Platelet to white blood cell ratio predicts 30-day postoperative infectious complications in patients undergoing radical nephrectomy for renal malignancy

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Abstract

Introduction: We sought to examine the relationship between preoperative platelet to white blood cell ratio (PLT/WBC), a hematological marker of the systemic inflammatory response, and postoperative infectious complications following radical nephrectomy for localized renal cell carcinoma.

Methods: We performed a retrospective cohort study of patients treated with radical nephrectomy for localized kidney cancer between January 1, 2005 and December 31, 2014 (n=6235) using the American College of Surgeons (ACS) National Surgical Quality Improvement Program (NSQIP) database. Univariate and multivariate analyses were used to assess the association between PLT/WBC ratio and 30-day infectious complications, including surgical site infection, urinary tract infection (UTI), pneumonia, and sepsis. Secondarily, we examined major complications and bleeding requiring transfusion.

Results: A lower PLT/WBC ratio was associated with an increased risk of sepsis, pneumonia, and UTI rates (p<0.05 for all). Furthermore, there was a significant trend of decreasing rates of sepsis and pneumonia with increasing PLT/WBC ratio across quintiles (p<0.05 for all). On multivariate analysis, patients with the lowest PLT/WBC ratios (Quintile 1) had a two-fold risk of having a postoperative infectious complication compared to patients in the highest quintile (odds ratio [OR] 2.01; 95% confidence interval [CI] 1.42–2.86; p<0.0001). Patients in Quintile 5 had a higher risk of requiring blood transfusion than those in Quintiles 2–4 (p<0.05 for all).

Conclusions: The PLT/WBC ratio represents a widely available and novel index to predict risk of infectious and bleeding complications in patients undergoing radical nephrectomy. External validation is required and the biological underpinning of this phenomenon requires further study.

Introduction

Renal cell carcinoma is the ninth most common cancer in the U.S. and its incidence has been increasing in developed countries. ^{1,2} The vast majority of patients present with localized disease. ^{1,2} Surgical extirpation, by radical or partial nephrectomy, remains the standard of care for these patients. ³ Postoperative complications and re-admission rates following radical nephrectomy have been reported to be as high as 30%. ^{4,5} We previously identified bleeding (requiring transfusion) and infectious complications to be the most common postoperative complications. ⁶ While patient comorbidities are known to predict postoperative complications, ^{7,8} the risk is difficult to quantify depending on number and type. A simple test that could quickly assess perioperative risk in the clinic would be useful in counselling patients.

The systemic inflammatory response may be involved in kidney cancer progression and kidney tumour cell biology, and may have utility as a prognostic biomarker. 9,10 These scoring systems are calculated using hematological tests, which are routinely collected preoperatively.1 While their usefulness in predicting prognosis in localized renal cell carcinoma has recently been demonstrated,11 the small number and heterogeneity of studies, as well as the lack of large multi-institutional data, limits wider adoption.1 The platelet to white blood cell count (PLT/WBC) has demonstrated prognostic ability for mortality and long-term outcomes following myocardial infarction, 12,13 but its association with perioperative outcomes has not been fully assessed. 14 The PLT/WBC ratio would be an ideal marker for perioperative outcomes, as it is easily calculated and routinely collected prior to major surgery.

We hypothesized that there may be a relationship between the systemic inflammatory response and postoperative infections. We sought to evaluate the association between preoperative PLT/WBC ratio and postoperative infectious complications among patients undergoing radical nephrectomy for localized kidney cancer. Secondarily, we examined the relationship between the preoperative PLT/WBC ratio and postoperative bleeding complications. In order to do so, we used the National Surgical Quality Improvement Program (NSQIP), a multi-institutional registry developed by the American College of Surgeons (ACS).¹⁵ Studies have found NSQIP to more accurately identify 30-day postoperative patient outcomes compared to single-centre or administrative data sources.¹⁶⁻¹⁸

Methods

Study design and population

We performed a retrospective cohort study of patients aged 18 years and older who underwent a radical nephrectomy for kidney cancer between January 1, 2005 and December 31, 2014 using the ACS NSQIP database. NSQIP employs surgical clinical reviewers who collect validated preoperative and perioperative data until 30 days after surgery on patients at participating hospitals. ¹⁵ Its registry includes over 700 hospitals and is focused on reporting and improving perioperative outcomes. ¹⁹

We identified patients undergoing radical nephrectomy using Common Procedural Terminology (CPT) codes (CPT 50220, 50225, 50230, 50545, and 50546) who had complete data for preoperative PLT and WBC count. We subsequently restricted our cohort to patients with a postoperative diagnosis of kidney cancer (ICD-9 codes 189 or 189.0). Patients with a cancer diagnosis were chosen, as previous studies have linked the systemic inflammatory response to cancer prognosis. Patients younger than 18 years (n=2) and those missing data on weight (n=47), height (n=99), length of stay (n=3) or American Society of Anesthesiology (ASA) class (n=16) were excluded. Finally, patients with disseminated cancer (n=588), missing hematocrit (n=4) or gender (n=3) were excluded from our final analysis.

