

Prior Frequent Emergency Department Use as a Predictor of Emergency Department Visits After a New Cancer Diagnosis

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QUESTION ASKED: Among adults with newly diagnosed cancer, how strongly is emergency department (ED) visit volume before cancer diagnosis associated with ED visits after cancer diagnosis compared with other known predictors of ED use (social determinants of health [SDH] and clinical cancer-related factors)?

SUMMARY ANSWER: Frequent pre-cancer diagnosis ED use was strongly associated with post-cancer diagnosis ED visit volume, with a predictive impact nearly twice as high as those of the strongest SDH or clinical predictors.

WHAT WE DID: We identified adults diagnosed with cancer between 2008 and 2018 at an academic medical center and a safety-net hospital and matched them to a regional health information exchange of ED encounters at 98% of the nonfederal hospitals in North Texas. We used a multivariable negative binomial regression to model the number of ED visits in the first 6 months after an incident cancer diagnosis with the following prespecified predictors: ED visit history in the 6-12 months preceding cancer diagnosis, electronic health record proxy SDH (race, language preference, insurance type, and homelessness), and clinical severity (cancer type, stage at diagnosis, initial treatment modalities, and comorbidities).

WHAT WE FOUND: Among 35,090 patients with cancer (49% female and 50% non-White), 57% had ≥ 1 ED visit in the 6 months immediately following cancer

diagnosis and 20% had ≥ 1 ED visit in the 6-12 months prior to cancer diagnosis. The strongest predictor of postdiagnosis ED visits was frequent (≥ 4) prediagnosis ED visits (adjusted incidence rate ratio [aIRR]: 3.68; 95% CI, 3.36 to 4.02). Having 1-3 prediagnosis ED visits (aIRR: 1.32; 95% CI, 1.28 to 1.36), Hispanic (aIRR: 1.12; 95% CI, 1.07 to 1.17) and Black (aIRR: 1.21; 95% CI, 1.17 to 1.25) race, homelessness (aIRR: 1.95; 95% CI, 1.73 to 2.20), advanced-stage cancer (aIRR: 1.30; 95% CI, 1.26 to 1.35), and treatment regimens including chemotherapy (aIRR: 1.44; 95% CI, 1.40 to 1.48) were also associated with greater postdiagnosis ED use.

BIAS, CONFOUNDING FACTOR(S): This was an observational study, so there may be residual unmeasured confounders affecting our results. However, we did adjust for major SDH and measures of clinical severity such as cancer type and stage, comorbidities, and treatment modalities received. Furthermore, our findings were robust to a number of sensitivity analyses.

REAL-LIFE IMPLICATIONS: Frequent ED use prior to cancer diagnosis may serve as a surrogate marker that could be used to identify patients likely to have frequent postdiagnosis ED visits. Efforts to reduce ED visits among patients with cancer should consider tailoring interventions that target heavy prior ED users when offering alternative acute care options.

ASSOCIATED CONTENT

Appendix

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PURPOSE To determine whether emergency department (ED) visit history prior to cancer diagnosis is associated with ED visit volume after cancer diagnosis.

METHODS This was a retrospective cohort study of adults (≥ 18 years) with an incident cancer diagnosis (excluding nonmelanoma skin cancers or leukemia) at an academic medical center between 2008 and 2018 and a safety-net hospital between 2012 and 2016. Our primary outcome was the number of ED visits in the first 6 months after cancer diagnosis, modeled using a multivariable negative binomial regression accounting for ED visit history in the 6-12 months preceding cancer diagnosis, electronic health record proxy social determinants of health, and clinical cancer-related characteristics.

RESULTS Among 35,090 patients with cancer (49% female and 50% non-White), 57% had ≥ 1 ED visit in the 6 months immediately following cancer diagnosis and 20% had ≥ 1 ED visit in the 6-12 months prior to cancer diagnosis. The strongest predictor of postdiagnosis ED visits was frequent (≥ 4) prediagnosis ED visits (adjusted incidence rate ratio [aIRR]: 3.68; 95% CI, 3.36 to 4.02). Other covariates associated with greater postdiagnosis ED use included having 1-3 prediagnosis ED visits (aIRR: 1.32; 95% CI, 1.28 to 1.36), Hispanic (aIRR: 1.12; 95% CI, 1.07 to 1.17) and Black (aIRR: 1.21; 95% CI, 1.17 to 1.25) race, homelessness (aIRR: 1.95; 95% CI, 1.73 to 2.20), advanced-stage cancer (aIRR: 1.30; 95% CI, 1.26 to 1.35), and treatment regimens including chemotherapy (aIRR: 1.44; 95% CI, 1.40 to 1.48).

CONCLUSION The strongest independent predictor for ED use after a new cancer diagnosis was frequent ED visits before cancer diagnosis. Efforts to reduce potentially avoidable ED visits among patients with cancer should consider educational initiatives that target heavy prior ED users and offer them alternative ways to seek urgent medical care.

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INTRODUCTION

The literature on frequent emergency department (ED) use among the general patient population largely focuses on clinical characteristics (eg, mental health conditions)¹⁻³ and indicators of social disadvantage (eg, social determinants of health [SDH])⁴ as predictors of ED use. Currently, health care systems largely rely on patients to triage the severity and urgency of their illness when they get sick and decide between going to the ED versus less expensive alternative sites of care, something patients may be ill-equipped to judge.

How to decide when and where to seek acute care when sick is even more challenging for patients with

cancer. Adults newly diagnosed with cancer frequently visit the ED for nonemergent conditions because of side effects of their treatment and/or symptoms related to their underlying disease.⁵⁻¹⁰ This problem may be exacerbated by warnings to be alert to seemingly minor changes in condition: between 15% and 22% of patients with cancer have two or more ED visits made within 180 days of diagnosis.^{5,11}

Even privately insured patients encounter administrative barriers in accessing timely, appropriate-site, and appropriate-specialty care.¹² It follows that patients who have had poor prior experiences receiving a timely and complete evaluation in outpatient settings might learn to use the ED as a primary source of care.

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Other patients may not have enough health system savvy from the outset to be aware of and seek out less costly sites of acute care.

For a population-based cohort of patients with cancer in two health systems, we used longitudinally linked ED visits, including visits preceding the cancer diagnosis, to predict the factors influencing ED visits in the 6 months after a new cancer diagnosis. We hypothesized that ED visits post-cancer diagnosis would be associated with their pre-cancer diagnosis ED visit patterns, independent of other known drivers of acute care such as cancer- and treatment-related characteristics and SDH.

METHODS

Study Design, Population, and Setting

We conducted a retrospective analysis of comprehensive, longitudinally linked ED encounters from a regional health information exchange, the Dallas-Fort Worth Hospital Council Foundation (Foundation), which contains information on ED visits to > 80 hospitals in North Texas (all the nonfederal hospitals in a 100-mile radius of Dallas). We evaluated predictors of ED visits after a new cancer diagnosis across two different patient populations: patients with cancer treated at the University of Texas Southwestern Medical Center (University) and at the Parkland Health & Hospital System (Parkland), the safety-net integrated health system for Dallas County. We included patients who had an incident cancer diagnosis between 2008 and 2018 among the University population and between 2012 and 2016 among the Parkland population. For patients with synchronous diagnoses, we selected the highest-stage cancer. For patients with metachronous cancer diagnoses, we selected the first diagnosis and excluded subsequent diagnoses. We excluded patients with only nonmelanoma skin cancer because of lower ED use overall. We also excluded patients with leukemia because of prolonged inpatient stays for acute leukemia and frequent hospitalizations for relapse of disease,¹³ which make post-cancer diagnosis ED use difficult to compare with other cancers within a 6-month window.

Parkland is the primary provider of care for the under- and uninsured in Dallas County and is the sole provider of cancer care for the uninsured in the region. However, Parkland patients do visit EDs at other hospitals, which may not be captured by any single health system electronic health record (EHR). Among our University population, we have also found that two thirds of ED visits are made to other health systems and are not captured by the local EHR.¹⁴ Both health systems use the same EHR vendor (Epic, Verona, WI). Some University faculty physicians supervise trainees at Parkland, but the clinic staff providing acute care triage are exclusive to each health system. We linked patients from local cancer registries to the regional Foundation database, which collects information for approximately

12 million unique patients and their 65 million hospital encounters. Visit-level data are organized into a master patient index that assigns a unique identifier, allowing longitudinal tracking of patients over time to all hospitals in the Foundation database.

Using a combination of name, date of birth, zip code of residence, and medical record number, we matched patients to their unique Foundation identifiers. The Foundation database provided dates of ED arrival and discharge, name of hospital and health system, and discharge disposition (including died in ED, transferred to another hospital, observation stay, inpatient admission, and discharged home).

Outcome

Our primary outcome was the number of ED visits in the first 6 months (180 days) after a new diagnosis of cancer. ED visits that resulted in the initial pathology-confirmed cancer diagnosis were excluded.

Covariates

Our primary prespecified predictor of interest was ED use prior to cancer diagnosis, which we defined as the number of ED visits 6-12 months (181-365 days) before diagnosis. We categorized this variable as zero visits, 1-3 visits, or ≥ 4 visits. To be conservative, we did not use ED visits in the 0-6 months immediately before diagnosis, as they could potentially reflect early symptoms of the subsequently diagnosed cancer.

