Staphylococcus aureus Skin and Soft Tissue Infections at a Tertiary Hospital in Botswana

Hong Truong
Samir S. Shah
University of Pennsylvania

Johnathan Ludmir
University of Pennsylvania

Ephraim O. Tawanana

Margaret Bafana

See next page for additional authors

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Recommended Citation
Truong, Hong; Shah, Samir S.; Ludmir, Johnathan; Tawanana, Ephraim O.; Bafana, Margaret; Wood, Sarah M.; Moffat, Howard; and Steenhoff, Andrew P., "Staphylococcus aureus Skin and Soft Tissue Infections at a Tertiary Hospital in Botswana" (2011). Botswana-UPenn Scholarly Publications. Paper 6.
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Abstract

Objectives. To study the epidemiology of *Staphylococcus aureus* skin and soft-tissue infections (SSTIs) in hospitalised children and adults in Gaborone, Botswana, and to describe the changes in antimicrobial susceptibilities of *S. aureus* isolates over time.

Methods. A retrospective cohort study evaluated SSTI isolates from January 2000 to December 2007 at Princess Marina Hospital (PMH), a large tertiary referral centre in Gaborone. Eligible subjects were those hospitalised at PMH during the study period who had a skin or soft-tissue culture yielding a bacterial or fungal pathogen. The primary outcome measure was a skin or soft-tissue culture yielding *S. aureus*. Secondary outcomes were the organism’s antimicrobial susceptibilities.

Results. *S. aureus* was detected in 857 (35.8%) of single-organism SSTI cultures, and 194 (22.6%) of these isolates were methicillin resistant (MRSA). The proportion of MRSA isolates increased over time (linear test of trend: *p*=0.03 from 2000 to 2003), and MRSA isolates were more likely than methicillin-susceptible isolates to be resistant to commonly used antimicrobials recommended by the national SSTI treatment guideline.

Conclusions. We report a high and increasing proportion of MRSA SSTIs in Gaborone. This high rate of MRSA resistance to currently recommended empiric antibiotics for SSTIs dictates the need for revising national guidelines and ongoing prospective surveillance of SSTIs in this setting.

Keywords

staphylococcus aureus, infection, Botswana

Disciplines

Bacterial Infections and Mycoses | Skin and Connective Tissue Diseases

Author(s)

Hong Truong, Samir S. Shah, Johnathan Ludmir, Ephraim O. Tawanana, Margaret Bafana, Sarah M. Wood, Howard Moffat, and Andrew P. Steenhoff

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Over the past decades, methicillin-resistant *Staphylococcus aureus* (MRSA) has become a key pathogen in skin and soft-tissue infections (SSTIs). The emergence of MRSA in 1961 occurred predominantly in patients exposed to health care facilities; community-acquired MRSA (CA-MRSA) isolates were identified in the 1990s. From 1993 to 2005, the impact of CA-MRSA nearly tripled emergency department SSTI visits in the USA. However, there are limited data on the incidence and characteristics of SSTIs in sub-Saharan Africa, and knowledge of pathogen-specific attributable morbidity is lacking.

HIV-infected patients are at markedly increased risk for CA-MRSA SSTIs in hospitalised children and adults in Botswana’s capital city of Gaborone over an 8-year period. The secondary objective was to describe temporal changes in antimicrobial susceptibilities of *S. aureus* isolates over this period.

Method

This retrospective cohort study evaluated SSTI isolates from January 2000 to December 2007 at Princess Marina Hospital (PMH), a large tertiary referral centre in Gaborone. Eligible subjects were those hospitalised at PMH during the study period who had a skin or soft-tissue culture yielding a bacterial or fungal pathogen. The primary outcome measure was a skin or soft-tissue culture yielding *S. aureus*. Secondary outcomes were the organism’s antimicrobial susceptibilities.

Results. *S. aureus* was detected in 857 (35.8%) of single-organism SSTI cultures, and 194 (22.6%) of these isolates were methicillin-resistant (MRSA). The proportion of MRSA isolates increased over time (linear test of trend: \( p=0.03 \) from 2000 to 2003), and MRSA isolates were more likely than methicillin-susceptible isolates to be resistant to commonly used antimicrobials recommended by the national SSTI treatment guideline.

Conclusions. We report a high and increasing proportion of MRSA SSTIs in Gaborone. This high rate of MRSA resistance to currently recommended empiric antibiotics for SSTIs dictates the need for revising national guidelines and ongoing prospective surveillance of SSTIs in this setting.

Gaborone, Botswana. PMH is the largest tertiary care referral centre in Botswana, with 330 beds and 20 000 - 30 000 admissions annually. PMH provides virtually free health care to citizens of Botswana and charges a subsidised fee to non-citizens. Since January 2002, Botswana as a whole, and PMH in particular, have provided free highly active antiretroviral therapy (HAART) to citizens who are HIV positive and who meet specified clinical criteria. All patients hospitalised at PMH during the study period who had a skin or soft-tissue culture yielding a bacterial or fungal pathogen were included.

