ORIGINAL ARTICLE

Differential Effects of Chlorhexidine Skin Cleansing Methods on Residual Chlorhexidine Skin Concentrations and Bacterial Recovery

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BACKGROUND.   Bathing intensive care unit (ICU) patients with 2% chlorhexidine gluconate (CHG)–impregnated cloths decreases the risk of healthcare-associated bacteremia and multidrug-resistant organism transmission. Hospitals employ different methods of CHG bathing, and few studies have evaluated whether those methods yield comparable results.

OBJECTIVE.   To determine whether 3 different CHG skin cleansing methods yield similar residual CHG concentrations and bacterial densities on skin.

DESIGN.   Prospective, randomized 2-center study with blinded assessment.

PARTICIPANTS AND SETTING.   Healthcare personnel in surgical ICUs at 2 tertiary-care teaching hospitals in Chicago, Illinois, and Boston, Massachusetts, from July 2015 to January 2016.

INTERVENTION.   Cleansing skin of one forearm with no-rinse 2% CHG-impregnated polyester cloth (method A) versus 4% CHG liquid cleansing with rinsing on the contralateral arm, applied with either non–antiseptic-impregnated cellulose/polyester cloth (method B) or cotton washcloth dampened with sterile water (method C).

RESULTS.   In total, 63 participants (126 forearms) received method A on 1 forearm (n = 63). On the contralateral forearm, 33 participants received method B and 30 participants received method C. Immediately and 6 hours after cleansing, method A yielded the highest residual CHG concentrations (2500 µg/mL and 1250 µg/mL, respectively) and lowest bacterial densities compared to methods B or C ($P < .001$).

CONCLUSION.   In healthy volunteers, cleansing with 2% CHG-impregnated cloths yielded higher residual CHG concentrations and lower bacterial densities than cleansing with 4% CHG liquid applied with either of 2 different cloth types and followed by rinsing. The relevance of these differences to clinical outcomes remains to be determined.

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The antiseptic properties of chlorhexidine gluconate (CHG) have been known since the 1950s.[1] CHG has diverse clinical applications from oral hygiene[2] to preoperative surgical site skin preparation.[3] The use of no-rinse 2% CHG-impregnated cloths for routine patient bathing in the intensive care unit (ICU) has been shown to decrease the risk of

healthcare-associated bloodstream infections[4–6] and to reduce cross transmission of multidrug-resistant organisms (MDROs).[4,7] Other ICU-based studies in which patients were bathed daily with 4% CHG formulations instead of no-rinse 2% CHG-impregnated cloths have suggested similar decreases in MDRO transmission;[8,9] however, these studies have been

limited by lack of randomization and by varying bathing techniques.

Few studies have directly compared outcomes between skin cleansing with no-rinse 2% CHG-impregnated cloths versus 2% or 4% CHG liquid formulations. Despite the lack of comparative data, some hospitals assume equivalent efficacy for these techniques and bathe patients with 4% CHG liquid formulations rather than 2% CHG-impregnated cloths based on relatively lower cost.[10,11] The objective of the present study was to determine whether cleansing with 2% CHG-impregnated cloths versus cleansing with 4% CHG liquid delivered by 2 commonly used methods yielded comparable CHG concentrations and residual microbial densities on the skin of healthy volunteers.

METHODS

Study Population and Recruitment

Healthcare personnel from the surgical ICUs of Rush University Medical Center (RUMC) in Chicago, Illinois, and from Brigham and Women's Hospital/Harvard Medical School (BWH) in Boston, Massachusetts, were recruited between July 2015 and January 2016. Subjects with known allergies or prior adverse reactions to CHG and those who had nonintact skin or well-defined erythema (defined as erythema grade ≥2; erythema grade scale modified from Vernon et al[7]) on their forearms were excluded. All other healthcare personnel who worked in the surgical ICUs were eligible for study participation. The institutional review boards of RUMC and BWH reviewed and approved the study independently; written informed consent was required at RUMC and verbal assent was required at BWH.

