

# LOCAD-PTS: Operation of a new system for microbial monitoring aboard the International Space Station (ISS)

J. Maule<sup>1</sup> and N. Wainwright<sup>2</sup>

*British Aerospace (BAE) Systems, Huntsville, AL, 35806; and Charles River Laboratories, Charleston, SC 29407*

A. Steele<sup>3</sup>

*Geophysical Laboratory, Carnegie Institution for Science, Washington, DC, 20015*

D. Gunter, G. Flores and M. Effinger<sup>4</sup>

*NASA MSFC, Huntsville, AL 35812*

M. Damon and M. Wells<sup>5</sup>

*British Aerospace (BAE) Systems, Huntsville, AL, 35806; and University of Alabama at Huntsville (UAH), Huntsville AL 35806*

S. Williams<sup>6</sup>

*Astronaut Office, NASA Johnson Space Center, Houston, TX, 77058*

H. Morris and L. Monaco<sup>7</sup>

*Jacobs Technology Inc., Huntsville, AL 35806*

Microorganisms within the space stations *Salyut*, *Mir* and the International Space Station (ISS), have traditionally been monitored with culture-based techniques. These techniques involve growing environmental samples (cabin water, air or surfaces) on agar-type media for several days, followed by visualization of resulting colonies; and return of samples to Earth for ground-based analysis. This approach has provided a wealth of useful data and enhanced our understanding of the microbial ecology within space stations. However, the approach is also limited by the following: i) More than 95% microorganisms in the environment cannot grow on conventional growth media; ii) Significant time lags occur between onboard sampling and colony visualization (3-5 days) and ground-based analysis (as long as several months); iii) Colonies are often difficult to visualize due to condensation within contact slide media plates; and iv) Techniques involve growth of potentially harmful microorganisms, which must then be disposed of safely. This report describes the operation of a new culture-independent technique onboard the ISS for rapid analysis (within minutes) of endotoxin and  $\beta$ -1, 3-glucan, found in the cell walls of gram-negative bacteria and fungi, respectively. This technique involves analysis of environmental samples with the *Limulus* Amebocyte Lysate (LAL) assay in a handheld device. This handheld device and sampling system is known as the Lab-On-a-Chip Application Development Portable Test System (LOCAD-PTS). A poster will be presented that describes a comparative study between LOCAD-PTS analysis and existing culture-based methods onboard the ISS; together with an exploratory survey of surface endotoxin throughout the ISS. It is concluded that while a general correlation between LOCAD-PTS and traditional culture-based methods should not necessarily be expected, a combinatorial approach can be adopted where both sets of data are used *together* to generate a more complete story of the microbial ecology on the ISS.

<sup>1</sup> Project Scientist, LOCAD-PTS, British Aerospace (BAE) Systems, Huntsville, AL, 35806.

<sup>2</sup> Principal Investigator, LOCAD-PTS, Charles River Labs, Wappoo Road, Charleston, SC, 29407.

<sup>3</sup> Staff Scientist, Geophysical Laboratory, Carnegie Institution for Science, Washington DC, 20015.

<sup>4</sup> Lead Flight Operations Engineer and Project Managers (past and present), LOCAD Group, MSFC, Huntsville, AL, 35812.

<sup>5</sup> Scientist, BAE Systems, Huntsville, AL, 35806; Engineer, UAH, Huntsville, AL, 35806.

<sup>6</sup> Deputy Chief, Astronaut Office, NASA Johnson Space Center, Houston, TX, 77058.

<sup>7</sup> Project Scientist and Science Team Lead, Jacobs Technology Inc., Huntsville, AL, 35805.