Exposure and covariates

Our exposure of interest was preoperative PLT (10³/microliter) to WBC (number of WBC/microliter) ratio (PLT/WBC), categorized as quintiles. We abstracted data on *a priori* selected covariates based on a literature review to identify preoperative patient factors that may affect postoperative outcomes and may thus confound the relationship between PLT/WBC ratio and perioperative complications. These covariates included age, gender, race, ASA score, body mass index (BMI, kg/m²), smoking status (active smoker within one year of surgery), history of cardiac disease, history of diabetes (requiring oral

medication or insulin therapy), history of neurologic disease, history of chronic obstructive pulmonary disease (COPD), requirement for hemodialysis, functional status (dependent, partially dependent, independent, and unknown), chronic steroid use, surgical technique (laparoscopic vs. open) and preoperative anemia (hematocrit <39%).

Outcome

Our primary outcome was infectious complications (surgical site infections, pneumonia, urinary tract infection [UTI] or sepsis) within 30 days after surgery. Secondary outcomes included bleeding requiring transfusion and major complications within 30 days. Major complications included mortality, reoperation, neurological event (stroke or coma), and/or cardiac event (myocardial infarction or cardiac arrest). These outcome were chosen as they are major contributors to patient mortality, morbidity, and increased healthcare costs.²¹

Statistical analysis

Baseline demographic variables were assessed using frequencies and proportions for categorical variables, means and standard deviations for normally distributed continuous variables, and medians and interquartile ranges for non-normal continuous variables. We compared differences between PLT/WBC ratio quintiles using the Pearson chi-squared test for categorical variables and one-way analysis of variances (ANOVA) for continuous variables. For variables with ordinal structure, we assessed trends across the quintiles using the Cochran-Armitage test for trend.

We compared the proportion of cases resulting in a complication between each of our quintiles. We assessed for differences between the quintiles using the Pearson chi-squared test and examined trends across quintiles using the Cochran-Armitage test. Multivariable logistic regression modelling was used to assess the association between PLT/WBC ratio and each outcome while adjusting for age, gender, race, BMI category, ASA class, history of cardiac disease, diabetes, COPD, neurological disease, on dialysis, chronic use of steroids, functional status, anemia, and smoking status.

Statistical significance was set at p value equal to 0.05. All tests were two-tailed and all statistical analyses were performed using SPSS v24 (IBM Corp., Armonk, NY, U.S.).

Results

We identified 6235 eligible patients who underwent radical nephrectomy for localized kidney cancer during the study period. We found that both lower preoperative WBC as well as higher preoperative PLT counts were responsible for the observed variation in PLT/WBC ratio (Table 1). Patients in the lowest quintile were older, more likely to be male, have

medical comorbidities, to have undergone laparoscopic surgery, and be of non-African descent than patients in higher quintiles (Table 1).

An increasing PLT/WBC ratio was associated with a decreasing risk of 30-day postoperative infectious complications (p<0.0001, Table 2). Univariate analysis examining

Table 1. Demographic characteristics of patients treated with radical nephrectomy for localized kidney cancer, according to quintiles of platelet to white blood cell count (PLT/WBC) ratio