SSTI was defined as a culture of non-blood, non-stool or non-cerebrospinal fluid specimens that yielded a single bacterial or fungal pathogen. Most specimens were collected from pus, wound sites, eye, ear, throat, vaginal swabs, pleural fluid, pericardial fluid, joint fluid or brain abscesses. Polymicrobial skin and soft-tissue cultures were considered contaminants and were not included. Multiple cultures from an individual patient within 7 days of the original culture were considered part of the same episode of SSTI and therefore excluded. Clinical wards at PMH were categorised by three parameters: the clinical nature of the ward (medical, surgical, intensive care unit, accident and emergency, orthopaedic, oncology, ophthalmology, radiology); gender (male v. female medical or orthopaedic wards); and age of patients (neonatal, paediatric or adult).

Skin and soft-tissue cultures were subjected to microscopy, culture and sensitivity testing. Antimicrobial susceptibility patterns of isolates were determined by disc-diffusion methods according to Clinical Laboratory Standards Institute guidelines. Antimicrobial agents used were amikacin, ampicillin, amoxicillin–clavulanate, cefotaxime, ceftazidime, cephradine, chloramphenicol, ciprofloxacin, clindamycin, erythromycin, gentamicin, methicillin, penicillin G, piperacillin, tetracycline, trimethoprim, trimethoprim–sulfamethoxazole (TMP-SMX) and vancomycin. The D test for inducible clindamycin resistance was not available. Organisms that were resistant to all of the following were classified as multidrug-resistant organisms: ampicillin, amoxicillin–clavulanate, cefotaxime, gentamicin, tetracycline, trimethoprim and TMP-SMX. The proportion of resistance among *S. aureus* isolates was calculated by dividing the number of resistant isolates by the total number of isolates.

Electronic and manual records of the Botswana National Health Laboratory were reviewed to identify skin and soft-tissue samples that met the study criteria. Demographic data were abstracted from the records, including anatomical site of sample, age, gender, and the hospital ward at the time of specimen collection. Results of antimicrobial susceptibility testing were recorded. Owing to laboratory personnel changes, transition from manual to electronic records and hospital ward at the time of specimen collection. Results of antimicrobial susceptibility testing were recorded. Owing to laboratory personnel changes, transition from manual to electronic medical records and periodic shortages of antibiotics, there were missing laboratory records in 39 of the 96 months (41%) of the study period – May 2000 - July 2001, December 2002 - June 2003, April 2004 – February 2005, June 2005 - September 2005, and November 2005 - December 2005.

Data were analysed using STATA version 11.0 (Stata Corp., College Station, TX). Continuous variables such as age were described using mean, median, and range or interquartile range (IQR) values and compared using the Wilcoxon rank-sum test. Categorical variables were described using counts and percentages and compared using the chi-square test. We used univariable logistic regression to determine the association of individual covariates with MRSA infection and a linear test of trend to identify changes in MRSA prevalence over time.

**Results**

During the study period, 2 979 (31.6%) of 9 428 cultures from SSTIs were positive. A single organism was isolated from 2 404 (80.7%) of the positive cultures. *S. aureus* was detected in 857 (35.8%) of the single-organism cultures.

Table 1 describes the demographic characteristics of patients with *S. aureus* infections. MRSA accounted for 194 (22.6%) of *S. aureus* isolates. Of patients with an MRSA SSTI, 55.1% were male. Patients 13 - 49 years of age accounted for almost two-thirds of cases (Table 1). The most common sites of *S. aureus* infection or colonisation were vaginal swabs (N=53), wounds (N=42) and eyes (N=36). However, in the vast majority of *S. aureus* isolates the anatomical site was unspecified ‘pus’ and ‘unspecified fluid, yielding 2 083 (86.6%) and 60 (2.5%) of isolates, respectively.

The male orthopaedic ward had the highest rate of MRSA (42 cases – 20.6% of MRSA SSTIs). Patients in the male medical ward (odds ratio (OR) 4.5, 95% confidence interval (CI) 1.8 - 18.2, p=0.01) were at increased odds of having a MRSA SSTI compared with SSTIs in the neonatal unit. Fig. 1 displays the annual prevalence of MRSA. The lowest proportion of MRSA SSTIs was in 2000 (15.9%) and the highest proportion was in 2003 (27%; linear regression p=0.03 from 2000 to 2003). Although there was an increase in the proportion of MRSA from 2000 to 2003, there were no further increases from 2003 to 2007 (p=0.48).

*S. aureus* isolates were routinely tested for susceptibility to amoxicillin–clavulanate, cephradine, chloramphenicol, clindamycin, erythromycin, gentamicin, methicillin, penicillin, piperacillin, tetracycline and vancomycin. Susceptibility to TMP-SMX was not routinely tested; results were available for only 18 isolates (Fig. 2). *S. aureus* was most susceptible to vancomycin, clindamycin and chloramphenicol, with susceptibility rates of 99.5%, 94.5% and 82.5%, respectively. No chloramphenicol susceptibility data were available for 2000.