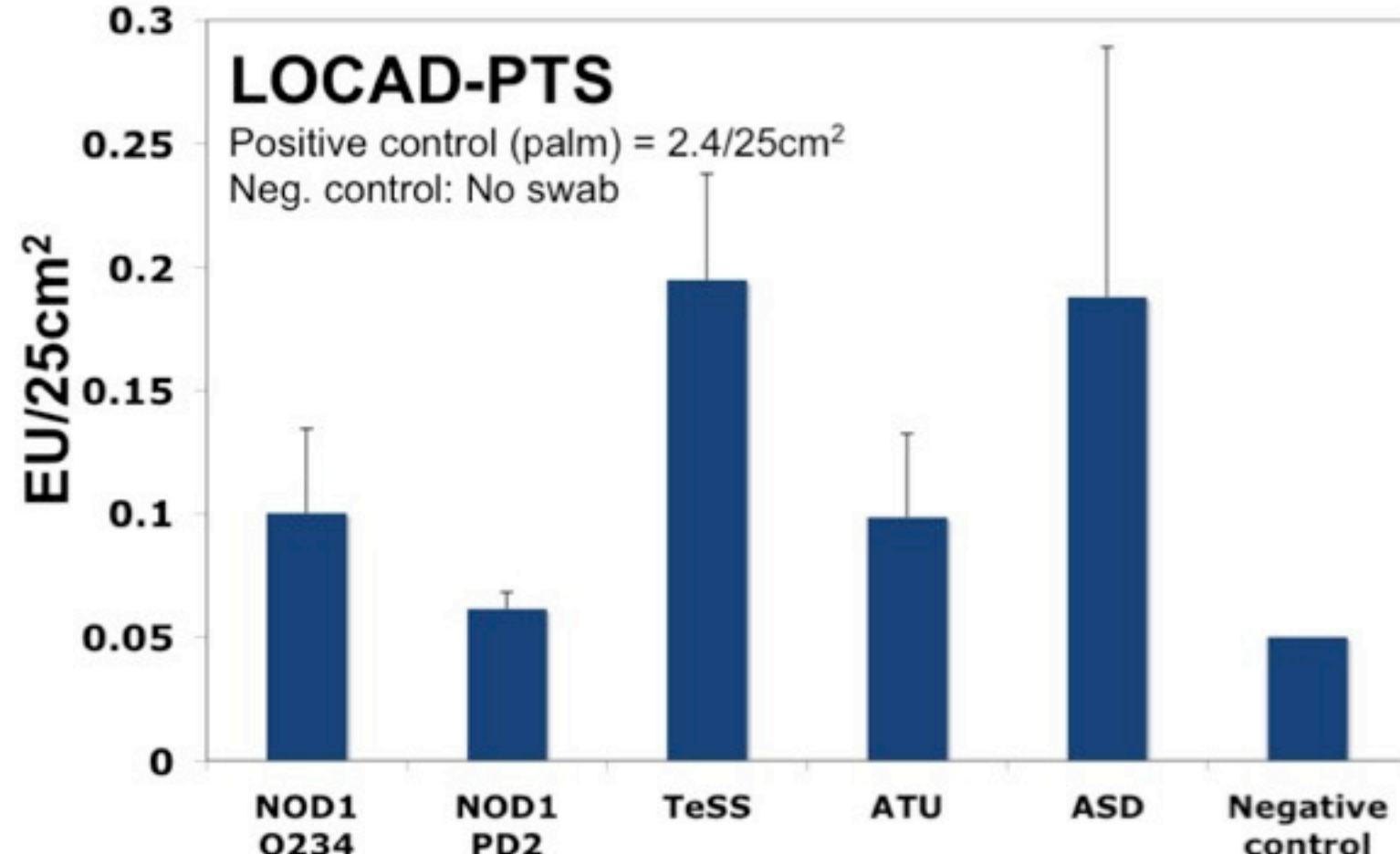
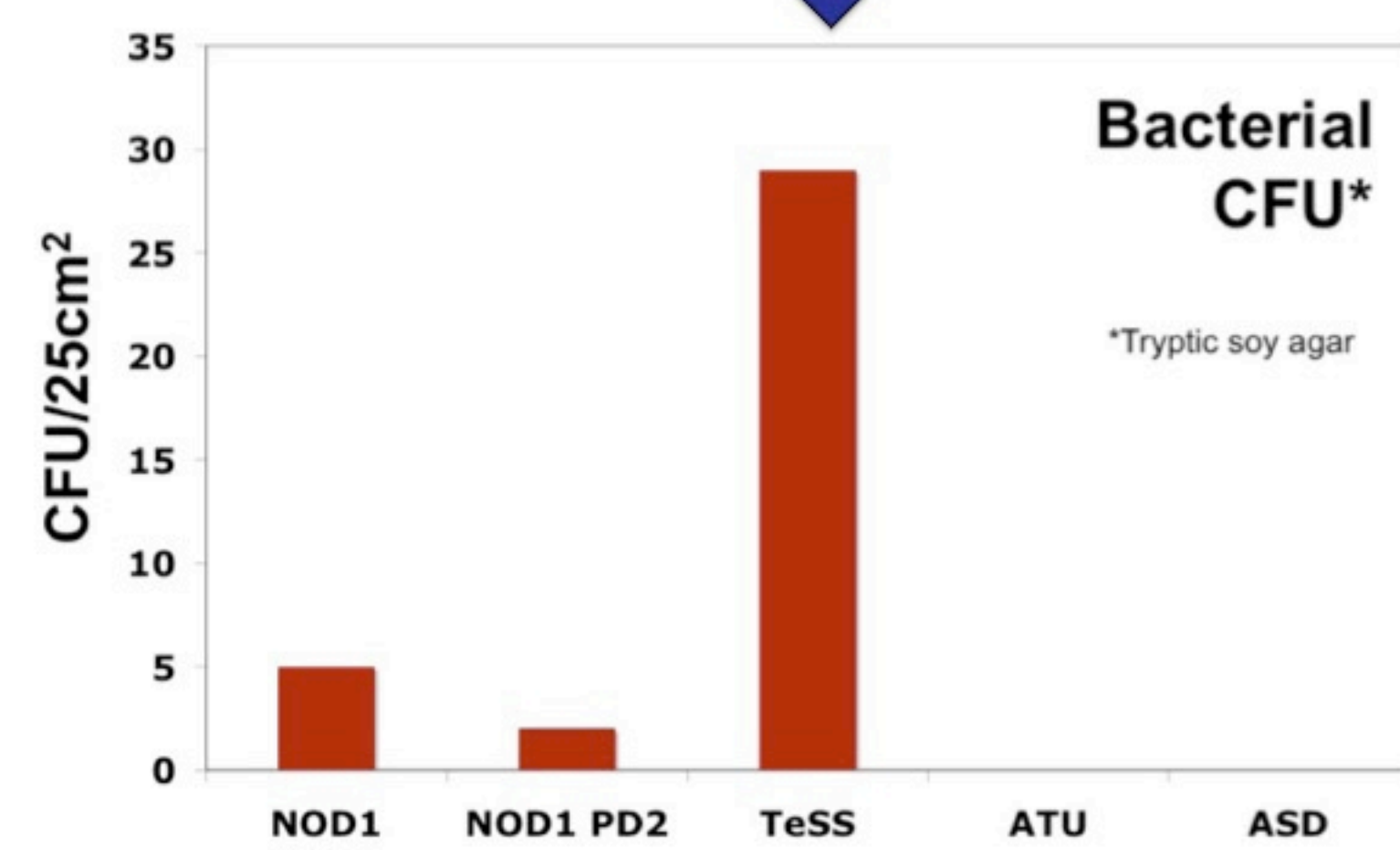
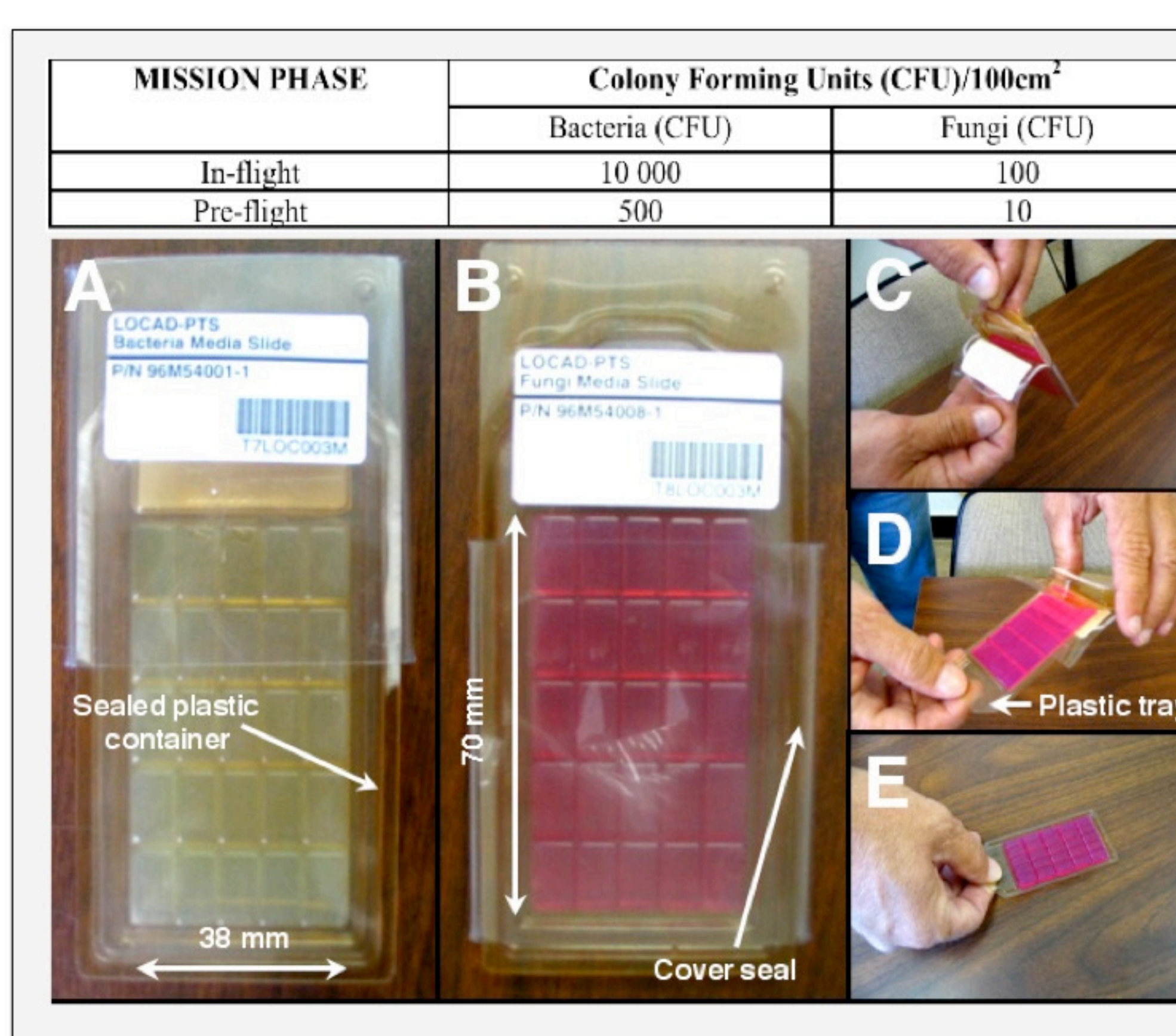
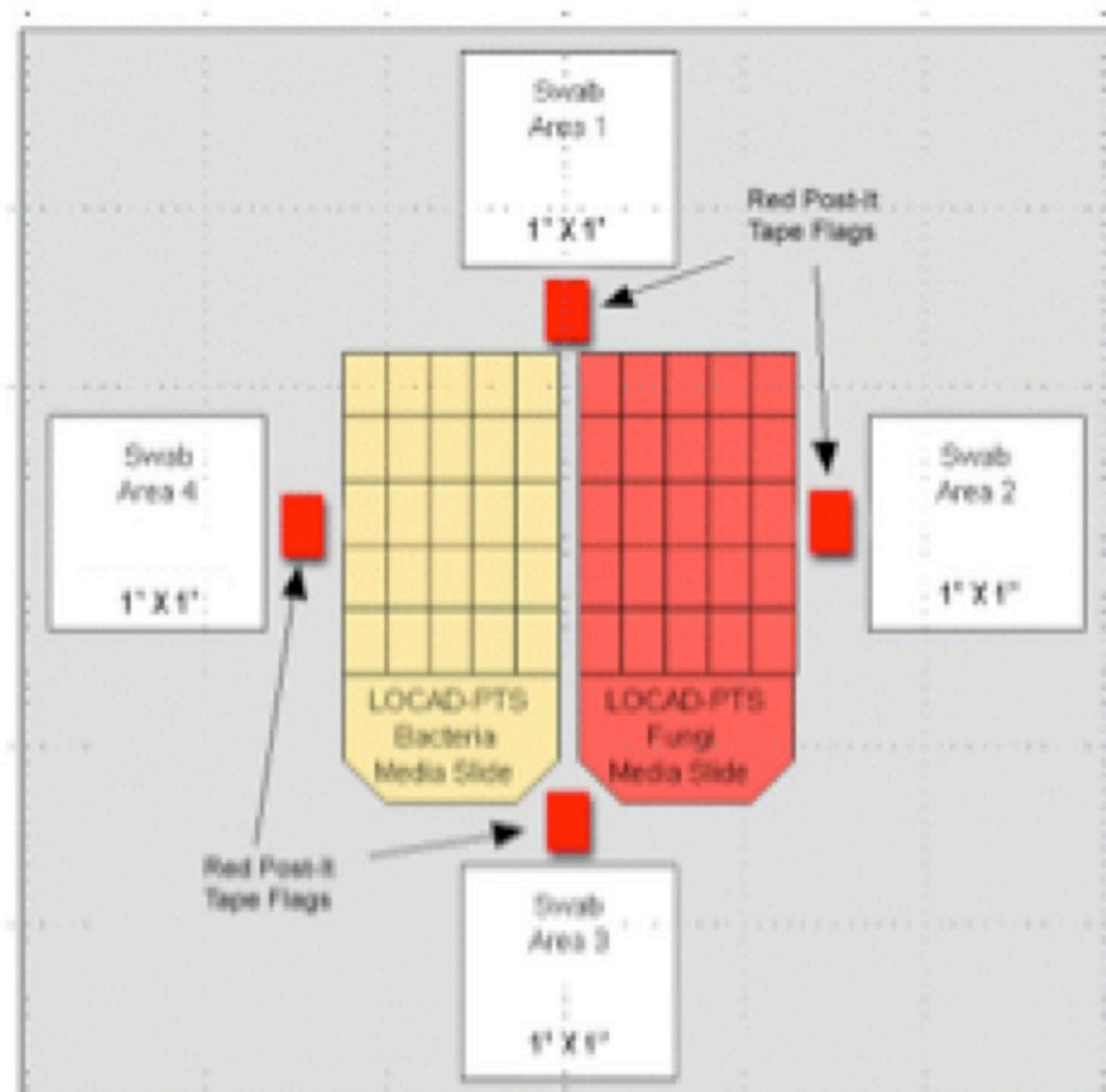
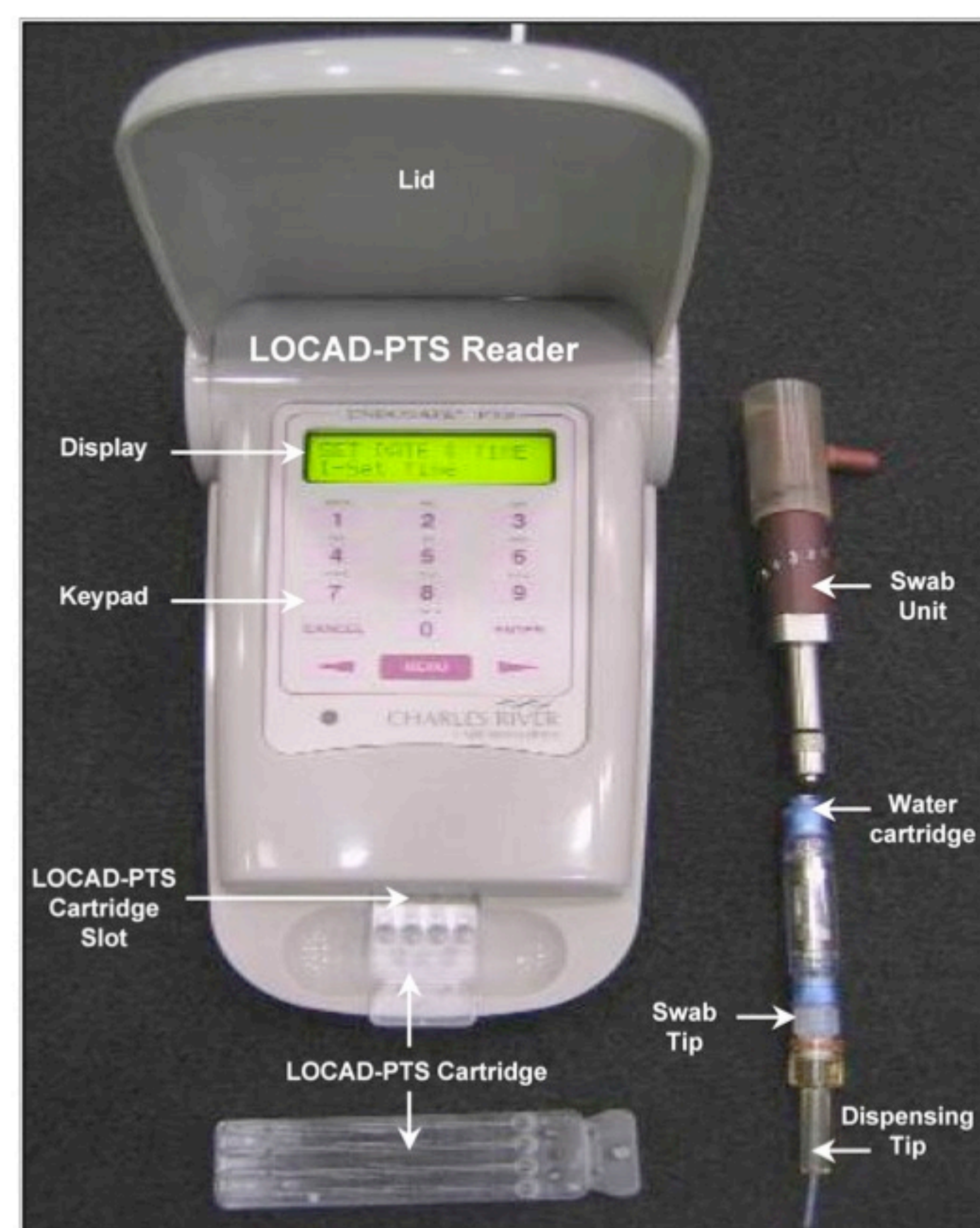
# LOCAD-PTS: Operation of a new system for microbial monitoring aboard the International Space Station (ISS)



