# Risk of mental health outcomes and peak risk periods posthematologic cancer diagnosis in the United States

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### Key Takeaway



The importance of early and long-term mental health prevention and support post cancer diagnosis is one key aspect underscored by these findings

#### Conclusions



There were consistently increased incidence rates of depressive and anxiety disorders at 3-years post-hematologic cancer diagnosis



Effect sizes were most pronounced at 30-days for the occurrence of any depressive and any anxiety disorder



Additional research should focus on improving mental health assessment and management in care of cancer patients



Il co-authors are employees of Johnson & Johnson. T Beaulieu, J Gendreau, and D Gifkins are also equity holders of Johnson & Johnso.

#### Introduction

- According to the American Cancer Society and estimates using data derived from the National Cancer Institute's Surveillance Epidemiology and End Results (SEER) database, the lifetime risk of a person developing cancer (any cancer) is approximately 39-40% in the United States
- The lifetime risk of a person developing leukemia is about 1-2%, and for non-Hodgkin lymphoma (NHL), it is roughly 2%
- A considerable proportion of individuals diagnosed with cancer will develop a subsequent mental health condition
- Building from the work of Forbes et al. (2024), we sought to quantify the real-world and contemporary incidence of mental health conditions post-cancer diagnosis

We also sought to identify populations and time periods where risk is more pronounced

#### Methods

- Data were derived from Optum's de-identified Clinformatics® Data Mart Database (January 2016 through December 2024)
- Individuals with a leukemia or non-Hodgkin lymphoma (NHL) diagnosis were matched to individuals without a cancer diagnosis (1:1 nearest neighbor without replacement)
- With a standardized mean difference threshold of 0.25, this approach yielded adequate balance for most covariates

- Primary outcomes of interest included any depressive disorder and any anxiety disorder\*
- A washout period of 365-days was used to help ensure that any observed outcomes were new-onset depressive and anxiety disorders
- Incidence rates (IRs) of depressive and anxiety disorders were calculated
- Adjusted hazard ratios (aHRs) with 95% confidence intervals (CIs) were derived from multivariate cox models
- We sought to examine risk conferred by cancer diagnosis at 30days, 180-days, 1-year, and 3-years post-cancer diagnosis

#### Results

- For the outcome of depressive disorders, each matched cohort was comprised of 80,224 individuals (**Table 1**)
- For the outcome of anxiety disorders 79,480 individuals remained in each cohort post-match

Table 1. Baseline characteristics post matching

	Depressive disorders N=80,224	Anxiety disorders N=79,480
Age		
median (IQR)	72 (65, 79)	72.00 (65, 79)
18-39	3,704 (4.6)	3,618 (4.6)
40-49	3,658 (4.6)	3,587 (4.5)
50-59	7,226 (9.0)	7,140 (9.0)
60-69	17,080 (21.3)	16,970 (21.4)
70-79	28,849 (36.0)	28,519 (35.9)
80-89	19,707 (24.6)	19,646 (24.7)
Sex at birth		
Female	34,006 (42.4)	33,581 (42.3)
Male	46,218 (57.6)	45,899 (57.7)
Race/Ethnicity		
Asian	2,276 (2.8)	2,269 (2.9)
Black or African American	7,815 (9.7)	7,783 (9.8)
No matching concept	26,384 (32.9)	26,069 (32.8)
White	43,749 (54.5)	43,359 (54.6)
Hispanic or Latino	6,582 (8.2)	6,506 (8.2)
No matching concept	25,619 (31.9)	25,317 (31.9)
Not Hispanic or Latino	48,023 (59.9)	47,657 (60.0)
Region		
Midwest	18,159 (22.6)	18,104 (22.8)
Northeast	11,831 (14.7)	11,566 (14.6)
South	32,473 (40.5)	32,088 (40.4)
Unknown	241 (0.3)	236 (0.3)
West	17,520 (21.8)	17,486 (22.0)
Diabetes	24,149 (30.1)	24,199 (30.4)
Dementia	2,724 (3.4)	2,797 (3.5)
Heart disease	36,469 (45.5)	36,174 (45.5)
Tobacco use disorder	287 (0.4)	301 (0.4)
Alcohol use disorder	743 (0.9)	759 (1.0)
Substance use disorder*	20,733 (25.8)	20,542 (25.8)

 At each time point (30-days, 180-days, 1-year, 3-years) incidence of depressive disorders was consistently elevated, and consistently higher among those with hematologic cancer as compared to those without cancer (Figure 1)

