

Welcome to Austin



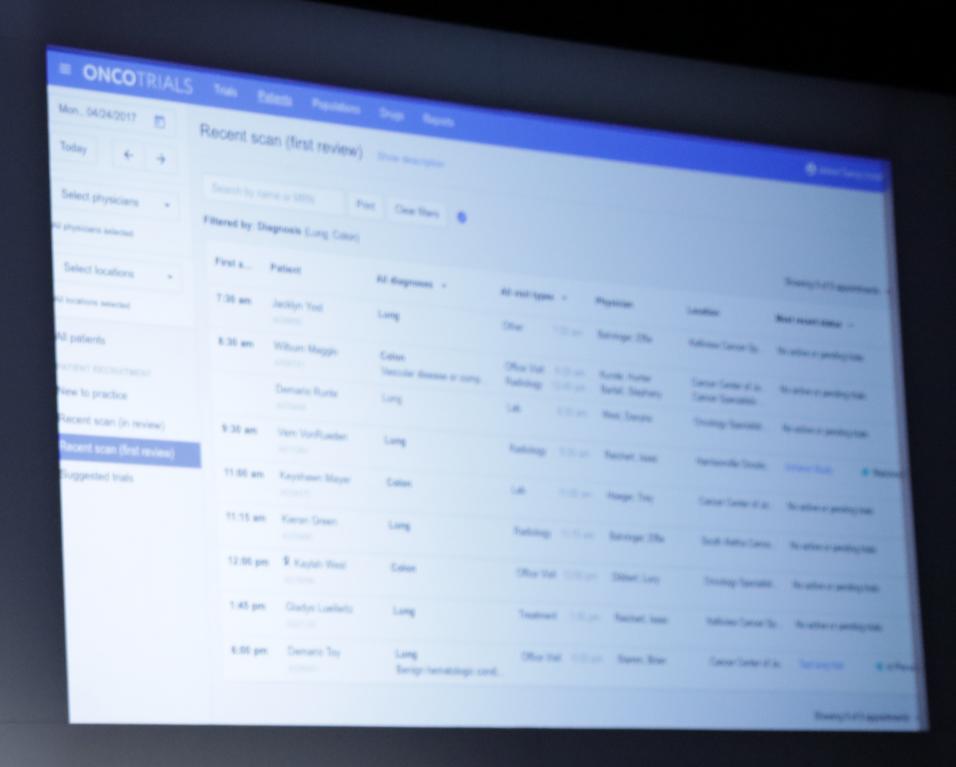
















500+

practice attendees

THANK YOU TO OUR SPONSORS:







































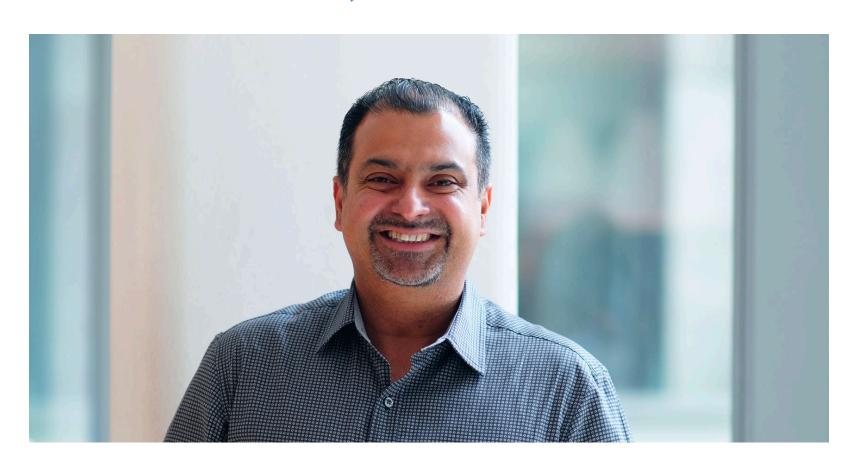


From: Tesh Khullar [mailto:tesh@flatiron.com]

Sent: Thursday, June 06, 2019 12:06 PM

Subject: [External]Flatiron Provider News - Tesh Khullar





Friends-

After what will be an amazing five and a half year journey at Flatiron, I wanted to let you all know that I have decided to move on from Flatiron at the end of 2019. It was a very tough decision for me but ultimately came down to me wanting to prioritize my family more and travel less. What made this decision so difficult was that I love working with you and will always consider community oncology my home. Although I do not know what I plan to do next, I am confident it will be something that will allow me to be home more.





Thank you.





Carolyn Starrett
Senior Vice President,
Provider Solutions



Goals for this weekend

Goals for this weekend

Bring practice partners together to create new connections

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Bring practice partners together to create new connections

Learn how you can get more from Flatiron's software, services and teams

Conference Agenda

General Sessions

State of the Union

Product Development

Tips and Trends for the Future of Community Practice

Workshop Tracks

Advancing the Business of Oncology

Streamlining Clinical Workflows

Empowering Operational Teams

Research Innovation



Askthe Experts

Level 5

SATURDAY

7:45am - 1:00pm

2:45pm - 5:45pm

SUNDAY

9:00am - 12:00pm

New Features

Level 5

SATURDAY

7:45am - 1:00pm

2:45pm - 5:45pm

SUNDAY

9:00am - 12:00pm



Communities





Lounge Level 5

SATURDAY

7:45am - 1:00pm

2:45pm - 5:45pm

SUNDAY

9:00am - 12:00pm

Roundtable Discussions Level 4

SUNDAY

10:00am - 12:00pm





Evening Evening Events

FRIDAY

Palm Park Terrace Fairmont Austin, Level 7 6:00pm - 9:00pm

SATURDAY

Banger's Sausage House & Beer Garden Historic Rainey Street 7:00pm - 11:00pm







Questions?

