Economic Burden of Opioid-Induced Constipation Among Patients With or Without Cancer in the United States

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Background

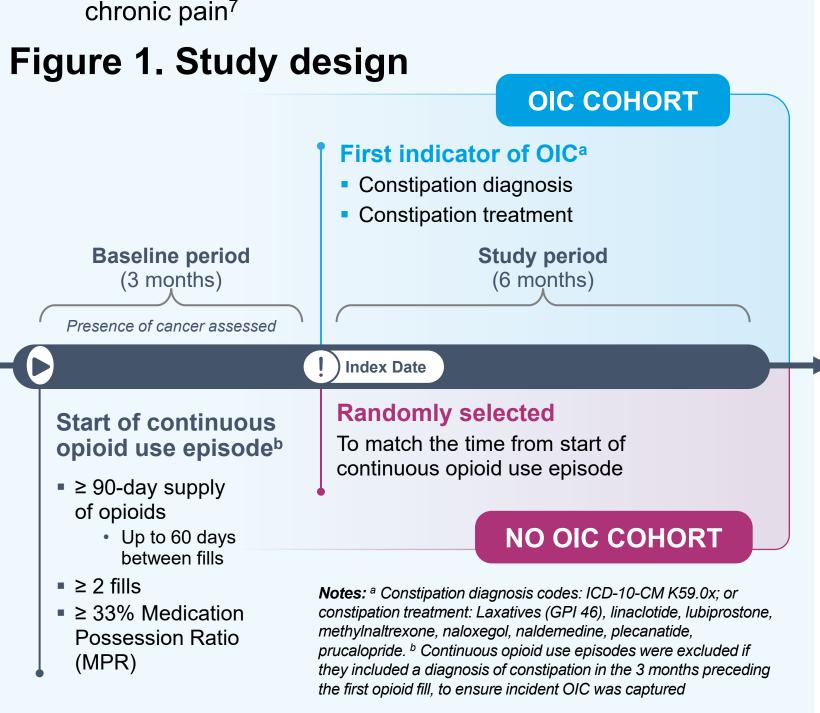
- Opioid use can cause opioid-induced constipation (OIC), often leading to difficult bowel movements with substantial discomfort^{1,2}
- Previous studies have highlighted incremental healthcare costs linked to OIC³⁻⁶, but they are limited by issues like outdated data and small sample sizes, with scant evidence on OIC among patients with long-term opioid use to manage chronic pain⁷
- OIC can negatively affect daily activities and productivity in patients managing non-cancer pain with opioids^{1,2}
- Patients with cancer commonly suffer from pain caused by their condition, resulting in increased opioid consumption and a higher likelihood of developing OIC⁸

Objective

To compare healthcare costs between commercially insured continuous opioid users with and without opioid-induced constipation (OIC) in the United States (US), separately among patients with and without cancer