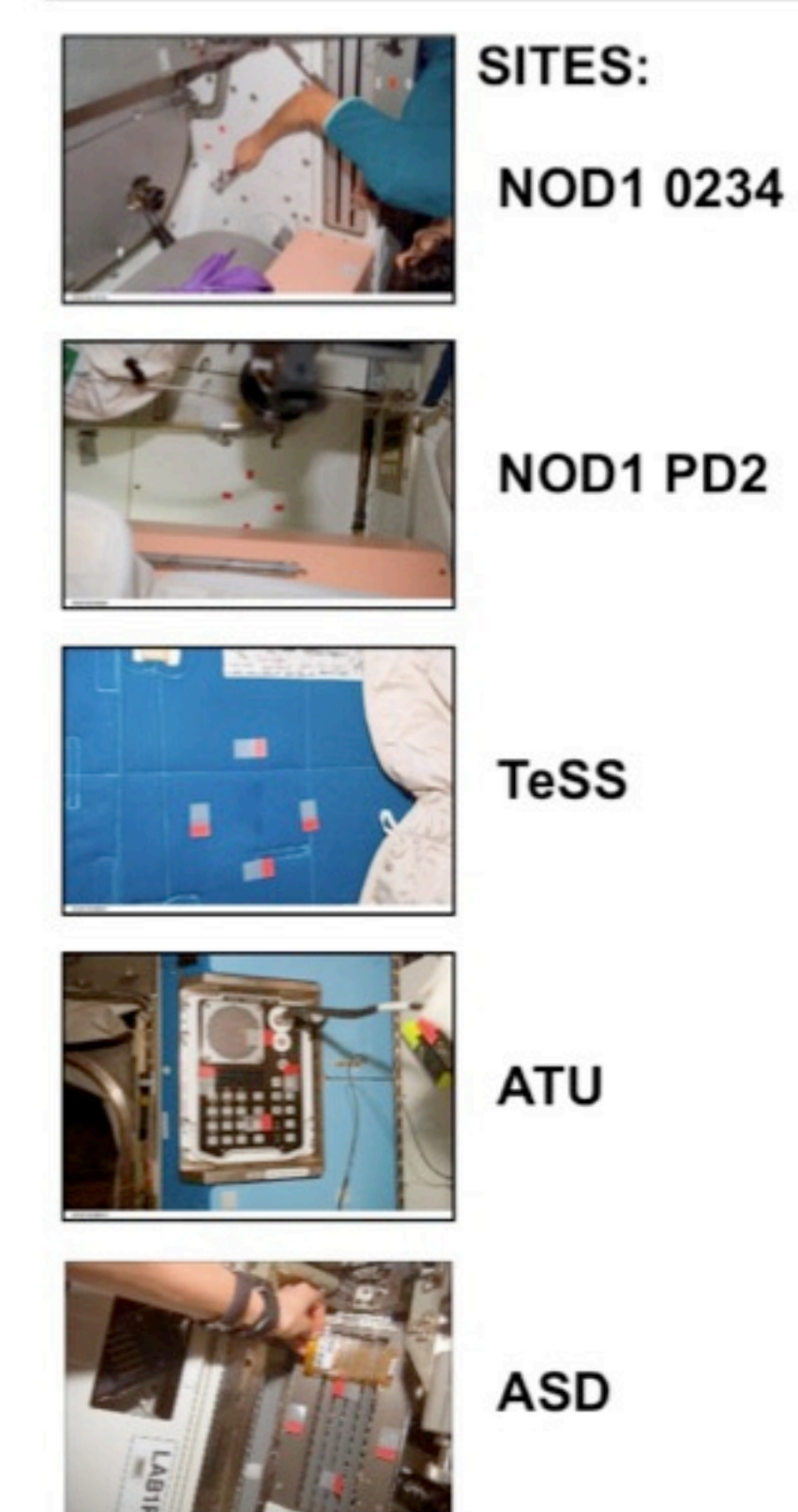
J. Maule<sup>1</sup>, N. Wainwright<sup>2</sup>, A. Steele<sup>3</sup>, D. Gunter<sup>4</sup>, G. Flores<sup>4</sup>, M. Effinger<sup>4</sup>, M. Damon<sup>1</sup>, M. Wells<sup>5</sup>, S. Williams<sup>5</sup>, H. Morris<sup>6</sup> and L. Monaco<sup>6</sup>  
<sup>1</sup>British Aerospace (BAE) Systems, Huntsville, AL, 35806. <sup>2</sup>Charles River Laboratories Inc., Charleston, SC 29407. <sup>3</sup>Carnegie Institution for Science, Geophysical Laboratory, Washington DC 2005. <sup>4</sup>NASA MSFC, Huntsville AL 35812. <sup>5</sup>University of Alabama at Huntsville (UAH), AL 35806. <sup>6</sup>Astronaut Office, NASA Johnson Space Center, Houston, TX 77058. <sup>6</sup>Jacobs Technology Inc., Huntsville, AL 35806.