Download the conference app: search for "Flatiron Health" on the app store

Ask anyone with a purple staff badge

Visit the Registration Desk

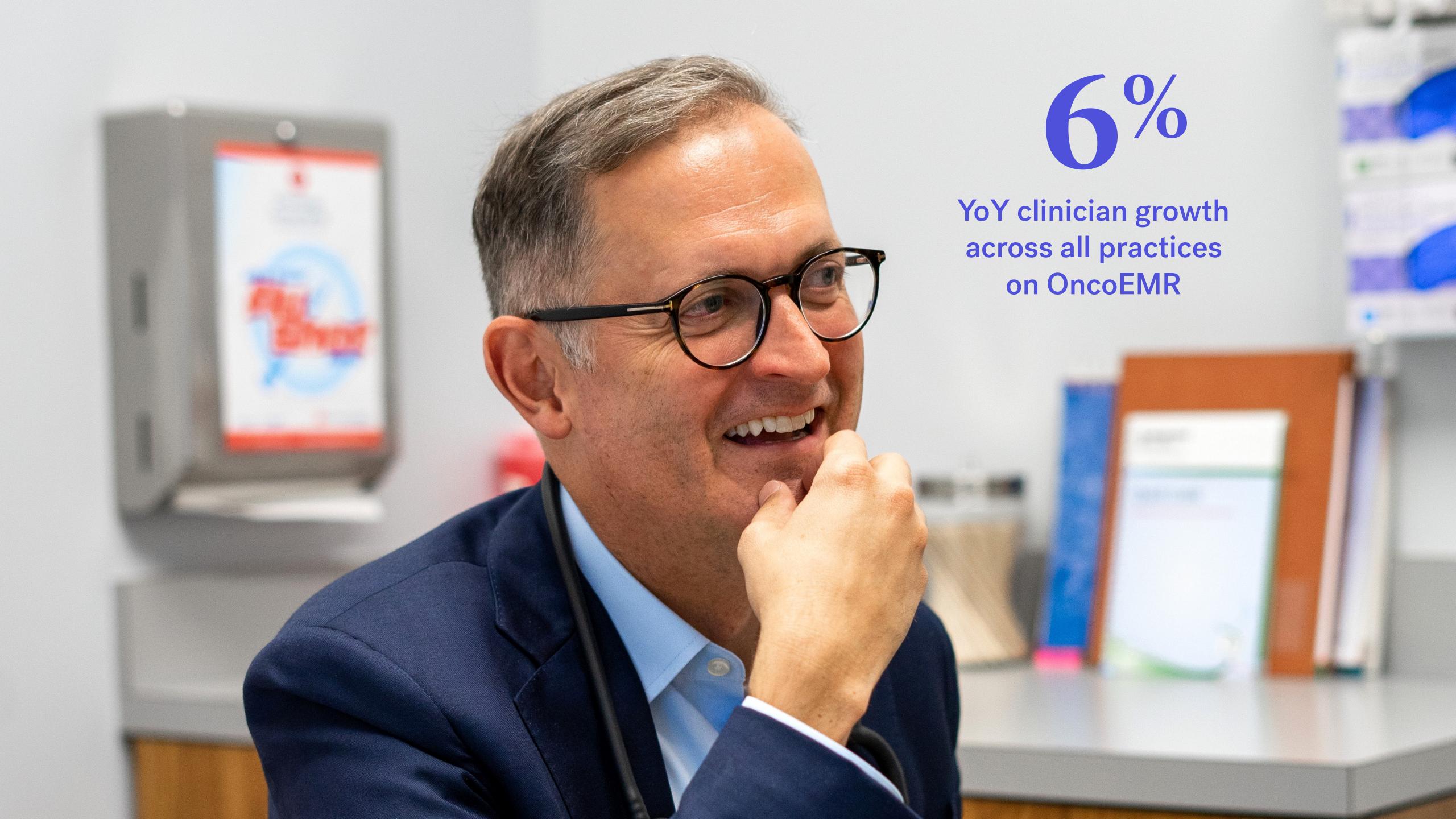
Email eventdesk@flatiron.com

State of the Union













ONCOLOGY

Pillars of Community Oncology







Perseverance & Flexibility





Perseverance & Flexibility



