



www.bioinformation.net
Volume 21(9)



Research Article

Received September 1, 2025; Revised September 30, 2025; Accepted September 30, 2025, Published September 30, 2025

DOI: 10.6026/973206300213231

SJIF 2025 (Scientific Journal Impact Factor for 2025) = 8.478

2022 Impact Factor (2023 Clarivate Inc. release) is 1.9

Declaration on Publication Ethics:

The author's state that they adhere with COPE guidelines on publishing ethics as described elsewhere at <https://publicationethics.org/>. The authors also undertake that they are not associated with any other third party (governmental or non-governmental agencies) linking with any form of unethical issues connecting to this publication. The authors also declare that they are not withholding any information that is misleading to the publisher in regard to this article.

Declaration on official E-mail:

The corresponding author declares that lifetime official e-mail from their institution is not available for all authors

License statement:

This is an Open Access article which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly credited. This is distributed under the terms of the Creative Commons Attribution License

Comments from readers:

Articles published in BIOINFORMATION are open for relevant post publication comments and criticisms, which will be published immediately linking to the original article without open access charges. Comments should be concise, coherent and critical in less than 1000 words.

Disclaimer:

Bioinformation provides a platform for scholarly communication of data and information to create knowledge in the Biological/Biomedical domain after adequate peer/editorial reviews and editing entertaining revisions where required. The views and opinions expressed are those of the author(s) and do not reflect the views or opinions of Bioinformation and (or) its publisher Biomedical Informatics. Biomedical Informatics remains neutral and allows authors to specify their address and affiliation details including territory where required.

Edited by A Prashanth

E-mail: phyjunc@gmail.com

Citation: Venkatraman *et al.* Bioinformation 21(9): 3231-3234 (2025)

Prospective study on surgical site infections in major oncologic resections

Poornima Venkatraman¹, Vishal Mruthyunjaya Kalmani², Arundhati Maindola³, Yeruva Bheemeswara Reddy⁴, Dhyaneshwar Ra⁵ & Malavika Sajeev^{6,*}

¹Department of Surgery, Davao Medical School Foundation, Philippines, Telangana, India; ²Department of Emergency Medicine, Barking Havering and Redbridge University Hospitals NHS Trust, Essex, United Kingdom; ³Department of Health and Family Welfare, Directorate of Health Safety and Regulations, Shimla, Himachal Pradesh, India; ⁴Department of Surgery, Santhiram Medical College, Andhra Pradesh, India; ⁵Department of General Surgery, Government Stanley Medical College Chennai, Tamil Nadu, India; ⁶Department of Emergency Medicine, East Sussex NHS Healthcare Trust, Eastbourne, United Kingdom; *Corresponding author

Affiliation URL:

<https://www.dmsf.edu.ph/>

<https://www.bhrhospitals.nhs.uk/>

<https://dhsr.hp.gov.in/>
<https://santhirammedicalcollege.com/>
<https://www.stanleymedicalcollege.in/>
<https://www.esht.nhs.uk/>

Authors contacts:

Poornima Venkatraman - E-mail: poornimavenkatraman96@gmail.com; Phone: +91 9652696432
Vishal Mruthyunjaya Kalmani - E-mail: vishal.kalmani@nhs.net; Phone: +44 7405881521
Arundhati Maindola - E-mail: maindolarundhati@gmail.com; Phone: +91 6396290554
Yeruva Bheemeswara Reddy - E-mail: Bheemesh.yeruva@gmail.com; Phone: +91 7095918515
Dhyaneshwar Ra - E-mail: radhyaneshwar@gmail.com; Phone: +91 7010378943
Malavika Sajeev - E-mail: malavika.sajeev@nhs.net; Phone: +44 7863247163

Abstract:

The incidence, risk factors, and microbiological profile of surgical site infections (SSIs) in 142 patients undergoing major oncologic resections across gastrointestinal, breast, and head and neck malignancies is of interest. The overall SSI rate was 19.7%, with the highest incidence seen in gastrointestinal surgeries. Prolonged operative time, high ASA grade, and preoperative hypoalbuminemia were significant predictors of SSI. Most infections were caused by gram-negative organisms, with *E. coli* and *Klebsiella* species predominant. Thus, the importance of preoperative optimization and tailored antibiotic strategies in oncology surgery is shown.