| | PLT/WBC ratio quintiles | | | | | | | | | |
|-----------------------------------|-----------------------------------------|---------------------------|-----------------------------------------|----------------------------|-------------------|----------|------------|--|--|--|
| Variable | Quintile 1 <24.42 | Quintile 2 24.43–30.22 | Quintile 3 30.23–36.20 | Quintile 4 36.21– 44.55 | Quintile 5 >44.55 | р | p (trend)* | | | |
| N | 1265 | 1272 | 1257 | 1245 | 1196 | | | | | |
| Age, median (IQR) | 64 (56–73) | 64(55-72) | 65 (55–72) | 63 (54–71) | 62 (54–71) | <0.0001 | n/a | | | |
| WBC, mean (SD) | 9.96 (4.0) | 8.1 (2.0) | 7.4 (1.9) | 6.9 (1.8) | 6.2 (1.9) | <0.0001 | n/a | | | |
| PLT count, mean (SD) | 187.4 (60.4) | 221.2 (55.7) | 244.8 (63.2) | 274.2 (72.6) | 341.6 (119.0) | <0.0001 | | | | |
| Gender | | | | | | < 0.0001 | < 0.001 | | | |
| Male | 920 (72.7%) | 873 (68.6%) | 811 (64.6%) | 710 (57.1%) | 602 (50.3%) | | | | | |
| Female | 345 (27.3%) | 399 (31.4%) | 445 (35.4%) | 533 (42.9%) | 594 (49.7%) | | | | | |
| Preoperative diabetes | | | | | | < 0.0001 | < 0.0001 | | | |
| Yes | 339 (26.8%) | 281 (22.1%) | 268 (21.3%) | 239 (19.2%) | 203 (17.0%) | | | | | |
| Smoker within 1 year | | | | | | < 0.0001 | < 0.0001 | | | |
| Yes | 313 (24.7%) | 274 (21.5%) | 237 (18.9%) | 184 (14.7%) | 182 (15.2%) | | | | | |
| Preoperative COPD | | | | | | < 0.0001 | < 0.0001 | | | |
| Yes | 115 (9.1%) | 86 (6.8%) | 66 (5.3%) | 46 (3.7%) | 46 (3.8%) | | | | | |
| Preoperative MI, CHF, or angina | | | | | | 0.002 | < 0.0001 | | | |
| Yes | 68 (5.4%) | 63 (5.0%) | 54 (4.3%) | 43 (3.5%) | 30 (2.5%) | | | | | |
| ASA classification | | | | | | < 0.0001 | n/a | | | |
| 1 | 12 (1.1%) | 10 (0.8%) | 24 (1.9%) | 24 (1.9%) | 26 (2.2%) | | | | | |
| 2 | 307 (24.3%) | 409 (32.2%) | 445 (35.4%) | 460 (36.9%) | 455 (38.0%) | | | | | |
| 3 | 787 (62.2%) | 737 (57.9%) | 695 (55.3%) | 692 (55.6%) | 653 (54.6%) | | | | | |
| 4 | 157 (12.4%) | 116 (9.1%) | 93 (7.4%) | 69 (5.5%) | 62 (5.2%) | | | | | |
| Preoperative stroke or neurologic | cal history | | | | | 0.271 | 0.473 | | | |
| Yes | 24 (1.9%) | 23 (1.6%) | 27 (2.1%) | 29 (2.3%) | 14 (1.2%) | | | | | |
| Preoperative BMI | | | | | | < 0.0001 | n/a | | | |
| <18.5 | 14 (1.1%) | 7 (0.6%) | 9 (0.7%) | 13 (1.0%) | 25 (2.1%) | | | | | |
| 18.5–24.9 | 181 (14.3%) | 191 (15.0%) | 213 (16.9%) | 262 (21.0%) | 326 (27.3%) | | | | | |
| 25–29.9 | 399 (31.5%) | 419 (32.9%) | 452 (36.0%) | 429 (34.5%) | 416 (34.8%) | | | | | |
| 30+ | 671 (53.0%) | 655 (51.5%) | 583 (46.4%) | 541 (43.5%) | 429 (35.9%) | | | | | |
| Functional status | | | | | | 0.073 | n/a | | | |
| Independent | 1217 (96.2%) | 1245 (97.9%) | 1220 (97.1%) | 1218 (97.8%) | 1170 (97.8%) | | | | | |
| Partially dependent | 40 (3.2%) | 21 (1.7%) | 29 (2.3%) | 21 (1.7%) | 19 (1.6%) | | | | | |
| Totally dependent | 2 (0.2%) | 0 | 0 | 3 (0.2%) | 3 (0.3%) | | | | | |
| Not reported | 6 (0.5%) | 6 (0.5%) | 8 (0.6%) | 3 (0.2%) | 4 (0.3%) | | | | | |
| Race | ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | | ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | | , | < 0.0001 | n/a | | | |
| African | 76 (6.0%) | 98 (7.7%) | 98 (7.8%) | 122 (9.8%) | 149 (12.5%) | | .,. | | | |
| Caucasian | 1046 (82.7%) | 1017 (80.0%) | 1032 (82.1%) | 972 (78.1%) | 859 (71.8%) | | | | | |
| Other | 143 (11.3%) | 157 (12.3%) | 127 (10.1%) | 151 (12.1%) | 188 (15.7%) | | | | | |
| Dialysis-dependent | | | , (10.170) | | .55 (1017,07 | < 0.0001 | <0.0001 | | | |
| Yes | 78 (6.2%) | 61 (4.8%) | 55 (4.4%) | 45 (3.6%) | 36 (3.0%) | | .3.000 | | | |
| Chronic steroid use | , 5 (5.2/0) | 01 (1.070) | 00 (1.470) | 10 (3.070) | 00 (0.070) | < 0.0001 | <0.0001 | | | |
| Yes | 81 (6.4%) | 53 (4.2%) | 43 (3.4%) | 41 (3.3%) | 38 (3.2%) | \0.000 i | \J.0001 | | | |
| Anemia | 01 (0.470) | OO (-T.2 /0) | -0 (0. -70) | - 1 \J.J/0/ | 00 (0.270) | < 0.0001 | <0.001 | | | |
| Yes | 486 (38.4%) | 430 (33.8%) | 486 (38.7%) | 596 (47.9%) | 736 (61.6%) | \0.000 i | \0.00T | | | |