Innovation & Advocacy





Perseverance & Flexibility



Innovation & Advocacy







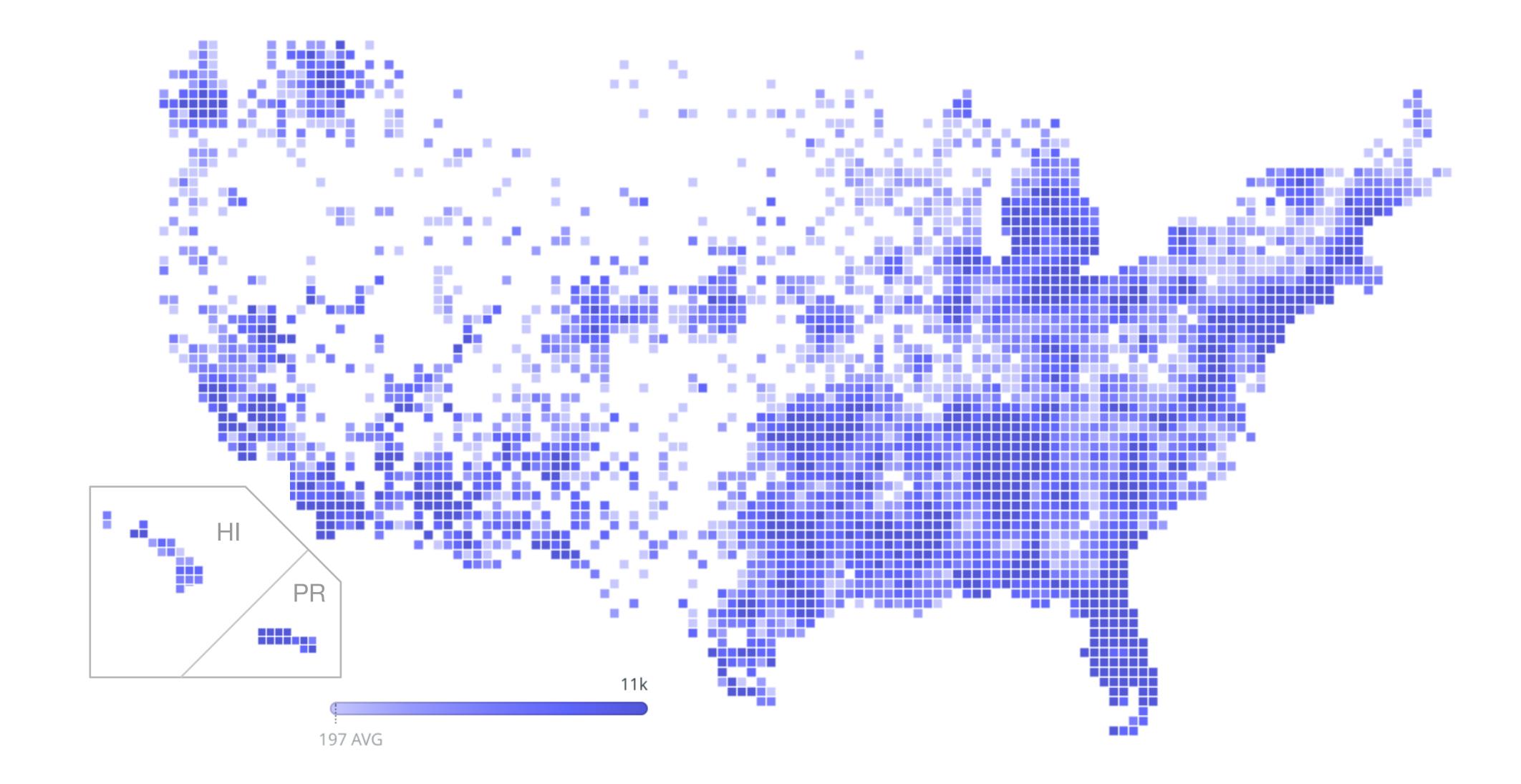


"There is a huge difference between a willingness to work on patients quickly and cheerfully, versus doing so with eye rolls and sighs—it's a very different experience for staff who have to ask a doctor to overbook."