## ABSTRACT

Microorganisms within the space stations *Salyut*, *Mir* and most recently, the International Space Station (ISS), have traditionally been monitored with culture-based techniques. These techniques involve growing environmental samples (cabin water, air or surfaces) on agar-type media for several days, followed by visualization of resulting colonies; and the return of samples to Earth for ground-based analysis. This approach has provided a wealth of useful data and enhanced our understanding of the microbial ecology within space stations. However, the approach has been limited by the following: i) As many as 95% of the microorganisms in the environment cannot grow on conventional growth media; ii) Significant time lags often occur between onboard sampling and colony visualization (3-5 days) and ground-based analysis (as long as several months); iii) Colonies are often difficult to visualize due to condensation within contact slide media plates; and iv) Culture-based techniques involve growth of potentially harmful microorganisms, which must then be disposed of safely. This report describes the operation of a new culture-independent technique onboard the ISS for rapid analysis (within minutes) of endotoxin and b-1, 3-glucan, found in the cell walls of gram-negative bacteria and fungi, respectively. This technique involves analysis of environmental samples with the *Limulus* Amebocyte Lysate (LAL) assay in a handheld device. This handheld device and sampling system is known as the Lab-On-a-Chip Application Development Portable Test System (LOCAD-PTS). We describe a comparative study between LOCAD-PTS analysis and existing culture-based methods onboard the ISS; together with an exploratory survey of surface endotoxin throughout the ISS.



## Phase 1 LAL on ISS



## Phase 2 LAL Results

Phase 2 Session	Date	Operator	Module	Location	EU/25cm <sup>2</sup>	
1	Dec. 20, 2007	DT	US Lab	Laptop palm rest	0.239	
				Hygiene station wall	0.434	
				TeSS Door	0.149	
2	Dec. 21, 2007	DT	US Lab	CEVIS (Bogun arm)	0.469	
				Node 1	Handrail (PMA1)	0.140
				Bathroom Mirror	3.07	
				CEVIS	0.271	
				LAB1 D5 Rack Face	> 0.050	
3	Dec. 24, 2007	PW	US Lab	SM	Water Dispenser	0.139
				US Lab	Microphone	0.124
				SM	SM table top	0.564
				Air diffuser (P50)	< 0.05	
				TeSS wall	0.054	
4	Jan. 26, 2008	PW	US Lab	Behind panel (D2)	< 0.05	
				Airlock	A/L vent	0.05
				N/A	Negative control	0.05
				US Lab	ACY mirror	0.188
				US Lab	CEVIS (Bogen arm)	< 0.05
5	Feb 2, 2008	DT	Airlock	FGB	Wall opposite mirror	> 0.543
				N/A	Negative control	< 0.05
				N/A	Negative control	< 0.05
				Mfg label at EV hatch	< 0.05	
				EMU glove	< 0.057	
PGT Handle	< 0.051					
Docking Compartment Hatch Handle	> 0.788					

DT = Dan Tani, PW = Peggy Whitson (Increment 16), FGB = функционально-грузовой блок, or "Functional Cargo Block", also known as *Zarya*; SM = Service Module.

## CONCLUSION

It is concluded that while a general correlation between LOCAD-PTS and traditional culture-based methods should not necessarily be expected, a combinatorial approach should be adopted where both sets of data are used *together* to generate a more complete story of the microbial ecology on the ISS.

## References and Acknowledgements

Astronauts Peggy Whitson, Dan Tani and Greg Chamitoff; LOCAD Team Payload Integration Managers Lorrie Hellier and Jeff Durham; ISS Program Scientist Julie Robinson; Increment Scientists Vic Cooley and Kenol Jules, NASA Johnson Space Center (JSC); Duane Pierson and Mark Ott, Microbiology Group, NASA JSC.

# LOCAD-PTS: Operation of a new system for microbial monitoring aboard the International Space Station (ISS)

J. Maule\* and N. Wainwright†

*British Aerospace (BAE) Systems, Huntsville, AL, 35806; and Charles River Laboratories, Charleston, SC 29407*

A. Steele‡

*Geophysical Laboratory, Carnegie Institution for Science, Washington, DC, 20015*

D. Gunter, G. Flores and M. Effinger§

*NASA MSFC, Huntsville, AL 35812*

M. Damon and M. Wells\*\*

*British Aerospace (BAE) Systems, Huntsville, AL, 35806; and University of Alabama at Huntsville (UAH), Huntsville AL 35806*

S. Williams††

*Astronaut Office, NASA Johnson Space Center, Houston, TX, 77058*

H. Morris and L. Monaco‡‡

*Jacobs Technology Inc., Huntsville, AL 35806*

**Microorganisms within the space stations *Salyut*, *Mir* and the International Space Station (ISS), have traditionally been monitored with culture-based techniques. These techniques involve growing environmental samples (cabin water, air or surfaces) on agar-type media for several days, followed by visualization of resulting colonies; and return of samples to Earth for ground-based analysis. This approach has provided a wealth of useful data and enhanced our understanding of the microbial ecology within space stations. However, the approach is also limited by the following: i) More than 95% microorganisms in the environment cannot grow on conventional growth media; ii) Significant time lags occur between onboard sampling and colony visualization (3-5 days) and ground-based analysis (as long as several months); iii) Colonies are often difficult to visualize due to condensation within contact slide media plates; and iv) Techniques involve growth of potentially harmful microorganisms, which must then be disposed of safely. This report describes the operation of a new culture-independent technique onboard the ISS for rapid analysis (within minutes) of endotoxin and  $\beta$ -1, 3-glucan, found in the cell walls of gram-negative bacteria and fungi, respectively. This technique involves analysis of environmental samples with the *Limulus* Amebocyte Lysate (LAL) assay in a handheld device. This handheld device and sampling system is known as the Lab-On-a-Chip Application Development Portable Test System (LOCAD-PTS). A poster will be presented that describes a comparative study between LOCAD-PTS analysis and existing culture-based methods onboard the ISS; together with an exploratory survey of surface endotoxin throughout the ISS. It is concluded that while a general correlation between LOCAD-PTS and traditional culture-based methods should not necessarily be expected, a combinatorial approach can be adopted where both sets of data are used *together* to generate a more complete story of the microbial ecology on the ISS.**

---

\* Project Scientist, LOCAD-PTS, British Aerospace (BAE) Systems, Huntsville, AL, 35806.

† Principal Investigator, LOCAD-PTS, Charles River Labs, Wappoo Road, Charleston, SC, 29407.

‡ Staff Scientist, Geophysical Laboratory, Carnegie Institution for Science, Washington DC, 20015.

§ Lead Flight Operations Engineer and Project Managers (past and present), LOCAD Group, MSFC, Huntsville, AL, 35812.

\*\* Scientist, BAE Systems, Huntsville, AL, 35806; Engineer, UAH, Huntsville, AL, 35806.

†† Deputy Chief, Astronaut Office, NASA Johnson Space Center, Houston, TX, 77058.

‡‡ Scientist and Science Team Lead, Jacobs Technology Inc., Huntsville, AL, 35805.

## I. Introduction

The International Space Station (ISS) is an enclosed habitat, where crews of three to six astronauts live and work for several months at a time. It was recognized early on in the history of the space program that microorganisms might pose a health hazard to the crew if left unmonitored<sup>1</sup>. Since the first space flight, by Yuri Gagarin in 1961, microorganisms within spacecraft have traditionally been analyzed with culture-based techniques, both in the United States<sup>2-5</sup> and Russia<sup>6-13</sup>.