We prespecified additional predictors of postdiagnosis ED visits on the basis of our team's multidisciplinary expertise and from prior literature that were available in the EHR. These included a set of previously validated SDH variables (race and/or ethnicity, non-English language preference, insurance type at time of diagnosis, and homelessness),^{3,15-19} comorbidities at diagnosis (organized into Charlson Comorbidity Index),^{20,21} and clinical cancer-specific characteristics obtained from local cancer registries maintained by health systems that are Commission on Cancer-designated Academic Comprehensive Care, and National Cancer Institute-designated Comprehensive Cancer Programs (cancer type, whether advanced stage at diagnosis, and initial treatment modalities [chemotherapy, radiation therapy, surgery, and/or immunotherapy]).^{5,22} We categorized a patient as homeless if their address was listed as homeless shelter in the EHR at any point between 1 year (365 days) before or 6 months (180 days) after cancer diagnosis or if they received care with the homeless health care program during that period. Advanced stage was defined as stage IIIB or higher for lung cancer, stage III or higher for pancreatic cancer, and stage IV for all others except for brain cancer.^{5,23} Because our overall cohort consisted of patients from two distinct health systems spanning different time periods, we included the patient's health system (University or Parkland) and year of cancer diagnosis as covariates. The model was also adjusted for age, sex, and whether the

patient died within 6 months (180 days) after cancer diagnosis.

Statistical Analysis

We first grouped ED visits and ED visit history into a patient-level data set and used descriptive statistics to characterize the patient cohort. Next, we applied a multivariable negative binomial regression to model the number of ED visits within the first 6 months after a new cancer diagnosis. We chose negative binomial regression because of overdispersion of the outcome variable. For improved interpretability, we used marginal effects methods to estimate adjusted postdiagnosis ED visit counts across key strata of interest.²⁴

Sensitivity Analyses

To assess the robustness of our findings, we conducted a series of sensitivity analyses for the University and Parkland cohorts separately. First, we reran our analyses for each health system cohort separately. For both cohorts, we repeated analyses with (1) ED visit history 12-18 months prior to cancer diagnosis as the primary predictor, (2) added interaction terms between initial treatment modalities and cancer stage at diagnosis, (3) neighborhood education level included as a binary covariate (low v not low), and (4) excluded pre- and post-cancer diagnosis ED visits that resulted in hospitalization.

For the University cohort, we repeated analyses with neighborhood poverty level included as a binary covariate (high v not high). This was not done for the Parkland cohort because many patients were enrolled in low-income insurance programs at the county (charity assistance) or state level (Medicaid). We used validated measures to characterize census tracts as low education ($\geq 25\%$ of individuals older than age 25 years did not graduate high school) or high poverty ($\geq 10\%$ of households below poverty level).²⁵⁻²⁷

For the Parkland cohort, we repeated analyses with the addition of whether the patient listed a missing or non-sensical social security number as a proxy for undocumented immigration status as we hypothesized this could influence ED utilization. Social security numbers were not reliably identifiable in the University EHR.

Statistical analyses were performed using SAS 9.4 (SAS Institute, Cary, NC) and Stata/SE 15.0 (StataCorp, College Station, TX). The University of Texas Southwestern Medical Center institutional review board approved this study (STU 112017-026 and 122017-042).

RESULTS

Patient Characteristics

We matched 35,090 patients from the University and Parkland cancer registries to the Foundation database. Half (49.4%) were female, one-third (33.7%) were ≥ 65 years old, 18.3% had a non-English language preference, 30.0%

were uninsured or enrolled in Medicaid at the time of diagnosis, and 0.8% were homeless. The most common cancers were breast (14.2%), lung (10.9%), GI (10.6%), prostate (10.2%), and colorectal (5.9%). Overall, one quarter (24.2%) of individuals had advanced-stage cancer at diagnosis, 41.6% had initial treatment regimens that included chemotherapy, and 5.4% died within 6 months after diagnosis.

More than half of the cohort (57.2%) had at least one ED visit in the 6 months after cancer diagnosis, whereas one-fifth (19.5%) had at least one ED visit in the 6-12 months prior to cancer diagnosis. Complete demographics, cancer-related characteristics, and ED visit counts are shown in [Table 1](#).

Predictors of ED Visits After Cancer Diagnosis

In our multivariable adjusted regression analysis, the strongest independent predictor of postdiagnosis ED visits was frequent (≥ 4) prediagnosis ED visits (adjusted incidence rate ratio [aIRR]: 3.68; 95% CI, 3.36 to 4.02) ([Fig 1](#), [Appendix Table A1](#), online only). Patients with 1-3 prediagnosis ED visits also had more ED visits following cancer diagnosis, although to a lesser extent (aIRR: 1.32; 95% CI, 1.28 to 1.36).

SDH variables associated with greater postdiagnosis ED use included Hispanic (aIRR: 1.12; 95% CI, 1.07 to 1.17) and Black (aIRR: 1.21; 95% CI, 1.17 to 1.25) race and/or ethnicity, Medicaid (aIRR: 1.45; 95% CI, 1.39 to 1.52) and county charity assistance (aIRR: 1.32; 95% CI, 1.27 to 1.38) insurance types, and homelessness (aIRR: 1.95; 95% CI, 1.73 to 2.20). Non-English language preference was not significantly associated with postdiagnosis ED visits.

Among the clinical covariates, we observed greater postdiagnosis ED use for individuals with lung cancer (aIRR: 1.26; 95% CI, 1.19 to 1.34) and colorectal cancer (aIRR: 1.19; 95% CI, 1.11 to 1.28), advanced-stage cancer at diagnosis (aIRR: 1.30; 95% CI, 1.26 to 1.35), and initial treatment regimens that included chemotherapy (aIRR: 1.44; 95% CI, 1.40 to 1.48). Breast cancer was associated with fewer postdiagnosis ED visits (aIRR: 0.91; 95% CI, 0.86 to 0.97). Full model results for the overall cohort are shown in [Appendix Table A1](#). [Appendix Tables A2](#) and [A3](#) (online only) contain full model results stratified by health system.

Marginally Adjusted ED Visit Counts

In our adjusted marginal effects analysis, patients with infrequent (1-3) and frequent (≥ 4) prediagnosis ED visits had 0.33 (95% CI, 0.29 to 0.37) and 2.77 (95% CI, 2.43 to 3.10) more postdiagnosis ED visits, respectively, than patients with zero prediagnosis ED visits ([Table 2](#)).

Sensitivity Analyses

Our findings were not substantively changed when modeled for the two health system cohorts separately, nor in

TABLE 1. Characteristics of Adults With Newly Diagnosed Cancer Across Two Health Systems^a

Characteristic	University Cohort (n = 26,493)	Parkland Cohort (n = 8,597)	Overall Cohort (N = 35,090)
Age ≥ 65 years	38.3	19.3	33.7
Female	47.6	54.9	49.4
Race and/or ethnicity			
Non-Hispanic White	59.2	22.6	50.2
Hispanic	15.3	38.3	20.9
Black	19.4	33.7	22.9
Others	6.1	5.5	6.0
Non-English language preference	13.4	33.4	18.3
Insurance type			
Commercial or other government payer	47.7	4.2	37.1
Medicare	36.5	14.4	31.1
Medicaid	6.7	34.2	13.4
County charity assistance (uninsured)	8.6	41.3	16.6
Unknown	0.5	5.9	1.8
Homeless	0.1	3.1	0.8
Mean (SD) Charlson Comorbidity Index at diagnosis	1.7 (3.0)	0.8 (1.4)	1.5 (2.7)
Cancer type			
Lung	11.4	9.3	10.9
Breast	13.6	16.0	14.2
Colorectal	6.0	5.5	5.9
GI	9.7	13.5	10.6
Head and neck	7.3	4.6	6.6
Brain	4.5	1.8	3.8
Kidney	5.5	4.8	5.4
Prostate	11.4	6.7	10.2
Cervical, uterine, ovarian, and vaginal	6.1	11.3	7.4
Lymphoma	8.3	7.2	8.1
Others	16.2	19.2	16.9
Advanced cancer stage at diagnosis	23.6	25.9	24.2
Died within 180 days after diagnosis	4.0	9.8	5.4
Initial cancer treatment modality			
Surgery	54.4	50.5	53.5
Radiation therapy	35.8	27.9	33.9
Chemotherapy	41.7	41.2	41.6
Immunotherapy	5.9	6.3	6.0
No. of ED visits 6-12 months before cancer diagnosis			
0	81.7	76.8	80.5
1-3	17.2	21.0	18.2
≥ 4	1.1	2.2	1.4
No. of ED visits 6 months after cancer diagnosis			
0	43.0	42.4	42.8
1-3	48.9	45.7	48.1
≥ 4	8.2	11.8	9.1

Abbreviations: ED, emergency department; SD, standard deviation.

^aData presented as % unless otherwise specified.