Study Design

With randomized laterality, each participant had 1 forearm cleansed using method A: 2% CHG-impregnated polyester cloth, ~ 500 mg CHG per cloth as per manufacturer (Sage 2% Chlorhexidine Gluconate Cloths, Sage Products [now part of Stryker], Cary, IL). Each participant's contralateral forearm was then randomized to cleansing with 5 mL of undiluted 4% CHG (Hibiclens, Mölnlycke Health Care, Norcross, GA) liquid suspension applied with either a cellulose/polyester cloth impregnated with a nonantiseptic solution (Comfort Bath, Sage Products) (Method B) or with a cotton washcloth dampened with sterile water (method C). Each cleansing method was applied to the skin for 20 seconds. For 4% CHG products, this was followed by a 2-minute dwell time,[12,13] then a 20-second wipe off with a clean cloth of the same type used for application. The wipe-off technique for the 4% CHG liquid was consistent with manufacturer recommendations for general skin cleansing.[14] At each participating hospital, skin cleansing protocols were standardized to be identical and were conducted by a single investigator to ensure uniformity.

Research staff, who were blinded to cleansing assignment, collected swab samples within a 5×5-cm^2 area on the forearm at 3 time points: immediately prior to, immediately after, and 6 hours after skin cleansing. Throughout the 6 hours, participants performed their routine clinical work. To test for residual CHG on the skin of each participant, sterile swabs were moistened with sterile water (Bio-Swab, Arrowhead Forensics, Lenexa, KS), and forearms were swabbed for 10 seconds. For bacterial cultures, flocked swabs (FLOQSwabs, Copan, Murrieta, CA) were moistened in sterile water and a 5×5-cm^2 area near the area swabbed for residual CHG was swabbed for 10 seconds. To avoid the wipe-off effect of CHG sample collection, the same area was not swabbed twice. A short survey was administered to each study participant to collect demographic data, information on use of skin products, and reports of skin reactions after cleansing. Subject-level data were deidentified before analysis.

Laboratory Methods

Chlorhexidine gluconate concentration was measured using a colorimetric method described previously.[13,15] Swab samples for culture were placed immediately into 500 μL neutralizing agent[16,17] without ether sulfate, as this reagent was unavailable for purchase in the United States at the time of testing. Serially diluted 100 μL volumes were aliquoted to 5% sheep's blood agar plates in duplicate and incubated at 35 ± 2°C in ambient air for up to 48 hours. Colonies were counted and transformed to colony forming units (CFUs) after correcting for any dilution factor. Presumptive identifications were performed using standard microbiologic methods. All laboratory testing was conducted by research personnel blinded to study assignments.

Statistical Design and Analysis

Sample size estimates were based on an earlier study[15] that found an effect size of $d = 0.62$[18] for the reduction in CFUs when the CHG concentration was increased from 37.5–150 μg/mL to 300–600 μg/mL. Assuming an effect size of 0.62, a sample of 60, and a 1-tailed α of 0.05, a power of 0.96 was obtained for the current study design.

Deidentified results were recorded in an electronic database and shared between institutions. The Kruskall-Wallis test was used to compare differences in bacterial density, CHG skin concentrations, and other ordinal variables. For comparison of categorical variables, χ^2 analysis or the Fischer exact test was used as appropriate for expected values. All statistical tests were 2-tailed; an α level of 0.05 was considered significant. Testing was performed using SPSS version 22 software (IBM, Armonk, NY), SAS version 9.2 software (SAS Institute, Cary, NC), or R software (R Foundation for Statistical Computing, Vienna, Austria).

RESULTS

Survey Results

In total, 32 participants were enrolled at RUMC and 31 participants were enrolled at BWH (63 participants and 126 forearms in total). The median age was 35 years (range, 30–45

years). Most participants were female (86%) and right-handed (87%). Nurses and patient care assistants made up the largest category of healthcare personnel (70%). Also, 2 participants reported having eczema, 1 participant reported having psoriasis, and 1 participant reported a remote history of idiopathic urticaria. When stratified by CHG cleansing intervention, no statistically significant differences were detected among groups in age, body mass index, gender, medical team role, dominant arm cleansed, presence of watch or bracelet, underlying dermatologic conditions affecting the forearms, use of topical skin products in the 24 hours prior to skin cleansing, or sleeve length prior to cleansing or throughout the 6-hour period (Table 1). Most participants reported cleansing to be relatively pleasant (≥4/5 on pleasantness scale). However, 2 participants reported minimal erythema at 6 hours after cleansing, 1 of whom had no change from baseline erythema that was caused by bracelet irritation.