Method

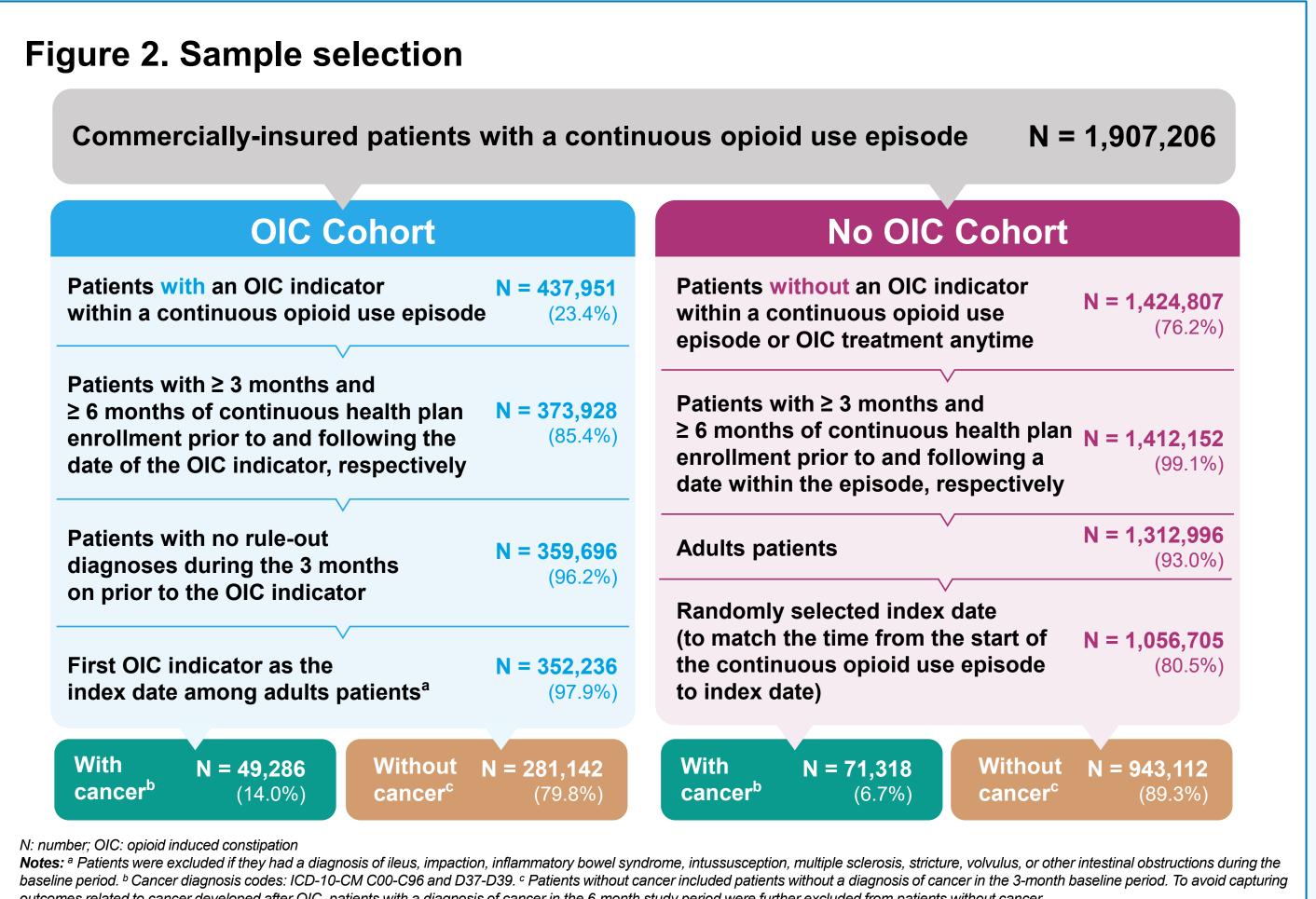
- Data source: Komodo Research Data (KRD+) from 01/2016 - 02/2024
- Adult patients with ≥1 continuous opioid use episode were grouped into two distinct cohorts based on the presence of an OIC indicator during an episode (Figure 1)
- Continuous opioid use episodes were defined as opioid use for ≥ 90 days, to align with the CDC's definition of chronic pain⁷