^{*}Cochrane Armitage test for trend . 'Statistical test performed using analysis of variance (ANOVA) with Dunnett's C post-hoc testing. ASA: American Society of Anesthesiologists; BMI: body mass index; CHF: congestive heart failure; COPD: chronic obstructive pulmonary disease; IQR: interquartile range; MI: myocardial infarction; SD: standard deviation.

each site of infection demonstrated a significant difference between quintiles for sepsis, pneumonia, and UTI, with significant trend for decreasing complication rate with increasing PLT/WBC quintile for both sepsis and pneumonia (Table 2). While there appeared to be a decreasing trend for UTIs with increasing PLT/WBC quintiles, this was not significant (p=0.14). There was no significant difference in surgical site infection rates between quintiles (p=0.38). After multivariable regression, patients with low PLT/WBC ratio (Quintiles 1 and 2) had a significantly higher risk of infectious complication compared to patients in Quintile 5 (Table 3).

Among our secondary outcomes, there was a significant difference in transfusion rates, but not major complication rates between PLT/WBC quintiles (Table 2). Further, there was a significant trend for higher transfusion rates among patients in higher quintiles (Cochran Armitage test for trend, p=0.0009). On multivariable analysis, patients in Quintiles 2, 3, and 4 had significantly lower odds of requiring transfusion compared to Quintile 5 (Table 3). There was no significant difference in major complication rates between quintiles on either univariate or multivariable analyses (Tables 2 and 3).

Discussion

Using a large, prospectively collected and validated multiinstitutional registry, we identified PLT/WBC ratio as a novel, independent predictor of infectious and bleeding complications following radical nephrectomy for kidney cancer. While PLT/WBC ratio was strongly related to baseline patient characteristics, including age and comorbidity, a lower PLT/ WBC ratio (Quintile 1) was independently associated with a two-fold higher risk of infectious complications compared to patients in the highest PLT/WBC quintile (Quintile 5) after adjusting for these characteristics. Further, a higher PLT/WBC ratio was independently associated with higher risk of requiring blood transfusion compared to lower PLT/WBC ratios (Quintiles 2, 3, and 4). The major complication rate for the whole cohort was low, consistent with rates reported in the literature,²² and not associated with the PLT/WBC ratio.

To the best of our knowledge, this is the first study to identify PLT/WBC ratio as a predictor of postoperative complications in surgical patients with localized cancer. Previous studies have used other hematological markers as surrogates to the inflammatory response to predict prognosis in patients with metastatic and localized disease; 1,11,23 however, the use of such markers to predict short-term outcomes has not been previously studied. Furthermore, these calculations usually

Table 2. Univariate analysis of 30-day postoperative complications in patients undergoing radical nephrectomy for localized kidney cancer (n=6235)

