

Aparna Krishnan¹, Jilcha Feyisa MD^{1,2}, Nirav S. Kapadia MD^{1,2,3}, James B. Yu MD^{1,2,3}

1. Geisel School of Medicine at Dartmouth, NH, USA

2. Dartmouth Cancer Center, Department of Radiation Oncology & Applied Sciences, NH, USA

3. The Dartmouth Institute for Clinical Practice and Health Policy, NH, USA

INTRODUCTION

Prostate cancer ranges from indolent to aggressive disease.

Changes in tumor biology and diagnostics (e.g., MRI-targeted biopsy) may contribute to **increased high-grade diagnoses.**

We aim to evaluate temporal trends in high-grade prostate cancer in the United States from 2004-2022.

METHOD

Cohort: 2.26 million men with prostate cancer (18.8% with high-grade) from 2004-2022 (2025 NCDB)

High-grade: Gleason score ≥ 8 (Grade Group ≥ 4)

Descriptive analyses:

- % of high-grade diagnoses by biopsy practice eras (2004–2012, 2013–2017, 2018–2022)
- Annual median PSA change (high- vs. lower-grade diagnoses)

Multivariable analysis:

- Logistic regression modeling odds of high-grade diagnosis
- Covariates: age, race, treatment facility type

RESULTS

Fig. 1: High-Grade Diagnoses Over Time

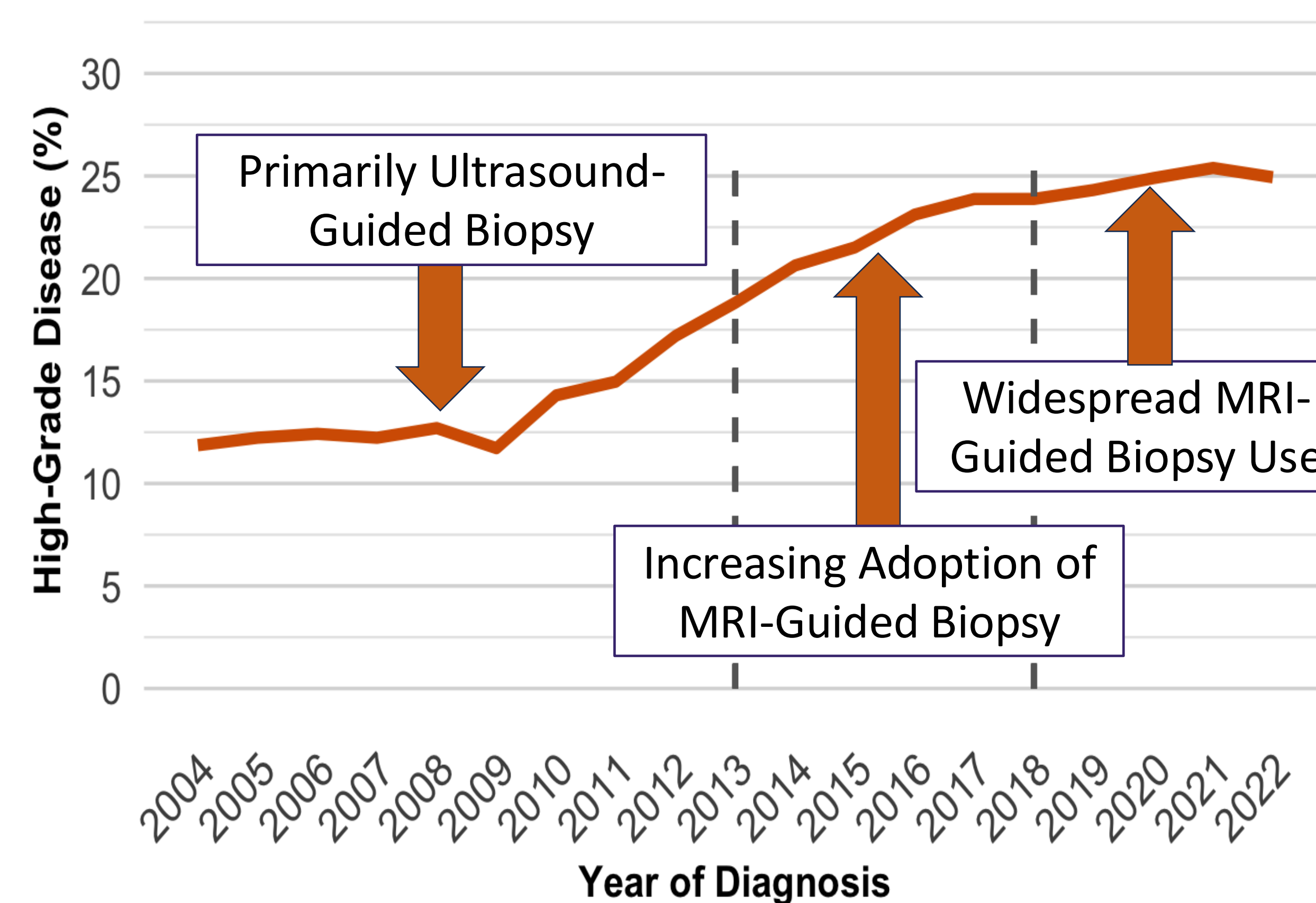


Fig. 1: Rising high-grade trend relative to ultrasound-guided to MRI-guided biopsy shift. Proportion **doubled from 2004 to 2022 (11.9% to 24.9%).**

