Incidence of cachexia and health care resource use (HCRU) in patients with breast, colorectal, lung, pancreatic, and prostate cancers

Introduction

- Cachexia in patients with cancer is characterized by involuntary weight loss, muscle atrophy, increased lipolysis and energy expenditure, and reduced appetite.¹
- It can lead to a decreased quality of life, reduced cancer treatment effectiveness, increased susceptibility to cancer treatment side effects, and increased cancer-related mortality.^{2,3}
- Based on available scientific literature from clinical studies, cachexia affects ~50–80% of patients with advanced cancer and ~20% of cancer deaths are attributable to cachexia.^{3,4}
- Cachexia is likely to be under-diagnosed due to inconsistent application of diagnostic criteria in clinical practice as well as challenges when interpreting changes in bodyweight (for example, due to obesity, edema, and/or ascities).^{5,6}
- There are also limited data on how cachexia may impact health care resource use (HCRU) in patients with cancer.⁷

Objectives

- To estimate the incidence of cachexia in patients newly diagnosed with breast, colorectal, lung, pancreatic, and prostate cancer.
- To compare measures of health care resource utilization in patients diagnosed with cancer who do and do not have cachexia.

Methods

 This retrospective, observational, cohort study integrated electronic health record and administrative claims data using the Optum[®] Market Clarity database (Figure 1).



≥12 months after cancer diagnosis, until death, loss to follow-up, or end of study period (30 September 2023), whichever occured first

- Inclusion and exclusion criteria are shown in Table 1.
- Cachexia was defined as $\geq 5\%$ loss in bodyweight calculated from 2 bodyweight measurements ~6 months apart.⁸
- The cachexia index date was the earliest date after the cancer index date on which \geq 5% loss of bodyweight was recorded.
- In the control group, the cachexia index date was the date after the cancer index date on which <5% loss of bodyweight was recorded.
- The primary outcome was the number of health care visits per-patient-per-month (PPPM; 30 days) during the 12 months post-cachexia index date.
- Inverse probability of treatment weighting was used to adjust for confounding between cohorts (cachexia vs no cachexia) at the cachexia index date.
- Student's *t*-test was used to compare HCRU between cohorts (cachexia vs no cachexia) by tumor type.

REFERENCES

1. Nishie K, et al. Drug Discov Today 2023;28:103689. 2. Dev R. Ann Palliat Med 2019;8:24-32

3. von Haehling S, Anker SD. J Cachexia Sarcopenia Muscle 2014:261-3. 4. Argilés JM, et al. Nat Rev Cancer 2014;14:754-62. 5. Nishikawa H, et al. Int J Mol Sci 2021;22. 6. Bianchini C, et al. Cancer Treat Rev 2024;125:102717.

7. Tarricone R, et al. Crit Rev Oncol Hematol 2016;99:49-62. 8. Fearon K, et al. Lancet Oncol 2011;12:489-95.

Table 1: Eligibility criteria

Inclusion criteria

- Diagnosed with ≥1 malignant solid tumor of interest between 1 October 2016 and 30 September 2022, as identified by the following ICD-10-CM codes in ≥ 1 claim or encounter in a hospital inpatient setting, or ≥ 2 claims or encounters in selected outpatient setting \geq 30 days apart:
- Breast cancer (ICD-10 codes C50.x)
- Colorectal cancer (ICD-10 codes C18.x-C21.x, C49.x, C7A.x)
- Lung cancer (ICD-10 codes C34.x, C7A.090)
- Pancreatic cancer (ICD-10 codes C25.x)
- Prostate cancer (ICD-10 codes C61.x)
- \geq 19 years of age at cancer index date (ie, \geq 18 years of age at the beginning of the 12 months before cancer index date)
- EHR enrollment (ie, EHR activity) for 12 months before cancer index date
- ≥ 2 bodyweight assessments 150–210 days apart in the 12 months before cancer index
- ≥ 2 bodyweight assessments 150–210 days apart with ≥ 1 of those assessments occurring in the 12 months after cancer index
- EHR enrollment for 12 months after cancer index date or death, whichever occured first

For analyses related to HCRU:

- Continuous health plan enrollment for 12 months before cachexia index date
- Continuous health plan enrollment for ≥ 1 month after cachexia index or until death, whichever occured first

Exclusion criteria

- In the 12 months before the cancer index date, ≥ 1 claim or encounter in any setting with a diagnosis of:
- Any solid tumor other than the index (incident cancer) diagnosis (ICD-10 codes C00.x-C75.x or C7A.x)

Any hematologic malignancy (ICD-10 codes C81.x-C96.x)

- Any cancer with metastasis (ICD-10 codes C77.x, C78.x, C79.x, C7B.x)
- Any other cancer type (ICD-10 codes C45.x, C46.x, C76.x, C80.x, C96.x)
- History of cachexia in the pre-index period (ie, patient met criteria for cachexia in the 12 months before the cancer index date, as defined below) but not in the post-index period:
- 2 bodyweight measurements within 150–210 days of each other - 2nd bodyweight was $\leq 95\%$ of the first bodyweight measurement
- Unknown gender
- Prostate cancer cohort only: Female gender

EHR=electronic health record; HCRU=health care resource use; ICD-10-CM=International Classification of Diseases 10th Revision, Clinical Modification

Results INCIDENCE

BREA



Demo Femal

Age, m Race, Africar

Asian White

Other/ _____



Demo Female Age, m Race, Africar Asian White Other/



Demo Female Age, m Race, Africar Asian White Other/

DISCLOSURES

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- A total of 27,428 patients with a malignant solid tumor were included.
- Within 12 months of receiving a cancer diagnosis, cachexia was observed in 38% of patients.