| | PLT/WBC ratio quintiles | | | | | | | |
|----------------------------|-------------------------|--------------|--------------|--------------|--------------|-------------------------|-------------------------------------|--|
| | Quintile 1 | Quintile 2 | Quintile 3 | Quintile 4 | Quintile 5 | Pearson chi-square p | Cochrane-Armitage test for trend | |
| Infectious complications | | | | | | < 0.0001 | < 0.0001 | |
| No | 1155 (91.3%) | 1186 (93.2%) | 1196 (95.1%) | 1195 (96.0%) | 1138 (95.2%) | | | |
| Yes | 110 (8.7%) | 86 (6.8%) | 61 (4.9%) | 50 (4.0%) | 58 (4.8%) | | | |
| Sepsis | | | | | | < 0.0001 | < 0.0001 | |
| No | 1210 (95.7%) | 1248 (98.1%) | 1238 (98.5%) | 1225 (98.4%) | 1180 (98.7%) | | | |
| Yes | 55 (4.3%) | 24 (1.9%) | 19 (1.5%) | 20 (1.6%) | 16 (1.3%) | | | |
| Surgical site infections | | | | | | 0.383 | Not performed | |
| No | 1237 (97.6%) | 1241 (98.2%) | 1234 (98.4%) | 1225 (98.4%) | 1178 (98.5%) | | | |
| Yes | 28 (2.2%) | 31 (2.4%) | 23 (1.8%) | 20 (1.6%) | 18 (1.5%) | | | |
| Pneumonia | | | | | | | | |
| No | 1239 (97.9%) | 1247 (98.0%) | 1240 (98.6%) | 1236 (99.3%) | 1182 (98.8%) | 0.028 | 0.005 | |
| Yes | 26 (2.1%) | 25 (2.0%) | 17 (1.4%) | 9 (0.7%) | 14 (1.2%) | | | |
| Urinary tract infection | | | | | | | | |
| No | 1239 (97.9%) | 1251 (98.3%) | 1243 (98.9%) | 1236 (99.3%) | 1175 (98.2%) | 0.042 | 0.144 | |
| Yes | 26 (2.1%) | 21 (1.7%) | 14 (1.1%) | 9 (0.7%) | 21 (1.8%) | | | |
| Major complications | | | | | | 0.655 | Not performed | |
| No | 1215 (96.0%) | 1231 (96.8%) | 1219 (97.0%) | 1206 (96.9%) | 1160 (97.0%) | | | |
| Yes | 50 (4.0%) | 41 (3.2%) | 38 (3.0%) | 39 (3.1%) | 36 (3.0%) | | | |
| Bleeding requiring transfu | usion | | | | | < 0.0001 | 0.0009 | |
| No | 1078 (85.2%) | 1133 (89.1%) | 1109 (88.2%) | 1095 (88.0%) | 962 (80.4%) | | | |
| Yes | 187 (14.8%) | 139 (10.9%) | 148 (11.8%) | 150 (12.0%) | 234 (19.6%) | | | |

Table 3. Multivariate analysis assessing the association between PLT/WBC ratio and major, bleeding, and infectious complications 30 days postoperatively in patients who underwent radical nephrectomy for kidney cancer (n=6235)

| Variable | Odds ratio* | Lower 95% CI | Upper 95% CI | р | | | | |
|--------------------------------|----------------|-----------------|-----------------|----------|--|--|--|--|
| Infectious complication | าร | | | | | | | |
| Quintile 1 | 1.88 | 1.33 | 2.66 | < 0.0001 | | | | |
| Quintile 2 | 1.58 | 1.10 | 2.25 | 0.013 | | | | |
| Quintile 3 | 1.11 | 0.76 | 1.62 | 0.60 | | | | |
| Quintile 4 | 0.87 | 0.59 | 1.29 | 0.48 | | | | |
| Quintile 5 | REFERENCE | | | | | | | |
| Major complications | | | | | | | | |
| Quintile 1 | 1.04 | 0.66 | 1.65 | 0.87 | | | | |
| Quintile 2 | 0.94 | 0.58 | 1.50 | 0.78 | | | | |
| Quintile 3 | 0.89 | 0.55 | 1.43 | 0.63 | | | | |
| Quintile 4 | 0.98 | 0.61 | 1.56 | 0.91 | | | | |
| Quintile 5 | REFERENCE | | | | | | | |
| Bleeding requiring transfusion | | | | | | | | |
| Quintile 1 | 0.83 | 0.66 | 1.05 | 0.114 | | | | |
| Quintile 2 | 0.65 | 0.51 | 0.83 | 0.001 | | | | |
| Quintile 3 | 0.67 | 0.53 | 0.85 | 0.001 | | | | |
| Quintile 4 | 0.62 | 0.49 | 0.79 | < 0.0001 | | | | |
| Quintile 5 | REFERENCE | | | | | | | |

^{*}Adjusted for the effect of age, diabetes history, smoking history, chronic obstructive pulmonary disease history, cardiac history, American Society of Anesthesiologists classification, neurological history, body mass index, gender, mobility, race, undergoing dialysis, chronic steroid use, and anemia history. Cl: confidence interval; PLT/WBC: platelet to white blood cell count

require neutrophil and lymphocytes counts, data that is not uniformly collected in registry databases.

It is notable that PLT/WBC ratio was significantly associated with sepsis, pneumonia, and UTI, but not wound infections. Wound infections are much more likely to be influenced by technical considerations, including skin preparation, shaving, and wound closure, ²⁴ in addition to patient factors, as compared to other sites of infection. Further, because radical nephrectomy incisions are considered clean, surgical site infections are rare. ²⁵