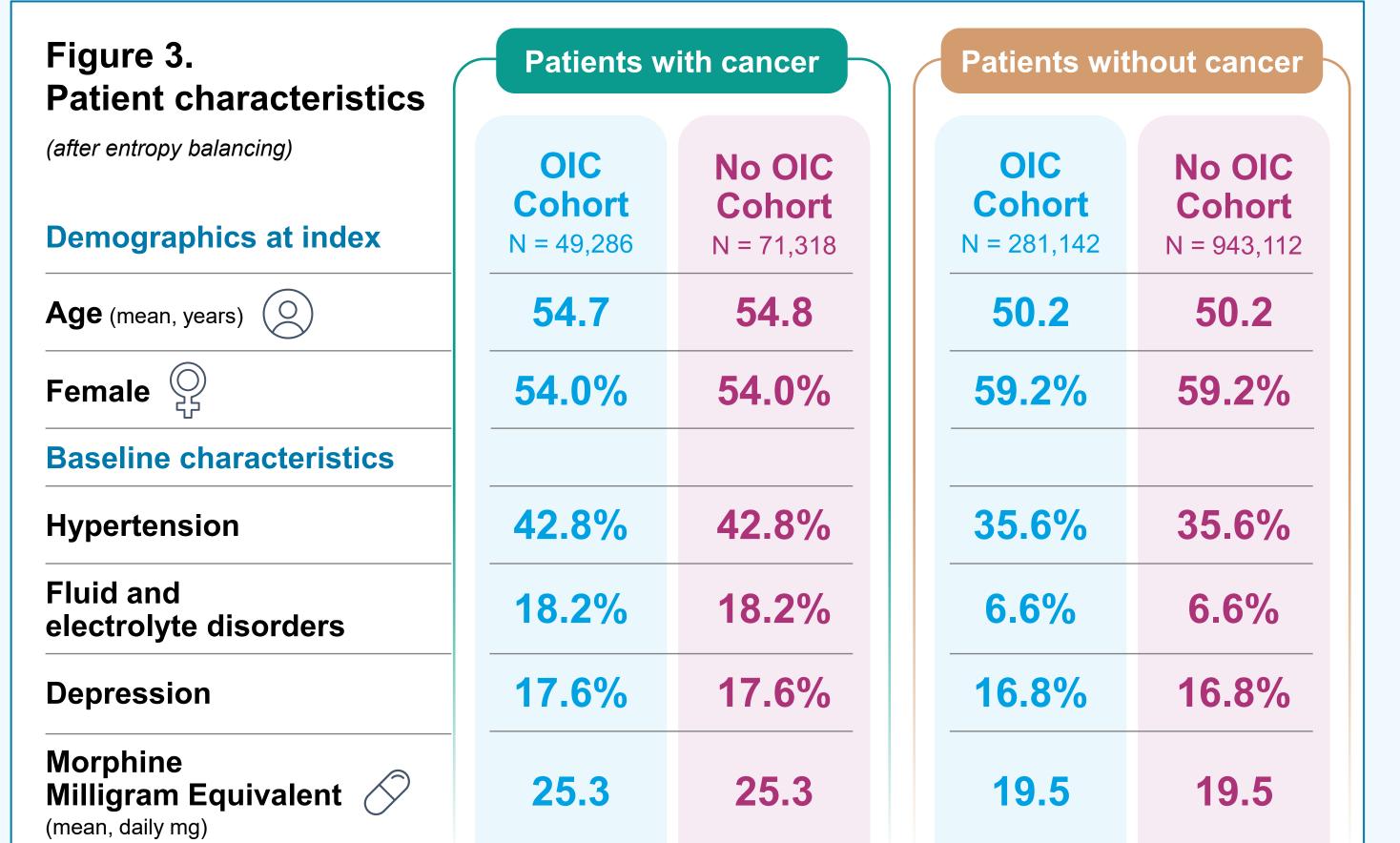


- Entropy balancing was applied to adjust for baseline characteristics including demographics (e.g., age, sex, region, insurance type), duration of opioid use, baseline morphine milligram equivalent (MME), and comorbidity profile, between the OIC and no OIC cohorts for patients with and without cancer, separately
- Outcomes included all-cause healthcare costs (estimated allowed amounts from medical claims, including inpatient, outpatient, and emergency department visits, and pharmacy claims; 2024 USD) during the study period, reported as per patient per year (PPPY)
- Weighted GLM with a log link and Gamma distribution was used to compare healthcare costs between the OIC and no OIC cohorts for patients with and without cancer, separately