Fig. 2: Median PSA at Diagnosis Over Time (by Grade)

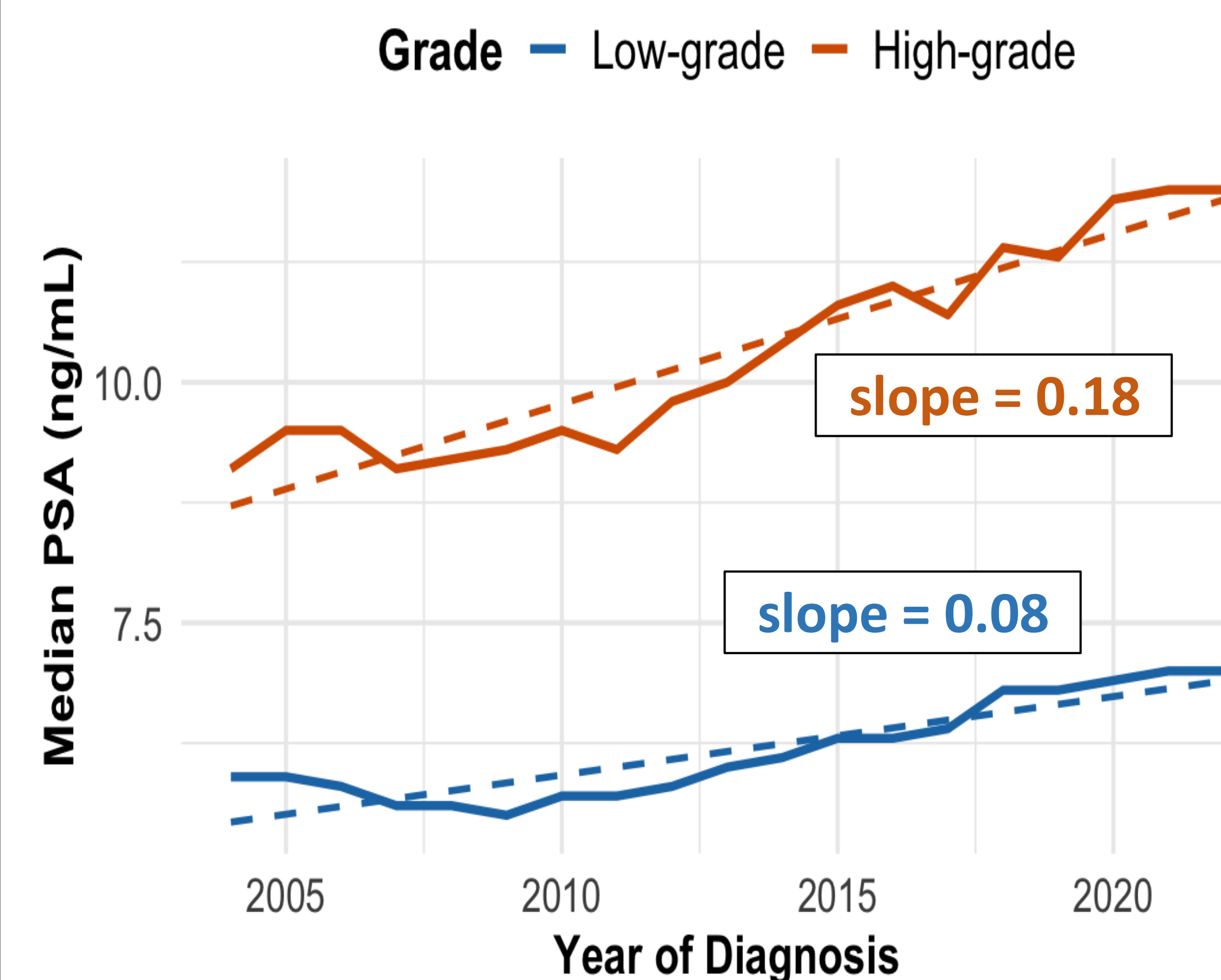


Fig. 2: Median PSA rose overall. Rate of increase is **steeper for high-grade tumors (0.18 vs 0.08 per year).**

