Disparities in receipt of radiotherapy and survival by age, sex, and race among patients with non-metastatic squamous cell carcinoma of the anus

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Background: Combination chemoradiotherapy is the standard of care for treatment of non-metastatic squamous cell carcinoma of the anus (SCCA). This population based study evaluated disparities in receipt of radiotherapy (RT) as well as comparative survival rates for SCCA patients in the United States.

Methods: The Surveillance, Epidemiology, and End Results (SEER) 18 database was used to identify patients with non-metastatic SCCA diagnosed between 1998 and 2008. Multivariate logistic regression was used to model the relationships between age, sex, and race and the receipt of RT, adjusting for marital status and stage of disease. Relative survival (RS) rates were compared by each factor, with added adjustment for RT status, using Cox proportional hazards model.

Results: A total of 3,885 patients with localized or regional SCCA as the only primary malignancy were included in the study, of which, 3,192 (82%) received RT. In our multivariate analysis, lower rates of RT were found for those 65+ years old [adjusted odds ratio (OR) 0.71; P<0.001], males (adjusted OR 0.65; P<0.001), and blacks (adjusted OR 0.78; P=0.049). Multivariate survival analysis showed worse survival among those 65+ years old (adjusted HR 1.65; P<0.001), males (adjusted HR 1.53; P<0.001), and blacks (adjusted HR 1.35; P=0.001).

Conclusions: This population based study identified older patients, males, and blacks as less likely to receive RT. Worse survival was also found in these groups.

Keywords: Anal cancer; seer; survival; radiotherapy (RT)

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Introduction

Squamous cell carcinoma of the anus (SCCA) constitutes 2.4% of gastrointestinal cancers. An estimated 7,210 new cases and 950 deaths in the United States are predicted for 2014 (1). Both the incidence and mortality of anal cancer has been increasing in the U.S. with an annual percentage change of 1.9% and 3.6%, respectively (1).

Current standard of care for patients with non-metastatic SCCA is chemoradiotherapy with mitomycin and 5-fluorouracil (1). This is based on findings of the EORTC trial that investigated the addition of mitomycin/5-fluorouracil to the traditional radiotherapy (RT) regimen and found an increased rate of complete remission (80% vs. 54%) (2). Ajani et al. (3) investigated the replacement of mitomycin with cisplatin, but found no improvement in disease-free survival and higher rates of colostomy in the cisplatin group.
Blacks have higher annual percentage increases in incidence and mortality of SCCA compared to whites (2.3 vs. 2.0 and 4.4 vs. 3.6, respectively) (1). Furthermore, blacks with SCCA suffer from lower survival rates compared to whites [5-year relative survival (RS) 56% vs. 67%]. It is unclear whether the inferior survival rates are due to disparities in cancer care in these patients. Thus, we conducted this study to evaluate disparities in receipt of RT and survival in patients with non-metastatic squamous cell carcinoma.

**Methods**

**Data source**

We used the Surveillance, Epidemiology, and End Results (SEER) program database to identify patients with SCCA diagnosed between 1998 and 2008 (4). The SEER 18 database is a population-based cancer database sponsored by the National Cancer Institute that collects high-quality incidence and survival data from several cancer registries throughout the United States (4). The SEER program’s standard for case ascertainment is 98%, and this goal is achieved by conducting rigorous quality control studies every other year (5). The SEER database covers approximately 28 percent of the US population overall and 26 percent of African Americans (4). We gathered data using registries from the following areas: Atlanta, Connecticut, Detroit, Hawaii, Iowa, New Mexico, San Francisco-Oakland, Seattle-Puget Sound, Utah, Los Angeles, San Jose-Monterey, Rural Georgia, greater California, Kentucky, Louisiana, New Jersey, and greater Georgia.

**Study population**

We identified a total of 6,695 patients with non-metastatic (localized or regional) carcinoma of the anus diagnosed between January 1, 1998 and December 31, 2008. We defined SCCA as cases with ICD-O-3 histology codes 8051 through 8081. Inclusion criteria were non-Hispanic black or white race and no prior malignancies. American Indian/Alaska Native and Asian/Pacific Islander patients were excluded because of insufficient numbers for meaningful analysis. Patients were excluded if their disease was not histologically confirmed, their diagnosis was made from death certificate or autopsy report, or they had missing data for age, marital status, or RT status. Patients receiving radioactive implants, radioisotopes, or with unspecified mode of radiation were also excluded. The final study population was N=3,885 (Figure 1).

**Study variables**

We assessed the variables age, sex, race, stage, marital status,
and RT status. We dichotomized age as <65 or 65+ years old. Marital status separated patients into three groups; married (including common law), single (never married), and S/D/W/U (separated, divorced, widowed, unmarried or domestic partner). Stage was either localized or regional based on the LRD stage variable in SEER (6). We divided treatment groups into those who received RT (beam radiation only) and those who did not (including those who refused RT).

### Statistical analysis

We used univariate logistic regression to assess the relationship between independent variables and receipt of RT and reported unadjusted odds ratios (ORs). To adjust for covariates we used a multivariate logistic regression model and reported adjusted ORs. These statistical analyses were performed using Stata 13.0 (StataCorp, College Station, TX, USA).