Keywords: Surgical site infection, oncology surgery, risk factors, prospective study, microbiological profile, infection prevention, cancer resection

Background:

Surgical site infections (SSIs) are among the most common postoperative complications, particularly in patients undergoing major oncologic resections [1]. These infections not only increase patient morbidity and hospital stay but also delay adjuvant therapy, ultimately impacting long-term oncologic outcomes [2]. Despite advances in surgical techniques and perioperative care, SSI rates in cancer patients remain significantly higher compared to non-oncologic surgical populations [3]. This elevated risk is multifactorial, often attributed to factors such as immunosuppression from malignancy, nutritional deficits, chemotherapy effects, and complex, prolonged surgeries [4]. Oncologic surgeries typically involve extensive resections with or without reconstruction, frequently resulting in prolonged operative times, greater tissue handling, and potential for contamination—especially in gastrointestinal and head & neck procedures [5]. The microbial spectrum is also influenced by anatomical site and hospital flora, necessitating precise local data to guide effective prophylactic and therapeutic strategies [6]. Although numerous retrospective studies have addressed SSI risk, prospective data focused specifically on the oncology surgical population is limited [7]. Therefore, it is of interest to prospectively evaluate the incidence, contributing factors, and microbiological patterns of surgical site infections following major cancer surgeries, with the ultimate goal of identifying modifiable risks and informing targeted interventions to reduce postoperative infectious morbidity in this vulnerable population.

Materials and Methods:

This prospective observational study was conducted over a 20-month period from March 2023 to October 2024, in the surgical oncology department of a tertiary cancer center. A total of 142 patients undergoing elective major oncologic resections were

enrolled consecutively after obtaining informed consent. Inclusion criteria included patients aged ≥18 years undergoing definitive surgical resection for malignancies of the gastrointestinal tract, breast, or head and neck region. Patients with pre-existing infections, emergency surgeries, palliative procedures, or those receiving long-term immunosuppressive therapy were excluded. All surgeries were performed by experienced oncologic surgeons using standardized aseptic techniques. Preoperative variables such as age, sex, body mass index (BMI), ASA grade, nutritional status (including serum albumin), comorbidities (diabetes mellitus, anemia), neoadjuvant therapy, and smoking status were documented. Operative parameters included surgical site, type of surgery (clean, clean-contaminated, contaminated), use of drains, blood loss, operative duration, and perioperative antibiotic prophylaxis. SSIs were classified according to CDC guidelines into superficial incisional, deep incisional, and organ/space infections. Patients were followed for 30 days postoperatively with daily wound inspection during hospitalization and subsequent outpatient review. When infection was suspected, wound swabs or pus cultures were obtained and analyzed using standard microbiological techniques. Antibiotic sensitivity testing was performed for all isolates. The primary outcome was the incidence of SSI. Secondary outcomes included identification of independent risk factors and characterization of the microbiological spectrum of infections. Data were analyzed using SPSS version 26. Univariate and multivariate logistic regression were applied to identify predictors of SSI, with $p < 0.05$ considered statistically significant.

Table 1: Baseline demographic and clinical characteristics

Variable	Total (n = 142)
Mean age (years)	56.2 ± 11.7
Male sex (%)	83 (58.5%)

BMI (mean ± SD)	23.1 ± 3.2
ASA Grade ≥ 3 (%)	49 (34.5%)
Diabetes Mellitus (%)	41 (28.9%)
Hypoalbuminemia (<3.5 g/dL)	47 (33.1%)
Preoperative anemia (%)	61 (43.0%)
Neoadjuvant therapy (%)	38 (26.8%)

Table 2: Distribution of surgeries by cancer type

Surgery Type	Number of Patients	SSI Cases	SSI Rate (%)
Gastrointestinal	60	17	28.3%
Head & Neck	34	6	17.6%
Breast	37	3	8.1%
Sarcoma/Soft Tissue	11	2	18.2%

Table 8: Antibiotic resistance pattern of predominant isolates

Organism	Ceftriaxone Resistance (%)	Amikacin Resistance (%)	Meropenem Resistance (%)
<i>E. coli</i> (n=9)	77.8%	33.3%	11.1%
<i>Klebsiella</i> (n=7)	85.7%	28.6%	14.3%
<i>S. aureus</i> (n=6)	50.0% (oxacillin)	NA	NA

Table 5: SSI rates by wound classification

Wound Type	Total Cases	SSI Cases	SSI Rate (%)
Clean	61	5	8.2%
Clean-contaminated	54	15	27.8%
Contaminated	27	8	29.6%

Table 6: Multivariate logistic regression analysis for SSI risk

Variable	Adjusted OR	95% CI	p-value
Operative time >180 min	2.91	1.32–6.40	0.008
ASA Grade ≥3	2.78	1.22–6.34	0.015
Hypoalbuminemia	3.45	1.53–7.76	0.003