Chlorhexidine Gluconate Concentration and Bacterial Density on Skin

Chlorhexidine gluconate was not detected on any participant's forearm immediately prior to cleansing (Figure 1). Cleansing with no-rinse 2% CHG yielded the highest residual CHG skin concentrations immediately and 6 hours after cleansing; this difference remained statistically significant when compared to each of the two 4% CHG cleansing methods independently ($P < .001$ for each comparison) or combined ($P < .001$). Cleansing with 4% CHG liquid and non–antiseptic-impregnated cloths resulted in higher residual CHG concentrations than cleansing with 4% CHG with cotton washcloth immediately and 6 hours after cleansing ($P < .001$ and $P = .002$, respectively).

Bacterial density on forearms varied by participant prior to cleansing, but the differences were not statistically significant ($P = .29$) (Table 2). Immediately and 6 hours after cleansing, bacterial densities decreased after all 3 cleansing methods. The no-rinse 2% CHG cloth cleansing yielded the lowest bacterial densities at both time points when compared to both 4% CHG cleansing methods combined ($P < .001$). Differences in bacterial densities between 4% CHG cleansing methods were not statistically significant at either time point after cleansing. Residual bacteria isolated after cleansing were commensal skin microbiota such as coagulase-negative staphylococci, micrococcus, and *Bacillus* species.

DISCUSSION

We found that cleansing the skin of healthy volunteers with no-rinse 2% CHG-impregnated cloths yielded the highest residual CHG concentrations and lowest bacterial densities compared to cleansing with 4% CHG liquid applied with non–antiseptic-impregnated cellulose/polyester cloths or with cotton washcloths dampened with sterile water, followed by rinsing. The strengths of our study design include the use of highly standardized cleansing methods and incorporation of

blinded sample collection and outcome assessment. More importantly, we compared residual CHG concentrations and microbial densities on the skin directly between participants. Our findings thus extend those of 2 other separate investigations in which the application of 2% CHG-impregnated cloths yielded higher residual CHG concentrations[13] or greater microbial reductions at skin sites[19] in healthy volunteers compared to cleansing with 4% CHG liquid.

The CHG concentration needed to reduce microbial bioburden on skin appears to be dependent at least in part on microbial susceptibility to CHG.[20] In a study of ICU patients, Popovich et al[15] found that a CHG skin concentration ≥18.75 µg/mL was inversely associated with gram-positive bacterial colony counts, including *Staphylococcus aureus* and *Enterococcus* species. Lin et al[21] calculated that the relative risk of skin contamination with *Klebsiella pneumoniae* carbapenemase (KPC)-producing *Enterobacteriaceae* was decreased by approximately half in long-term acute-care hospital patients who had CHG skin concentrations ≥128 µg/mL. In both studies, the optimal concentration of CHG on skin exceeded the minimum inhibitory concentrations (MICs) of the targeted bacteria. Notably, some strains of bacteria have been reported to have CHG MICs as high as 2500 µg/mL.[22] In our study, cleansing with no-rinse 2% CHG-impregnated cloth was the only method that achieved CHG concentrations as high as 2500 µg/mL.

Several factors may account for the observed differences in effects of 2% CHG-impregnated cloth and 4% CHG liquid skin cleansing methods. First, 2% CHG-impregnated cloth bathing is "no-rinse," whereas manufacturers of 4% liquid CHG solutions instruct users to rinse off the solution after skin cleansing, an instruction that we followed.[14,23] Few data have been published on the safety of bathing patients with undiluted CHG liquid without rinsing. In a study of patients admitted to a long-term acute-care hospital, daily bathing with 2% CHG liquid solution without rinsing was discontinued in 3 of 405 patients due to mild and reversible generalized erythema or pruritus.[24] Swan et al[25] published a study of 350 surgical ICU patients who were bathed every other day with CHG liquid without rinsing, alternating with soap and water. Their results showed the same incidence of adverse skin reactions compared to bathing patients daily with soap and water only. More recently, Alserehi et al[26] demonstrated that bathing patients in a trauma ICU with diluted 4% CHG solution without rinsing resulted in near equivalent proportions of skin samples that had adequate CHG skin concentrations (defined as ≥18.75 µg/mL) compared to bathing with CHG-impregnated cloths. However, only 10 patients were studied, and adverse skin reactions related to the no-rinse solution were not evaluated.

