

# Association Between Emergency Department (ED) Involvement in Cancer Diagnosis and Survival Across Cancer Types in the Medicare Population

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## BACKGROUND

- 20–50% of cancers are diagnosed following emergency department (ED) presentation, rising to 60% for some malignancies (e.g., pancreatic).<sup>1,2,4</sup>
- ED diagnosis is associated with advanced stage, poorer survival, higher comorbidity, and worse patient experience.<sup>3</sup>
- ED presentation can reflect underlying barriers to timely diagnosis (e.g., limited access).<sup>5,6</sup>
- Evidence on the independent impact of diagnosis route on survival, especially in Medicare populations, remains limited.

## OBJECTIVE

To quantify the association between ED involvement at diagnosis and overall survival across different cancer types in the SEER-Medicare population.

## METHODS

Figure 1. Study design.

**METHODS OVERVIEW**  
*Retrospective Cohort Study Using SEER-Medicare Data (2009-2020)*

**EXCLUSION CRITERIA**

- Age <18 years at index
- Prior cancer during 12-month baseline
- In situ or non-malignant disease
- Diagnosed after death autopsy

Final Analytic Cohort: n = 818,120

**INCLUSION CRITERIA**

- Medicare FFS beneficiaries linked SEER-Medicare database with SEER-confirmed primary cancer
- 12+ months cont. enrollment pre-index, 1+ post-index
- Cancer site matches study ICD-0-3 groupings

**INDEX DATE**

- Earliest ICD-9/10 claim for cancer site with +1 month of SEER diagnosis month
- Imputed to the 15<sup>th</sup> of SEER diagnosis month of not found

**BASELINE PERIOD**  
12 months pre-index

- Demographics, CCI, prior history

**FOLLOW-UP PERIOD**

From index date until first of death, disenrollment or subsequent primary cancer

- End of Study Period: December 31, 2020

**PRIMARY OUTCOME:** Overall survival

- Time from index date to death from any cause (days)

**STATISTICAL ANALYSIS**

- Kaplan-Meier:** unadjusted survival by route of diagnosis/log rank tests
- Cox proportional hazard models:** ED as reference, adjusted for age, race/ethnicity, dual eligibility, CCI, year of diagnosis, & AJCC staging (across study period)
- Overall cohort and cancer-specific analyses based on independent models

## KEY RESULTS: ED INVOLVEMENT WAS COMMON, ROSE WITH STAGE, AND ASSOCIATED WITH 2-4X HIGHER MORTALITY RISK

### Study Population

- 818,120 Medicare beneficiaries (mean age 74) diagnosed with invasive cancer between 2010-2020.
- Cohort was mostly non-Hispanic White (82.8%), followed by non-Hispanic Black (8.9%), Hispanic (2.2%), and other/unknown (6.0%). 54.6% were female and 45.4% were male (Table 1).

Table 1. Baseline characteristics among patients with invasive cancer diagnosis between 2010-2020.

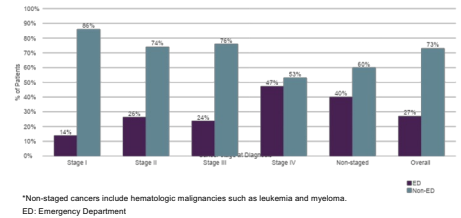
Description	Overall Sample
<b>Number of Patients</b>	818,120
<b>Age at Index</b>	
Mean (years, SD)	74 (9.1)
<b>Sex</b>	
Female	54.6%
<b>Race/Ethnicity</b>	
Non-Hispanic White	82.8%
Non-Hispanic Black	8.9%
Hispanic	2.2%
Other/Unknown	6.0%
<b>Original Reason for Entitlement</b>	
Age	80.3%
Disability or ESRD	19.3%
Both	0.4%
<b>Dual Eligibility</b>	
Full	19.9%
Partial	6.1%
No or Unknown	74.0%
<b>Stage*</b>	
I	36.3%
II	16.7%
III	14.1%
IV	24.9%
Unknown/Missing	8.0%
<b>Charleston Comorbidity Index (CCI)</b>	
Mean	3.5
0	16.6%
1	15.0%
2	16.2%
3+	52.2%

