Washington University School of Medicine

Digital Commons@Becker

Open Access Publications

2020

Clinical implications of restrictions in criteria for defining surgical site infections after mastectomy

Antoinette A A Bediako-Bowan

Washington University School of Medicine in St. Louis

David K Warren

Washington University School of Medicine in St. Louis

Katelin B Nickel

Washington University School of Medicine in St. Louis

Victoria J Fraser

Washington University School of Medicine in St. Louis

Margaret A Olsen

Washington University School of Medicine in St. Louis

Follow this and additional works at: https://digitalcommons.wustl.edu/open_access_pubs

Please let us know how this document benefits you.

Recommended Citation

Bediako-Bowan, Antoinette A A; Warren, David K; Nickel, Katelin B; Fraser, Victoria J; and Olsen, Margaret A, "Clinical implications of restrictions in criteria for defining surgical site infections after mastectomy." *Infection Control & Hospital Epidemiology*. 1 - 3. (2020).
https://digitalcommons.wustl.edu/open_access_pubs/10001

This Open Access Publication is brought to you for free and open access by Digital Commons@Becker. It has been accepted for inclusion in Open Access Publications by an authorized administrator of Digital Commons@Becker. For more information, please contact vanam@wustl.edu.

Concise Communication

Clinical implications of restrictions in criteria for defining surgical site infections after mastectomy

Antoinette A. A. Bediako-Bowan MBChB^{1,2} , David K. Warren MD, MPH¹ , Katelin B. Nickel MPH¹ ,
Victoria J. Fraser MD¹ and Margaret A. Olsen PhD, MPH^{1,3}

¹Division of Infectious Diseases, Washington University School of Medicine, St Louis, Missouri, United States, ²Department of Surgery, University of Ghana, Accra, Ghana and ³Division of Public Health Sciences, Washington University School of Medicine, St Louis, Missouri, United States

Abstract

More than 50% of women with clinically apparent infection after mastectomy did not meet the 2020 National Healthcare Safety Network (NHSN) definition for surgical site infection (SSI). Implant loss was similar whether the 2020 NHSN SSI definition was met or not, suggesting equivalent adverse outcomes regardless of restriction to the surveillance definition.

(Received 10 June 2020; accepted 18 October 2020)

Surgical site infections (SSIs) are the most common healthcare-associated infection in the United States.¹ Accurate SSI surveillance and feedback to surgeons is essential to devise and implement strategies to prevent postoperative infections. The Centers for Disease Control and Prevention (CDC) National Healthcare Safety Network (NHSN) definitions for deep-incisional and organ-space SSIs have changed over time. For example, in 2013 the surveillance period for deep and organ space infections was reduced from 1 year to 90 days for procedures including implants.^{2,3} We sought to determine the impact of underreporting infections after mastectomy using the 2020 NHSN SSI definition.

Methods

We conducted a retrospective cohort study using electronic health record and billing data from 1 academic hospital and 1 community hospital. We identified mastectomy admissions among women aged ≥ 18 years from July 1, 2010 to June 30, 2015, using the *International Classification of Diseases, 9th Revision* (ICD-9) procedure codes (Supplementary Table 1 online) and verified mastectomy by surgeon description.

Potential SSIs were identified by selecting records to review based on performance of a microbiology culture or diagnosis or procedure code suggestive of a wound complication within 180 days after mastectomy (Supplementary Table 1 online). Infections were verified by review of outpatient and hospital records for signs and symptoms of infection, intervening procedures, expansion history, and microbiology data.

We defined clinically apparent infections as infections that met the pre-2013 NHSN definition or documented signs consistent with SSI (eg, cellulitis necessitating implant removal). The 180-day SSI rates were calculated using the pre-2013, 2020

NHSN, and clinical definitions (Supplementary Table 2 online). The Fisher's exact test was used to compare implant removal rates and physician documentation of infection using SAS version 9.4 software (SAS Institute, Cary, NC). The study was approved by the Washington University Human Research Protection Office with a waiver of informed consent.