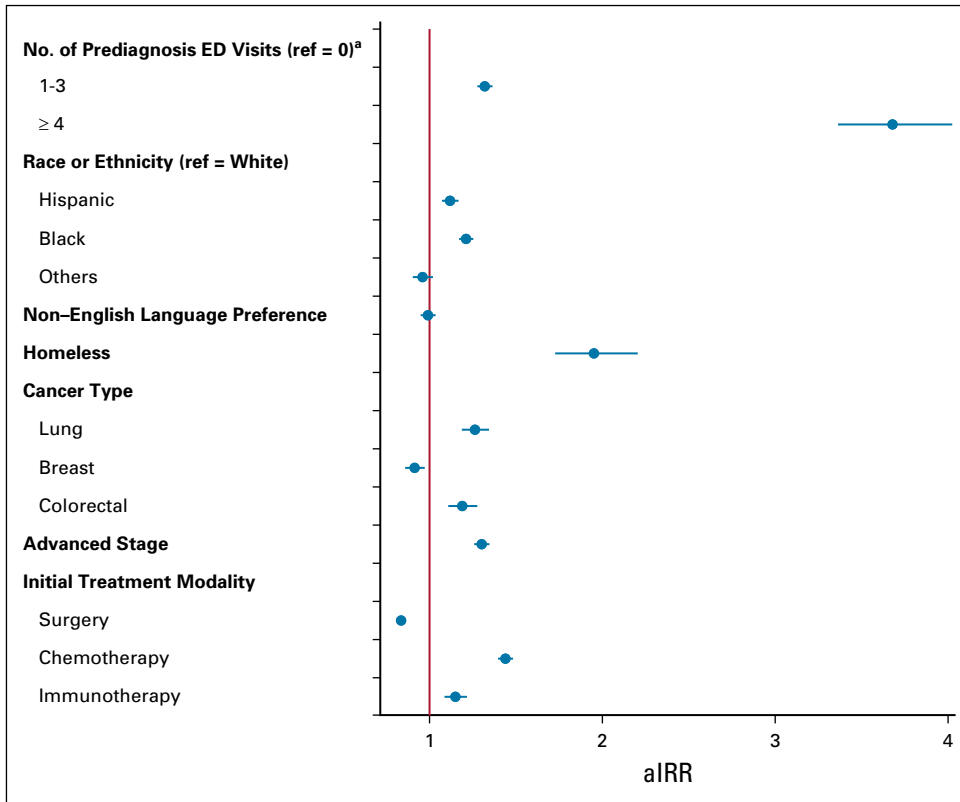


FIG 1. Key predictors of number of ED visits 6 months after cancer diagnosis. Shown are incidence rate ratios and 95% CIs of ED visits within 6 months after cancer diagnosis generated from a multivariable negative binomial regression adjusted for health system (Parkland v University), age, sex, race and/or ethnicity, language preference, insurance type, homelessness, Charlson Comorbidity Index at time of diagnosis, year of cancer diagnosis, cancer type, advanced cancer stage at diagnosis, death within 6 months of diagnosis, and initial cancer treatment modalities. Full model results are shown in Appendix Table A1. ^aNumber of ED visits within 6-12 months before cancer diagnosis. aIRR, adjusted incidence rate ratio; ED, emergency department; ref, reference.

sensitivity analyses that (1) used ED visit history 12-18 months prior to cancer diagnosis as the primary predictor, (2) included interaction terms between initial treatment modalities and cancer stage, (3) adjusted for neighborhood education level, (4) excluded pre- and post-cancer diagnosis ED visits that resulted in hospitalization, (5) adjusted for neighborhood poverty level, and (6) adjusted for whether a social security number was missing in the EHR.

See Appendix Tables A2 and A3 for complete sensitivity analysis results.

DISCUSSION

In this retrospective analysis of 35,090 adults with newly diagnosed cancer across two health systems with diverse populations, we identified a strong association between ED use before cancer diagnosis and ED use after cancer diagnosis. Patients with frequent (≥ 4) ED visits in the 6 months prior to cancer diagnosis averaged nearly three more postdiagnosis ED visits than patients with zero pre-diagnosis ED visits. Our estimates are likely conservative because we excluded ED visits in the 6 months leading up to cancer diagnosis. Furthermore, our results were robust to a number of sensitivity analyses. We also confirmed known risk factors of postdiagnosis ED visits such as minority race and/or ethnicity, homelessness, advanced-staged cancer, and treatment regimens including chemotherapy and radiation therapy.²⁸⁻³⁰

Our most practice- and policy-relevant finding is the strength of a patient’s prior ED visit history in predicting ED

TABLE 2. Predicted Number of ED Visits Within 6 Months After Cancer Diagnosis Adjusted Difference in No. of Postdiagnosis ED Visits (95% CI)

No. of Prediagnosis ED Visits ^a	Univariate	Multivariable
	Ref	Ref
0	Ref	Ref
1-3	0.38 (0.33 to 0.43)	0.33 (0.29 to 0.37)*
≥ 4	3.11 (2.69 to 3.53)	2.77 (2.43 to 3.10)*

NOTE. * $P < .001$.

Abbreviations: ED, emergency department; ref, reference.

^aNumber of ED visits within 6-12 months before cancer diagnosis.

visits in the first 6 months after an incident cancer diagnosis. Adding to prior studies that identified prior acute care use as a predictor of acute care use after initiation of chemotherapy or radiation therapy,²⁹⁻³³ our study sampled a diverse population of patients (half non-White), including patients without insurance and who had not undergone chemotherapy or radiation therapy, and accounted for SDH variables. Frequent prediagnosis ED visits was by far the strongest predictor of postdiagnosis ED visit volume, with a predictive impact that nearly doubled those of the strongest clinical or SDH predictors. Recognizing that sociodemographic and cancer-related factors are often beyond the control of a patient or health system, and that many ED visits may be clinically necessary, frequent prediagnosis ED use may be a marker for potentially modifiable ED-seeking behavior. Although we account only for certain SDH measurable in the EHR, prediagnosis ED use was also strongly predictive for the University cohort. We would not expect the prevalence of only the unmeasured proxy SDH to be disproportionately higher in the University cohort. Although some faculty physicians practice at the University and supervise at Parkland as well, we have low concern for correlation between the health systems given that clinic staff that provide acute care triage advice work exclusively at one health system.

We suspect that patients with prior frequent ED use are exhibiting a behavior reinforced by interactions with the health care system. Patients that frequently visit the ED may find certain features of ED visits favorable to the alternatives (eg, urgent clinic visit), and some may be unaware of alternatives altogether. Although ED visits are accompanied by longer visit times³⁴ and higher cost-sharing,^{35,36} the ED is open 24 hours, does not require a scheduled appointment, does not rely on patients to determine the severity of their own medical condition, and is the only site of guaranteed evaluation and clinical stabilization regardless of ability to pay.³⁷ Health systems might use this information to develop educational initiatives that target heavy prior ED users and offer them alternative ways to seek urgent medical advice.

To date, policy efforts discouraging patients from inappropriate ED visits have relied on post hoc financial penalties, with some insurers requiring higher cost-sharing for ED visits that, with the benefit of hindsight, are deemed avoidable.^{38,39} Health care providers and policymakers keen on reducing ED visits among patients with cancer should be careful not to ignore that the patient's decision to visit the ED is made under a great deal of uncertainty. Efforts to reduce ED use by patients with cancer should include strategies to educate patients about the best ways to judge how and where to receive care when sick (eg, urgent telephone advice and ED v outpatient cancer urgent care visit) with a sensitivity to understanding a patient's prior experiences. Although cancer centers often provide instructions about when to call, come to the clinic, or go to the ED, more in-depth educational approaches may be needed for the subgroup of patients who are avid ED users

prior to cancer diagnosis. Existing 24-hour telephone triage lines could be emphasized, as well as ED alternatives such as oncology urgent care clinics.⁴⁰ Such an effort may be more effective and patient-centered than issuing post hoc financial penalties.

Our findings should be interpreted in the context of several limitations. First, this is an observational study, and there may be unmeasured confounders affecting our results. We were unable to adjust for presenting complaint, which prior studies have shown is closely tied to acute care use.^{28,41,42} However, we carefully adjusted for many of the important predictors of illness severity and reliably measurable SDH. Second, of the original 9,050 Parkland patients and 31,048 University patients, 448 Parkland (5.0%) and 4,555 University (14.7%) patients were not matched to the Foundation database. Most patients were likely unmatched because of data entry errors and discrepancies. Some patients may not have matched because of recent in-migration to the region. However, a selective lack of prediagnosis ED visits would tend to underestimate the magnitude of our prediagnosis ED visit predictor. It is also possible that the matching process combined two different patients with the same names, dates of birth, and zip codes of residence, which would bias our results in both directions. We suspect the frequency of this type of mismatch to be exceedingly low given the multiple levels of detail used to match patients. Third, we did not distinguish which ED visits after cancer diagnosis could have been avoidable—addressable in either an outpatient or specialized oncology urgent care clinic. As to the generalizability of our findings, we captured a broad array of cancer types within two distinct health systems with diverse patient populations over a 10-year time span. Conducting similar analyses across a wider geographic range would only be possible with a specific subset of insured enrollees (eg, Medicare fee-for-service) rather than our all-ages, population-level analysis and would exclude uninsured patients. Additionally, the EHR-based measures of SDH that we used are much less widely available in national-level administrative data sets. However, our findings may not be completely reflective of patients who had synchronous or metachronous cancers.