The current microbial requirements for the ISS are shown in Table 1 below. These requirements are currently verified by the Crew Health Care System (CHeCS), an onboard kit consisting of various culture-based methods for the analysis of air and water, as well as surfaces. Surfaces are analyzed with ‘Contact Slides’ (Biotest, Rocaway, NJ), applied to a surface, incubated for 3-5 days, followed by visualization of colonies. More sophisticated analyses are performed on the ground following the return of flight samples to NASA Johnson Space Center.

**Table 1.** Current Microbial Requirements for Internal Surfaces of the International Space Station (ISS)

MISSION PHASE	Colony Forming Units (CFU)/100cm <sup>2</sup>	
	Bacteria (CFU)	Fungi (CFU)
In-flight	10 000	100
Pre-flight	500	10

Requirements from <sup>1</sup>ISS Mission Operations Requirements Document (MORD) section “7.4.4: In-flight environmental microbiology specifications and monitoring of air and surfaces” and “7.4.3 Preflight Environmental Microbiology Specifications and Monitoring of Air and Surfaces”.

These culture-based techniques have provided extremely useful data regarding the type and distribution of microorganisms in space stations, from *Salyut*<sup>13</sup> to the ISS<sup>2</sup>. However, as with all techniques, there are several limitations; in this case, of using a system to detect microorganisms based on their ability to grow on culture media. Four of these limitations include: i) More than 95% microbial species cannot be cultured in conventional growth media<sup>14-15</sup>; ii) Significant time lags often occur between onboard sampling and colony visualization (3-5 days) and ground-based analysis (as long as several months); iii) Colonies are often difficult to visualize due to condensation within contact slide media plates (see Fig. 1); iv) Culture-based techniques involve growth of potentially harmful microorganisms, which must then be disposed of safely.

There are a variety of techniques used to detect microorganisms, each suitable to a particular purpose. These techniques may test parameters such as a microorganism’s ability to grow, metabolic activity, or the presence of characteristic biological molecules. The Lab-On-a-Chip Application Development

Portable Test System (LOCAD-PTS) detects the latter i.e., the presence of biological molecules found in the cell walls of bacteria and fungi. These molecules include: i) Endotoxin, ii)  $\beta$ -1, 3-glucan and iii) lipoteichoic acid/peptidoglycan.

A common classification in microbiology is to divide bacteria into two groups: Gram-negative and Gram-positive. The ‘Gram’ refers to the Gram stain, developed by Hans Christian Gram in 1884, to differentiate *Klebsiella pneumoniae* from pneumococci<sup>16</sup>. With this method, Gram-negative bacteria stain pink and Gram-positive bacteria stain purple, and is related to structure of their cell walls. Endotoxin - the bacterial form of which is also known as lipopolysaccharide (LPS) - is found predominantly in the cell wall outer membrane of Gram-negative bacteria<sup>17</sup>, whereas lipoteichoic acid and peptidoglycan are found in the cell walls of Gram-positive bacteria. The polysaccharide  $\beta$ -1, 3-glucan is not found in either group of bacteria, but only in the cell walls of fungi such as the baker’s yeast, *Saccharomyces cerevisiae*.

LOCAD-PTS detects endotoxin and  $\beta$ -1, 3-glucan with the *Limulus* amoebocyte lysate (LAL) assay. The LAL assay is derived from early discoveries that microbial infection causes blood clots in the horseshoe crab *Limulus polyphemus*<sup>18</sup> and that the causative agent is endotoxin<sup>19-21</sup>. In 1977, the LAL assay replaced the rabbit pyrogen test for endotoxin<sup>22</sup> and in 1983, the Food and Drug Administration (FDA) certified the LAL assay as a standard test for endotoxins.



**Figure 1. Condensation within a ‘Contact Slide’ onboard ISS**

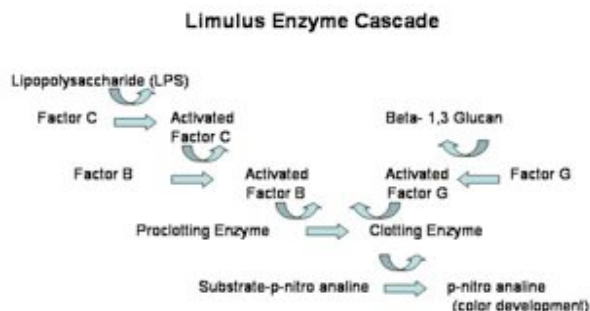
The LAL assay is now used routinely to monitor endotoxin, as an important entity by itself and as a proxy for Gram-negative bacteria, in a wide range of products and applications. These include: dairy products<sup>23</sup>, eye surgery<sup>24</sup>, heart surgery<sup>25</sup>, dental surgery<sup>26</sup>, diagnosis of gram-negative peritonitis<sup>27</sup>, biofilms in dialysis systems<sup>28</sup>, organs for transplantation<sup>29</sup>, microbial contamination of metal-working fluids<sup>30</sup>, air monitoring in factories<sup>31</sup>, research of cyanobacterial blooms<sup>32</sup>, in plasmid-DNA based gene therapy<sup>33,34</sup> to verify that gene vectors are endotoxin-free<sup>33,34</sup>.

While the LAL assay can be coupled to a turbidimetric reaction, this is time-consuming and labor-intensive. To address this issue, the LAL assay was coupled to a colorimetric reaction (see Fig. 2) by Charles River Laboratories Inc. (Charleston, SC) to generate the Endosafe™ Portable Test System (PTS). The Endosafe PTS is a portable spectrophotometer coupled with a range of thumb-sized cartridges in which the colorimetric LAL reaction takes place. Each LAL-containing cartridge also contains p-nitroalanine, which is converted into a yellow/green product if endotoxin is present in the original sample.

The Endosafe™ PTS was then modified by the LOCAD Team at NASA Marshall Space Flight Center (MSFC) to deem it flight qualified, and ensure it could withstand the rigors of launch, and the space environment. The instrument, along with a novel sample collection device, became known as the LOCAD-PTS. LOCAD-PTS therefore consists of a modified Endosafe™ PTS, plus a surface sampling/processing kit that enables astronauts to collect surface samples, dilute them with endotoxin-free water, mix, and then dispense as a liquid into a LOCAD-PTS cartridge.

LOCAD-PTS was launched to the International Space Station (ISS) aboard Space Shuttle Discovery in December 2006 and remains onboard the ISS. This report describes preliminary results from eleven operations of LOCAD-PTS onboard the ISS from March 2007 until the spring of 2008.

This report describes LOCAD-PTS operations performed on the ISS to address five main objectives: i) Demonstrate correct instrument function in a mission environment (success criterion: mid/high reading for positive control); ii) Demonstrate crew handling of LOCAD-PTS supplies without biological contamination e.g. from skin (success criterion: <0.05EU/ml reading for negative control); iii) Demonstrate consistent and reliable crew operations of LOCAD-PTS in a mission environment; iv) Conduct general survey of surface endotoxin within Node 1 and the US Lab; and iv) Compare and contrast endotoxin data with colony forming unit (CFU) data obtained from adjacent areas at each site.



**Figure 2. Colorimetric LAL cascade**

## II. Hardware



Figure 3. LOCAD-PTS

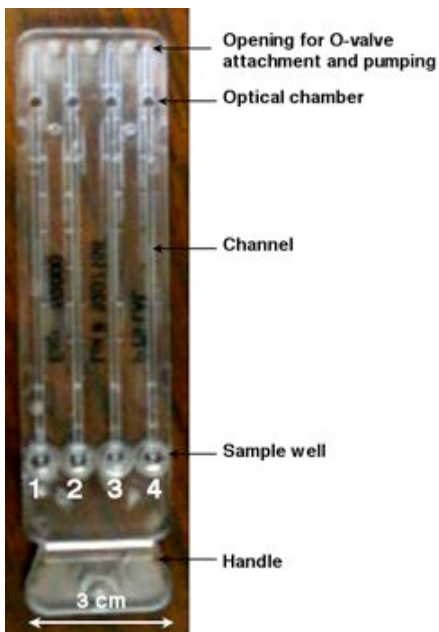


Figure 4. Cartridge

swabbing kit consisted of three components: a water cartridge, a swab tip and a dispensing tip. The water cartridge was constructed with two Luer-lok valves at either end. The swabbing tip consisted of a polycarbonate Luer-lok, covered with Dacron material to form a circular tip with a diameter of 8mm. The capacity of each dispensing tip was 150 $\mu$ l. Each component of the swabbing kit was cleaned to remove endotoxin by boiling in 6%

### A. Reader

The LOCAD-PTS Reader is a handheld integrated spectrophotometer, heater, pump with built-in electronics and software to perform various assays with LOCAD-PTS Cartridges (see Fig. 3). Two Readers were chosen for space flight (one as back-up) and both were manufactured and modified for space flight by Charles River Laboratories. Modifications of the commercially available Readers were minor as far as assay function was concerned and included removal of the battery from the Reader to comply with ISS safety regulations.