AST CAN	ICER				Population size: 10,500 Incidence of cachexia: 34%			
graphics	With cachexia n=3595	Without cachexia n=6905	P value	HCRU	With cachexia n=3595	Without cachexia n=6905	P value	
e, %	99	99	0.31	Inpatient	0.98	0.80	0.05	
nean (SD), y	65 (12)	65 (12)	0.74	Outpatient	5.11	3.46	<0.0001	
%			0.87	Office	3.69	3.44	0.02	
n American	13	12		Home	0.67	0.83	0.05	
	2	2		Telehealth	0.03	0.03	0.51	
	81	82		Other	4.75	3.85	<0.0001	
/Unknown	4	4		Overall	15.22	12.41	<0.0001	

COLORECTAL CANCER

Population size: 3304 **Incidence of cachexia:** 51%

Population size: 3345

Incidence of cachexia: 55%

	With cachexia	Without cachexia	Ρ		With cachexia	Without cachexia	Ρ
graphics	n=1683	n=1621	value	HCRU	n=1683	n=1621	value
e, %	52	52	0.92	Inpatient	2.72	2.32	0.21
nean (SD), y	66 (13)	66 (13)	0.83	Outpatient	5.36	3.93	<0.0001
%			0.98	Office	4.38	3.55	<0.001
n American	11	11		Home	1.51	0.84	<0.0001
	1	1		Telehealth	0.02	0.02	0.37
	83	83		Other	6.49	5.63	0.07
Únknown	5	5		Overall	20.48	16.30	<0.0001

LUNG CANCER

	With	Without			With	Without	
	cachexia	cachexia	Ρ		cachexia	cachexia	Ρ
graphics	n=1845	n=1500	value	HCRU	n=1845	n=1500	value
e, %	57	56	0.55	Inpatient	6.92	5.45	0.03
ean (SD), y	69 (10)	69 (10)	0.94	Outpatient	6.89	6.14	0.07
%			0.93	Office	4.36	4.29	0.78
n American	10	10		Home	1.89	1.89	1.00
	1	1		Telehealth	0.02	0.02	0.74
	86	86		Other	9.52	7.53	<0.001
Unknown	3	3		Overall	29.59	25.32	<0.001



Dem



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HEALTH CARE RESOURCE USE

• Across all tumor types, hospital outpatient and office visits PPPM were numerically greater in patients with vs without cachexia, with statistical significance varying by tumor type. • Other differences in hospital inpatient, home, telehealth, and other visits were also observed.

PANCREATIC CANCER

Population size: 475 **Incidence of cachexia**: 74%

1822P

	With	Without			With	Without	
Demographics	n=350	n=125	P value	HCRU	n=350	n=125	P value
Female, %	52	54	0.72	Inpatient	8.25	12.50	0.13
Age, mean (SD), y	67 (12)	67 (13)	0.90	Outpatient	10.80	8.72	0.18
Race, %			0.95	Office	6.41	2.66	<0.0001
African American	15	17		Home	2.49	1.47	0.09
Asian	1	1		Telehealth	0.05	0.02	0.21
White	79	79		Other	12.60	14.08	0.58
Other/Unknown	5	4		Overall	40.59	39.45	0.83

PROSTATE CANCER

Population size: 9804 **Incidence of cachexia:** 29%

qp	With	Without			With	Without	
	cachexia	cachexia	Ρ		cachexia	cachexia	Ρ
Demographics	n=2865	n=6939	value	HCRU	n=2865	n=6939	value
Male, %	100	100	-	Inpatient	1.46	0.69	<0.0001
Age, mean (SD), y	69 (9)	69 (9)	0.25	Outpatient	3.14	2.50	<0.0001
Race, %			0.94	Office	3.27	3.06	0.03
African American	15	15		Home	0.64	0.43	<0.01
Asian	1	1		Telehealth	0.02	0.01	0.15
White	79	80		Other	4.53	3.20	<0.0001
Other/Unknown	4	4		Overall	13.04	9.89	<0.0001

CONCLUSIONS

O Incidence of cachexia was highest in patients with pancreatic cancer followed by lung and colorectal cancer, and lowest in breast and prostate cancers. • Patients with cachexia used more health care resources, highlighting a high clinical and economic burden in this population.

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