The cloth that was used to apply 4% CHG solution to the skin may have also affected residual CHG concentrations and bacterial densities. In our study, the application of 4% CHG with cotton washcloths yielded consistently lower CHG concentrations than application with cellulose/polyester cloths that were impregnated with a nonantiseptic solution. Cotton fibers

TABLE 1. Demographics, Study Characteristics, Survey Feedback, and Observations by Chlorhexidine Gluconate (CHG) Skin Cleansing Intervention Groups

Characteristic	2% CHG Impregnated, No-Rinse Cloth (n = 63), No. (%)[a]	4% CHG Applied With Non–Antiseptic-Impregnated Cloth (n = 33), No. (%)[a]	4% CHG Applied With Cotton Washcloth (n = 30), No. (%)[a]	P Value
Demographics				
Age, median (IQR), y	35 (30–45)	36 (30.5–45.5)	33 (28.8–45.3)	.73
BMI, median (IQR)[b]	24 (21.8–27.8)	24.2 (22.2–27.2)	23.3 (21.6–29.4)	.98
Sex				.88
Male	9 (14.3)	4 (12.1)	5 (16.7)	
Female	54 (85.7)	29 (87.9)	25 (83.3)	
Occupation				.99
Nurse	39 (61.9)	21 (63.6)	18 (60)	
Patient care assistant	5 (7.9)	2 (6.1)	3 (10)	
Physician	7 (11.1)	4 (12.1)	3 (10)	
Physical therapist	1 (1.6)	0 (0)	1 (3.3)	
Respiratory therapist	6 (9.5)	4 (12.1)	2 (6.7)	
Other[c]	5 (7.9)	2 (6.1)	3 (10)	
Study characteristics				
Laterality of arm cleansed				.91
Right	32 (50.8)	17 (51.5)	14 (46.7)	
Left	31 (49.2)	16 (48.5)	16 (53.3)	
Dominant arm cleansed				.81
Yes	32 (50.8)	15 (45.5)	16 (53.3)	
No	31 (49.2)	18 (54.5)	14 (46.7)	
Survey results as reported by study participants				
Eczema on forearm	2 (3.2)	2 (6.1)	0 (0)	.56
Psoriasis on forearm	1 (1.6)	0 (0)	1 (3.3)	.49
Other dermatologic conditions on forearm[d]	1 (1.6)	1 (3)	0 (0)	.99
≥ 1 topical product used on forearm in the last 24 h	55 (87.3)	29 (87.9)	26 (86.7)	.99
Sleeve length				
Prior to cleansing				.86
Long	32 (50.8)	18 (54.5)	14 (46.7)	
Three-quarters	6 (9.5)	4 (12.1)	2 (6.7)	
Short	25 (39.7)	11 (33.3)	14 (46.7)	
Throughout most of 6 h				.30
Long	32 (50.8)	19 (57.6)	13 (43.3)	
Three-quarters	10 (15.9)	7 (21.2)	3 (10)	
Short	21 (33.3)	7 (21.2)	14 (46.7)	
Pleasantness of cleansing method[e]				.86
Level 2	1 (1.6)	0 (0)	1 (3.3)	
Level 3	19 (30.2)	9 (27.3)	8 (26.7)	
Level 4	10 (15.9)	5 (15.2)	4 (13.3)	
Level 5	33 (52.4)	19 (57.6)	17 (56.7)	
Gross contamination on the forearm within 6 hours of cleansing	3 (4.8)	0 (0)	0 (0)	.43
Researcher observations				
Hair on forearm	25 (39.7)	14 (42.4)	11 (36.7)	.89
Wearing a watch[f]	7 (11.1)	1 (3)	5 (16.7)	.18
Wearing a bracelet[f]	4 (6.3)	4 (12.1)	4 (13.3)	.45
Erythema on forearm[g]				
Prior to cleansing				.63
Grade 0	62 (98.4)	32 (97)	30 (100)	
Grade 1	1 (1.6)	1 (3)	0 (0)	