### B. Cartridges

LOCAD-PTS LAL cartridges (manufactured by CRL) were approximately 10 cm long, 3 cm wide and inserted into the Reader to perform each assay (see Fig. 4). Each cartridge was manufactured from polycarbonate plastic and consisted of four sample wells (each designed to receive 25 $\mu$ l of liquid sample), with each sample well connected to a channel that leads to an optical chamber. Beyond each of the four optical chambers is a small opening that interfaces with O-rings and a pump in the Reader. This pump was programmed by the Reader software to move fluid up and down the channels of the cartridge as required.

### C. Swabbing Unit

The Swabbing Unit was designed and manufactured by the University of Huntsville in Alabama (UAH) (see Fig. 5). It consists of a hand-held pipette and swabbing device. Several notches were included on the swabbing unit, labeled (from the top): P, W, 1, 2, 3, 4, 5 and 6. Notch W was the largest notch and moving the knob up and down in this position enabled sample mixing. Notches 1-6 were of identical size and enabled dispensing of uniform 25 $\mu$ l droplets into the sample wells (see below for a more detailed description of these procedures). Two swabbing units were chosen to launch on STS-116 (one as a back-up).

### D. Swabbing Kits

Each swabbing kit consisted of a water cartridge, swabbing tip and dispensing tip, and was designed and manufactured at NASA MSFC (see Fig. 6). Each

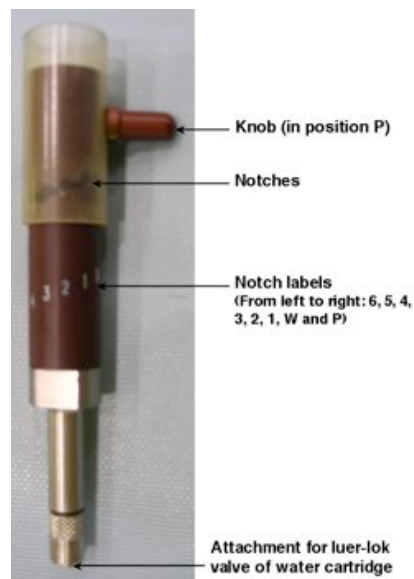


Figure 5. Swabbing Unit

hydrogen peroxide for at least 20 minutes. The components and Teflon-980 material were then dried in a clean, sterile, laminar flow hood. After drying, water cartridges were filled with endotoxin-free water (LAL reagent water, LRW, product number W110; CRL, Charleston, SC) under a laminar flow hood. The swabbing kits were then packaged to include: one full water cartridge, one swabbing tip and one dispensing tip. To ascertain cleanliness of these swabbing kits, 5 out of every 100 produced were analyzed with LOCAD-PTS on the ground before launch. A total of 84 swabbing kits were launched on Space Shuttle STS-116.

#### E. Media “Contact” slides.

This investigation used the term ‘media slides’ interchangeably to refer to the ‘contact slides’ manufactured by Biotest USA (Denville, NJ). Media slides consisted of microbial growth media in the form of a solid gel rectangular pad, supported by a plastic (PVC) tray (see Fig. 7 A and B). This media slide was stored in sealed plastic container until use. When required, the container was peeled open (see Fig. 7, C), the media slide removed (see Fig. 7, D) and applied to a surface of interest (see Fig. 7, E). Following surface application, the media slide was returned to the container, which was secured with a plastic cover seal (see Fig. 7 B).



Figure 6. Swabbing Kit

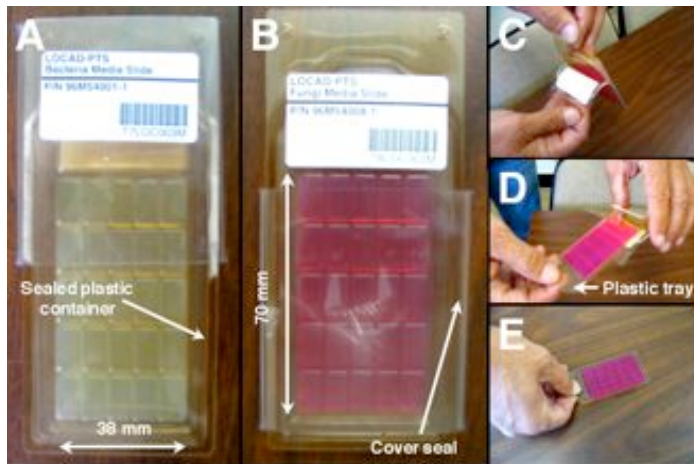


Figure 7. Media ‘Contact’ Slides

Two types of media slide were used in this investigation and referred to as ‘Bacteria’ and ‘Fungi’ media slides. Bacteria media slides corresponded to and were identical to the TC contact slides from Biotest and consisted of tryptic soy agar (product number 931 250 100, Biotest USA). Fungi media slides corresponded to and were identical to YM contact slides and consisted of Rose Bengal Agar with gentamicin sulphate (product number 931 280 100, Biotest USA). The surface area of each media slide was 25cm<sup>2</sup> (to meet International Regulations USP24 and EP3).

### III. Procedures

#### A. Crew Training

In general, crewmembers were trained in LOCAD-PTS operations approximately 12-18 months before launch. In this experiment, procedures onboard ISS were performed by crewmember Sunita Williams, Peggy Whitson, Dan Tani and Greg Chamitoff. Each received a 1-hour training session in the mock-up ISS U.S. Laboratory facility at NASA Johnson Space Center (Houston, TX) several months preceding their mission. In addition, some crewmembers attended a 2-day training session in molecular biology at the Marine Biological Lab (Woods Hole, MA) during 2004, where they were familiarized with LOCAD-PTS. In the JSC training session, Williams was trained to perform a complete end-to-end analysis of a surface swab with LOCAD-PTS, with special attention paid to operation of the swabbing unit to perform mixing and dispensing operations.

#### B. Shipping and pre-launch

All LOCAD-PTS equipment was packaged by Teledyne Brown Engineering (TBE) at MSFC into a Half Cargo Transfer Bag (CTB). The ‘Half’ refers to middeck locker equivalents (MLEs) i.e. a Half CTB will fill half of a Space Shuttle middeck locker. A Half-CTB has a 30 pounds maximum carrying capacity, with the dimensions of

16.75 by 9.75 by 9.25 inches. In order to support and protect equipment against vibrations and shock during ground transport and launch, major components (such as the two Readers) were separated by a solid foam material.

### C. Launch and transfer to ISS

LOCAD-PTS was launched aboard Space Shuttle Discovery (STS-116) on December 9<sup>th</sup> 2006, which docked with the ISS on December 12<sup>th</sup> 2006. Following docking, the half-CTB containing LOCAD-PTS was transferred to the US Lab of the ISS, and stowed at ambient pressure (15 psi) and temperature (76°F) until use on March 31<sup>st</sup> 2007.

### D. Overview of ISS Operations

Each operating session was defined as either a ‘Phase 1’ session or ‘Phase 2’ session. Each Phase 1 session was performed according to the surface sampling layout in Fig. 8, with four surface swabs performed around two centrally placed media slides (for the culture-based detection of fungi and bacteria). Each of these four swabs was analyzed with LOCAD-PTS, so that data from LOCAD-PTS culture-based methods were obtained for each site. The duration of a typical phase 1 session was approximately 3 hours. Each Phase 2 session consisted of five individual surface swabs of separate sites within the ISS; these sessions were slightly shorter than Phase 1 sessions, running approximately 2-3 hours. The selection of these sites were sometimes guided by the LOCAD-PTS operations team, but often left to crew discretion. These Phase 2 sessions constituted an exploratory survey of surface endotoxin throughout the ISS.

A total of five Phase 1 sessions and five Phase 2 sessions were performed for LOCAD-PTS LAL analysis of surface endotoxin. Each Phase 1 session was performed by astronaut Suni Williams on March 31<sup>st</sup>, April 30<sup>th</sup>, May 5<sup>th</sup>, May 8<sup>th</sup> and May 15<sup>th</sup>, 2007. The surface sites chosen for each of these five sessions are shown in Fig. 9. Each of the Phase 2 LAL sessions were performed by astronauts Peggy Whitson and Dan Tani on December 20<sup>th</sup>, 21<sup>st</sup> and 24<sup>th</sup> 2007 and February 2<sup>nd</sup> 2008. Phase 1 Glucan sessions began on June 21<sup>st</sup> 2008, performed by astronaut Greg Chamitoff.

The LOCAD-PTS operations team supported each session from mission control at the Payload Operations Integration Center (POIC) at MSFC. Prior to each astronaut’s first session, a science conference was scheduled so that the crew could discuss procedures and scientific objectives with the LOCAD-PTS team. Prior to on-orbit operations, all crew procedures were tested first by the LOCAD-PTS operations team in the microgravity environment of parabolic flight (Maule et al., 2004).