The mechanistic relationship between PLT/WBC ratio and infection remains to be elucidated. One possibility is that the PLT/WBC ratio is a surrogate for a patient's baseline health status. Patients with low ratios were more likely to have more comorbidities (diabetes, COPD, dialysis-dependence, use of chronic steroids) and to be current smokers, which are all known to alter the inflammatory response and are risk factors for postoperative infections. Furthermore, other studies have found poor health status (high BMI, >5% weight loss, low albumin, lymphocyte count) of patients to result in worse postoperative outcomes in patients undergoing nephrectomy for renal cell carcinoma. Notably, PLT/WBC ratio remained an important predictor of infectious complications after adjusting for these factors. Further, this ratio is likely also an indicator of metabolic syndrome, a

constellation of physiological and biochemical abnormalities, resulting in inappropriate activation of inflammatory pathways.³⁰ Metabolic syndrome is a well-known risk factor for postoperative complications, including infection.^{27,31,32}

In addition to the association with infectious complications, PLT/WBC ratio was significantly associated with rates of bleeding requiring transfusion. Patients in the lowest quintile were overall in poorer general health compared to patients in higher quintiles and were more likely to have undergone laparoscopic surgery compared to patients in Quintile 5. These higher transfusion rates may be explained by the fact that a greater proportion of patients underwent open surgery in Quintile 5. Many studies have found higher transfusion rates with open compared to laparoscopic nephrectomy.^{33,34} Indeed, in our model, open surgical technique was a strong predictor of increased transfusion risk (data not shown). Furthermore, it is unknown if patients undergoing open surgery had increased tumour complexity (data is not captured), which could increase risk of transfusion. Another explanation is that there are a higher proportion of female patients in Quintile 5 (49.7%) compared to Quintile 1 (27.3%). Biologically, females have lower hemoglobin levels compared to males.³⁵ As these patients were treated at many different hospitals, it is unknown what parameters were used for initiating transfusion and how these may have differed between institutions.

We excluded patients with disseminated cancer at time of surgery, as studies have found these patients to be at increased risk of complications compared to patients with localized disease. ^{6,36} Further, we excluded patients for whom the postoperative diagnosis was not explicitly recorded as kidney cancer. Thus, these results may not directly be extrapolated to patients with metastatic disease, nor those undergoing nephrectomy for benign indications.

Despite the strengths of the NSQIP database, there are limitations to this analysis. As we used retrospective data, there are likely other confounders that were not included in our model that could be affecting complication rates. In addition, as NSQIP only captures complications within the first 30 postoperative days, the results may not apply to complications that occur beyond this period. NSQIP does not collect information on tumour staging and renal nephrometry scores. As a result, we cannot comment on stage migration over time; however, epidemiological data suggest that due to imaging studies being performed for other indications, renal cell carcinoma is commonly found incidentally.³⁷ This has led to a stage migration overall towards lower stage of disease at time of diagnosis.³⁸

In order to provide clinically meaningful interpretations of PLT/WBC ratio, we used quintiles; however, future research may identify more informative ways to operationalize this variable, such as a dichotomous threshold. Currently, no such cutoff has been identified. As this study only exam-

ined PLT/WBC ratio in patients with renal cell carcinoma, its results may not be applicable to patients undergoing other procedures. Finally, we cannot assess whether neutrophil to lymphocyte ratio would perform similarly to PLT/WBC ratio.

It was our hypothesis that the PLT/WBC, a measure of the systemic inflammatory response, would reflect the overall health status of the patient, taking into account their medical comorbidities, cancer status, and lifestyle. We found that the PLT/WBC ratio was associated with postoperative infections both in univariate analysis and independently in multivariable analyses accounting for these factors. Thus, this simple ratio may prove useful for busy clinicians, as it is easier to interpret this result than to integrate the risks of each predictive factor.

Conclusion

Our study is the first to examine usefulness of the PLT/WBC ratio in predicting short-term 30-day postoperative outcomes in patients undergoing surgery for renal malignancy. Patients who had low PLT/WBC ratios had significantly higher odds of infectious complications compared to patients with high PLT/WBC ratios. Further studies need to be performed to verify these findings in other populations.

Competing interests: The authors report no competing personal or financial interests.

This paper has been peer-reviewed.