Results

SAMPLE SELECTION AND PATIENT CHARACTERISTICS

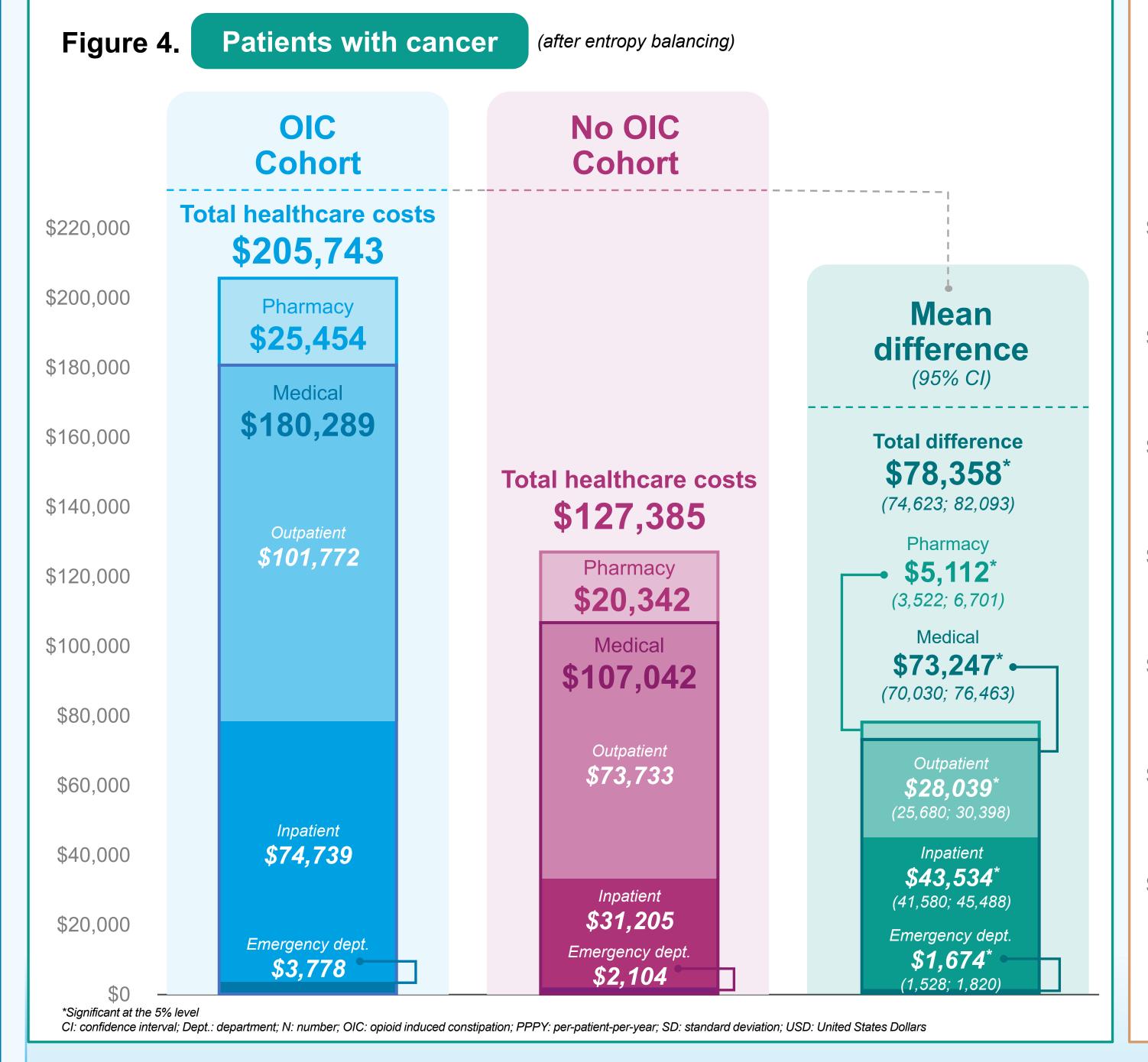


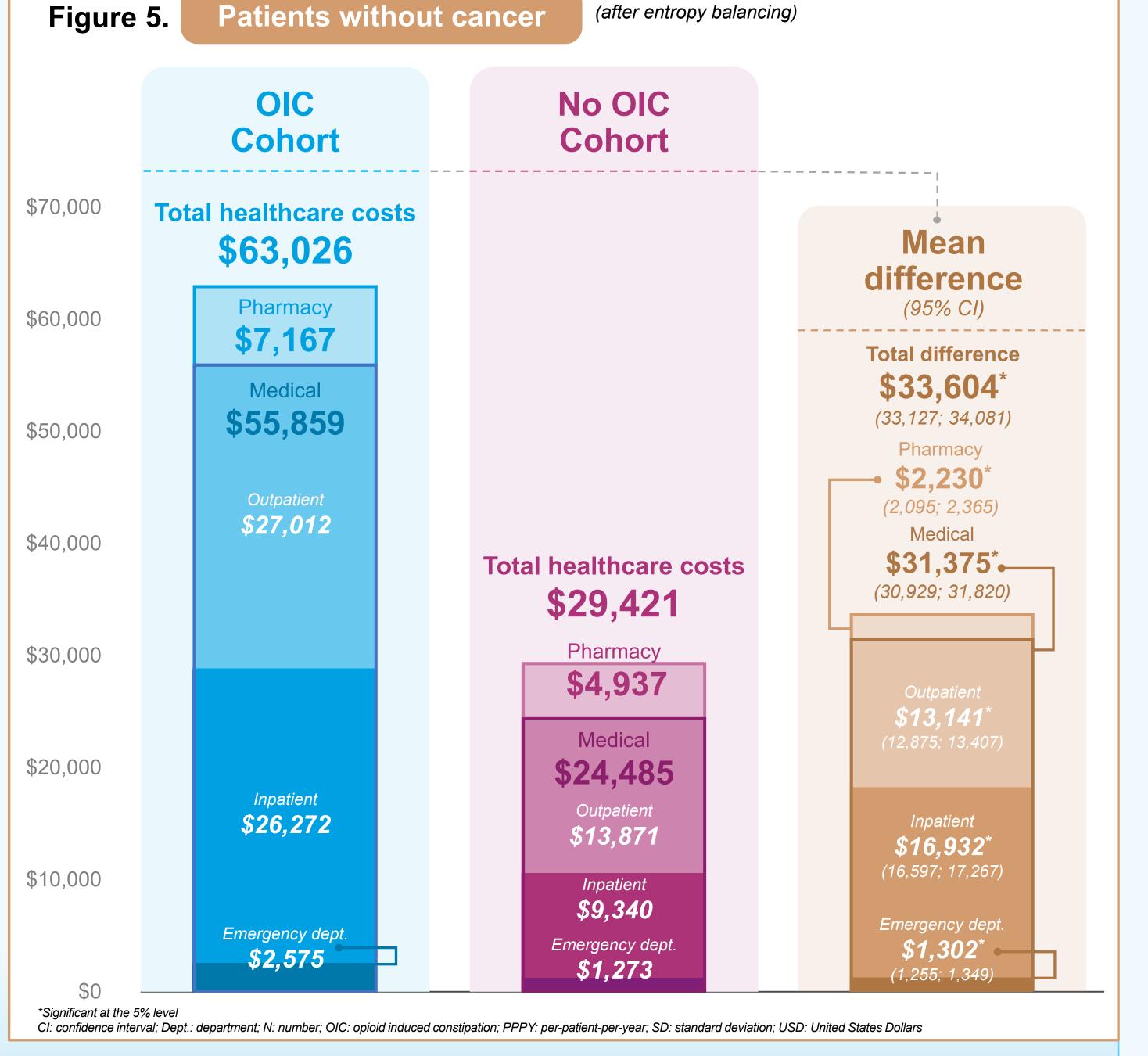


CCI: Charlson Comorbidity Index; MME: morphine milligram equivalent; MPR: medication possession ratio; N: number; OIC: opioid induced constipation; SD: standard deviation; std. diff.: standardized difference

Note: ^a The top 3 most frequent comorbidities observed in patients with cancer were reported.

HEALTHCARE COSTS





For both patients with and without cancer, separately, patients in the OIC Cohort experienced higher healthcare costs than patients in the No OIC Cohort

Conclusions

Compared to the No OIC Cohort, the OIC Cohort had significantly higher all-cause annual healthcare costs, driven by increased medical and pharmacy costs, for both patients with and without cancer

Limitations

- Since this is a claims-based study, over-the-counter treatments for constipation were not captured
- Results are based on a commercially insured population and may not reflect the broader U.S. adult population with OIC
- This study faces typical limitations associated with claims data, including possible billing discrepancies and incomplete data

References

- 1. Argoff CE. Clin J Pain. **2020**;36(9):716-722.
- Andresen V, et al. United European Gastroenterol J. 2018;6(8):1254-1266.
- 3. Fine PG, et al. Support Care Cancer. **2019**;27(2): 687-696.
- Fernandes AW, et al. Am Health Drug Benefits.
 2016;9(3):160-170.
- 79-86.
 6. Wan Y, et al. Am Health Drug Benefits. **2015**;8(2):

5. Olufade T, et al. Am Health Drug Benefits. 2017;10(2):

- 93-102.
 7. Dowell D, et al. MMWR. Recommendations and
- 8. ALMouaalamy N. Cureus. **2021**;13(4):e14386.

Sponsorship

reports. 2022;71.

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Disclosures

PGS, RB, RB, NG, and AG are employees of Analysis Group, Inc., a consulting company that has provided paid consulting services to Bausch Health Companies, Inc., which funded the development and conduct of this study. AL is an employee of Salix Pharmaceuticals. SKS and OO are employees of and have stock ownership in Bausch Health. AS was a postdoctoral fellow with Rutgers Pharmaceutical Industry Fellowship Program at the time of study completion.