Fig. 3: Odds of High-Grade Diagnosis

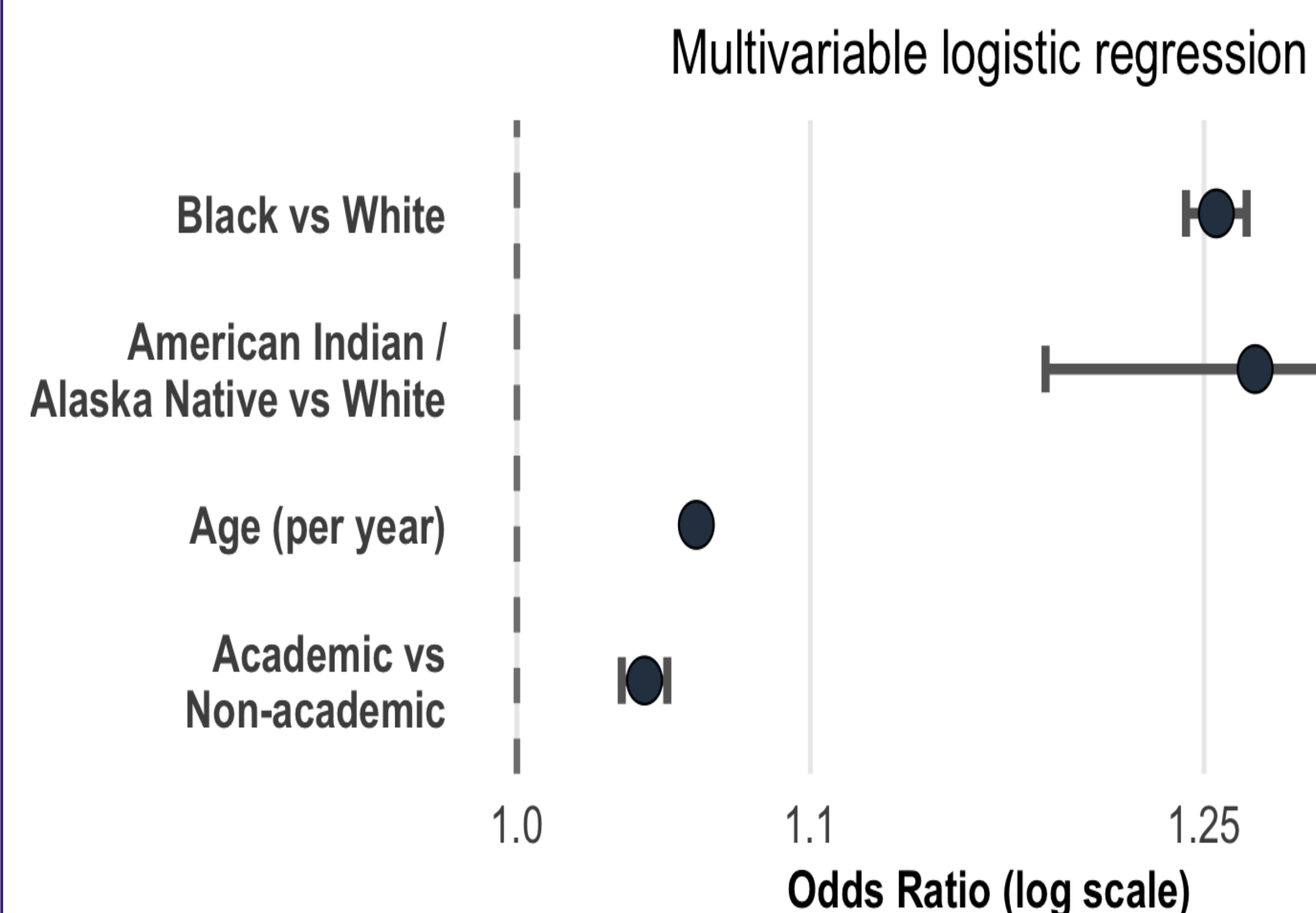


Fig. 3:

- **African American vs White: OR 1.25 (95% CI 1.24–1.27)***
- **American Indian/Alaska Native vs White: OR 1.27 (95% CI 1.19–1.36)***
- **Increasing age: OR 1.06 per year (95% CI 1.058–1.059)***
- **Diagnosis at an academic center vs non-academic: OR 1.04 (95% CI 1.03–1.05)***

*95% CI has $p < 0.001$

CONCLUSIONS

- High-grade diagnoses **more than doubled** from 2004–2022, coinciding with MRI-guided biopsy adoption & potentially reflecting **evolving pathology grading or diagnostic thresholds.**
- Steeper PSA rise in high-grade disease may suggest **improved detection** rather than biologic change.
- Higher odds of high-grade in **African American and AI/AN men**; risk rises **~6% per year of age.**
- Modest effect sizes may have large population impact. Benefits of advances may be unevenly distributed.

REFERENCES

1. Goldberg H, et al. J Urol. 2020;203(6):1085–93.
2. Ahmed HU, et al. Lancet. 2017;389:815–22.
3. Ahdoot M, et al. N Engl J Med. 2020;382:917–28.
4. Epstein JI, et al. Am J Surg Pathol. 2016;40:244–52.

ACKNOWLEDGEMENTS

The National Cancer Database (NCDB) is a joint project of the Commission on Cancer (CoC) of the American College of Surgeons and the American Cancer Society. The CoC's NCDB and the hospitals participating in the CoC's NCDB are the source of the de-identified data used herein; they have not verified and are not responsible for the statistical validity of the data analysis or the conclusions derived by the authors.

CONTACT INFORMATION

Aparna.Krishnan.Med@dartmouth.edu