Figure 1. Incidence of depressive disorders per 100 persons

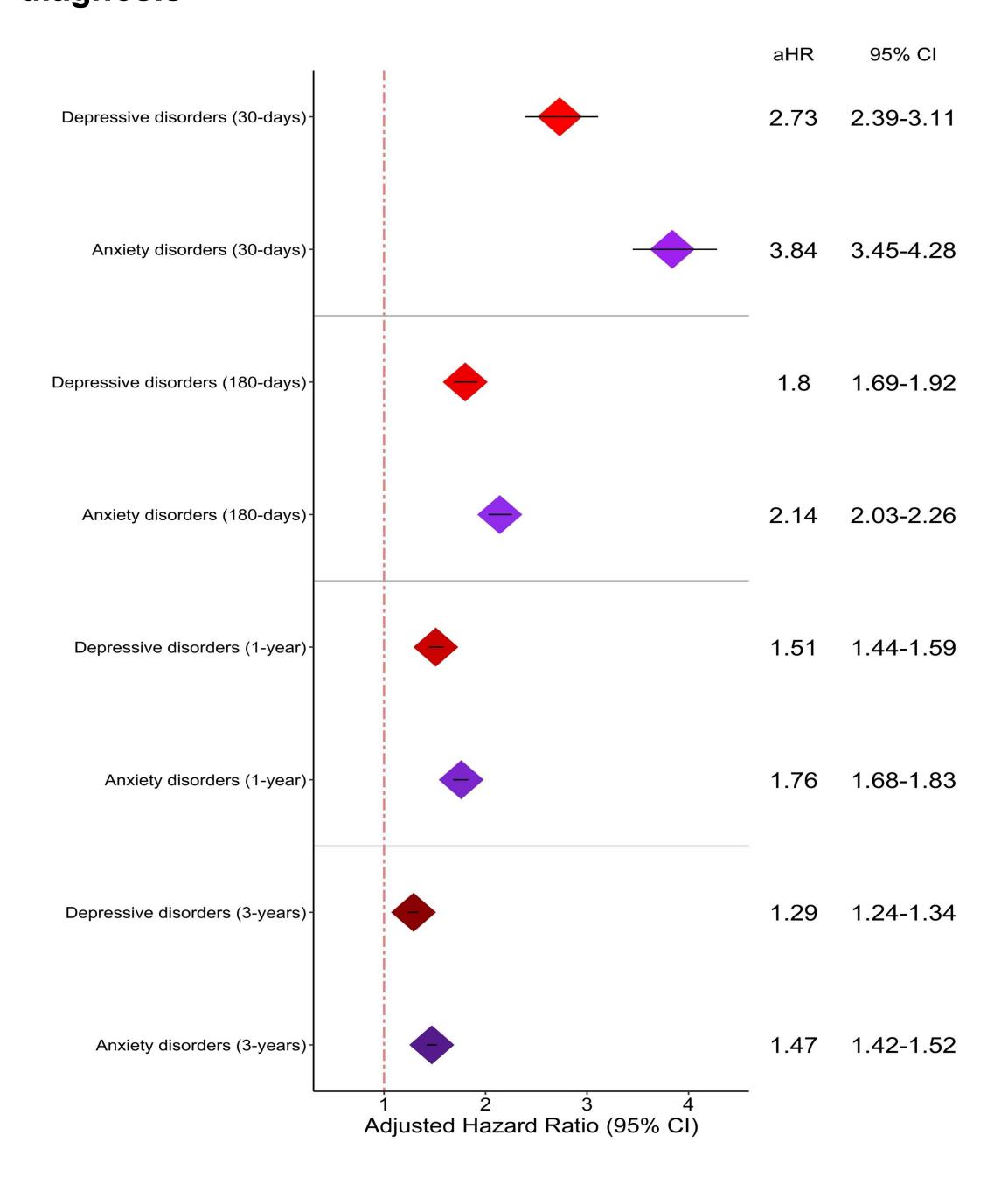
	Hematologic cancer diagnosis	No cancer diagnosis
30-days	2.18	0.54
180-days	12.64	5.21
1-year	21.16	12.25
3-years	42.11	33.15

 A similar trend was observed for incidence of anxiety disorders (Figure 2)

## Figure 2. Incidence of anxiety disorders per 100

persons	Hematologic cancer diagnosis	No cancer diagnosis
30-days	3.67	0.76
180-days	17.24	6.48
1-year	27.01	14.46
3-years	48.03	37.02

#### Figure 3. A forest plot of adjusted hazard ratios for depressive and anxiety disorders post hematologic cancer diagnosis



- For depressive disorders, there was a stepwise decrease in effect size, with effect size appearing most pronounced at 30days post cancer diagnosis in the hematologic cancer cohort (Figure 3).
- The effect size was greater for anxiety disorders than was observed for depressive disorders at each time point.
- These findings should be viewed in the context of several limitations including but not limited to the potential for misclassification, unmeasured confounding and bias, coding and programming inaccuracies, and so forth.

Forbes H, Carreira H, Funston G, Andresen K, Bhatia U, Strongman H, Abrol E, Bowen L, Giles C, Bhaskaran K. Early, medium and long-term mental health in cancer survivors compared with cancer-free comparators: matched cohort study using linked UK electronic health records.

Hennessy S, Atsuta Y, Hill S, Rägo L, Juhaeri J; Council for International Organizations of Medical Sciences (CIOMS) Working Group XIII. Real-World Evidence in Regulatory Decision Making: Report Summary From the Council for International Organizations of Medical Sciences (CIOMS) Working Group XIII. Pharmacoepidemiol Drug Saf. 2025 Mar;34(3):e70117. doi: 10.1002/pds.70117. PMID: 40070104; PMCID: PMC118970 Tyree PT, Lind BK, Lafferty WE. Challenges of using medical insurance claims data for utilization analysis. Am J Med Qual. 2006 Jul-Aug:21(4):269-75. doi: 10.1177/1062860606288774.

Hematologic Malignancies