As part of efforts to reduce potentially avoidable ED visits among a population at high risk, we should take care to understand the behavioral and experiential components of patients seeking acute care. Frequent ED use prior to cancer diagnosis may serve as a surrogate marker that we can use to identify patients likely to have frequent postdiagnosis ED visits. Further research is needed to understand the reasons that underpin patient decisions to seek ED care prior to cancer diagnosis, including qualitative studies exploring the philosophy of seeking on-demand care, cultural and/or community norms, and previous experiences with the health care system. Health systems will also need to provide more timely and patient-friendly sick care telephone advice and accessible ED alternatives such as specialized oncology urgent care clinics.

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DISCLAIMER

The content is solely the responsibility of the authors and does not necessarily represent the official views of Texas Health Resources, the University of Texas Southwestern Medical Center, the National Institutes of Health, or the Agency for Healthcare Research and Quality. The funders had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; and preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

EQUAL CONTRIBUTION

A.S.H. and D.Q.N. contributed equally to this work as first co-authors.

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST**Prior Frequent Emergency Department Use as a Predictor of Emergency Department Visits After a New Cancer Diagnosis**

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APPENDIX

TABLE A1. Negative Binomial Regression Analysis for the Overall Cohort (N = 35,090)

Covariate	Univariate IRR (95% CI)	Multivariable aIRR (95% CI)
Age, years		
18-24	Ref	Ref
25-34	1.06 (0.92 to 1.21)	0.98 (0.86 to 1.11)
35-44	1.05 (0.93 to 1.19)	0.98 (0.87 to 1.10)
45-54	1.06 (0.94 to 1.20)	0.94 (0.84 to 1.06)
55-64	0.99 (0.88 to 1.11)	0.90 (0.80 to 1.00)*
≥ 65	0.82 (0.73 to 0.93)	0.83 (0.74 to 0.93)***
Female	1.07 (1.04 to 1.10)	1.02 (0.99 to 1.06)
Race or ethnicity		
Non-Hispanic White	Ref	Ref
Hispanic	1.28 (1.24 to 1.33)	1.12 (1.07 to 1.17)***
Black	1.45 (1.40 to 1.50)	1.21 (1.17 to 1.25)***
Others	0.97 (0.91 to 1.03)	0.96 (0.90 to 1.02)
Non-English language preference	1.07 (1.03 to 1.11)	0.99 (0.95 to 1.04)
Insurance type		
Commercial	Ref	Ref
Medicare	1.11 (1.08 to 1.15)	1.17 (1.12 to 1.22)***
Medicaid	1.71 (1.64 to 1.78)	1.45 (1.39 to 1.52)***
County charity assistance (uninsured)	1.47 (1.41 to 1.53)	1.32 (1.27 to 1.38)***
Unknown	1.21 (1.09 to 1.34)	1.18 (1.06 to 1.30)***
Homeless	2.63 (2.30 to 3.01)	1.95 (1.73 to 2.20)***
Charlson Comorbidity Index at diagnosis, per 1 unit	1.03 (1.03 to 1.04)	1.02 (1.02 to 1.03)***
Year of cancer diagnosis		
2008	Ref	Ref
2009	1.11 (1.01 to 1.23)	1.04 (0.95 to 1.14)
2010	1.22 (1.12 to 1.34)	1.18 (1.08 to 1.29)***
2011	1.23 (1.12 to 1.35)	1.16 (1.07 to 1.27)***
2012	1.32 (1.22 to 1.44)	1.19 (1.10 to 1.29)***
2013	1.36 (1.25 to 1.48)	1.23 (1.13 to 1.33)***
2014	1.35 (1.24 to 1.47)	1.22 (1.12 to 1.32)***
2015	1.33 (1.22 to 1.44)	1.19 (1.10 to 1.29)***
2016	1.37 (1.27 to 1.49)	1.21 (1.11 to 1.31)***
2017	0.93 (0.86 to 1.01)	0.94 (0.87 to 1.02)
2018	0.77 (0.70 to 0.84)	0.86 (0.78 to 0.94)***
Cancer type		
Lung	1.46 (1.40 to 1.52)	1.26 (1.19 to 1.34)***
Breast	0.79 (0.76 to 0.83)	0.91 (0.86 to 0.97)***
Colorectal	1.14 (1.08 to 1.21)	1.19 (1.11 to 1.28)***
Head and neck	1.23 (1.17 to 1.30)	1.20 (1.12 to 1.28)***
Melanoma	0.49 (0.44 to 0.53)	0.85 (0.77 to 0.94)***
Brain	1.02 (0.95 to 1.10)	1.34 (1.23 to 1.45)***

(continued on following page)

TABLE A1. Negative Binomial Regression Analysis for the Overall Cohort (N = 35,090) (continued)

Covariate	Univariate IRR (95% CI)	Multivariable aIRR (95% CI)
Kidney	0.81 (0.76 to 0.86)	1.21 (1.12 to 1.30)***
Prostate	0.48 (0.45 to 0.50)	0.78 (0.73 to 0.84)***
Other GU	0.90 (0.84 to 0.97)	1.35 (1.24 to 1.46)***
Lymphoma	1.10 (1.04 to 1.15)	1.05 (0.98 to 1.12)
Cervical, uterine, ovarian, and vaginal	1.27 (1.20 to 1.33)	1.30 (1.22 to 1.39)***
Pancreas	1.39 (1.29 to 1.51)	1.36 (1.24 to 1.48)***
Other GI	1.33 (1.27 to 1.40)	1.31 (1.23 to 1.40)***
Advanced cancer stage at diagnosis	1.71 (1.65 to 1.76)	1.30 (1.26 to 1.35)***
Died within 180 days after diagnosis	1.58 (1.49 to 1.67)	1.30 (1.24 to 1.38)***
Initial cancer treatment modality		
Surgery	0.65 (0.63 to 0.67)	0.84 (0.81 to 0.86)***
Radiation therapy	1.40 (1.36 to 1.44)	1.18 (1.15 to 1.22)***
Chemotherapy	1.72 (1.68 to 1.77)	1.44 (1.40 to 1.48)***
Immunotherapy	1.20 (1.13 to 1.27)	1.15 (1.09 to 1.22)***
No. of prediagnosis ED visits ^a		
0	Ref	Ref
1-3	1.32 (1.28 to 1.37)	1.32 (1.28 to 1.36)***
≥ 4	3.67 (3.33 to 4.05)	3.68 (3.36 to 4.02)***
Cohort		
University	Ref	Ref
Parkland	1.21 (1.17 to 1.25)	0.84 (0.81 to 0.88)***

NOTE. Boldface indicates a statistically significant aIRR: * $P < .05$, *** $P < .001$.

Abbreviations: aIRR, adjusted incidence rate ratio; ED, emergency department; GU, genitourinary; IRR, incidence rate ratio; ref, reference.

^aNumber of ED visits within 6-12 months before cancer diagnosis.

TABLE A2. Negative Binomial Regression and SAs for the University Cohort (n = 26,493)