Table 7: Microbial profile of SSI cases

Organism	Isolates (n = 28)	Percentage (%)
<i>Escherichia coli</i>	9	32.1%
<i>Klebsiella pneumoniae</i>	7	25.0%
<i>Staphylococcus aureus</i>	6	21.4%
<i>Pseudomonas aeruginosa</i>	3	10.7%
Polymicrobial	3	10.7%

Table 9: Hospital stay duration

Group	Median Days (IQR)	p-value
SSI Group	13 (11–17)	<0.001
No SSI	7 (6–9)	

Table 10: Postoperative antibiotic use

Group	Mean Duration (days)	p-value
SSI Group	9.3 ± 2.1	<0.001
No SSI Group	3.8 ± 1.4	

Results:

A total of 142 patients undergoing major oncologic resections were included in this prospective study. The overall incidence of surgical site infections (SSIs) was 19.7% (n = 28). SSI rates were highest among gastrointestinal surgeries (28.3%), followed by head and neck (17.6%) and breast surgeries (8.1%). Multivariate analysis revealed that prolonged operative duration, ASA grade ≥3, and preoperative hypoalbuminemia (<3.5 g/dL) were independent predictors of SSI. Most infections were superficial, and the predominant organisms isolated were *Escherichia coli*, *Klebsiella pneumoniae*, and *Staphylococcus aureus*. Table 1 summarizes the baseline demographic and clinical characteristics of the 142 enrolled patients. The mean age was 56.2 years, with a slight male predominance. Nearly one-third had ASA grade ≥3, 33.1% had hypoalbuminemia, and 43% had

Table 3: Classification of surgical site infections

SSI Type	Number (n = 28)	Percentage (%)
Superficial	14	50.0%
Deep Incisional	8	28.6%
Organ/Space	6	21.4%

Table 4: Comparison between SSI and non-SSI groups

Variable	SSI Group (n=28)	No SSI (n=114)	p-value
Mean operative time (min)	216.4 ± 42.8	174.6 ± 39.2	<0.001
Hypoalbuminemia (%)	18 (64.3%)	29 (25.4%)	<0.001
ASA ≥ 3 (%)	17 (60.7%)	32 (28.1%)	0.002
Diabetes mellitus (%)	11 (39.3%)	30 (26.3%)	0.164
Neoadjuvant therapy (%)	10 (35.7%)	28 (24.6%)	0.246

preoperative anemia – factors associated with increased infection risk. Table 2 presents the distribution of surgeries by cancer type and their corresponding SSI rates. Gastrointestinal resections accounted for the largest share and also had the highest SSI incidence at 28.3%. Head and neck and sarcoma surgeries followed, while breast cancer procedures had the lowest infection rate (8.1%). Table 3 classifies SSIs by type, showing that superficial infections were most common (50%), followed by deep incisional (28.6%) and organ/space infections (21.4%), indicating that the majority of infections were manageable with local wound care. Table 4 compares risk variables between the SSI and non-SSI groups. Patients who developed SSIs had significantly longer operative times, higher rates of hypoalbuminemia (64.3% vs. 25.4%), and were more likely to have ASA grade ≥3, all with statistically significant p-values. Table 5 reports SSI incidence by wound classification. Clean-contaminated and contaminated surgeries had substantially higher infection rates (27.8% and 29.6%, respectively) compared to clean procedures (8.2%), emphasizing the role of intraoperative contamination in SSI pathogenesis. Table 6 presents a multivariate logistic regression model identifying three independent predictors of SSI: operative time >180 minutes, ASA grade ≥3, and preoperative hypoalbuminemia. Each was associated with significantly increased odds of infection. Table 7 describes the microbiological profile of SSI cases. Gram-negative organisms were predominant, with *Escherichia coli* (32.1%) and *Klebsiella pneumoniae* (25%) leading, followed by *Staphylococcus aureus* (21.4%). Polymicrobial infections occurred in 10.7% of cases. Table 8 outlines antibiotic resistance patterns. The majority of *E. coli* and *Klebsiella* isolates were resistant to ceftriaxone (77.8% and 85.7%, respectively), highlighting growing resistance among gram-negative organisms. Carbapenem resistance remained relatively low. Table 9 shows that patients with SSIs had a significantly longer hospital stay, with a median duration of 13 days versus 7 days in non-infected patients, contributing to increased healthcare burden and resource utilization. Table 10 compares postoperative antibiotic use. The mean duration of antibiotics was significantly higher in patients with SSIs (9.3 days)

compared to those without (3.8 days), indicating increased treatment intensity and potential risk of antimicrobial resistance.