References

- Grimes N, Tyson M, Hannan C, et al. A systematic review of the prognostic role of hematological scoring systems in patients with renal cell carcinoma undergoing nephrectomy with curative intent. Clin Genitourin Cancer 2016;14:271-6. https://doi.org/10.1016/j.clgc.2016.01.006
- Rendon RA, Kapoor A, Breau R, et al. Surgical management of renal cell carcinoma: Canadian Kidney Cancer Forum Consensus. Can Urol Assoc J 2014;8:E398-412. https://doi.org/10.5489/cuaj.1894
- Ljungberg B, Bensalah K, Canfield S, et al. EAU guidelines on renal cell carcinoma: 2014 update. Eur Urol 2015;67:913-24. https://doi.org/10.1016/j.eururo.2015.01.005
- Ebbing J, Wiebach T, Kempkensteffen C, et al. Evaluation of perioperative complications in open and laparoscopic surgery for renal cell cancer with turnour thrombus involvement using the Clavien—Dindo classification. Eur J Surg Oncol 2015;41:941-52. https://doi.org/10.1016/j.ejso.2015.02.009
- Hadjipavlou M, Khan F, Fowler S, et al. Partial vs. radical nephrectomy for T1 renal tumours: An analysis from the British Association of Urological Surgeons nephrectomy audit. BJU Int 2016;117:62-71. https://doi.org/10.1111/bju.13114
- Wallis CJ, Bjarnason G, Byrne J, et al. Morbidity and mortality of radical nephrectomy for patients with disseminated cancer — an analysis of the National Surgical Quality Improvement Program Database. *Urology* 2016;95:95-102. https://doi.org/10.1016/j.urology.2016.04.055
- Novitsky YW, Orenstein SB. Effect of patient and hospital characteristics on outcomes of elective ventral hemia repair in the United States. Hemia 2013;17:639-45. https://doi.org/10.1007/s10029-013-1088-5
- Scrutinio D, Giannuzzi P. Comorbidity in patients undergoing coronary artery bypass graft surgery: Impact on outcome and implications for cardiac rehabilitation. Eur J Cardiovasc Prev Rehabil 2008;15:379-85. https://doi.org/10.1097/HJR.0b013e3282fd5c6f
- Roxburgh CS, McMillan DC. Role of systemic inflammatory response in predicting survival in patients with primary operable cancer. Future Oncol 2010;6:149-63. https://doi.org/10.2217/fon.09.136

- Ramsey S, Lamb GW, Aitchison M, et al. Evaluation of an inflammation-based prognostic score in patients with metastatic renal cancer. Cancer 2007;109:205-12. https://doi.org/10.1002/cncr.22400
- Lucca I, de Martino M, Hofbauer SL, et al. Comparison of the prognostic value of pretreatment measurements of systemic inflammatory response in patients undergoing curative resection of clear-cell renal cell carcinoma. World J Urol 2015;33:2045-52. https://doi.org/10.1007/s00345-015-1559-7
- Dehghani MR, Rezaei Y, Taghipour-Sani L. White blood cell count to mean platelet volume ratio as a novel non-invasive marker predicting long-term outcomes in patients with non-ST elevation acute coronary syndrome. Cardiol J 2015;22:437-45. https://doi.org/10.5603/CJ.a2015.0015
- Çiçek G, Açıkgöz SK, Yayla Ç, et al. White blood cell count to mean platelet volume ratio: A novel and promising prognostic marker for ST-segment elevation myocardial infarction. Cardiol J 2016;23:225-35. https://doi.org/10.5603/CJ.a2016.0001
- Ashrafganjoei T, Mohamadianamiri M, Farzaneh F, et al. Investigating preoperative hematological markers for prediction of ovarian cancer surgical outcome. Asian Pac J Cancer Prev 2016;17:1445-8. https://doi.org/10.7314/APJCP.2016.17.3.1445
- Cohen ME, Liu Y, Ko CY, et al. Improved surgical outcomes for ACS NSQIP hospitals over time: Evaluation of hospital cohorts with up to 8 years of participation. *Ann Surg* 2016;263:267-73. https://doi.org/10.1097/SLA.000000000001192
- Davenport DL, Holsapple CW, Conigliaro J. Assessing surgical quality using administrative and clinical data sets: A direct comparison of the University HealthSystem Consortium Clinical Database and the National Surgical Quality Improvement Program data set. Am J Med Qual 2009;24:395-402. https://doi.org/10.1177/1062860609339936
- Cima RR, Lackore KA, Nehring SA, et al. How best to measure surgical quality? Comparison of the Agency for Healthcare Research and Quality Patient Safety Indicators (AHRQ-PSI) and the American College of Surgeons National Surgical Quality Improvement Program (ACS-NSQIP) postoperative adverse events at a single institution. Surgery 2011;150:943-49. https://doi.org/10.1016/j. surg.2011.06.020
- Koch CG, Li L, Hixson E, et al. What are the real rates of postoperative complications: Elucidating inconsistencies between administrative and clinical data sources. J Am Coll Surg 2012;214:798-805. https://doi.org/10.1016/i.jamcollsurg.2011.12.037
- Ingraham AM, Richards KE, Hall BL, et al. Quality improvement in surgery: The American College of Surgeons National Surgical Quality Improvement Program approach. Adv Surg 2010;44:251-67. https://doi.org/10.1016/j.yasu.2010.05.003
- Ramsey MT, Fabian TC, Shahan CP, et al. A prospective study of platelet function in trauma patients. J Trauma Acute Care Surg 2016;80:726-32; discussion 32-3. https://doi.org/10.1097/ TA.000000000001017
- Eappen S, Lane BH, Rosenberg B, et al. Relationship between occurrence of surgical complications and hospital finances. JAMA 2013;309:1599-606. https://doi.org/10.1001/jama.2013.2773
- De P, Otterstatter MC, Semenciw R, et al. Trends in incidence, mortality, and survival for kidney cancer in Canada, 1986—2007. Cancer Causes & Control 2014;25:1271-81. https://doi.org/10.1007/ s10552-014-0427-x
- Grimes N, Tyson M, Hannan C, et al. A systematic review of the prognostic role of hematological scoring systems in patients with renal cell carcinoma undergoing nephrectomy with curative intent. Clin Genitourin Cancer 2016;14:271-6. https://doi.org/10.1016/j.clgc.2016.01.006
- Gottrup F MA, Hollander DA. An overview of surgical site infection: Etiology, incidence, and risk factors. Eur Wound Manag Assoc J 2005;5:11-5.
- Kortram K, Ijzermans JN, Dor FJ. Perioperative events and complications in minimally invasive live donor nephrectomy: A systematic review and meta-analysis. *Transplantation* 2016;100:2264-75. https://doi.org/10.1097/TP.000000000001327
- Klinger MH, Jelkmann W. Role of blood platelets in infection and inflammation. J Interferon Cytokine Res 2002;22:913-22. https://doi.org/10.1089/10799900260286623
- Dandona P, Aljada A, Chaudhuri A, et al. Metabolic syndrome: A comprehensive perspective based on interactions between obesity, diabetes, and inflammation. *Circulation* 2005;111:1448-54. https://doi.org/10.1161/01.CIR.0000158483.13093.9D
- Morgan TM, Tang D, Stratton KL, et al. Preoperative nutritional status is an important predictor of survival in patients undergoing surgery for renal cell carcinoma. Eur Urol 2011;59:923-8. https://doi.org/10.1016/j.eururo.2011.01.034
- Jeon HG, Choi DK, Sung HH, et al. Preoperative prognostic nutritional index is a significant predictor of survival in renal cell carcinoma patients undergoing nephrectomy. *Ann Surg Oncol* 2016;23:321-7. https://doi.org/10.1245/s10434-015-4614-0
- Odrowaz-Sypniewska G. Markers of pro-inflammatory and pro-thrombotic state in the diagnosis of metabolic syndrome. Adv Med Sci 2007;52:246-50.
- Doyle SL, Lysaght J, Reynolds JV. Obesity and postoperative complications in patients undergoing nonbariatric surgery. Obes Rev 2010;11:875-86. https://doi.org/10.1111/j.1467-789X.2009.00700.x