TABLE 1.   *Continued*

Characteristic	2% CHG Impregnated, No-Rinse Cloth (n = 63), No. (%)[a]	4% CHG Applied With Non−Antiseptic-Impregnated Cloth (n = 33), No. (%)[a]	4% CHG Applied With Cotton Washcloth (n = 30), No. (%)[a]	P Value
Immediately after cleansing				.20
Grade 0	63 (100)	33 (100)	29 (96.7)	
Grade 1	0 (0)	0 (0)	1 (3.3)	
6 hours after cleansing				.04
Grade 0	63 (100)	33 (100)	28 (93.3)	
Grade 1	0 (0)	0 (0)	2 (6.7)	

NOTE. IQR, interquartile range.
[a]Unless otherwise specified.
[b]For BMI, data were missing for 2 participants for 2% CHG intervention group and 1 participant for each 4% CHG intervention group.
[c]Other occupations included clerk, occupational therapist, and pharmacist.
[d]Other dermatologic conditions includes one patient with remote idiopathic urticaria.
[e]Level corresponds to degree of pleasantness and ranges from 1 to 5, with 1 indicating very unpleasant and 5 indicating very pleasant.
[f]Subjects who wore a watch or bracelet at the wrist or distal forearm prior to cleansing were allowed to keep the jewelry on throughout the 6 hours.
[g]Grade of erythema corresponds to severity and ranges from 0 to 4 with a grade of 0 indicating no erythema, grade 1 indicating very slight erythema (barely perceptible), grade 2 indicating well-defined erythema, grade 3 indicating moderate to severe erythema, and grade 4 indicating severe erythema (beet redness).
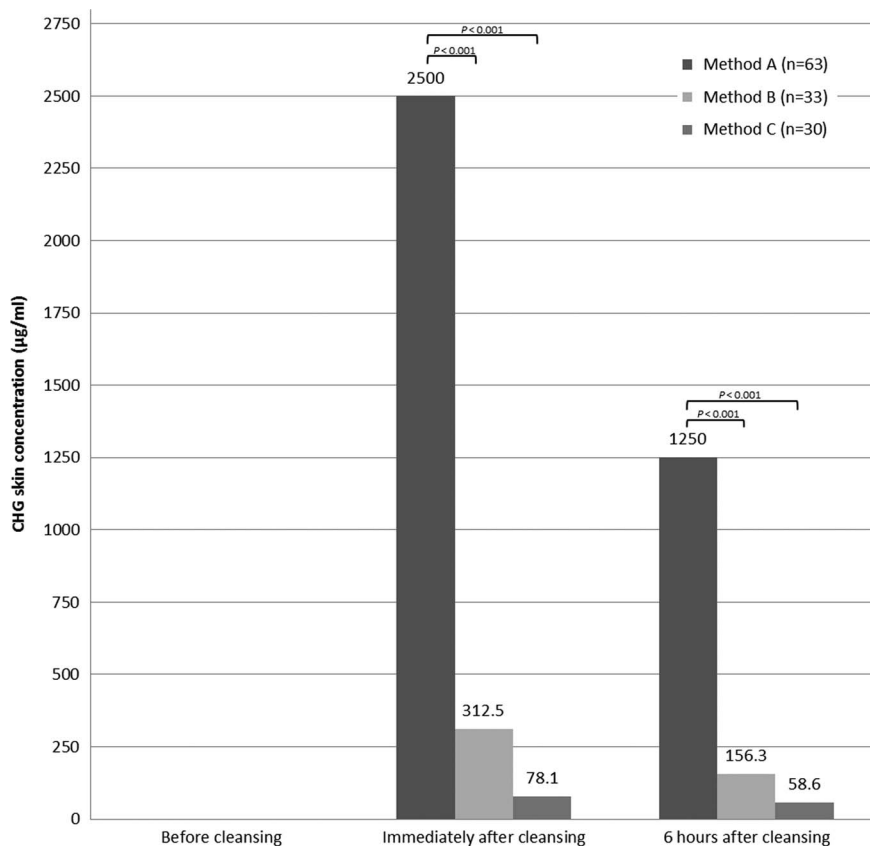

