

# Antibacterial effect of a novel material "earth-plus" for resistant nosocomial bacteria

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# BACKGROUND

With regard to hospital-associated infections, bacterial contamination of fabrics such as doctor's coats, ties, and curtains is problematic and non-negligible. However, it is difficult to disinfect these fabrics.

## PURPOSE

The earth-plus<sup>TM</sup> (Shinshu Ceramics Co. Ltd., Nagano, Japan) is a novel paint, which contains a hydroxyapatite-binding Ag/TiO<sub>2</sub> ceramic composite. The bactericidal effect of cotton / polyester fabric coated with the earth-plus<sup>TM</sup> was investigated. In addition, the duration of the bactericidal effect was evaluated when the fabric was repeatedly washed.

# RESULTS

Table : Bactericidal effects for fabric with or without e+

Bacterial strains	objects -	Incubation time		Inhibitation rate
		Oh	18h	of growth
HA-MRSA	fabric without e+	4.3	6.7	
(ATCC BAA-1699)	fabric with e+	4.4	2.3	99.99
CA-MRSA	fabric without e+	4.7	6.9	
(ATCC BAA-1680)	fabric with e+	4.7	3.2	99.99
VRE (ATCC 51299)	fabric without e+	4.7	5.9	
	fabric with e+	4.1	2.7	99.92
ESBL E. coli (clinical	fabric without e+	2.4	6.3	
isolate)	fabric with e+	3.3	<1.3	99.99
MDRP (clinical isolate)	fabric without e+	2.4	6.3	
	fabric with e+	3.3	<1.3	99.99
MDRA (clinical isolate)	fabric without e+	4.4	5.9	
	fabric with e+	4.3	<1.3	99.99

