contributing to a compromised skin barrier leading to increased itch and skin odour. Atopic dogs are also at a significantly increased risk of developing skin infections (pyoderma). This pilot study was aimed to evaluate the effects of regular topical application of Enviro-Biotics[™], on dogs with atopic dermatitis and their sleeping area.

Furthermore, the assessment of specific bacteriological cultures, pruritus (itch) scores and owner's perception of odour, on skin and bedding before and during the use of the product over a 21 day period was evaluated. Pruritis assessment:

For the pruritis variable we found a reduction in value of -0.6% per day. The reduction was found to be statistically significant (F1,11.01=7.05, p=0.022, Figure 1a).

Odour assessment:



Fig.1: a. Mean pruritis assessment values with error bars representing the standard deviation across four repetitions. b. Mean odour assessment values with error bars representing the standard deviation across four repetitions.

Field household study to evaluate Enviro-Biotics™ on atopic dogs

continued



Australia 2018

Bacterial culture assessment:

The number of bacteriological cultures positive for S. pseudintermedius declined for these dogs over Day 7-21, while the number of cultures positive for Enviro-Biotics™ remained constant over this period





Conclusion

Overall, results are very encouraging considering the pilot data set and the relatively short assessment time frame for a chronic condition.

We can see a statistically significant decrease in Pruritis scoring, together with a clear decrease in odour, furthermore, we can see a decrease in S. pseudintermedius (primarily a pathogen for domestic animals, but has been known to affect humans as well) while Enviro-Biotics™ remained constant.

This study was conducted by Invetus, the largest Australasian veterinary contract research organisation (CRO) with sites throughout Australia and New Zealand and managed by Orivet genetic pet care, Australia.

Israeli Police Forces, Israel



Improved productivity associated with the reduction of airborne irritants.

Reduction of headaches and noticeable improvement in air and surface quality, additional symptoms associated with SBS.

1 - Before treatment

HMC ID Number: 1500527	3 - 2 Samp	ole Media: Sv	wab	Volume: 1.	00 Limit of detection: 100 cfu / cm2
Sample ID Number: 2	Sam	ple Name: Zi	on 1		
Organism	Туре	Raw Count	cfu/cm2	% of Total	Note
Alternaria	Fungi	9	900	4%	
Aspergillus	Fungi	11	1100	4%	
Rhodotorula	Fungi	14	1400	6%	
Bacillus	Bacteria	28	2800	11%	
Corynebacterium	Bacteria	13	1300	5%	
Micrococcus	Bacteria	6	600	2%	
Pseudomonas sp.	Bacteria	150	15000	60%	

2 - After 12 weeks of treatment

HMC ID Number: 1600481	3 - 3 Samj	ole Media: Sv	wab	Volume: 1	.00
Sample ID Number: 3	Sam	ple Name: Sv	wab		
Organism	Туре	Raw Count	cfu / cm2	% of Total	
No Fungi Detected	Fungi				
Bacillus	Bacteria	40	4000	100%	



2012 Hayes Microbial, Testing US

Caspi Sror Law Offices, Tel Aviv, Israel



Outbreak of rashes and itchy eyes associated with poor indoor air **and surface** quality .

Reduction in high levels of 3 dangerous strains, of allergy causing bacteria, and specific reduction of Staph, a pathogenic bacteria to non detectable levels. Return to a balanced indoor air and surface microbiome. Reduced headaches.



Altman Offices, Israel



Presence of strains of fungi and bacteria that are associated with health

Levels of disease causing bacteria and fungi returned to balanced levels. High levels of betterair Enviro-biotics™ replaced the allergy causing microbes.

Before treatment

HMC ID Number: 15008044 - 2 Sar		ple Media: Sv	wab	Volume: 1	.00			
Sample ID Number: 2	Sam	Sample Name: Surface / Table						
Organism	Туре	Raw Count	cfu / cm2	% of Total				
Rhodotorula	Fungi	15	1500	12%				
Unspecified Mold	Fungi	2	200	2%				
Yeast	Fungi	33	3300	26%				
Bacillus	Bacteria	11	1100	9%				
Staphylococcus sp.	Bacteria	68	6800	53%				

After treatment

HMC ID Number: 150091	36 - 3	Samp	ole Media: Sv	vab	Volume: 1.00	
Sample ID Number: 3		Sam	rface / Table			
Organism	Ту	ре	Raw Count	cfu / cm2	% of Total	
No Fungi Detected	Fu	ngi				
Bacillus	Bac	teria	61	6100	100%	



El AI - Boeing 777, International Flights



Unhygienic in flight conditions on surfaces, in the air and in toilets, with associated high levels of infection causing microbes present.

