The Impact of Socioeconomic Status, Insurance Type, and Treatment Facility on Survival in Metastatic Prostate Cancer: A Systematic Literature Review

# Objective



To systematically identify real-world evidence (RWE) that examined the impact of social determinants of health (SDOH), including socioeconomic status (SES), insurance status, and treatment facility (academic/non-academic/community). on survival in metastatic prostate cancer (mPC).

# Conclusions



Findings from this systematic literature review (SLR) suggest that higher SES, having health insurance, particularly private insurance, and treatment in an academic facility are associated with improved overall survival (OS) in patients with mPC. Focused efforts are needed to understand this survival advantage and to expand the benefit to all patients.



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2021. <sup>7</sup>GA Wells BS et al.

References: <sup>1</sup>Cackowski FC et al. Am Soc Clin Oncol Educ Book. 2021;41:1-1. <sup>2</sup>Hinata et al. World J Mens Health. 2022;40(2):217-227. <sup>3</sup>Coughlin SS. Prostate Int. 2020; 8(2):49-54 <sup>4</sup>Page, et al. *BMJ*. 2021;372:n160 <sup>5</sup>Page, et al. *BMJ*. 2021;372:n71. <sup>6</sup>Higgins JPT TJ et al.

Disclosures: SJF is a consultant for Pfizer, Astellas, Janssen, Bayer, Merck, Astra Zeneca Wovant, and Sanofi and a speaker for Astra Zeneca and Sanofi, AL, AH, MR, ER, and IAS are employees of EVERSANA, Canada, which was contracted by Pfizer to work on this project. AN and BA are employees of Pfizer and own Pfizer stock. JN is supported by the 2021-2023 University of Illinois Chicago-Pfizer Fellowship

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# Background

 Despite recent advancements in cancer treatment, prevention and care, significant disparities in incidence, treatment, and survival by SDOH persist.<sup>1-3</sup>

· We previously reported the association between race, a key SDOH, and OS in mPC. Herein, we examined additional SDOH (i.e., SES, insurance status, and treatment facility) on survival in mPC.

## Materials and Methods

### SYSTEMATIC LITERATURE REVIEW

• MEDLINE®. Embase, and Cochrane Library were searched for full-texts from Jan 2012 to July 2022 and for conference abstracts from Jan 2019 to July 2022 using Ovid®. A supplemental search of key congresses was also conducted. Article inclusion was based on a pre-defined PICOS criteria (Table 1).

• Review implementation and reporting followed the PRISMA statement<sup>4-5</sup> and Cochrane guidelines<sup>6</sup> (PROSPERO registration: CRD42022350888).

Quality assessment was performed on full-text publications using Newcastle-Ottawa Scales (NOS).7

# Results

### LITERATURE SEARCH

 Identified 3228 unique records: 95 records (representing 86 studies) were included. Here, we present results on the 14 studies (12 US, 1 ex-US, 1 mixed) which assessed the impact of SES, insurance status, and treatment facility on OS (Figure 1).

• Findings were from 12 full-text publications (10 US, 1 ex-US, 1 mixed) and 2 conference abstracts (both US).

#### • ≥90% studies scored ≥7/9 on the NOS, indicating high quality. Figure 1: PRISMA Flow Diagram Identification of studies via databases and registers Identification of studies via other methods Records identified from: Records removed before screening: Records identified from Databases Duplicate records removed (n = Conferences (n = 173) (n = 4,496) Records screened Records excluded (n = 2,879) (n = 3,228) Records screened Records excluded (n = 173) (n = 166) Reports not retrieved Reports sought for retrieval (n = 0)(n = 349) Reports excluded Reports assessed for eligibility (n=261)<sup>.</sup> (n = 349) Non-English (n = 2) Population (n = 187 Outcome (n = 52)Studies included in review Study Design (n = 12) (n = 86) Date (full text pre-2012 or abstract pre-Reports of included studies 2019 (n = 2) (n = 95) Incomplete/insufficient/partial data (n = 1) Duplicate (n = 5)Studies that assessed the SDOH of interest (n = 14) Reports of included studies (n = 14)

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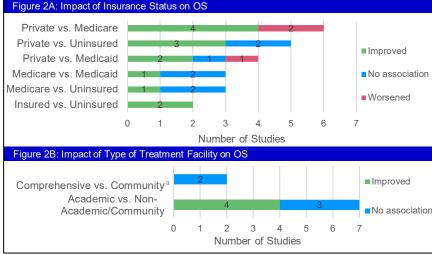
Table 1: PICOS Criteria	
Population	<ul> <li>Study populations or subgroups of patients (humans only; men) with:</li> <li>Age ≥18 years</li> <li>Histologically or cytologically confirmed adenocarcinoma of the prostate</li> <li>Metastatic disease</li> <li>Castration-sensitive OR castration-resistant</li> </ul>
Interventions/ Comparators	Any except alternative medicines
Outcomes	<ul> <li>Studies that mention a relationship between SDOH and target outcome:</li> <li>Clinical outcomes of interest:</li> <li>Real-world progression-free survival, OS, objective response rate, treatment duration, real-world patient-reported outcomes, time to subsequent therapy/chemotherapy</li> <li>Adherence/access outcomes of interest:</li> <li>Medication access (payer rejection rates, etc), medication persistence (abandonment rates, etc), medication adherence, treatment intensity/ intensification, initiation of life-prolonging therapies, treatment consistent with guidelines, health insurance access</li> <li>Note: Full-text publications were further filtered by those reporting the impact of SES, insurance status, and treatment facility on OS.</li> </ul>
Study Design	Real-world data

Impact of Insurance Status and Treatment Facility on OS

- A majority of studies showed that patients with private insurance had improved survival compared to other insurance types (Figure 2A; all US).
- Most studies (all US) showed patients in an academic facility had improved survival versus a non-academic facility (Figure 2B; all US).

## Impact of SES on OS

• Two studies (1 ex-US, 1 mixed) investigated the impact of SES, and both reported that higher SES was associated with significantly improved OS.



Comprehensive treatment facility was defined as those that treated ≥500 cancer patients per year, based on the National Cancer Database used in both studies