Ted Arrowsmith, MDTennessee Oncology

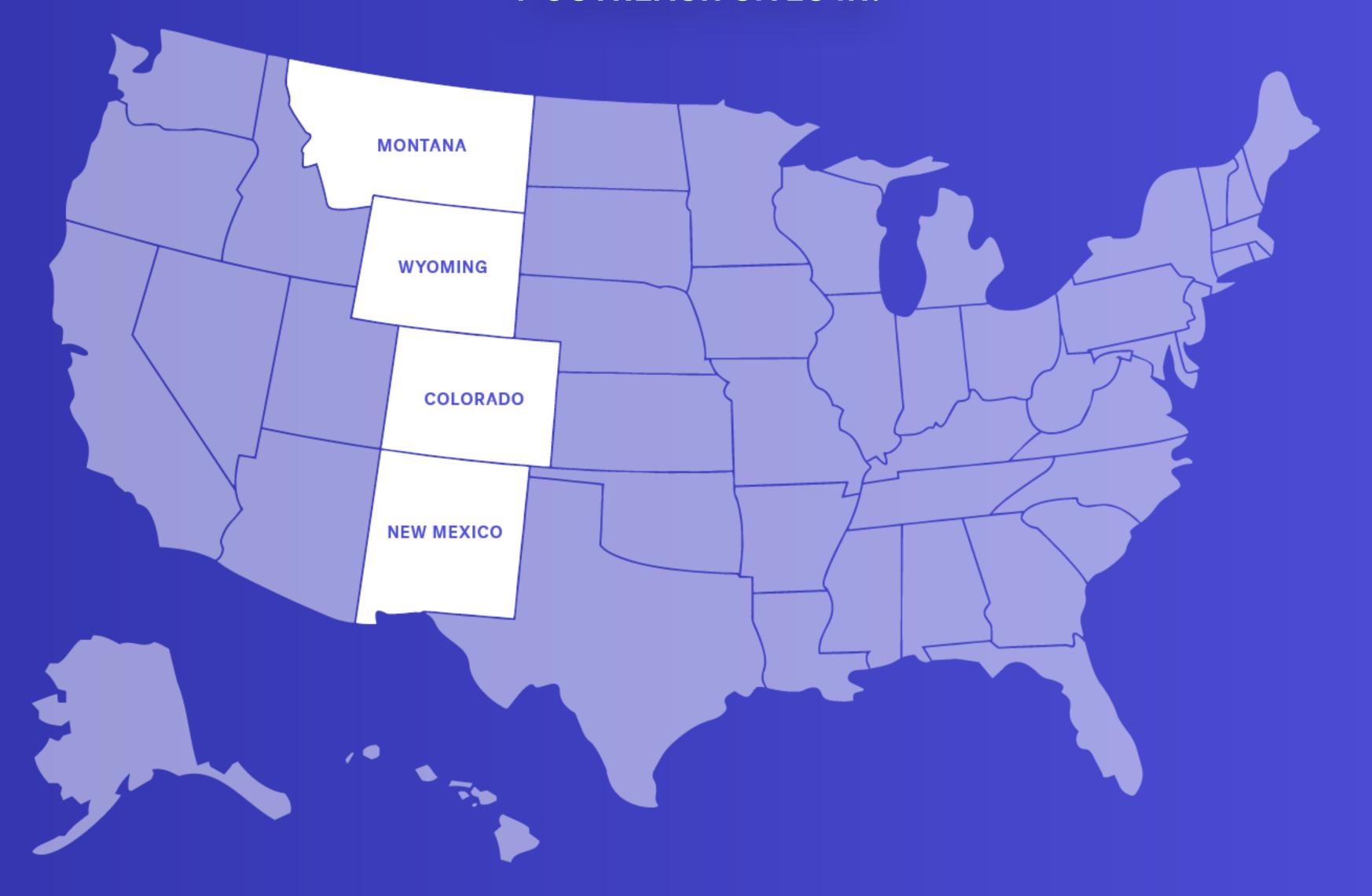








7 OUTREACH SITES IN:









Perseverance & Flexibility



Innovation & Advocacy





Perseverance & Flexibility



Innovation & Advocacy



Julia Morton
Senior Director,
Customer Success





Next





Edit

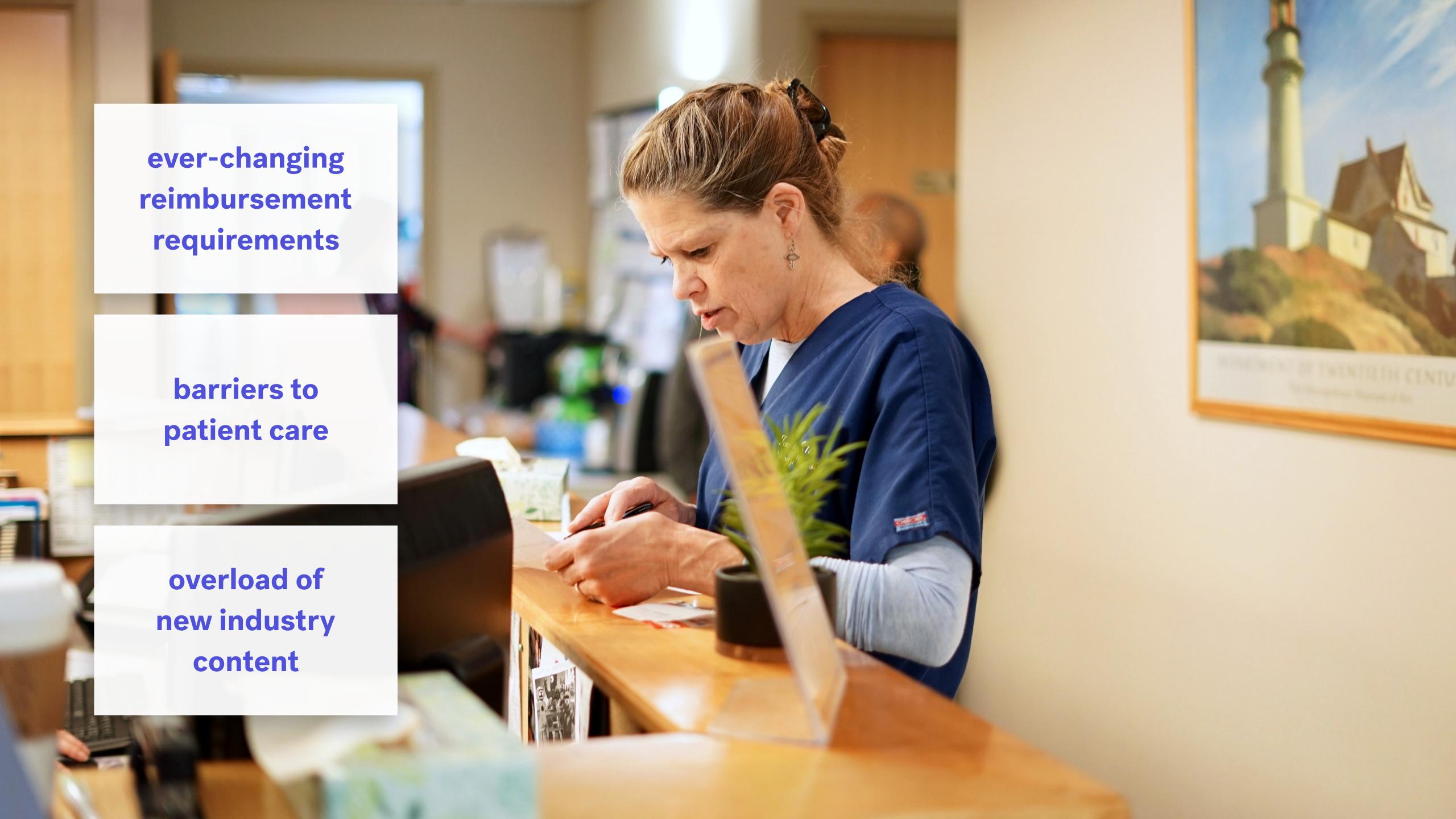
Filter













OCM standards for every patient



OCM standards for every patient

expanded triage system



OCM standards for every patient

- expanded triage system
- 24/7 CCM services



OCM standards for every patient