Results

In total, 1,902 women underwent mastectomy, of whom 148 developed clinically apparent infection at the surgical site within 180 days after operation. Infections in 69 women (46.6%) met the 2020 NHSN SSI criteria. Infections in 102 women (68.9%) met the pre-2013 NHSN criteria, due to 33 infections after implant reconstruction meeting the earlier SSI criteria (Table 1).

Of the 148 women with clinically apparent infections, 23 (15.5%) underwent mastectomy only and 125 (84.5%) underwent immediate reconstruction. Moreover, 100 women (67.6%) had placement of tissue expander(s), 10 (6.8%) had permanent implant(s), and 15 (10.1%) underwent autologous flap reconstruction.

The reasons for exclusion of clinically apparent infections based on 2020 NHSN criteria (not mutually exclusive) included diagnosis of a superficial incisional SSI >30 days after mastectomy ($n = 22$), diagnosis of a deep incisional or organ-space SSI >90 days after mastectomy ($n = 19$; 2 deep incisional, 17 organ-space), manipulation of the surgical site after mastectomy in the absence of signs/symptoms of infection [ie, surgical debridement ($n = 18$), needle aspiration of seroma ($n = 9$), tissue expander access ($n = 40$)], and negative intraoperative cultures ($n = 14$) (Table 2). Methicillin-sensitive *Staphylococcus aureus* was the most commonly isolated etiologic organism, regardless of the onset timing of infection (Supplementary Table 3 online).

Tissue expanders were accessed in 40 women before the onset of clinically apparent infection, a median of 2 times (range, 1–7), resulting in exclusion of 13 infections (32.5%) using the 2020

Author for correspondence: Margaret A. Olsen, E-mail: molsen@wustl.edu

Cite this article: Bediako-Bowan AAA, et al. (2020). Clinical implications of restrictions in criteria for defining surgical site infections after mastectomy. *Infection Control & Hospital Epidemiology*, <https://doi.org/10.1017/ice.2020.1288>

Table 1. Impact of Changes in the National Healthcare Safety Network Surgical Site Infection (SSI) Definitions on the Number and Rate of Infections After Mastectomy

Procedure	Total Surgeries	Total No. of Clinically Apparent Infections (n=148) ^a	No. of SSI Meeting the Pre-2013 NHSN Definition (n=103)	No. of SSI meeting the 2020 NHSN definition, (n=69), No. (No. Identified By Surgeons)	Total No. of Clinically Apparent Infections Excluded by 2020 NHSN Definition (n=79), No. (No. identified by surgeons)	SSI Rate by Clinically Apparent Infections	SSI Rate by Pre-2013 NHSN Definition	SSI Rate by 2020 NHSN Definition
Mastectomy only	683	23	12	12 (11)	11 (9)	3.37	1.76	1.76
Mastectomy + implant reconstruction	1,122	110	86	53 (46)	57 (48)	9.80	7.66	4.72
Mastectomy + flap reconstruction	97	15	4	4 (3)	11 (7)	15.46	4.12	4.12

^aClinically apparent infections at the surgical site includes infections that met the pre-2013 NHSN definition, plus any of the following within 180 d after surgery: infections with onset after a prior diagnostic/therapeutic procedure, superficial infections >30 d, deep-incisional and organ-space infections >90 d with no implant, physician opened the deep wound in the presence of signs/symptoms of infection (eg, erythema, warmth), or the physician described the wound as infected, despite negative culture results.