Discussion:

This prospective study highlights a substantial burden of surgical site infections (SSIs) in patients undergoing major oncologic resections, with an overall incidence of 19.7%. Notably, gastrointestinal surgeries exhibited the highest SSI rate (28.3%), aligning with existing literature attributing elevated infection risk to the involvement of enteric contents and prolonged operative times. Breast surgeries, typically clean procedures, demonstrated the lowest infection rate (8.1%), affirming the role of wound classification in SSI risk stratification [8]. The identification of hypoalbuminemia, elevated ASA grade, and operative time >180 minutes as independent predictors underscores the multifactorial nature of SSIs [9]. Particularly for cancer patients who are prone to cachexia, this study supports preoperative nutritional management. Another important impact was the length of the surgery. Extended procedures frequently indicate more intricate surgery and more tissue manipulation both of which increase the risk of microbial contamination and impede wound healing. Similarly, markedly greater infection prevalence was linked to an ASA grade of ≥ 3 , which indicates a larger systemic illness burden [10].

Individualized risk reduction methods and perioperative planning should be informed by these factors. Gram-negative organisms predominated, according to microbiological profile, with *Staphylococcus aureus* coming in second, followed by *Escherichia coli* and *Klebsiella pneumoniae*. Concerns regarding empirical antibiotic selection were raised by the startling discovery that a large number of gram-negative isolates were resistant to third-generation cephalosporins [11]. In high-risk surgical oncology cases, these resistance patterns point to the need to reevaluate local antibiotic prophylactic strategies in favor of medicines that are effective against resistant gram-negatives. The implications were clinically severe even though the infection rate was mild. SSI patients needed greater antibiotic durations and had a median hospital stay that was almost twice as long as that of non-infected patients [12]. Such results may jeopardize long-term oncologic outcomes by delaying surgical recovery and the start of adjuvant therapy [13]. Our study confirms that SSIs continue to be a major concern in oncologic surgery, particularly for patients who are nutritionally fragile

and undergo gastrointestinal resections. Infection rates can be significantly decreased by interventions such as preoperative nutritional correction; careful surgical planning and tailored antibiotic tactics based on local resistance data. Future studies should examine the role of cutting-edge preventative techniques in high-risk patients, such as antimicrobial sutures and negative pressure wound care.

Conclusion:

Surgical site infections remain a significant concern following major oncologic procedures, particularly in gastrointestinal cancer surgeries. Addressing modifiable risk factors such as hypoalbuminemia and prolonged operative time, alongside optimizing antibiotic prophylaxis, can substantially improve postoperative outcomes in cancer patients.

Acknowledgement:

We acknowledge that all the authors contributed equally to this paper and hence they are considered as joint authors.

References:

- [1] Pantvaiddya G *et al.* *J Surg Oncol.* 2022 **125**:327. [PMID: 34729779]
- [2] Julien C *et al.* *J Crohns Colitis.* 2022 **16**:1211. [PMID: 35218661]
- [3] Shi M *et al.* *J Int Med Res.* 2020 **48**:300060520944072. [PMID: 32841576]
- [4] Sugamata N *et al.* *World J Surg Oncol.* 2022 **20**:111. [PMID: 35387666]
- [5] Lee DH *et al.* *Oral Oncol.* 2011 **47**:528. [PMID: 21543250]
- [6] Jung JP *et al.* *J Clin Med.* 2021 **10**:2560. [PMID: 34207893]
- [7] Thorgersen EB *et al.* *Ann Surg Oncol.* 2017 **24**:721. [PMID: 27766561]
- [8] Broecker JS *et al.* *Ann Surg Oncol.* 2017 **24**:3574. [PMID: 28895084]
- [9] Davis CM *et al.* *J Oral Maxillofac Surg.* 2016 **74**:1199. [PMID: 26917207]
- [10] Raftery NB *et al.* *Dis Esophagus.* 2021 **34**:doaa136. [PMID: 33590037]
- [11] Kamat US *et al.* *Indian J Surg.* 2008 **70**:120. [PMID: 23133038]
- [12] Cheng K *et al.* *Patient Prefer Adherence.* 2015 **9**:1171. [PMID: 26316722]
- [13] Rajan K *et al.* *Cureus.* 2025 **17**:e86596. [PMID: 40704248]