FIGURE 1.   Chlorhexidine gluconate (CHG) concentrations measured on the skin of participants randomized to 3 different CHG cleansing methods. NOTE. Median values are reported. Data are also significant at $P < .001$ between all 3 groups immediately and 6 hours after cleansing. Method A: 2% CHG-impregnated cloth; method B: 4% CHG liquid with non−antiseptic-impregnated cloth; method C: 4% CHG liquid with cotton washcloth. No participants had CHG detected on skin at the baseline measurement.

TABLE 2. Bacterial Density Measured on the Skin of Participants Randomized to 3 Different Chlorhexidine Gluconate (CHG) Cleansing Methods

CHG Cleansing Method	Bacterial Density (CFU/25 m^2)[a]		
	Before Cleansing	Immediately After Cleansing	6 Hours After Cleansing
Method A: 2% CHG-impregnated cloth (n = 63)	465	0	0
Method B: 4% CHG liquid with non–antiseptic-impregnated cloth (n = 33)	442.5	2.5	15
Method C: 4% CHG liquid with cotton washcloth (n = 30)	685	5	16.3
P value[b]	.29	<.001	<.001

NOTE. CFU, colony-forming units.

[a]Median values are reported.

[b]P values shown test the null hypothesis that all 3 groups are equal immediately and 6 hours after cleansing. Data also significant at $P = .002$ for method A vs method B immediately after cleansing, and $P < .001$ for method A vs method B 6 hours after cleansing. $P < .001$ for method A vs method C immediately after and 6 hours after cleansing.

are known to bind CHG[27] and may release less CHG to patients' skin during bathing, in comparison to noncotton cloths. Finally, the differential effects of the 3 bathing methods may have been due to the different quantities of CHG that were applied to skin: 200 mg CHG per 5 mL 4% CHG liquid versus 500 mg CHG per 2% CHG-impregnated cloth. Our skin cleansing protocols were developed to simulate actual patient bathing practices at our institutions.[7] Other studies of patient bathing with 4% CHG liquid have used various approaches, including different dwell times[9,12] and CHG dilutions.[8,25,28] The optimal method of 4% liquid CHG bathing remains to be determined.

Our study has limitations. First, we measured CHG concentrations and bacterial skin densities up to 6 hours from time of CHG cleansing (partly due to availability of healthcare professionals in an average shift), yet patients in most hospitals are bathed only once daily. We do not know whether the differences in CHG skin concentration and microbial densities observed for the different bathing methods persisted longer than 6 hours. In an earlier study, we found that reductions in microbial bioburden on the skin of ICU patients after cleansing with no-rinse 2% CHG-impregnated cloths persisted for up to 24 hours.[15] Second, we elected to study healthy healthcare personnel because they comprise a more homogenous population than do hospital patients. Skin microbial communities of healthy volunteers are likely different from skin microbial communities of hospital patients; the latter group may be more prone to harboring MDROs with high CHG minimum inhibitory concentrations (MICs).[21,29] Although residual CHG concentrations in our study were greater than CHG MICs reported for many MDROs,[20] additional studies of hospital patients are needed to determine whether the relative differences in CHG concentration and bacterial densities that we observed are present over a longer time. Finally, the relation between the outcomes we studied— CHG concentration and microbial density on skin—and clinically relevant outcomes such as bacteremia is unknown.

In conclusion, our findings using a randomized study design with blinded assessment demonstrated that cleansing with no-rinse 2% CHG-impregnated cloths yielded significantly higher residual skin concentrations and lower bacterial density in healthy volunteers for at least 6 hours after application compared to cleansing with 4% CHG liquid using 2 alternative cloth delivery vehicles, followed by rinsing. Reasons for poorer performance of 4% CHG liquid bathing methods in our study are likely multifactorial, including rinsing after cleansing, cloth material, and lower absolute quantity of CHG applied. Clinical studies that compare standardized CHG formulations and bathing techniques are needed to evaluate the effect of different CHG bathing methods on patient outcomes.

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