Covariate	Multivariable aIRR (95% CI)								
	Unadjusted IRR (95% CI)	Adjusted Model ^a	SA 1*	SA 2*	SA 3*	SA 4*	SA 5*	SA 6*	SA 7*
Age, years	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
18-24									
25-34	1.03 (0.88 to 1.20)	0.97 (0.84 to 1.12)	0.99 (0.85 to 1.14)	0.97 (0.80 to 1.17)	0.97 (0.84 to 1.12)	0.98 (0.85 to 1.13)	0.98 (0.85 to 1.13)	0.97 (0.84 to 1.12)	0.99 (0.83 to 1.19)
35-44	1.09 (0.94 to 1.26)	1.00 (0.87 to 1.14)	1.03 (0.90 to 1.18)	1.04 (0.87 to 1.24)	1.00 (0.87 to 1.14)	1.00 (0.88 to 1.15)	1.01 (0.88 to 1.15)	1.00 (0.87 to 1.14)	1.01 (0.85 to 1.19)
45-54	1.05 (0.91 to 1.21)	0.94 (0.83 to 1.08)	0.97 (0.85 to 1.11)	0.95 (0.79 to 1.13)	0.94 (0.83 to 1.08)	0.95 (0.83 to 1.08)	0.95 (0.83 to 1.08)	0.94 (0.83 to 1.08)	0.95 (0.80 to 1.12)
55-64	0.94 (0.81 to 1.08)	0.87 (0.77 to 1.00)**	0.89 (0.78 to 1.02)	0.90 (0.76 to 1.07)	0.87 (0.77 to 1.00)**	0.88 (0.77 to 1.00)**	0.88 (0.77 to 1.00)**	0.87 (0.77 to 1.00)**	0.84 (0.71 to 0.98)**
≥ 65	0.83 (0.73 to 0.96)	0.84 (0.73 to 0.96)**	0.85 (0.74 to 0.98)**	0.88 (0.74 to 1.05)	0.84 (0.73 to 0.96)**	0.84 (0.74 to 0.96)**	0.84 (0.74 to 0.97)**	0.84 (0.73 to 0.96)**	0.77 (0.65 to 0.92)**
Female	1.12 (1.08 to 1.15)	1.03 (0.99 to 1.07)	1.03 (0.99 to 1.07)	1.01 (0.96 to 1.05)	1.03 (0.99 to 1.07)	1.03 (0.99 to 1.07)	1.03 (0.99 to 1.07)	1.03 (0.99 to 1.07)	1.03 (0.98 to 1.08)
Race or ethnicity	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
Non-Hispanic White									
Hispanic	1.50 (1.43 to 1.56)	1.15 (1.10 to 1.21)**	1.16 (1.10 to 1.22)**	1.10 (1.04 to 1.18)**	1.15 (1.10 to 1.21)**	1.15 (1.10 to 1.21)**	1.15 (1.10 to 1.21)**	1.15 (1.10 to 1.21)**	1.28 (1.20 to 1.36)**
Black	1.51 (1.45 to 1.57)	1.23 (1.19 to 1.28)**	1.25 (1.20 to 1.30)**	1.19 (1.13 to 1.25)**	1.23 (1.19 to 1.28)**	1.23 (1.18 to 1.28)**	1.23 (1.19 to 1.28)**	1.23 (1.19 to 1.28)**	1.30 (1.24 to 1.37)**
Others	1.00 (0.93 to 1.07)	0.95 (0.89 to 1.01)	0.95 (0.89 to 1.02)	0.99 (0.91 to 1.09)	0.95 (0.89 to 1.01)	0.95 (0.89 to 1.01)	0.95 (0.89 to 1.01)	0.95 (0.89 to 1.01)	0.95 (0.87 to 1.03)
Non-English language preference	1.31 (1.26 to 1.37)	1.12 (1.06 to 1.18)**	1.12 (1.06 to 1.18)**	1.14 (1.07 to 1.21)**	1.12 (1.06 to 1.18)**	1.12 (1.06 to 1.18)**	1.11 (1.05 to 1.17)**	1.12 (1.06 to 1.18)**	1.05 (0.98 to 1.12)
Insurance type	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
Commercial									
Medicare	1.09 (1.06 to 1.13)	1.16 (1.10 to 1.21)**	1.16 (1.11 to 1.21)**	1.09 (1.03 to 1.15)**	1.16 (1.10 to 1.21)**	1.16 (1.11 to 1.21)**	1.15 (1.10 to 1.21)**	1.16 (1.10 to 1.21)**	1.15 (1.08 to 1.22)**
Medicaid	2.05 (1.93 to 2.17)	1.49 (1.41 to 1.58)**	1.52 (1.43 to 1.60)**	1.42 (1.32 to 1.51)**	1.49 (1.41 to 1.58)**	1.49 (1.41 to 1.58)**	1.48 (1.40 to 1.57)**	1.49 (1.41 to 1.57)**	1.55 (1.44 to 1.66)**
Uninsured	1.64 (1.55 to 1.73)	1.32 (1.26 to 1.40)**	1.31 (1.25 to 1.39)**	1.27 (1.19 to 1.35)**	1.32 (1.26 to 1.40)**	1.32 (1.26 to 1.40)**	1.32 (1.25 to 1.39)**	1.32 (1.26 to 1.40)**	1.38 (1.29 to 1.48)**
Unknown	1.16 (0.93 to 1.45)	1.09 (0.88 to 1.34)	1.06 (0.86 to 1.31)	1.11 (0.85 to 1.45)	1.09 (0.88 to 1.34)	1.09 (0.88 to 1.34)	1.08 (0.88 to 1.34)	1.09 (0.88 to 1.34)	1.12 (0.85 to 1.46)
Homeless	2.92 (1.73 to 4.93)	2.31 (1.48 to 3.63)**	2.34 (1.49 to 3.69)**	2.59 (1.46 to 4.60)**	2.32 (1.48 to 3.64)**	2.32 (1.48 to 3.63)**	2.25 (1.44 to 3.53)**	2.32 (1.48 to 3.64)**	2.84 (1.67 to 4.84)**
Charlson Comorbidity Index, per 1 unit	1.04 (1.03 to 1.05)	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.02 to 1.03)**	1.02 (1.01 to 1.02)**
Year of cancer diagnosis	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
2008									
2009	1.11 (1.01 to 1.22)	1.06 (0.97 to 1.16)	1.07 (0.97 to 1.17)	1.10 (0.98 to 1.22)	1.06 (0.97 to 1.16)	1.06 (0.97 to 1.16)	1.06 (0.97 to 1.16)	1.06 (0.97 to 1.16)	1.10 (0.97 to 1.23)
2010	1.22 (1.12 to 1.34)	1.22 (1.12 to 1.33)**	1.21 (1.11 to 1.32)**	1.25 (1.13 to 1.39)**	1.22 (1.12 to 1.33)**	1.22 (1.12 to 1.33)**	1.21 (1.11 to 1.32)**	1.22 (1.12 to 1.33)**	1.29 (1.15 to 1.44)**
2011	1.23 (1.13 to 1.35)	1.21 (1.11 to 1.32)**	1.24 (1.14 to 1.35)**	1.23 (1.11 to 1.37)**	1.22 (1.12 to 1.32)**	1.21 (1.11 to 1.32)**	1.20 (1.10 to 1.31)**	1.21 (1.11 to 1.32)**	1.33 (1.19 to 1.49)**
2012	1.26 (1.16 to 1.38)	1.22 (1.12 to 1.33)**	1.23 (1.13 to 1.34)**	1.21 (1.09 to 1.34)**	1.22 (1.12 to 1.33)**	1.22 (1.12 to 1.33)**	1.21 (1.11 to 1.32)**	1.22 (1.12 to 1.33)**	1.32 (1.19 to 1.48)**
2013	1.32 (1.21 to 1.44)	1.30 (1.20 to 1.42)**	1.32 (1.21 to 1.44)**	1.34 (1.21 to 1.49)**	1.30 (1.20 to 1.42)**	1.30 (1.20 to 1.42)**	1.29 (1.19 to 1.41)**	1.30 (1.20 to 1.42)**	1.37 (1.23 to 1.52)**
2014	1.29 (1.18 to 1.40)	1.24 (1.14 to 1.35)**	1.25 (1.15 to 1.36)**	1.33 (1.20 to 1.48)**	1.24 (1.14 to 1.35)**	1.24 (1.14 to 1.35)**	1.24 (1.14 to 1.35)**	1.24 (1.14 to 1.35)**	1.38 (1.24 to 1.54)**
2015	1.32 (1.21 to 1.44)	1.26 (1.16 to 1.37)**	1.28 (1.17 to 1.39)**	1.27 (1.15 to 1.41)**	1.26 (1.16 to 1.37)**	1.26 (1.16 to 1.37)**	1.25 (1.15 to 1.36)**	1.26 (1.16 to 1.37)**	1.41 (1.27 to 1.57)**
2016	1.35 (1.24 to 1.47)	1.26 (1.16 to 1.37)**	1.30 (1.20 to 1.41)**	1.30 (1.18 to 1.44)**	1.26 (1.16 to 1.37)**	1.26 (1.16 to 1.37)**	1.26 (1.16 to 1.37)**	1.26 (1.16 to 1.37)**	1.42 (1.28 to 1.58)**
2017	0.93 (0.86 to 1.01)	0.99 (0.91 to 1.07)	0.99 (0.91 to 1.08)	1.03 (0.93 to 1.14)	0.99 (0.91 to 1.07)	0.99 (0.91 to 1.07)	0.98 (0.90 to 1.06)	0.99 (0.91 to 1.07)	1.08 (0.97 to 1.20)
2018	0.77 (0.70 to 0.84)	0.90 (0.83 to 0.99)**	0.91 (0.83 to 1.00)**	0.93 (0.83 to 1.04)	0.90 (0.83 to 0.99)**	0.90 (0.83 to 0.99)**	0.91 (0.83 to 0.99)**	0.90 (0.83 to 0.99)**	1.01 (0.90 to 1.13)
Cancer type	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
Lung	1.46 (1.40 to 1.54)	1.30 (1.21 to 1.41)**	1.30 (1.20 to 1.40)**	1.28 (1.16 to 1.41)**	1.30 (1.21 to 1.41)**	1.31 (1.21 to 1.41)**	1.29 (1.19 to 1.39)**	1.30 (1.20 to 1.41)**	1.24 (1.12 to 1.37)**
Breast	0.85 (0.81 to 0.89)	0.95 (0.88 to 1.03)	0.95 (0.88 to 1.03)	0.97 (0.87 to 1.07)	0.96 (0.88 to 1.03)	0.95 (0.88 to 1.03)	0.94 (0.86 to 1.01)*	0.96 (0.88 to 1.03)	1.10 (0.99 to 1.21)*
Colorectal	1.20 (1.13 to 1.28)	1.24 (1.13 to 1.35)**	1.25 (1.15 to 1.36)**	1.21 (1.09 to 1.35)**	1.24 (1.14 to 1.35)**	1.24 (1.14 to 1.35)**	1.21 (1.11 to 1.32)**	1.24 (1.14 to 1.35)**	1.18 (1.05 to 1.32)**
Head and neck	1.24 (1.17 to 1.32)	1.22 (1.12 to 1.33)**	1.24 (1.14 to 1.35)**	1.18 (1.06 to 1.31)**	1.22 (1.12 to 1.33)**	1.23 (1.13 to 1.34)**	1.22 (1.12 to 1.32)**	1.22 (1.12 to 1.32)**	1.45 (1.30 to 1.61)**
Melanoma	0.49 (0.45 to 0.54)	0.86 (0.77 to 0.96)**	0.86 (0.77 to 0.96)**	0.97 (0.84 to 1.11)	0.86 (0.77 to 0.96)**	0.87 (0.78 to 0.97)**	0.87 (0.78 to 0.97)**	0.86 (0.77 to 0.96)**	0.96 (0.84 to 1.11)
Brain	1.02 (0.95 to 1.10)	1.33 (1.21 to 1.46)**	1.35 (1.23 to 1.48)**	1.26 (1.12 to 1.42)**	1.33 (1.22 to 1.46)**	1.34 (1.22 to 1.48)**	1.31 (1.19 to 1.44)**	1.33 (1.22 to 1.46)**	1.38 (1.23 to 1.55)**
Kidney	0.79 (0.73 to 0.85)	1.22 (1.11 to 1.34)**	1.21 (1.11 to 1.33)**	1.16 (1.02 to 1.30)**	1.22 (1.11 to 1.34)**	1.24 (1.13 to 1.36)**	1.23 (1.12 to 1.35)**	1.22 (1.11 to 1.34)**	1.28 (1.14 to 1.44)**
Prostate	0.47 (0.44 to 0.49)	0.81 (0.74 to 0.88)**	0.81 (0.74 to 0.88)**	0.79 (0.71 to 0.89)**	0.81 (0.74 to 0.88)**	0.81 (0.74 to 0.88)**	0.82 (0.75 to 0.89)**	0.81 (0.74 to 0.88)**	1.04 (0.93 to 1.16)