- expanded triage system
- 24/7 CCM services
- survivorship and EOL planning
- social worker partnerships
- / clinical standards committee













ENTERS FOR CANCER CARE AND BLOOD DISORDERS





Perseverance & Flexibility



Innovation & Advocacy





Perseverance & Flexibility



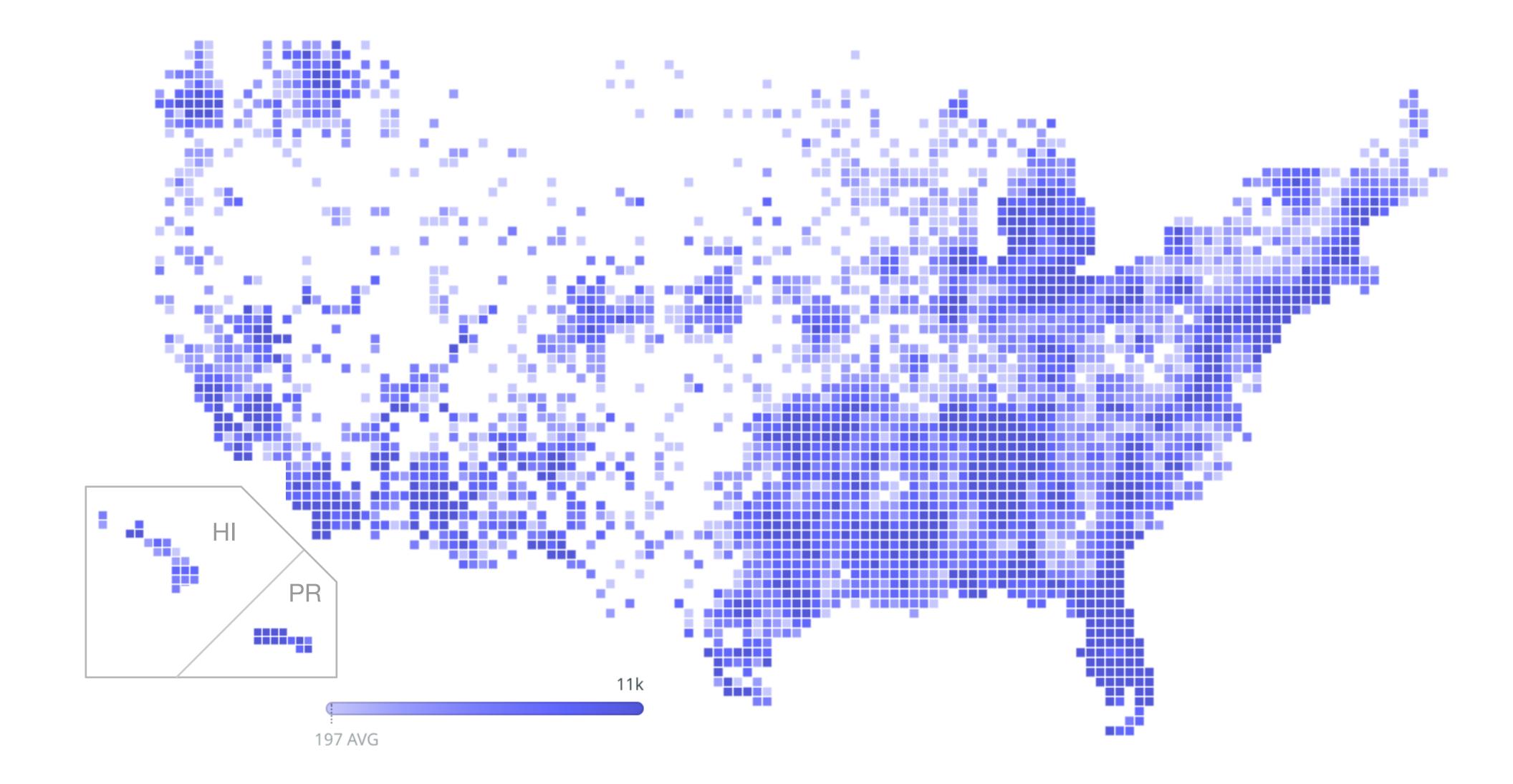
Innovation & Advocacy



Bobby Green, MD
Chief Medical Officer











OPINION | COMMENTARY

How I Was Wrong About ObamaCare



The law's drafters wanted consolidation: 112 hospital mergers last year. But smaller practices have improved care better.