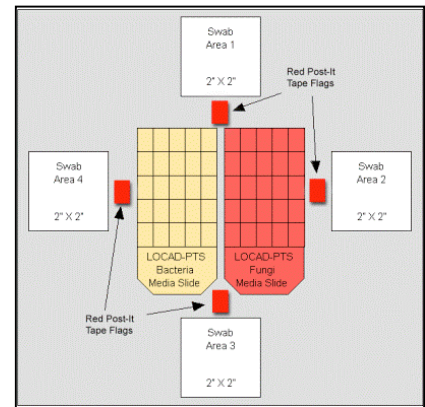


Figure 8. Surface sampling layout

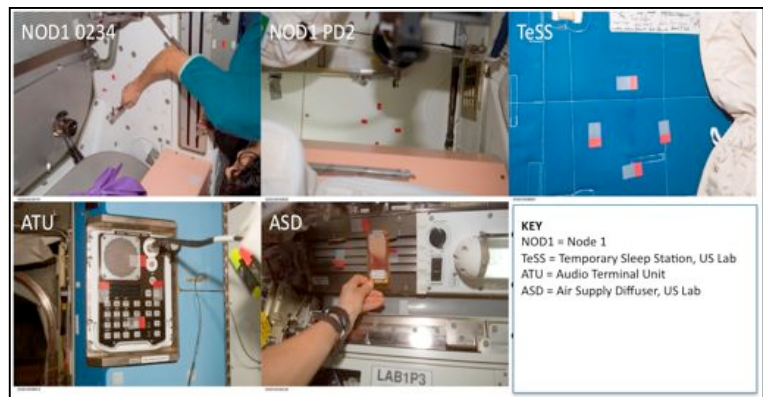


Figure 9. Surface site locations during Phase 1 LAL operations

### E. Swabbing

As can be seen from Fig. 8, each LOCAD-PTS swab covered a surface area of 25cm<sup>2</sup> (or 2” x 2”), with all four swabs surrounding two centrally placed contact slides (for the culture-based detection of bacteria and fungi) during Phase 1. Swabs performed during Phase 2 similarly covered 25cm<sup>2</sup> (but contact media slides were not used in Phase 2). In this way, both culture-independent and culture-based methods were used together to analyze an approximate 10” x 10” area (i.e., the grey box in Fig. 8). The crew prepared to swab in this layout by placing four red post-it flags on the surface site (see red rectangles in Fig. 8); applying contact slides within those four post-it flags; followed by four swabs at the positions indicated. The swabbing protocol began with direct attachment of the water cartridge of the swabbing kit via a push-and-turn action (while the knob of the swabbing unit was at position P). The water cartridge was then attached to the swabbing tip by another push-and-turn action. At this point the swab unit was in



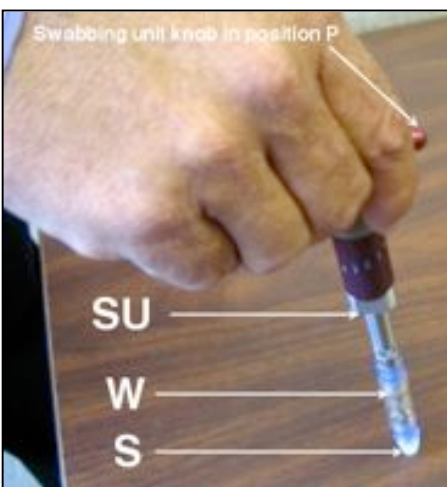
the ‘swabbing configuration’ i.e., a surface sample could be obtained by swabbing a surface with the swabbing tip (see Fig. 10).

#### F. Mixing

Following swabbing, the dispensing tip was attached to the swab tip via a push-on O-ring seal. This is called the ‘mixing/dispensing configuration’ and at this point, mixing and dispensing took place. The knob on the swabbing unit was manipulated up and down in notch W (see Fig. 5) five times at a rate of once per 2 seconds to mix the swab sample with 1.2ml of endotoxin-free water in the water cartridge. Following mixing, the crew performed the dispensing procedure (see below).

#### G. Dispensing

Following mixing, the knob of the swabbing unit was left in position 1. A further droplet was dispensed, and then droplets were dispensed into an inserted LOCAD-PTS cartridge by cycling through notches 2-6. Moving the knob down each of the subsequent notches (labeled 1-6) dispensed droplets of approximately 25 $\mu$ l each. Fig. 11 shows Suni Williams dispensing a sample onboard the ISS during Phase 1 LAL operations.



**Figure 10. Swabbing configuration.** SU = swab unit; W = water cartridge; S = swab tip



**Figure 11. Astronaut Suni Williams dispensing a mixed surface sample during Phase 1 LAL operations (March, 2007)**

### IV. Discussion

This report describes the operation of a new type of biochemical microbial analysis performed aboard the International Space Station (ISS) that builds upon the collective experience and excellent previous studies performed in both the United States and Russia, from the early programs of *Apollo*<sup>4,5</sup> and *Salyut*<sup>13</sup>, to space station *Mir*<sup>10-12</sup>, the Shuttle-*Mir* program<sup>3</sup> and most recently, the ISS<sup>2,6-9</sup>.

It was the first demonstration of complete biochemical analysis of environmental samples onboard a space station from sampling to onboard data. While the analytical test performed was relatively simple - the *Limulus* Amebocyte Lysate (LAL) assay for detection of endotoxin and  $\beta$ -1, 3-glucan - the demonstration of the technology and procedures serves as a foundation for future generations of more specific tests. The ease with which a variety of crewmembers – pilots and scientists – performed procedures was especially important, given that a relatively high level of hand-eye coordination and control was required, especially for the dispensing phase of operations. While this procedure might be second nature to a biochemist, it may not so familiar to others, especially in a microgravity environment; however, all crewmembers performed procedures equally well throughout each session.

A potential concern prior to this study was the absence of gloves from crew procedures, which could lead to skin-derived biological contamination of the swab tip and false positive readings. The negative control was designed to address this concern (where procedures were performed without a surface swab). No positive readings were obtained during any of the three negative controls performed on orbit – indicating that procedures did not introduce contamination to the swab tip (and that the reagents remained clean following extended storage on orbit).

The waived requirement for gloves is a major advantage given the extra up-mass of clean gloves (and onboard storage requirements), down-mass of used gloves, and greater crew time required for gloved operations. The robust design of the cartridges, which enabled handling with bare hands, was a major benefit to operations within the distinctly non-sterile ISS cabin environment.

While a general correlation should not be expected between data obtained with LOCAD-PTS and existing culture-based methods, it is concluded that the two types of data should be used *together* to generate a more complete story of microbial ecology on the ISS. This approach - using a *combination* of culture-independent and traditional methods, rather than one at the expense of another - has been implemented in many other areas from hygiene monitoring on Earth<sup>35</sup> to field biology investigations of extreme environments on Earth, such as Kamchatka<sup>36</sup> and the Arctic<sup>37</sup>.

### Acknowledgments

Many thanks to astronauts Suni Williams, Peggy Whitson, Dan Tani and Greg Chamitoff, who operated LOCAD-PTS on the ISS; LOCAD Team Payload Integration Managers Lorrie Hellier and Jeff Durham for their hard work preparing LOCAD-PTS for launch and during operations; ISS Program Scientist Julie Robinson, Increment Scientists Vic Cooley and Kenol Jules, NASA Johnson Space Center (JSC), for facilitating science sessions within the constraints of ISS mission operations; Duane Pierson, Mark Ott, Victoria Castro, Willy Wong, Anthony Jeevarajan and Bekki Bruce, Microbiology Group, NASA JSC, and Jeff Jones, Flight Surgeon, NASA JSC, for their support in the transition of LOCAD-PTS to a spaceflight instrument.