(continued on following page)

TABLE A2. Negative Binomial Regression and SAs for the University Cohort (n = 26,493) (continued)

Covariate	Multivariable aIRR (95% CI)								
	Unadjusted IRR (95% CI)	Adjusted Model ^a	SA 1 ^b	SA 2 ^c	SA 3 ^d	SA 4 ^e	SA 5 ^f	SA 6 ^g	SA 7 ^h
Other GU	0.89 (0.83 to 0.96)	1.37 (1.25 to 1.51)***	1.38 (1.26 to 1.52)***	1.39 (1.23 to 1.57)***	1.37 (1.25 to 1.51)***	1.38 (1.25 to 1.51)***	1.36 (1.24 to 1.50)***	1.37 (1.25 to 1.51)***	1.46 (1.30 to 1.65)***
Lymphoma	1.08 (1.02 to 1.14)	1.11 (1.02 to 1.21)**	1.12 (1.03 to 1.21)**	1.12 (1.01 to 1.24)**	1.11 (1.02 to 1.21)**	1.10 (1.01 to 1.20)**	1.08 (0.99 to 1.17)*	1.11 (1.02 to 1.21)**	1.02 (0.91 to 1.13)
Cervical, uterine, ovarian, and vaginal	1.36 (1.28 to 1.45)	1.33 (1.22 to 1.45)***	1.35 (1.24 to 1.47)***	1.30 (1.17 to 1.44)***	1.33 (1.22 to 1.45)***	1.34 (1.23 to 1.46)***	1.30 (1.20 to 1.42)***	1.33 (1.22 to 1.45)***	1.52 (1.36 to 1.69)***
Pancreas	1.34 (1.22 to 1.46)	1.39 (1.25 to 1.54)***	1.40 (1.26 to 1.55)***	1.30 (1.14 to 1.48)***	1.39 (1.25 to 1.54)***	1.38 (1.25 to 1.54)***	1.37 (1.23 to 1.52)***	1.39 (1.26 to 1.54)***	1.27 (1.10 to 1.45)***
Other GI	1.36 (1.28 to 1.44)	1.42 (1.30 to 1.54)***	1.44 (1.32 to 1.56)***	1.41 (1.27 to 1.57)***	1.42 (1.31 to 1.54)***	1.42 (1.30 to 1.54)***	1.39 (1.28 to 1.51)***	1.42 (1.30 to 1.54)***	1.31 (1.18 to 1.46)***
Advanced cancer stage at diagnosis	1.73 (1.67 to 1.79)	1.31 (1.26 to 1.36)***	1.30 (1.25 to 1.35)***	1.32 (1.26 to 1.38)***	1.31 (1.26 to 1.36)***	1.30 (1.25 to 1.35)***	1.59 (1.50 to 1.68)***	1.29 (1.22 to 1.35)***	1.19 (1.13 to 1.25)***
Died within 180 days of diagnosis	1.61 (1.49 to 1.73)	1.37 (1.28 to 1.47)***	1.37 (1.28 to 1.47)***	1.34 (1.22 to 1.46)***	1.37 (1.28 to 1.47)***	1.37 (1.28 to 1.47)***	1.34 (1.25 to 1.44)***	1.37 (1.28 to 1.47)***	0.99 (0.90 to 1.09)
Initial cancer treatment modality									
Surgery	0.67 (0.65 to 0.69)	0.87 (0.84 to 0.90)***	0.86 (0.83 to 0.90)***	0.85 (0.81 to 0.89)***	0.88 (0.84 to 0.92)***	0.87 (0.84 to 0.90)***	0.87 (0.84 to 0.90)***	0.87 (0.84 to 0.90)***	0.99 (0.95 to 1.04)
Radiation therapy	1.55 (1.50 to 1.60)	1.21 (1.17 to 1.25)***	1.21 (1.17 to 1.25)***	1.19 (1.14 to 1.25)***	1.21 (1.17 to 1.25)***	1.27 (1.21 to 1.33)***	1.19 (1.15 to 1.23)***	1.19 (1.15 to 1.24)***	1.26 (1.20 to 1.31)***
Chemotherapy	1.77 (1.71 to 1.82)	1.39 (1.34 to 1.44)***	1.38 (1.34 to 1.43)***	1.39 (1.33 to 1.45)***	1.41 (1.34 to 1.47)***	1.45 (1.39 to 1.52)***	1.55 (1.48 to 1.61)***	1.39 (1.35 to 1.44)***	1.33 (1.27 to 1.39)***
Immunotherapy	1.20 (1.12 to 1.28)	1.17 (1.09 to 1.24)***	1.16 (1.08 to 1.23)***	1.15 (1.06 to 1.25)***	1.17 (1.09 to 1.24)***	1.16 (1.09 to 1.24)***	1.16 (1.09 to 1.24)***	1.17 (1.09 to 1.24)***	1.20 (1.10 to 1.30)***
No. of prediagnosis ED visits ⁱ		Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
0	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
1-3	1.25 (1.20 to 1.30)	1.24 (1.19 to 1.28)***	1.23 (1.18 to 1.28)***	1.24 (1.18 to 1.30)***	1.24 (1.19 to 1.28)***	1.24 (1.19 to 1.28)***	1.24 (1.20 to 1.29)***	1.24 (1.19 to 1.28)***	1.43 (1.36 to 1.51)***
≥ 4	3.83 (3.39 to 4.33)	4.09 (3.66 to 4.56)***	4.46 (3.89 to 5.12)***	4.58 (4.00 to 5.24)***	4.08 (3.66 to 4.56)***	4.08 (3.66 to 4.56)***	4.13 (3.70 to 4.61)***	4.08 (3.66 to 4.56)***	7.17 (6.08 to 8.46)***
Additional covariates									
Low neighborhood education	—	—	—	1.08 (1.02 to 1.13)***	—	—	—	—	—
High neighborhood poverty	—	—	—	1.03 (0.98 to 1.08)	—	—	—	—	—
Chemotherapy × surgery	—	—	—	—	0.98 (0.92 to 1.04)	—	—	—	—
Chemotherapy × radiation	—	—	—	—	—	0.91 (0.86 to 0.97)**	—	—	—
Advanced stage × chemotherapy	—	—	—	—	—	—	0.73 (0.68 to 0.78)***	—	—
Advanced stage × radiation	—	—	—	—	—	—	—	1.04 (0.97 to 1.12)	—

NOTE. Boldface indicates a statistically significant aIRR: * $P < .05$, ** $P < .01$, and *** $P < .001$.

Abbreviations: aIRR, adjusted incidence rate ratio; ED, emergency department; GU, genitourinary; IRR, incidence rate ratio; ref, reference; SA, sensitivity analysis.

^aaIRRs of ED visits within 6 months after cancer diagnosis generated from a multivariable negative binomial regression adjusting for all covariates listed in this table except for additional covariates.

^bNo. of ED visits within 12-18 months before cancer diagnosis as the primary predictor.

^cIncludes neighborhood education and poverty level as binary covariates (low v high); 9,764 patients whose listed addresses could not be mapped to neighborhood education and poverty levels were excluded.

^dIncludes an interaction term between treatment with chemotherapy and surgery.

^eIncludes an interaction term between treatment with chemotherapy and radiation.

^fIncludes an interaction term between advanced-stage cancer and treatment with chemotherapy.

^gIncludes an interaction term between advanced-stage cancer and treatment with radiation therapy.