By Bob Kocher July 31, 2016 4:35 pm ET

I was wrong. Wrong about an important part of ObamaCare.

When I joined the Obama White House to advise the president on health-care policy as the only physician on the National Economic Council, I was deeply committed to developing the best health-care reform we could to expand coverage, improve quality and bring down costs. We worked for months to pass this landmark legislation, and I still count celebrating the passage of the Affordable Care Act with the president one balmy spring night in 2010 as one of my greatest Washington memories.





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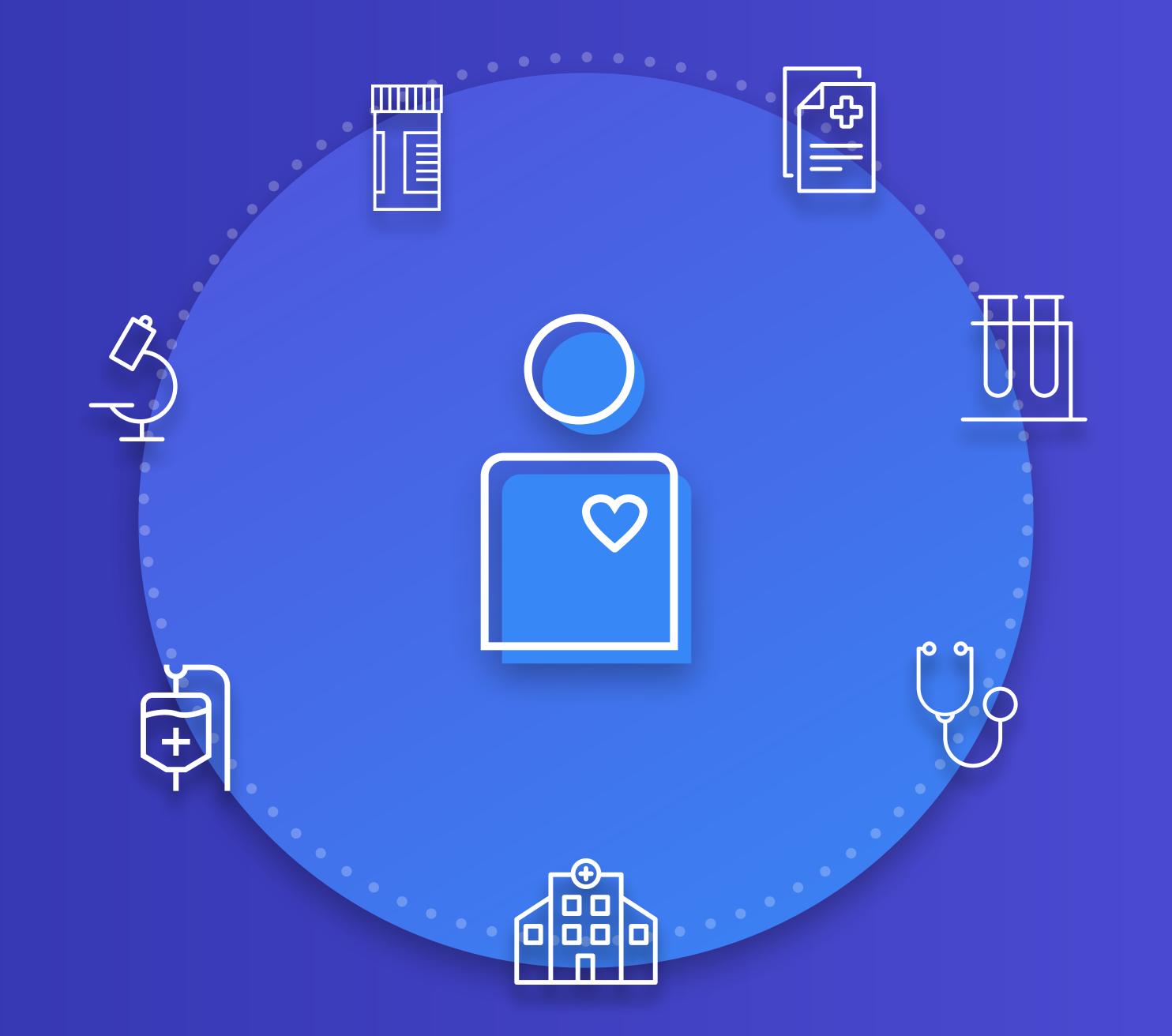
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ORIGINAL ARTICLE

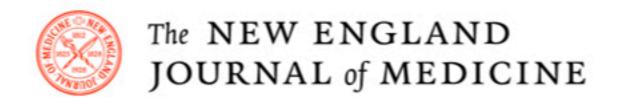
Nivolumab versus Docetaxel in Advanced Squamous-Cell Non–Small-Cell Lung Cancer

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M.D., Wilfried E.E. Eberhardt, M.D., Elena Poddubskaya, M.D., Scott Antonia,
M.D., Ph.D., Adam Pluzanski, M.D., Ph.D., Everett E. Vokes, M.D., Esther
Holgado, M.D., Ph.D., David Waterhouse, M.D., Neal Ready, M.D., Justin Gainor,
M.D., Osvaldo Arén Frontera, M.D., Libor Havel, M.D., Martin Steins, M.D.,
Marina C. Garassino, M.D., Joachim G. Aerts, M.D., Manuel Domine, M.D., Luis
Paz-Ares, M.D., Martin Reck, M.D., Christine Baudelet, Ph.D., Christopher T.
Harbison, Ph.D., Brian Lestini, M.D., Ph.D., and David R. Spigel, M.D.