Levels of disease causing bacteria and fungi returned to acceptable levels. High levels of betterair Enviro-biotics[™] replaced the allergy causing microbes.

1 - Before treatment

HMC ID Number: 17007210 - 1			Samp	ole Media: S	wab	Volume:	1.00
Sample ID Number:	1		Sam	ple Name: La	avatory		
Organism		Ту	ре	Raw Count	cfu / cm2	% of Total	
Yeast		Fu	ngi	4	400	2%	
Gram Negative Fermenter		Bac	teria	145	14500	78%	
Gram Negative non-Fermen	ter	Bac	teria	30	3000	16%	
Staphylococcus sp.		Bac	teria	8	800	4%	
	1700701		Com	la Madia: C		Valumas	1.00
HMC ID Number:	1/00/21	0-2	Samp	ble Media: 5		volume:	1.00
Sample ID Number:	2	.	Sam	Die Name: C	omp Fuselage	9/ of Total	_
Organism		I I Y	pe	Raw Count	ciu / cm2	% of Total	+
Aureobasidium		Fungi		3	300	2%	—
Rhodotorula		Fu	ngi	14	1400 7%		_
Gram Negative Fermenter		Bac	teria	50	5000	27%	_
Gram Negative non-Fermen	ter	Bac	teria	120	12000	64%	_
HMC ID Number:	1700721	0 - 3	Samp	ole Media: A	ir	Volume:	300.00
Sample ID Number:	3		Sam	ple Name: C	omp Fuselage B	ac	
Organism		Ту	pe	Raw Count	cfu / M3	% of Total	
Bacillus		Bac	teria	10	33	26%	
Micrococcus		Bac	teria	12	40	32%	
Staphylococcus sp.		Bac	teria	16	53	42%	
HMC ID Number:	1700721	0 - 4	Samp	ole Media: A	ir	Volume:	300.00
Sample ID Number:	4		Sam	ple Name: C	omp Fuselage M	old	
Organism		Ту	pe	Raw Count	cfu / M3	% of Total	
Cladosporium		Fu	ngi	28	93	100%	

2 - After treatment

HMC ID Number:	1701091	1 - 1	Samp	ole Media:	Swab	Volume:	1.00
Sample ID Number:	1		Sam	ole Name:	Lavatory		
Organism		Ту	pe	Raw Coun	t cfu / in2	% of Total	
Bacillus		Bac	teria	240	24000	100%	
No Fungi Detected							
HMC ID Number:	1701091	1 - 2	Samp	ble Media:	Swab	Volume:	1.00
Sample ID Number:	2		Sam	ole Name:	Composite Fusela	ge Swab	
Organism		Ту	ре	Raw Coun	it cfu / in2	% of Total	
Bacillus		Bac	teria	255	25500	100%	
No Fungi Detected							
HMC ID Number:	1701091	1 - 3	Samr	le Media:	Δir	Volume	500.00
Sample ID Number:	3		Sam	ole Name:	Composite Fusela	ge Air Ba	
Organism	•	Tv	ne	Baw Coun	t cfu / M3	% of Total	
Bacillus		Bac	teria	190	380	100%	
Daomas		Duo	tona	100	000	10070	_
							<u> </u>
HMC ID Number:	1701091	1 - 4	Samp	ole Media:	Air	Volume:	500.00
Sample ID Number:	4		Sam	ole Name:	Composite Fusela	ge Air Mold	
Organism		Ту	pe	Raw Coun	it cfu / M3	% of Total	
Cladosporium		Fu	ngi	4	8	100%	

Beit Gadi Medical Center



High levels of bacteria and fungi found in recovery rooms and patient rooms - leading to high probability of hospital acquired infections.

Post treatment result showed non detectable level of associated microbes.

1 - Before treatment

Job Number: 241114 Collected by: Yuli Horesh Email: yuli@betterair.co.il	f la	J	ob Name: Beit	Gady Medical Ce	nter Date Collected: 11/24/2014 Date Received: 12/03/2014 Date Reported: 12/08/2014			
HMC ID Number: 1402066	2 - 1 Sar	nple Media: Air			Limit of detection: 10 cfu / M3			
Sample ID Number: 1	Sar	nple Name: Rm	302-BAC					
Organism	Type	Raw Count	cfu / M3	% of Total	Note			
Staphylococcus sp	Bacteria	1	10	5%				
Pseudomonas sp	Bacteria	20	200	96%				
					Pseudomonas in the Air			
HMC ID Number: 1402066	2 • 2 Sar	nple Media: Air			Limit of detection: 10 cfu / M3			
Sample ID Number: 2	Nar	me: Rm 302 Mol	d					
Organism	Type	Raw Count	ant cfu / plate % of Total		Note			
Cladosporium	Fungi	17	6	46%				
Rhizopus .	Fungi	7	7	54%				
HMC ID Number: 1402066	2 - 3 Sar	nple Media: Swa	ab Sample		Limit of detection: 100 cfu / cm2			
Sample ID Number: 3	Nar	ne: general sur	ace					
Organism	Type	Raw Count	cfu / cm2	% of Total	Note			
Cladosporium	Fungi	1	100	1%				
Yeast	Fungi	9	900	5%				
Pseudomonas sp.	Bacteria	168	16800	94%				
					Pseudomonas on the surface			