# MATERIALS AND METHODS

#### (1) Examination of the bactericidal effects

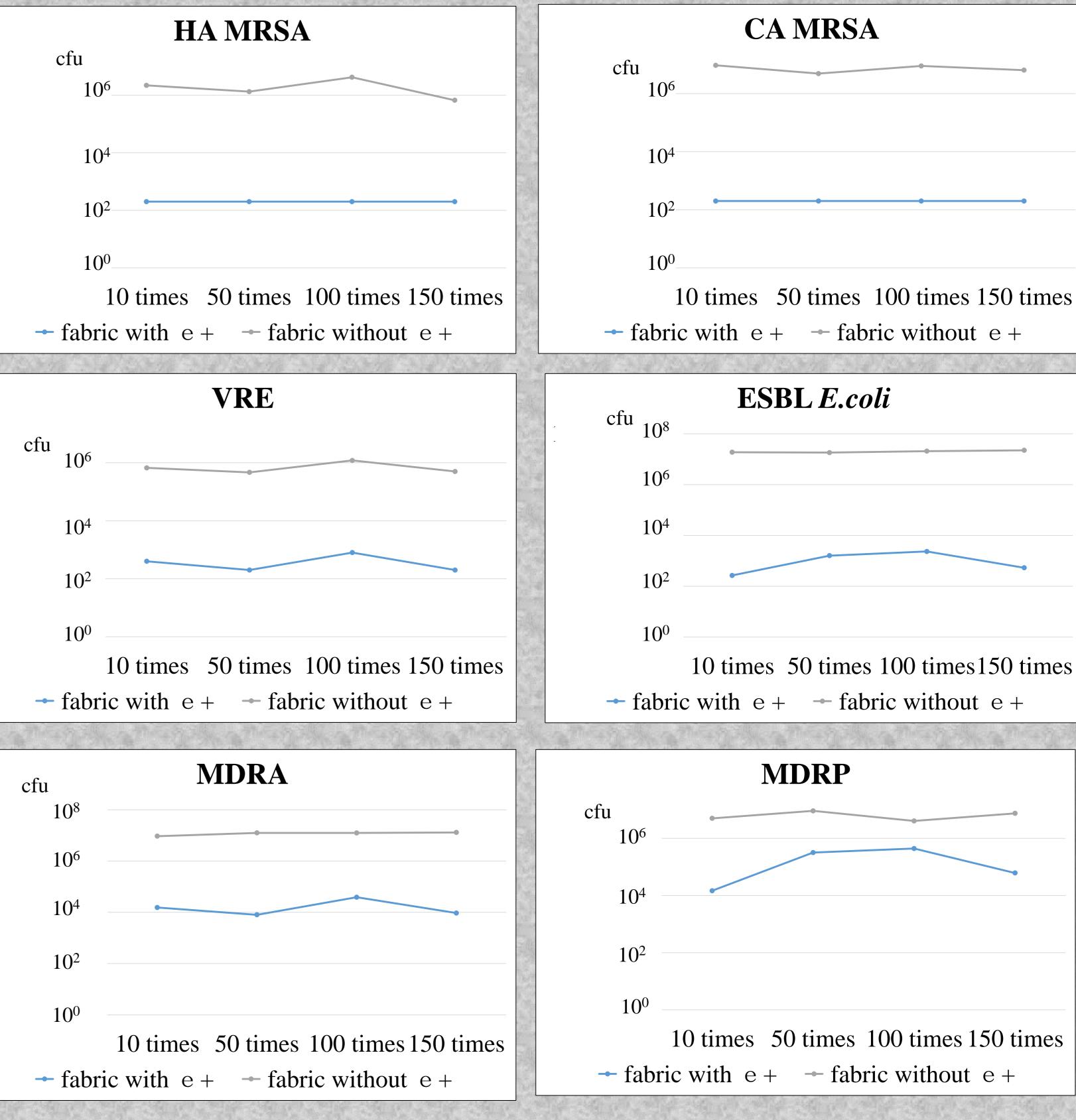
(i) Evaluated objects

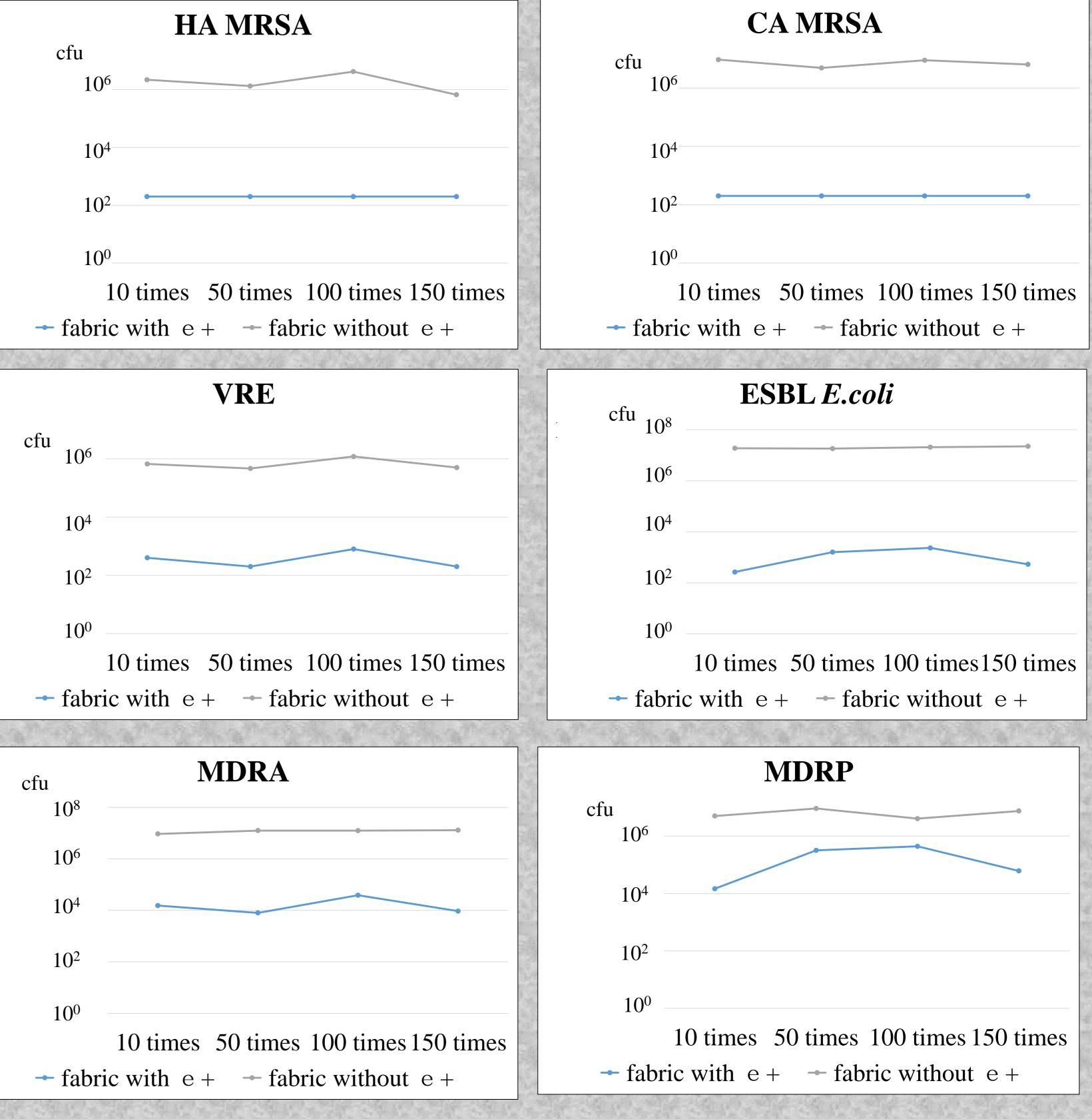
#1 cotton / polyester fabrics with the earth-plus  $^{TM}$  (e+) coating (4g/m)