Table 2. Reasons for Not Meeting the 2020 National Healthcare Safety Network Definition of Surgical Site Infection (SSI)

Reason for Not Meeting Criteria ^a	Mastectomy Only (n=11)	Mastectomy + Implant Reconstruction (n=57)	Mastectomy + Flap Reconstruction (n=11)	Total No. Of Patients With Signs Suggesting Surgical Site Infection ^a (n=79)
Superficial incisional SSI diagnosed >30 d postoperatively	10	5	7	22
Deep incisional or organ space SSI diagnosed > 90 d postoperatively	0	18	1	19
Prior procedure in the absence of signs/symptoms of infection				
Debridement of surgical site	1	12	5	18
Needle aspiration of breast	2	7	0	9
Accession of breast expanders	N/A	40	N/A	40
Culture negative microbiology from surgical site	0	14	0	14

^aAn infection may have been excluded for >1 of these reasons.

NHSN criteria. Expanders were last accessed a median 16 days (range, 5–132) before infection onset. Overall, 9 women had 1–3 documented needle aspirations of a seroma, with the last aspiration a median of 11 days (range, 1–48) before infection onset. Also, 18 women had debridement of necrotic skin flaps a median of 13.5 days (range, 3–111) before infection onset.

Of the 14 women who failed to meet the NHSN criteria for SSI due to a negative intraoperative culture, all had been treated with antibiotics (median, 3 days) in the 2 weeks prior to implant removal. Prior antibiotics before the negative intraoperative cultures included intravenous vancomycin, piperacillin/tazobactam, and/or clindamycin (n = 8 women), or oral clindamycin, doxycycline, cephalexin, and/or sulfamethoxazole-trimethoprim (n = 6).

Of the 33 women with infection after implant reconstruction that met the pre-2013 but not the 2020 NHSN SSI definition, 13 had an organ-space SSI >90 days after mastectomy, and 7 met the pre-2013 definition of deep-incisional or organ-space SSI (3 deep incisional and 4 organ space) solely due to surgeon diagnosis at the time of implant removal.

Of the 110 women who had expander or permanent implant reconstruction, 94 (85.5%) had an infection at the surgical site diagnosed by a surgeon and/or infectious diseases physician. The percentage of women with physician-documented infection was the same in women whose infection met the 2020 NHSN SSI criteria (86.8%, 46 of 53) or whose infection did not meet 2020 criteria (84.2%, 48 of 57) ($P = .790$).

Overall, 102 of 110 women (92.7%) who developed clinically apparent infection following immediate implant reconstruction had their implants removed. These rates were 92.5% (49 of 53) versus 93.0% (53 of 57) among those who met versus did not meet the 2020 NHSN SSI criteria ($P = 1.0$).

Discussion

We reviewed records of women after mastectomy with or without reconstruction and found that 53% of women with clinical infection did not meet the 2020 NHSN SSI definition. The number of infections after implant reconstruction that met NHSN criteria

decreased substantially after the criteria changes, which restricted the surveillance period in 2013 and definitions of deep-incisional and organ-space SSIs in 2014 and excluded infections after access of tissue expanders in 2017. The differential impact of the definition changes on implant reconstruction SSIs is important because immediate reconstruction has steadily increased during the past 20 years and tissue expanders account for the majority of reconstruction procedures.⁴ The SSI rate after mastectomy is typically at least twice as high with versus without reconstruction,⁵ which is obfuscated by changes to the surveillance definition that have differential impact on postimplant infections.

Manipulation of the surgical site post-mastectomy may contribute to the risk of infection, depending on the frequency and adherence to asepsis during manipulation. This is especially true for breast tissue expander reconstruction, since the port is accessed frequently for saline injection during expansion. Surveillance with feedback of all infections involving the surgical site post-mastectomy may alert surgeons and infection prevention specialists of suboptimal practices, particularly regarding tissue expansion, which could be targeted for infection prevention.^{6,7}

In our cohort, systemic antibiotic therapy was administered in women with implant reconstruction who presented with cellulitis, in hopes of salvaging the implant.⁸ This likely led to negative intra-operative cultures at the time of implant removal in the 14 women who met the clinical but not 2020 NHSN SSI definition. Most of these infections met the pre-2013 NHSN definition due to physician diagnosis. Re-evaluation of physician diagnosis to define SSI may need to be considered given the frequent empiric antibiotic treatment in this population.