We calculated 1- and 5-year RS rates using survival sessions in the SEER*Stat program (7). RS is a population-based measure of net-survival calculated as a ratio of the proportion of observed survivors (all causes of death) in our study population to the proportion of expected survivors in a group of individuals matched by age, sex, race, and year. The use of RS is based on the assumption that death from anal cancer is a negligible proportion of deaths in the general population, a reasonable assumption for a relatively rare cancer such as SCCA (8). We calculated hazard ratios (HRs) while adjusting for each covariate using Cox proportional hazards model in the CanSurv software package (9).

### Results

#### Patient characteristics and factors associated with receipt of RT

A total of 3,885 patients (Figure 1) were identified with SCCA as the first primary cancer that met our inclusion criteria. The median age at diagnosis was 56 years old, 30% were 65+ years old [1,176], 42% were male [1,633], 12% were black [469], and 38% had regional disease at the time of diagnosis [1,485]. See Table 1 for details.

#### Factors associated with receipt of RT

Of the patients studied, 3,192 (82%) received RT. A significant racial difference existed between whites and blacks with 83% of whites and 78% of blacks receiving RT. After adjusting for covariates age, sex, marital status, and stage, black patients were less likely to receive RT compared to whites (adjusted OR 0.781; 95% CI: 0.611–0.999; P=0.049). Older patients (65+ years) were less likely to receive RT compared to younger patients: 80% vs. 83% (adjusted OR 0.71; 95% CI: 0.59–0.86; P<0.001), and females were more likely to receive RT compared to males: 85% vs. 78% (adjusted OR 1.54; 95% CI: 1.28–1.85; P<0.001). Single patients and those with localized disease were also less likely to receive RT compared to their referent groups. See Table 2 for details.

#### Factors associated with RS

Among patients receiving RT, black patients had worse RS at 1 and 5 years compared to whites (1-year RS 88.4±1.8 vs. 93.0±0.5, P<0.005; 5-year RS 65.7±2.8 vs. 75.8±1.0, P<0.001). Older patients had worse RS than those <65 years old (1-year RS 88.0±1.2 vs. 94.3±0.5, P<0.001; 5-year RS 70.3±2.1 vs. 76.4±1.0, P<0.005). Males had worse RS at 1 and 5 years compared to females (1-year RS 90.0±0.9 vs. 94.2±0.6, P<0.001; 5-year RS 68.8±1.5 vs. 78.6±1.2, P<0.001).

#### Factors associated with survival: Cox model analysis

In order to assess RS while adjusting for covariates, Cox proportional hazards model was used. In this analysis, blacks had worse RS compared to whites (adjusted HR 1.35; 95% CI: 1.13–1.63; P<0.001). Older patients had worse RS compared to those <65 years old (adjusted HR 1.65; 95% CI: 1.41–1.94; P<0.001), and males had worse RS compared to males (adjusted HR 01.53; 95% CI: 1.31–1.78; P<0.001). See Table 3 for details.
Discussion

Receipt of RT

Since the release of the Institute of Medicine’s landmark report, Unequal Treatment: Confronting Racial and Ethnic Disparities in Health Care, increased attention has been directed toward healthcare disparities in the United States (10). Much of this attention has been aimed at disparities affecting blacks (11). Lower treatment rates are found among black patients for various cancers (11-14). These disparities may arise from a number of sources including lower SES (10). In the majority of studies that controlled for SES, disparities in treatment significantly diminished after adjustment, but persisted nonetheless (10). This makes it likely that, in addition to SES, sociocultural factors or provider biases are playing a role in their lower rates of treatment.

Additionally, older patients and men received lower rates of RT compared to younger patients and females. This is consistent with a study by Asch et al that assessed receipt of care according to age and sex and found that older patients (52% vs. 58%) and males (52% vs. 57%) were both less likely to receive recommended care (15). While older patients are eligible for Medicare and may see fewer access related barriers, they also have greater incidence of potentially RT-limiting comorbidities such as anemia (16). Age-related disparities in treatment of other cancers have been documented (17,18).

Survival

Our finding of worse RS among blacks matches the disparity noted in a recent study where blacks had a lower 5-year RS rate compared to whites (56% vs. 67%) (1).
Although we did not see a 5-year survival difference in our cohort, our multivariate analysis showed greater risk of death among blacks (adjusted HR 1.35; P=0.001). This same analysis found a greater risk of death among those not receiving RT (adjusted HR 1.22; P=0.036), and may be a reason black patients had worse survival.

The men in our study also had worse survival. Bartelink et al. identified male sex as a negative prognostic factor in SCCA (2). This poorer prognosis may be due to higher incidence of HIV/AIDS among males and their subsequent increased risk of developing SCCA (19). Thus, male sex may act as a proxy for HIV status, an important comorbidity leading to lower RS in that group.

Our study has several limitations. One of the major limitations is the lack of detail regarding therapeutic interventions such as dosage, regimens, missed appointments, and delays in initiation of treatment after diagnosis. Also, SEER does not collect data on comorbidities, socioeconomic status (except at the county level), or data regarding important risk factors such as HIV infection, HPV infection, and payer status.

In conclusion, there are significant disparities in the receipt of RT and survival in patients with non-metastatic SCCA. Identification of factors responsible for differences in treatment may lead to interventions to reduce disparities and improve SCCA outcomes.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: No formal approval of IRB is required as data were collected from a source that was publicly available and did not contain unique patient identifiers. Informed consent is not required as it was a SEER database study and data could not be tracked back to individuals.

References

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