The antibiotic susceptibilities of MRSA and MSSA are compared in Fig. 3. MRSA isolates were more likely to be resistant to clindamycin (OR 26.1, 95% CI 15.1 - 46.2, p<0.001), erythromycin (OR 25.5, CI 15.8 - 41.3, p<0.001) and tetracycline (OR 31.0, 95% CI 17.8 - 55.9, p<0.001), compared with MSSA isolates.

**Discussion**

The epidemiology and drug susceptibility of *S. aureus* SSTIs in Botswana are described. There is a high proportion of *S. aureus* SSTIs, particularly MRSA, at the tertiary hospital that serves the population of southern Botswana. Significant factors that increase the odds of MRSA infection among our cohort include age group 13 - 49 years, and hospitalisation in the intensive care unit, male orthopaedic and male medical wards.

An increasing proportion of MRSA SSTIs in Botswana between 2000 and 2003 and high levels of MRSA resistance to commonly used antibiotics are revealed. There is considerable variation in MRSA resistance profiles between regions and countries. In Australia, Singapore and Taiwan, MRSA is predominantly resistant to ciprofloxacin, clindamycin, erythromycin, gentamicin, TMP-SMX and tetracycline. In KwaZulu-Natal, South Africa, MRSA is resistant to clindamycin, erythromycin and TMP-SMX. Similarly, MRSA cases in this Botswana study displayed high resistance rates to clindamycin, erythromycin and tetracycline. This susceptibility pattern is of concern because patients with SSTIs in Botswana are routinely treated with one of these medications or penicillin. The empiric prescription of penicillin is probably targeting infection by *Streptococcus* group A, but our findings suggest that *S. aureus* is an under-recognised cause of SSTIs in Botswana.

Despite the Botswana Ministry of Health’s recommendation to administer co-trimoxazole as the drug of choice for cutaneous bacterial infections, which is supported by current US guidelines...
advocating TMP-SMX as the first-line therapy against community-acquired MRSA, *S. aureus* isolate susceptibilities to TMP-SMX were not routinely tested. Additionally, we found that vancomycin, the most effective antibiotic against MRSA, is not readily available at health centres other than the main hospital. Hence, we propose four improvements in managing SSTIs in Botswana: (i) revision of Botswana's national SSTI guidelines, including discontinuing advocating penicillin as empiric treatment for SSTIs in a setting where *S. aureus* accounts for more than a third of culture-positive SSTIs and virtually all of these *S. aureus* isolates are resistant to this class of antibiotics (this has been achieved, as Botswana’s national antibiotic guidelines, informed by information from this study, were revised in July 2010); (ii) TMP-SMX susceptibility testing of all *S. aureus* SSTI isolates; (iii) improved hospital infection control guidelines and practices to prevent MRSA transmission in the hospital setting; and (iv) vancomycin availability in district hospitals with strict guidelines governing its use.

Limitations of this study include: (i) missing laboratory data from our study period may have contributed to an underestimate of SSTI...
prevalence; (ii) the anatomical site was not known for most S. aureus isolates, making it impossible to define the exact nature of each SSTI; (iii) owing to the retrospective study design, standard laboratory techniques may have shifted over the 8-year study period; and (iv) our retrospective design makes our results susceptible to potential biases of studies of similar design.

In summary, we found a high and increasing proportion of MRSA SSTIs in Gaborone. Additionally, the high rate of MRSA resistance to currently recommended empiric antibiotics for SSTIs dictates the need for revising national guidelines and ongoing prospective surveillance of SSTIs in this setting.

We thank the Botswana Ministry of Health, Dr Harvey Friedman, Dr Rameshwari Thakur, Jack Rowe, the staff of the microbiology lab in the Botswana National Health Laboratory, Rudo Nthobatsang, Katherine Steele, Gill Jones and Nikki Jones, who made this project possible by providing invaluable guidance and/or assisting with the logistics.

Financial support. This project was supported in part by the University of Pennsylvania Center for AIDS Research IP30AI45008-01 to APS.

Conflict of interest. The contributing authors of this publication have no relationship or support that may constitute a conflict of interest.

References

Accepted 8 February 2011.

Fig. 3. Comparative susceptibility profiles of MRSA and MSSA.

### I love to watch the busy EGRET
(both the she-gret and the he-gret)
But what a way to earn your keep.
- Picking ticks off cows and sheep!

The egret has a graceful torso
And the HERON even more so.
See him, stone-still, in the creek,
S-shaped neck and dagger beak;
All-of-a-sudden with a swish!
Look – the heron has a fish!

Wheeling high up in the sky
The SEAGULL flies with raucous cry.
When underneath, you should be careful,
Or you’ll land up with a hairful.

I wish the awkward HA-DI-DAH
Would learn to whistle, tweet or baa.
A brownish bird, with fearsome beak
She calls her mate with rasping shriek
But someone loves the ha-di-dah
It is of course the ha-di-ma.

I’ve the greatest respect for the PLOVER
He is always so true to his lover
Her eggs he will shield
So exposed in the field
And he’ll never run off with anoher.

Maurice Kibel Children’s poems