We have demonstrated that implant loss was virtually the same in women with clinically diagnosed SSIs, regardless of whether the 2020 NHSN definition was met or not. Implant loss is important from a patient perspective because it results in additional morbidity, procedures, hospital costs, and delays in completion of adjuvant therapy.⁹

Our study of 2 academically affiliated hospitals may not reflect all community practices. The substantial changes in the NHSN SSI definitions in the past decade^{2,3,10} have resulted in underreporting of potentially preventable infections and underestimation of infectious morbidity after mastectomy. Comprehensive infection surveillance, particularly after breast implant reconstruction, is essential to provide women with accurate information about complication risks and to determine the need for additional infection prevention strategies.

Supplementary material. To view supplementary material for this article, please visit <https://doi.org/10.1017/ice.2020.1288>

Acknowledgments.

Financial support. Funding for this project was provided by the Centers for Disease Control and Prevention (grant no. U54CK000482 to V.J.F.).

Conflicts of Interest. M.A.O. reports consultant work with Pfizer and grant funding through Pfizer, Merck, and Sanofi Pasteur for work outside the submitted manuscript. V.J.F. reports her spouse is the Chief Clinical Officer at Cigna Corporation. D.K.W. reports consultant work with Centene, PDI, Pursuit Vascular, Homburg & Partner, and Carefusion/BD and as a subinvestigator for a Pfizer-sponsored study for work outside the submitted manuscript. All other authors report no conflicts of interest relevant to this article.

References

1. Magill SS, Edwards JR, Bamberg W, *et al.* Multistate point-prevalence survey of health care-associated infections. *N Engl J Med* 2014;370:1198–1208.
2. Mangram AJ, Horan TC, Pearson ML, Silver LC, Jarvis WR. Guideline for prevention of surgical site infection, 1999. Centers for Disease Control and Prevention (CDC) Hospital Infection Control Practices Advisory Committee. *Am J Infect Control* 1999;27:97–132.
3. National Healthcare Safety Network (NHSN) procedure-associated (PA) module: surgical site infection (SSI) event. Centers for Disease Control and Prevention website. <http://www.cdc.gov/nhsn/PDFs/pscManual/9pscSSICurrent.pdf>. Published 2019. Accessed June 17, 2019.
4. Nayyar A, Reddy KG, Strassle PD, *et al.* Temporal trends in immediate post-mastectomy breast reconstruction. *J Plast Reconstr Aesthet Surg* 2019; 72:513–527.
5. Olsen MA, Nickel KB, Fox IK, *et al.* Incidence of surgical site infection following mastectomy with and without immediate reconstruction using private insurer claims data. *Infect Control Hosp Epidemiol* 2015;36:907–914.
6. Sharara SL, Saunders HM, Fabre V, *et al.* Infection surveillance and prevention strategies to detect and prevent postaccess breast tissue expander infections. *Infect Control Hosp Epidemiol* 2019. doi: [10.1017/ice.2019.222](https://doi.org/10.1017/ice.2019.222).
7. Kraenzlin FS, Saunders H, Aliu O, *et al.* Classification of breast tissue expander infections: back to the basics. *J Surg Oncol* 2019;120:142–147.
8. Reish RG, Damjanovic B, Austen WG Jr, *et al.* Infection following implant-based reconstruction in 1952 consecutive breast reconstructions: salvage rates and predictors of success. *Plast Reconstr Surg* 2013;131:1223–1230.
9. Olsen MA, Nickel KB, Fox IK, Margenthaler JA, Wallace AE, Fraser VJ. Comparison of wound complications after immediate, delayed, and secondary breast reconstruction procedures. *JAMA Surg* 2017:e172338.
10. Dicks KV, Lewis SS, Durkin MJ, *et al.* Surveying the surveillance: surgical site infections excluded by the January 2013 updated surveillance definitions. *Infect Control Hosp Epidemiol* 2014;35:570–573.