### References

- <sup>1</sup>Fox, L., "The ecology of micro-organisms in a closed environment." *Life Sci Space Res.* 1971; 9: 69-74.
- <sup>2</sup>Castro, V.A., Thrasher, A.N., Healy, M., Ott, C.M., Pierson, D.L., "Microbial characterization during the early habitation of the International Space Station." *Microb. Ecol.* 2004 Feb; 47(2): 119-26.
- <sup>3</sup>Pierson, D.L., "Microbial contamination of spacecraft." *Gravit. Space Biol. Bull* 2001; 14(2):1-6.
- <sup>4</sup>Puleo, J.R., Oxborrow, G.S., Fields, N.D., Herring, C.M., Smith, L.S., "Microbiological profiles of four Apollo spacecraft." *Appl Microbiol.* 1973 Dec; 26(6): 838-45.
- <sup>5</sup>Puleo, J.R., Oxborrow, G.S., Fields, N.D., Hall, H.E., "Quantitative and qualitative microbiological profiles of the Apollo 10 and 11 spacecraft." *Appl Microbiol.* 1970 Sep; 20(3): 384-9.
- <sup>6</sup>Novikova, N., De Boever, P., Poddubko, S., Deshevaia, E., Polikarpov, N., Rakova, N., Coninx, I., Mergeay, M., "Survey of environmental biocontamination onboard the International Space Station." *Res Microbiol.* 2006 Jan-Feb; 157(1): 5-12.
- <sup>7</sup>Novikova, N.D., "Review of the knowledge of microbial contamination of the Russian manned spacecraft." *Microb Ecol.* 2004 Feb; 47(2):127-32.
- <sup>8</sup>Novikova, N.D., "Basic patterns of microflora development in the environment of orbital complex Mir." *Aviakosm Ekolog Med.* 2001; 35(4):32-40.
- <sup>9</sup>Rakova, N.M., Svistunova, I.V., Novikova, N.D., "Studies on the growth and reproduction of bacterial communities on structural materials of the international space station." *Aviakosm Ekolog Med.* 2005 Jan-Feb; 39(1): 44-6.
- <sup>10</sup>Viktorov, A.N., Novikova, N.D., Deshevaia, E.A., Bragina, M.P., Shnyreva, A.V., Sizova, T.P., D'iakov, I.T., "Residential colonization of orbital complex "Mir" environment by penicillium chrysogenum and problem of ecological safety in long-term space flight". *Aviakosm Ekolog Med.* 1998;32(5):57-62.
- <sup>11</sup>Viktorov, A.N., Novikova, N.D., Deshevaia, E.A., "The cabin microflora of manned space vehicles and the problem of the biological destruction of the construction materials used in them." *Aviakosm Ekolog Med.* 1992 May-Jun; 26(3): 41-8.
- <sup>12</sup>Shikina, M.I., Vinogradova, L.A., Kolesina, N.B., "Microflora of the drinking water regenerated from atmospheric moisture condensate in hermetically sealed quarters." *Kosm Biol Aviakosm Med.* 1988 Mar-Apr; 22(2): 53-6.
- <sup>13</sup>Nefedov, Y.G., Shilov, V.M., Konstantinova, I.V., Zaloguyev, S.N., "Microbiological and immunological aspects of extended manned space flights." *Life Sci Space Res.* 1971; 9:11-6.
- <sup>14</sup>Keep, N.H., Ward, J.M., Robertson, G., Cohen-Gonsaud M., Henderson, B., "Bacterial resuscitation factors: revival of viable but non-culturable bacteria". *Cell Mol Life Sci.* 2006 Nov; 63(22): 2555-9.
- <sup>15</sup>Byrd, J.J., Xu, H.S., Colwell, R.R., "Viable but nonculturable bacteria in drinking water." *Appl. Environ. Microbiol.* 1991 Mar; 57(3): 875-8.
- <sup>16</sup>Gram, H.C., "Über die isolierte Färbung der Schizomyceten in Schnitt- und Trockenpräparaten". *Fortschritte der Medizin,* 1884; 2: 185-89.
- <sup>17</sup>Tefsen, B., Geurtsen, J., Beckers, F., Tommassen, J., de Cock, H., "Lipopolysaccharide transport to the bacterial outer membrane in spheroplasts." *J. Biol. Chem.* 2005, Feb 11; 280(6):4504-9.
- <sup>18</sup>Bang, F.B., "A bacterial disease of *Limulus polyphemus*." *Bull Johns Hopkins Hosp.* 1956 May;98(5):325-51.
- <sup>19</sup>Levin, J., and Bang, F.B., "Clottable protein in *Limulus*; its localization and kinetics of its coagulation by endotoxin". *Thromb. Diath. Haemorrh.* 1968 Mar 31;19(1):186-97.
- <sup>20</sup>Levin, J., and Bang, F.B., "A description of cellular coagulation in the *Limulus*." *Bull Johns Hopkins Hosp.* 1964 Oct;115: 337-45.

- <sup>21</sup>Levin, J., and Bang, F.B., "The role of endotoxin in the extracellular coagulation of *Limulus* blood." *Bull Johns Hopkins Hosp.* 1964 Sep;115: 265-74.
- <sup>22</sup>Ronneberger, H.J., "Comparison of the pyrogen tests in rabbits and with *limulus* lysate." *Dev Biol Stand.* 1977;34: 27-36.
- <sup>23</sup>Mottar, J., De Block, J., Merchiers, M., Vantomme, K., Moermans, R., "Routine *limulus* amoebocyte lysate (LAL) test for endotoxin determination in milk using a Toxinometer ET-201." *J. Dairy Res.* 1993 May; 60(2): 223-8.
- <sup>24</sup>Spelsberg, H., Reinhard, T., Sokolovska, Y., Zysk, G., Böhringer, D., Sundmacher, R., "Endotoxin in storage medium of human corneal grafts and clinical course after penetrating normal-risk keratoplasty." *Eye.* 2004 Sep; 18(9):923-8.
- <sup>25</sup>Bowles, C.T., Ohri, S.K., Klangasuk, N., Keogh, B.E., Yacoub, M.H. and Taylor, K.M., "Endotoxaemia detected during cardiopulmonary bypass with a modified *Limulus* amoebocyte lysate assay." *Perfusion.* 1995 Jul; 10(4): 219-28.
- <sup>26</sup>Martinho, F.C. and Gomes, BP., "Quantification of endotoxins and cultivable bacteria in root canal infection before and after chemomechanical preparation with 2.5% sodium hypochlorite." *J. Endod.* 2008 Mar; 34(3): 268-72.
- <sup>27</sup>Pee, G., Serban, I., Blum, M., Cabili, S. and Iaina, A., "Early diagnosis of gram-negative peritonitis in continuous ambulatory peritoneal dialysis patients with the *Limulus* amoebocyte lysate assay." *Am. J. Nephrol.* 1992; 12(1-2):19-21.
- <sup>28</sup>Liu, H., Li, G., Zhang, X., Duan, X., Yao, Y., "A study on the endotoxin and bacterial contamination in reverse osmosis water and dialysis fluids." *Zhonghua Nei Ke Za Zhi.* 1999 Dec; 38(12): 806-9.
- <sup>29</sup>Pillay, S.P., Wynter, C., Lynch, S., Wall, D., Balderson, G., Strong, R., "Endotoxin levels in donors and recipients during orthotopic liver transplantation." *Aust N Z J Surg.* 1997 Apr; 67(4): 187-91.
- <sup>30</sup>Cyprowski, M., Piotrowska, M., Zakowska, Z., Szadkowska-Stańczyk, I., "Microbial and endotoxin contamination of water-soluble metalworking fluids." *Int J Occup Med Environ Health.* 2007; 20(4): 365-71.
- <sup>31</sup>Marchand, G., Lalonde, M., Beaudet, Y., Boivin, G., Villeneuve, S., Pépin, C., "Documentation of the endotoxins present in the ambient air of cotton fiber textile mills in Québec." *J. Environ. Monit.* 2007 Aug; 9(8): 869-76.
- <sup>32</sup>Bernardová, K., Babica, P., Marsálek, B. and Bláha, L.J., "Isolation and endotoxin activities of lipopolysaccharides from cyanobacterial cultures and complex water blooms and comparison with the effects of heterotrophic bacteria and green alga." *Appl Toxicol.* 2008 Jan; 28(1): 72-7.
- <sup>33</sup>Wells, D.J., Maule, J., McMahon, J., Mitchell, R., Damien, E., Poole, A., Wells, K.E., "Evaluation of plasmid DNA for in vivo gene therapy: factors affecting the number of transfected fibers." *J Pharm Sci.* 1998 Jun; 87(6):763-8.
- <sup>34</sup>Wells, K.E., Maule, J., Kingston, R., Foster, K., McMahon, J., Damien, E., Poole, A., Wells, D.J., "Immune responses, not promoter inactivation, are responsible for decreased long-term expression following plasmid gene transfer into skeletal muscle." *FEBS Lett.* 1997 Apr 28; 407(2):164-8.
- <sup>35</sup>Aycicek, H., Oguz, U., Karci, K., "Comparison of results of ATP bioluminescence and traditional hygiene swabbing methods for the determination of surface cleanliness at a hospital kitchen." *Int. J. Hyg Environ. Health.* 2006 Mar; 209(2): 203-6.
- <sup>36</sup>Maule, J., Morrill, P., Shaw, A. and Leuko, S., "International 2006 NAI Expedition to Kamchatka". *NASA Astrobiology Conference*, Santa Clara, CA. April 2008.
- <sup>37</sup>Steele, A., Amundsen, H.E.F., Conrad, P.G., Benning, L., and Fogel, M., on Behalf of the AMASE 07 team, "Arctic Mars Analogue Svalbard Expedition (AMASE) 2007". *Lunar and Planetary Science Conference XXXIX* (2008), Houston, TX.