^hExcludes pre- and post-cancer diagnosis ED visits that resulted in hospitalization.

ⁱNo. of ED visits within 6-12 months (or 12-18 months in SA 1) before cancer diagnosis.

TABLE A3. Negative Binomial Regression and SAs for the Parkland Cohort (n = 8,597)

Covariate	Unadjusted IRR (95% CI)		Adjusted Model ^a		SA 1 ^b	SA 2 ^c	SA 3 ^d	SA 4 ^e	SA 5 ^f	SA 6 ^g	SA 7 ^h
	Ref	Ref	Ref	Ref							
Age, years											
18-24	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
25-34	1.10 (0.85 to 1.42)	0.99 (0.78 to 1.26)	1.05 (0.82 to 1.33)	1.01 (0.80 to 1.29)	1.01 (0.80 to 1.29)	1.01 (0.80 to 1.29)	1.01 (0.80 to 1.29)	1.01 (0.80 to 1.29)	1.01 (0.80 to 1.29)	1.02 (0.80 to 1.29)	1.24 (0.94 to 1.63)
35-44	0.98 (0.77 to 1.24)	1.00 (0.79 to 1.25)	1.05 (0.83 to 1.31)	1.01 (0.80 to 1.27)	1.01 (0.80 to 1.27)	1.01 (0.80 to 1.27)	1.01 (0.80 to 1.27)	1.01 (0.81 to 1.27)	1.01 (0.81 to 1.27)	1.01 (0.81 to 1.27)	1.23 (0.95 to 1.60)
45-54	1.09 (0.87 to 1.38)	0.99 (0.80 to 1.24)	1.00 (0.80 to 1.25)	0.99 (0.80 to 1.24)	0.99 (0.80 to 1.24)	0.99 (0.80 to 1.24)	0.99 (0.80 to 1.24)	1.00 (0.80 to 1.24)	1.00 (0.80 to 1.24)	1.00 (0.80 to 1.25)	1.22 (0.94 to 1.58)
55-64	1.11 (0.88 to 1.40)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.25)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.26)	1.01 (0.81 to 1.26)	1.02 (0.81 to 1.27)	1.17 (0.90 to 1.51)
≥ 65	0.84 (0.67 to 1.07)	0.90 (0.71 to 1.14)	0.91 (0.72 to 1.16)	0.90 (0.71 to 1.14)	0.90 (0.71 to 1.14)	0.90 (0.71 to 1.14)	0.90 (0.71 to 1.14)	0.90 (0.71 to 1.14)	0.90 (0.71 to 1.14)	0.91 (0.72 to 1.15)	1.08 (0.82 to 1.41)
Female	0.91 (0.86 to 0.96)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.03 (0.96 to 1.10)	1.06 (0.98 to 1.15)
Race or ethnicity											
Non-Hispanic White	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
Hispanic	0.76 (0.70 to 0.82)	0.96 (0.90 to 1.07)	1.01 (0.93 to 1.10)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	0.98 (0.90 to 1.07)	1.06 (0.97 to 1.17)
Black	1.02 (0.94 to 1.10)	1.02 (0.94 to 1.09)	1.02 (0.95 to 1.10)	1.02 (0.94 to 1.09)	1.02 (0.94 to 1.09)	1.02 (0.94 to 1.09)	1.02 (0.94 to 1.09)	1.02 (0.94 to 1.09)	1.02 (0.95 to 1.10)	1.02 (0.95 to 1.10)	1.01 (0.93 to 1.10)
Others	0.72 (0.62 to 0.83)	0.91 (0.80 to 1.05)	0.89 (0.78 to 1.02)	0.92 (0.80 to 1.05)	0.91 (0.80 to 1.05)	0.91 (0.80 to 1.05)	0.92 (0.80 to 1.05)	0.92 (0.80 to 1.05)	0.92 (0.80 to 1.05)	0.92 (0.80 to 1.05)	0.77 (0.66 to 0.90) **
Non-English language preference	0.68 (0.64 to 0.72)	0.81 (0.74 to 0.87) **	0.88 (0.81 to 0.96) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.81 (0.74 to 0.87) **	0.94 (0.86 to 1.02)
Insurance type											
Commercial	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
Medicare	1.09 (0.92 to 1.28)	1.04 (0.88 to 1.23)	1.05 (0.89 to 1.24)	1.04 (0.89 to 1.23)	1.04 (0.89 to 1.23)	1.04 (0.89 to 1.23)	1.04 (0.89 to 1.23)	1.04 (0.89 to 1.23)	1.04 (0.88 to 1.22)	1.05 (0.89 to 1.23)	0.93 (0.77 to 1.12)
Medicaid	1.24 (1.06 to 1.45)	1.17 (1.02 to 1.35) **	1.19 (1.03 to 1.39) **	1.17 (1.02 to 1.35) **	1.17 (1.02 to 1.35) **	1.17 (1.02 to 1.35) **	1.17 (1.02 to 1.35) **	1.17 (1.02 to 1.35) **	1.17 (1.01 to 1.35) **	1.17 (1.01 to 1.35) **	1.17 (1.00 to 1.37) **
Uninsured	1.12 (0.96 to 1.30)	1.07 (0.93 to 1.24)	1.10 (0.95 to 1.26)	1.07 (0.93 to 1.24)	1.07 (0.93 to 1.24)	1.07 (0.93 to 1.24)	1.07 (0.93 to 1.24)	1.06 (0.92 to 1.23)	1.06 (0.92 to 1.23)	1.07 (0.93 to 1.24)	1.07 (0.91 to 1.25)
Unknown	1.00 (0.82 to 1.20)	1.01 (0.85 to 1.21)	1.05 (0.88 to 1.25)	1.01 (0.85 to 1.21)	1.01 (0.85 to 1.21)	1.01 (0.85 to 1.21)	1.01 (0.85 to 1.21)	1.00 (0.84 to 1.20)	1.00 (0.84 to 1.20)	1.01 (0.84 to 1.20)	0.93 (0.76 to 1.13)
Homeless	2.34 (2.01 to 2.71)	1.85 (1.61 to 2.12) **	1.84 (1.60 to 2.11) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.85 (1.61 to 2.12) **	1.83 (1.57 to 2.13) **
Charlson Comorbidity Index, per unit	1.05 (1.03 to 1.07)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.99 to 1.03)	1.01 (0.98 to 1.03)
Year of cancer diagnosis											
2012	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref
2013	1.00 (0.91 to 1.10)	1.00 (0.91 to 1.08)	0.99 (0.91 to 1.09)	1.00 (0.91 to 1.09)	1.00 (0.91 to 1.09)	1.00 (0.91 to 1.09)	1.00 (0.91 to 1.09)	1.00 (0.91 to 1.09)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.94 (0.85 to 1.03)
2014	1.03 (0.94 to 1.13)	1.04 (0.95 to 1.15)	1.04 (0.96 to 1.14)	1.04 (0.95 to 1.13)	1.04 (0.95 to 1.13)	1.04 (0.95 to 1.13)	1.04 (0.95 to 1.13)	1.04 (0.95 to 1.13)	1.03 (0.95 to 1.13)	1.04 (0.95 to 1.13)	1.11 (1.00 to 1.22) **
2015	0.95 (0.87 to 1.05)	0.97 (0.89 to 1.06)	0.98 (0.90 to 1.08)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	0.97 (0.89 to 1.06)	1.04 (0.95 to 1.15)
2016	0.99 (0.91 to 1.09)	0.99 (0.91 to 1.08)	1.00 (0.91 to 1.09)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	0.99 (0.91 to 1.08)	1.12 (1.02 to 1.23) **
Cancer type											
Lung	1.48 (1.34 to 1.62)	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.22 (1.09 to 1.37) **	1.20 (1.07 to 1.35) **	1.14 (1.00 to 1.31) *
Breast	0.65 (0.50 to 0.71)	0.84 (0.75 to 0.93) **	0.84 (0.75 to 0.94) **	0.84 (0.75 to 0.93) **	0.84 (0.75 to 0.93) **	0.84 (0.75 to 0.93) **	0.84 (0.75 to 0.93) **	0.84 (0.75 to 0.93) **	0.83 (0.74 to 0.92) **	0.84 (0.75 to 0.94) **	0.68 (0.79 to 1.00) **
Colorectal	0.99 (0.87 to 1.13)	1.11 (0.96 to 1.27)	1.10 (0.96 to 1.26)	1.10 (0.96 to 1.27)	1.11 (0.96 to 1.27)	1.11 (0.96 to 1.27)	1.10 (0.96 to 1.26)	1.10 (0.96 to 1.26)	1.10 (0.96 to 1.26)	1.10 (0.96 to 1.27)	0.95 (0.81 to 1.11)
Head and neck	1.31 (1.15 to 1.49)	1.23 (1.06 to 1.42) **	1.24 (1.07 to 1.43) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.24 (1.07 to 1.43) **	1.20 (1.03 to 1.39) **	1.41 (1.20 to 1.66) **
Melanoma	0.60 (0.45 to 0.81)	0.92 (0.70 to 1.22)	0.92 (0.70 to 1.21)	0.93 (0.70 to 1.22)	0.92 (0.70 to 1.22)	0.92 (0.70 to 1.22)	0.92 (0.70 to 1.22)	0.92 (0.70 to 1.22)	0.92 (0.69 to 1.21)	0.92 (0.69 to 1.22)	0.89 (0.65 to 1.22)
Brain	1.25 (1.01 to 1.54)	1.70 (1.38 to 2.08) **	1.64 (1.34 to 2.01) **	1.68 (1.37 to 2.09) **	1.70 (1.38 to 2.09) **	1.69 (1.38 to 2.09) **	1.69 (1.38 to 2.09) **	1.70 (1.38 to 2.09) **	1.70 (1.38 to 2.09) **	1.71 (1.39 to 2.10) **	1.48 (1.17 to 1.88) **
Kidney	0.88 (0.77 to 1.01)	1.23 (1.06 to 1.42) **	1.22 (1.05 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.23 (1.06 to 1.42) **	1.25 (1.08 to 1.44) **	1.22 (1.05 to 1.41) **	1.40 (1.19 to 1.63) **
Prostate	0.56 (0.49 to 0.64)	0.78 (0.68 to 0.90) **	0.78 (0.68 to 0.90) **	0.78 (0.68 to 0.91) **	0.78 (0.68 to 0.91) **	0.78 (0.68 to 0.91) **	0.78 (0.68 to 0.91) **	0.78 (0.68 to 0.91) **	0.79 (0.68 to 0.91) **	0.78 (0.68 to 0.91) **	0.99 (0.84 to 1.16)
Other GU	1.13 (0.93 to 1.38)	1.39 (1.14 to 1.70) **	1.35 (1.11 to 1.64) **	1.39 (1.14 to 1.70) **	1.39 (1.14 to 1.70) **	1.39 (1.14 to 1.70) **	1.39 (1.14 to 1.70) **	1.39 (1.14 to 1.70) **	1.38 (1.14 to 1.69) **	1.37 (1.13 to 1.67) **	1.51 (1.22 to 1.88) **
Lymphoma	1.18 (1.06 to 1.32)	0.96 (0.85 to 1.09)	0.96 (0.85 to 1.10)	0.96 (0.85 to 1.10)	0.96 (0.85 to 1.10)	0.96 (0.85 to 1.10)	0.96 (0.85 to 1.10)	0.96 (0.85 to 1.10)	0.95 (0.85 to 1.09)	0.96 (0.85 to 1.10)	0.83 (0.72 to 0.97) **
Cervical, uterine, ovarian, and vaginal	1.05 (0.96 to 1.15)	1.26 (1.13 to 1.41) **	1.28 (1.14 to 1.43) **	1.26 (1.13 to 1.41) **	1.26 (1.13 to 1.41) **	1.26 (1.13 to 1.41) **	1.26 (1.13 to 1.41) **	1.26 (1.13 to 1.41) **	1.25 (1.12 to 1.40) **	1.28 (1.13 to 1.41) **	1.40 (1.24 to 1.58) **
Pancreas	1.58 (1.33 to 1.87)	1.37 (1.15 to 1.63) **	1.39 (1.17 to 1.65) **	1.37 (1.15 to 1.62) **	1.37 (1.15 to 1.63) **	1.37 (1.15 to 1.63) **	1.37 (1.15 to 1.63) **	1.37 (1.15 to 1.63) **	1.34 (1.12 to 1.60) **	1.38 (1.16 to 1.64) **	1.30 (1.06 to 1.59) **
Other GI	1.22 (1.11 to 1.33)	1.15 (1.03 to 1.29) **	1.19 (1.06 to 1.33) **	1.15 (1.03 to 1.29) **	1.15 (1.03 to 1.29) **	1.15 (1.03 to 1.29) **	1.15 (1.03 to 1.29) **	1.15 (1.03 to 1.29) **	1.14 (1.02 to 1.27) **	1.15 (1.03 to 1.28) **	1.07 (0.95 to 1.22)