2 - After treatment

HMC ID Number: 150	001393 - 1	San	mple Media: Air		Volume: 100.00			Limit of detection: 10 cfu / M3	
Sample ID Number: 1		Sar	mple Name: Rm 3	02-BAC					
Organism	Тур	•	Raw Count	cfu / M3	% of Tota	Total Note			
Bacillus	Bacte	ria	6	60	100%				
<									
					0	nly	Better	Air Probiotics found	
			-					in the Air	
HMC ID Number: 15001393 - 2		Sar	mple Media: Air		Volum	e: 10	0.00	Limit of detection: 10 ofu / Ma	_
Sample ID Number: 2		San	mple Name: Rm 3	02-Mold					
Organism	Type		Raw Count	cfu / M3	% of Total		Note		
No Fungi Detected	Fun	gi							
<u>, </u>									
						1			
HMC ID Number: 150	01393 - 3	Sa	mple Media: Surf	ace Swab	Volum	ne: 10	0.00	Limit of detection: 10 cfu / cm2	
Sample ID Number: 3		Sa	mple Name: Gen	eral surface					
Organism	Typ	æ	Raw Count	cfu / M3	% of Tot	al		Note	
No Fungi Detected	Fungi								
No Bacteria Detected	Bad	teria							
								Construction of a second second	
							NO IN	rection found on surface	

Case Studies in US

Care Partners Assisted Living Facility 4 months study



Evaluate effect on a highly utilized living area. With dispersion of betterair probiotics through the HVAC system.



Figure 1. Vent interior Gram positive bacteria. Samples were taken from vent duct interior surfaces approximately 50cm distance from vent opening. Approximately 100cm 2 surface was swabbed and diluted in 1 ml saline with 0.1 ml subsamples plated on MSA medium. A ten fold dilution factor has been applied to CFU values to account for the dilution.



Figure2.Solid Surface Gram positive bacteria. Samples were taken on window stools (lowermost interior horizontal surface) directly above vents and on the upper surface of the door to the room. Approximately 100 cm2 surface was swabbed and diluted in 1 ml saline with 0.1 ml subsamples plated on MSA. A ten fold dilution factor has been applied to CFU values to account for the dilution.

Case Studies in US continued

Care Partners Assisted Living Facility 4 months study



In summary, this data shows that levels of Gram negative bacteria, including fecal indicator bacteria representing potential pathogens, were greatly reduced in numbers on the inner surface of the vent ducts in both the treated and untreated rooms following the application of the betterair Enviro-bioticsTM.



Replacement of gram negative bacteria was replaced with betterair gram positive bacteria.



Gram negative bacteria associated with fecal matter decreased on the vent surface eventually decreasing to levels of non detection.

SAFETY AND EFFICACY

In Vitro Lab, South Korea



Test efficacy of betterair individual strains of bacteria, against 6 forms of pathogenic bacteria and 3 strains of toxin producing mold

2017 HEM

1 - Inhalation infection test





2 - Lecithinase and hemolysis activity



Safety Studies



Safety test of our **product** in accordance with EPA protocols as part of a certification processs.



Effect on inhalation - Lung Challenge test



Antibiotics susceptibility test to determine resistance to antibiotics.



Passed the inhalation test and no residual presence of **betterair** bacteria was found in the lungs after testing. DNA sequencing used.



No breakdown of red blood cells - Lack of Hemolysis.



Do not produce Lecithinase enzyme which breaks down lecithin, an essential component of body cell membranes.

Summary

In this booklet, we have displayed study cases showing the efficacy of our products together with various conclusive safety tests completed at external laboratories.

We exhibited the ability of our strains in inhibition of various bacteria and mold ubiquitous to indoor environments, both in laboratory experiments and also in "real life" indoor environments, together with various allergens reduction.

We have also exhibited our strains safety by lack of Hemolysis and Lecithinase activity together with antibiotic susceptibility. in addition, our strains passed a stringent lung challenge test.



Team