July 9, 2015

N Engl J Med 2015; 373:123-135

DOI: 10.1056/NEJMoa1504627



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Journal of Thoracic Oncology

<u>J Thorac Oncol.</u> 2018 May;13(5):682-688. doi: 10.1016/j.jtho.2018.02.022. Epub 2018 Mar 6.

Phase 1/2 Study of the Safety and Tolerability of Nivolumab Plus Crizotinib for the First-Line Treatment of Anaplastic Lymphoma Kinase Translocation - Positive Advanced Non-Small Cell Lung Cancer (CheckMate 370).

Spigel DR¹, Reynolds C², Waterhouse D³, Garon EB⁴, Chandler J⁵, Babu S⁶, Thurmes P⁷, Spira A⁸, Jotte R⁹, Zhu J¹⁰, Lin WH¹⁰, Blumenschein G Jr¹¹.

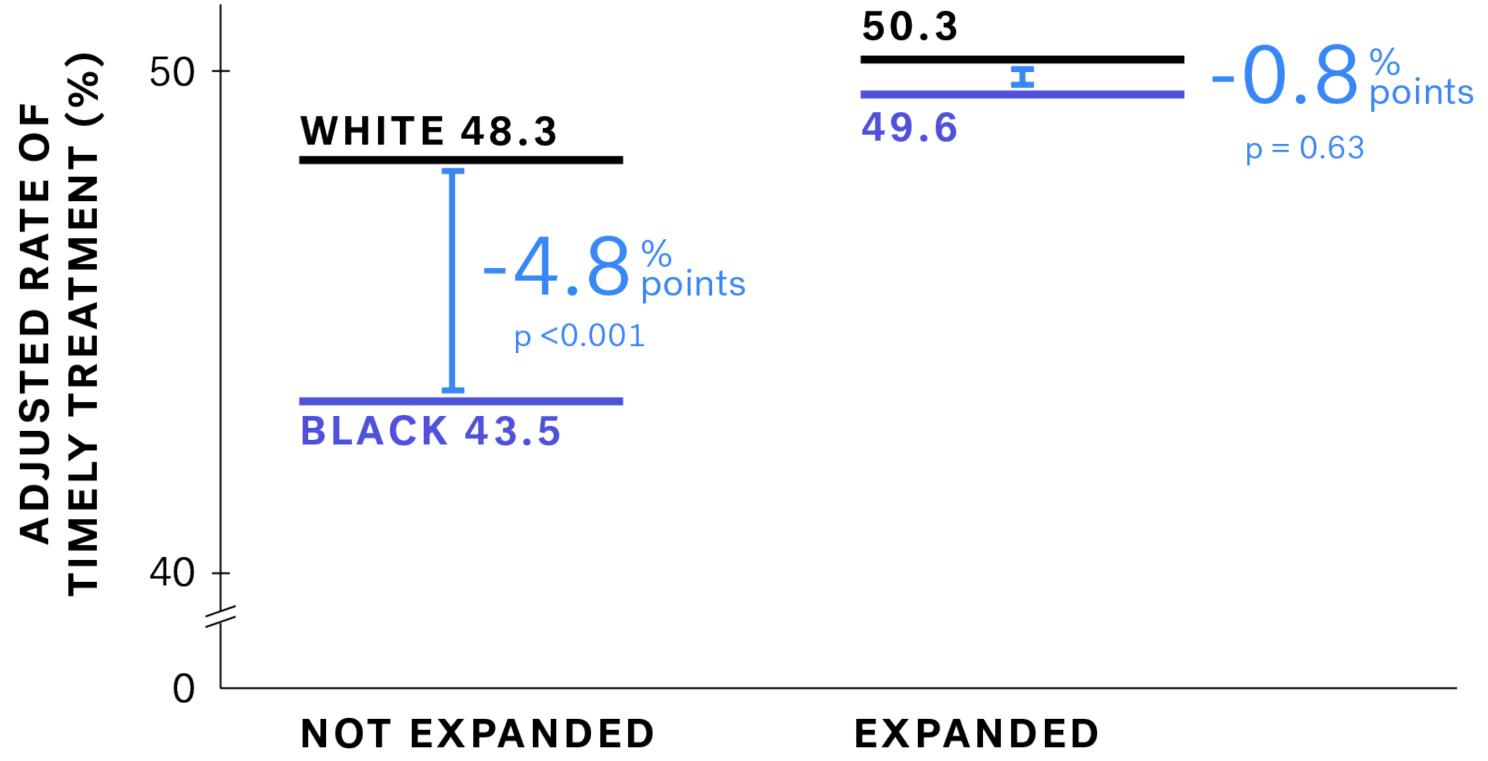
- 1 Sarah Cannon Research Institute, Nashville, Tennessee. Electronic address: dspigel@tnonc.com.
- 2 Florida Cancer Specialists, Ocala, Florida.
- OHC (Oncology Hematology Care), an affiliate of US Oncology Research, Cincinnati, Ohio; US Oncology Research, The Woodlands, Texas.
- 4 David Geffen School of Medicine at UCLA/Translational Research in Oncology-US Network, Santa Monica, California.
- 5 West Cancer Center, Memphis, Tennessee.
- 6 Fort Wayne Medical Oncology and Hematology, Fort Wayne, Indiana.
- 7 Minnesota Oncology, Minneapolis, Minnesota.
- 8 US Oncology Research, The Woodlands, Texas; Virginia Cancer Specialists, Fairfax, Virginia.
- 9 US Oncology Research, The Woodlands, Texas; Rocky Mountain Cancer Center, Denver, Colorado.
- 10 Bristol-Myers Squibb, Princeton, New Jersey.
- 11 Department of Thoracic/Head and Neck Medical Oncology, The University of Texas MD Anderson Cancer Center, Houston, Texas.