\*Among staged cancers; SD: Standard Deviation

### Emergency Department Involvement in Diagnosis

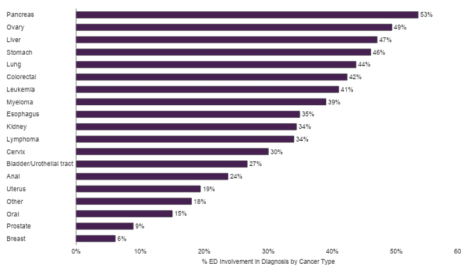
- 26.4% of cancer patients had ED involvement at diagnosis
- ED involvement rose with stage, reaching 47.3% in Stage IV; non-staged hematologic cancers had 40% ED involvement (Figure 2).
- Wide variation by cancer type (Figure 3):
  - Lowest: Breast (6%)
  - Highest: Ovarian (49%), and pancreatic (53%)
- Highest ED reliance observed in cancers with nonspecific or rapidly progressing symptoms (Figure 3).

Figure 2. Percent of patients diagnosed in the emergency department compared to non-emergency department settings, by stage.



\*Non-staged cancers include hematologic malignancies such as leukemia and myeloma.  
ED: Emergency Department

Figure 3. Percent of patients diagnosed in the emergency department, by cancer site.



Pancreas n = 26,342; Ovary n = 8,600; Liver n = 17,756; Stomach n = 10,229; Lung n = 158,706; Colorectal n = 86,078; Leukemia n = 25,317; Myeloma n = 17,805; Esophagus n = 7,294; Kidney n = 20,776; Lymphoma n = 28,844; Cervix n = 2,214; Bladder/Urthelial tract n = 33,331; Anal n = 4,118; Uterus n = 24,237; Other n = 16,746; Oral n = 18,499; Prostate n = 141,083; Breast n = 170,823  
\*Cancer groupings were defined according to SEER major cancer sites.  
\*\*Other cancer category includes thyroid, gallbladder, other digestive organs, and other ungrouped cancers. These cancers were combined to comply with SEER-Medicare cell suppression requirements.  
ED: Emergency Department.

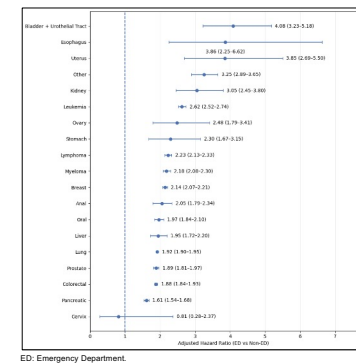
### Observed Mortality

- Follow-up mortality was substantially higher for ED-diagnosed patients (44%) vs. non-ED pathways (19%).
- Represents an approximate 2-fold higher crude mortality among ED-diagnosed patients.
- Adjusted Survival
  - After adjustment, route of diagnosis remained associated with hazard of death.
  - All non-ED pathways were associated with lower hazard of death vs ED diagnosis.
  - Screening-detected cancers showed the greatest survival advantage.

### Survival by Route of Diagnosis Across Cancer Types

- ED diagnosis was associated with higher hazard of death across most cancers (Figure 4).
- 3–4x: bladder/urothelial, uterus, esophagus
- 2x: breast, lung, colorectal, prostate, liver
- Overall, ED diagnosis was associated with 2–4x higher hazard, independent of stage and patient characteristics.
- Results were consistent in sensitivity analyses using SEER summary stage.

Figure 4. Mortality hazard ratios for ED vs non-ED by cancer site.



## CONCLUSIONS

- ED involvement accounted for a substantial share of overall mortality among cancer patients.
- ED involvement at diagnosis remained a strong independent predictor of mortality after adjusting for sociodemographics, comorbidities, and stage at diagnosis.
- These findings highlight the need for earlier, non-emergent diagnostic pathways.

## LIMITATIONS

- Residual confounding may remain; factors such as symptom duration, performance status, and access to or initiation of therapy were not captured.
- Route of diagnosis was claims-based and may be subject to misclassification.
- Stage adjustment may not fully account for disease severity or tumor biology.
- Findings may have limited generalizability to non-Medicare FFS populations.
- Observational study design limits causal inference.

## References

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## Disclosures

This study was sponsored by GRAIL, Inc.



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AJCC: American Joint Committee on Cancer; CCI: Charlson Comorbidity Index; CPT: Current Procedural Terminology; ED: Emergency Department; FFS: Fee-for-Service; HCPCS: Healthcare Common Procedures Coding System; HR: Hazard Ratio; ICD: International Classification of Diseases; IP: Inpatient; OP: Outpatient; SEER: Surveillance, Epidemiology, and End Results.