- Safranow K, Dziedziejko V, Rzeuski R, et al. Inflammation markers are associated with metabolic syndrome and ventricular arrhythmia in patients with coronary artery disease. *Postepy Hig Med Dosw* 2016;70:56-66. https://doi.org/10.5604/17322693.1194612
- Pak JS, Lee JJ, Bilal K, et al. Utilization trends and outcomes up to 3 months of open, laparoscopic, and robotic partial nephrectomy. J Robot Surg 2017;11:223-9. https://doi.org/10.1007/s11701-016-0650-4
- Ghani KR, Sukumar S, Sammon JD, et al. Practice patterns and outcomes of open and minimally invasive partial nephrectomy since the introduction of robotic partial nephrectomy: Results from the nationwide inpatient sample. J Urol 2014;191:907-13. https://doi.org/10.1016/j.juro.2013.10.099
- Bydon M, Abt NB, Macki M, et al. Preoperative anemia increases postoperative morbidity in elective cranial neurosurgery. Surg Neurol Int 2014;5:156. https://doi.org/10.4103/2152-7806.143754
- Moghadamyeghaneh Z, Hanna MH, Hwang G, et al. Outcomes of colon resection in patients with metastatic colon cancer. Am J Surg 2016;212:264-71. https://doi.org/10.1016/j.amjsurg.2016.01.025

- Sweeney JP, Thornhill JA, Graiger R, et al. Incidentally detected renal cell carcinoma: Pathological features, survival trends, and implications for treatment. Br J Urol 1996;78:351-3. https://doi.org/10.1046/ i.1464-410X.1996.00140.x
- Nason GJ, McGuire BB, Kelly ME, et al. Clinico-pathological analysis of renal cell carcinoma demonstrates decreasing tumour grade over a 17-year period. Can Urol Assoc J 2014;8:125-32. https://doi.org/10.5489/cuaj.1721

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