#2 cotton / polyester fabrics without the earth-plus<sup>TM</sup> ( $\alpha$ (e+) coating (ii) Bacterial strains

- #1 Hospital-acquired methicillin-resistant Staphylococcus aureus (ATCC BAA-1699)
- #2 Community-acquired methicillin-resistant *Staphylococcus aureus* (ATCC BAA-1680)
- #3 Vancomycin-resistant *Enterococcus faecalis* (ATCC 51299)
- #4 Extend spectrum  $\beta$ -lactamase-producing *Escherichia coli* (clinical isolate)
- #5 Multi-drug resistant *Pseudomonas aeruginosa* (clinical isolate)
- #6 Multi-drug resistant *Acinetobacter* species (clinical isolate)
- (iii) Bactericidal Procedure 1), 2)
- #1 According to Japanese Industrial Standards L1902 and ISO 20743

Figure ; Bactericidal effect for 10, 50, 100, and 150 washing times





#2 Bacterial suspensions

Density of approximately 1 x 10°CFU/mL  $(100\mu L=1 \times 10^{4} CFU \text{ was dropped})$ 

#3 Measurements at time points

0 and 18 hours

#### #4 Evaluation

- Bacterial solutions were extracted from each objects by shaking.
- The plate-colony count method is performed.
- Colonies were counted after incubation for 24 hours at 37 degree.
- The examination was performed three times.

#### (2) Examination of earthplus with various washing times

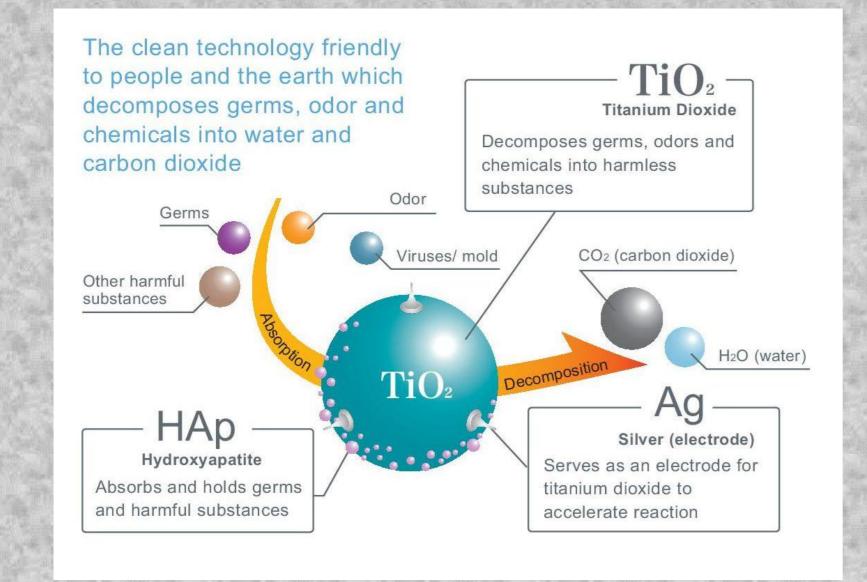
(i) Evaluated objects: The same objects (1)-(i) (ii) Bacterial strains: The same strains (1)-(ii) (iii) Washing Procedure

### #1 Washing method

- Hot water of 80 degrees was used
- Washing times are 10, 50, 100, and 150
- #2 Bacterial suspensions

After washing,  $1 \ge 10^{\circ}$  CFU was dropped on the objects and incubated at 37 degrees for 18 hours #3 Evaluation: Same as (1)-(iii)-#4

### **FUNCTIONS OF "earth-plus "TM**



• Non-elution feature

- Ceramics compound material
- The earthplus does not elute in water due to being a ceramics compound material, therefore, it is a safe, low burden to environments material.

### CONCLUSIONS

The earth-plus<sup>TM</sup> can be used to coat various materials. Cotton / polyester coated with the earthplus could contribute to a decrease of cross-contamination by problematic drug-resistant bacteria that cause hospital-associated infections due to its bactericidal effect and long-lasting antibacterial activity.

### **BIBLIOGRAPHY**

1. Japanese Industrial Standards Committee. JIS L1902. Available from: http://www.jisc.go.jp/index.html. Accessed August 26, 2011.

2. Swenson LM, Hindler JF, Jorgensen JH. Assessment of Bactericidal Activity by the Time-Kill Method. In Manual of Clinical Microbiology, 8th ed. Washington, DC: American Society of Microbiology; 2003.