

ORAL PRESENTATION



O006: Development of a method to simulate practical use conditions of hygienic handrubs

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From 2nd International Conference on Prevention and Infection Control (ICPIC 2013) Geneva, Switzerland. 25-28 June 2013

Introduction

The World Health Organization has recognized the shortcomings of current standards for evaluating the *in vivo* efficacy of hand hygiene products, and has called for methods which are "realistic under practical conditions".

Objectives

The objective of this study was to develop a method to evaluate the efficacy of alcohol-based handrubs, which reflects the mode of hand contamination typical in healthcare settings to provide an accurate assessment of product performance under in-use conditions.

Methods

Stainless steel discs 1 cm in diameter were contaminated with 10 μ l of a liquid suspension of *S. aureus* ATCC 6538 (8 log CFU/ml) and allowed to dry. Discs were stored in a humidity chamber at 50% RH for up to 72 hours prior to use. Hands were contaminated by firmly pressing each fingerpad to a contaminated disc for 2 seconds. Two fingers on each hand were sampled individually by kneading in a neutralizer solution for 30 seconds to obtain pre-treatment counts. A hand hygiene intervention was performed after which the remaining fingers were sampled to obtain post-treatment values. Recovered bacteria were quantified and mean log reductions per finger were calculated.

Results

S. aureus was stable on stainless steel discs for several days. Transfer and recovery of *S. aureus* from fingers was highly reproducible both between the fingers of individual subjects and between different subjects (mean recovery = $5.9\pm0.2 \log$ CFU per finger pad; *N*=65). The

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organism was stable on the fingers with no die off for at least 40 minutes. A 15 second non-antimicrobial handwash, 0.5 ml, and 1 ml of an alcohol based hand rub achieved log reductions of 3.2 ± 0.5 , 2.9 ± 1.3 and 3.7 ± 1.0 , respectively. Consistent with other hygienic hand rub methods, intra-subject variability was low and inter-subject variability was high.

Conclusion

Contamination of the fingers via contact with a dry surface appears to be a simple and highly reproducible means of evaluating the efficacy of hand hygiene products under practical use conditions. Furthermore, this method utilizes a relevant marker organism, and simulates the primary mode of hand contamination in healthcare settings. Finally, the sampling method may be applied to the clinical setting to perform Phase 3 field studies, to investigate prevention of cross-transmission of pathogens through use of a hygienic handrub.

Disclosure of interest

None declared.

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Published: 20 June 2013

doi:10.1186/2047-2994-2-S1-O6 **Cite this article as:** Rutter and Macinga: **O006**: **Development of a method to simulate practical use conditions of hygienic handrubs**. *Antimicrobial Resistance and Infection Control* 2013 **2**(Suppl 1):O6.



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