(continued on following page)

TABLE A3. Negative Binomial Regression and SAs for the Parkland Cohort (n = 8,597) (continued)

Covariate	Multivariable aIRR (95% CI)									
	Unadjusted IRR (95% CI)	Adjusted Model ^a	SA 1 ^b	SA 2 ^c	SA 3 ^d	SA 4 ^e	SA 5 ^f	SA 6 ^g	SA 7 ^h	
Advanced cancer stage at diagnosis	1.63 (1.53 to 1.74)	1.24 (1.16 to 1.34)***	1.21 (1.13 to 1.30)***	1.25 (1.17 to 1.34)***	1.25 (1.16 to 1.34)***	1.25 (1.16 to 1.34)***	1.45 (1.31 to 1.60)***	1.20 (1.10 to 1.30)***	1.08 (1.00 to 1.17)*	
Died within 180 days after diagnosis	1.43 (1.30 to 1.57)	1.24 (1.13 to 1.36)***	1.24 (1.13 to 1.36)***	1.24 (1.13 to 1.36)***	1.24 (1.13 to 1.36)***	1.24 (1.13 to 1.36)***	1.22 (1.11 to 1.33)***	1.24 (1.13 to 1.36)***	0.77 (0.68 to 0.86)***	
Initial cancer treatment modality										
Surgery	0.61 (0.58 to 0.65)	0.76 (0.71 to 0.81)***	0.76 (0.71 to 0.81)***	0.76 (0.71 to 0.81)***	0.75 (0.69 to 0.81)***	0.76 (0.71 to 0.81)***	0.76 (0.71 to 0.81)***	0.76 (0.71 to 0.81)***	0.88 (0.82 to 0.95)***	
Radiation therapy	1.11 (1.04 to 1.19)	1.04 (0.98 to 1.11)	1.04 (0.98 to 1.12)	1.04 (0.98 to 1.11)	1.04 (0.98 to 1.11)	1.03 (0.94 to 1.14)	1.03 (0.96 to 1.10)	1.00 (0.92 to 1.08)	1.02 (0.95 to 1.10)	
Chemotherapy	1.63 (1.54 to 1.72)	1.58 (1.48 to 1.68)***	1.56 (1.47 to 1.66)***	1.58 (1.48 to 1.68)***	1.56 (1.44 to 1.69)***	1.57 (1.46 to 1.69)***	1.71 (1.59 to 1.84)***	1.59 (1.49 to 1.69)***	1.57 (1.47 to 1.69)***	
Immunotherapy	1.19 (1.06 to 1.34)	1.12 (0.99 to 1.26)*	1.10 (0.98 to 1.24)	1.11 (0.99 to 1.25)*	1.12 (0.99 to 1.26)*	1.12 (0.99 to 1.26)*	1.12 (1.00 to 1.26)*	1.12 (0.99 to 1.26)*	1.09 (0.95 to 1.24)	
No. of prediagnosis ED visits ⁱ										
0	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	Ref	
1-3	1.47 (1.37 to 1.57)	1.51 (1.41 to 1.61)***	1.35 (1.26 to 1.45)***	1.49 (1.39 to 1.59)***	1.51 (1.41 to 1.61)***	1.50 (1.41 to 1.61)***	1.51 (1.42 to 1.62)***	1.51 (1.41 to 1.61)***	1.47 (1.37 to 1.59)***	
≥ 4	3.29 (2.78 to 3.89)	3.01 (2.57 to 3.53)***	3.46 (2.87 to 4.17)***	2.99 (2.56 to 3.50)***	3.01 (2.57 to 3.53)***	3.01 (2.57 to 3.53)***	3.01 (2.57 to 3.52)***	3.01 (2.57 to 3.53)***	4.65 (3.84 to 5.63)***	
Additional covariates										
Missing SSN	—	—	—	0.76 (0.69 to 0.83)***	—	—	—	—	—	
Chemotherapy × surgery	—	—	—	—	1.03 (0.92 to 1.16)	—	—	—	—	
Chemotherapy × radiation	—	—	—	—	—	1.01 (0.89 to 1.15)	—	—	—	
Advanced stage × chemotherapy	—	—	—	—	—	—	0.77 (0.68 to 0.87)***	—	—	
Advanced stage × radiation	—	—	—	—	—	—	—	1.15 (1.01 to 1.32)**	—	

NOTE. Boldface indicates a statistically significant aIRR: **P* < .05, ***P* < .01, and ****P* < .001.

Abbreviations: aIRR, adjusted incidence rate ratio; ED, emergency department; GU, genitourinary; IRR, incidence rate ratio; ref, reference; SA, sensitivity analysis; SSN, social security number.

^aaIRRs of ED visits within 6 months after cancer diagnosis generated from a multivariable negative binomial regression adjusting for all covariates listed in this table except for additional covariates.

^bNo. of ED visits within 12-18 months before cancer diagnosis as the primary predictor.

^cIncludes whether a SSN was available as a binary covariate.

^dIncludes an interaction term between treatment with chemotherapy and surgery.

^eIncludes an interaction term between treatment with chemotherapy and radiation.

^fIncludes an interaction term between advanced-stage cancer and treatment with chemotherapy.

^gIncludes an interaction term between advanced-stage cancer and treatment with radiation therapy.

^hExcludes pre- and post-cancer diagnosis ED visits that resulted in hospitalization.

ⁱNo. of ED visits within 6-12 months (or 12-18 months in SA 1) before cancer diagnosis.