OUR MISSION:

To improve lives by learning from the experience of every cancer patient.



Results: Change in Racial Disparity



Difference-in-Differences

4.0% points
Reduction in disparity
p = 0.042

Overall Mean: 47.7% Overall Mean: 49.8%





"We need to shift the data paradigm and understand how to safely and appropriately use our patients' data. This study is a leap forward and provides proof of concept that healthcare systems can use their data to improve outcomes and possibly reduce disparities."

Yousuf Zafar, MDDuke Cancer Institute

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Yousuf Zafar, MDDuke Cancer Institute





2019 World Conference on Lung Cancer

September 7–10, 2019 | Barcelona, Spain Conquering Thoracic Cancers Worldwide

MA14 THE ADEQUATE MTARGET IS STILL THE ISSUE MONDAY, SEPTEMBER 9 15:45–17:15

MA14.09 REAL-WORLD SURVIVAL OF RELAPSED COMPARED TO DE-NOVO STAGE IV DIAGNOSIS OF ADVANCED NON-SMALL CELL LUNG CANCER

K. Kerrigan¹, B. Haaland¹, B. Adamson², S. Patel¹, W. Akerley¹

Huntsman Cancer Institute, Salt Lake City/United States of America, ²Flatiron Health, New York City/United States of America

Background: Differences in tumor biology and cancer therapy in early stage lung cancer may affect overall survival (OS) of patients with relapsed stage IV disease compared to others with de-novo stage IV disease. This study aimed to compare real-world survival of these patients. Method: We selected patients with advanced NSCLC diagnosed between 2011 and 2017, who received at least one line of therapy, from the US Flatiron Health electronic health record-derived database. Patient data was collected through June 2018, providing at least 6 months of follow-up. OS was defined as time from advanced or metastatic diagnosis to the event of death,

censored at lass OS of patients fit multivariable hazard of death Overall survival

Progressed to Stage 4
De Novo Stage 4
Stratified Multivariate HR 0.76 (95% CI 0.73-0.79, p<0.001)

Future Medicine 1

JOURNAL OF COMPARATIVE EFFECTIVENESS RESEARCH, AHEAD OF PRINT | RESEARCH ARTICLE

Comparative effectiveness of first-line nab-paclitaxel versus paclitaxel monotherapy in triple-negative breast cancer

Patricia Luhn [™], Stephen Y Chui, "Angela" Fu-Chi Hsieh, Jingbo Yi, Almut Mecke, Preeti S Bajaj, Waseem Hasnain, Adeline Falgas, Thanh GN Ton & Allison W Kurian

Published Online: 9 Aug 2019 | https://doi.org/10.2217/cer-2019-0077

Aim: This observational study evaluated the effectiveness of nab-paclitaxel versus paclitaxel monotherapy as first-line (1L) treatment for metastatic triple-negative breast cancer (mTNBC). Materials & methods: 200 patients from the US Flatiron Health electronic health record-derived database (mTNBC diagnosis, January 2011–October 2016) who received 1L nab-paclitaxel (n = 105) or paclitaxel (n = 95) monotherapy were included. Overall survival and time to next treatment were evaluated. Results: The adjusted overall survival hazard ratio was 0.98 (95% CI: 0.67–1.44), indicating a similar risk of death between groups. Adjusted time to next treatment hazard ratio was 0.89 (95% confidence interval: 0.62–1.29). Conclusion: Nab-paclitaxel and paclitaxel monotherapy showed similar efficacy, suggesting their interchangeability as 1L treatments for mTNBC.

BMC Part of Springer Nature

BMC Medical Research Methodology

Validation of diagnosis codes to identify side of colon in an electronic health record registry

Patricia Luhn ™, Deborah Kuk, Gillis Carrigan, Nathan Nussbaum, Rachael Sorg, Rebecca Rohrer, Melisa G. Tucker,
Brandon Arnieri, Michael D. Taylor & Neal J. Meropol

Abstract

Background

The use of real-world data to generate evidence requires careful assessment and validation of critical variables before drawing clinical conclusions. Prospective

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Real-world progression, treatment, and survival outcomes during rapid adoption of immunotherapy for advanced nonsmall cell lung cancer

First published: 05 August 2019 | https://doi.org/10.1002/cncr.32383

We thank Nicole Lipitz, Julia Saiz Shimosato, Chris Gayer, and Sam Azaria, from Flatiron Health, for editing and administrative support.

Dr. Abernethy participated in this work prior to joining the FDA. This work and related conclusions reflect the independent work of study authors and does not necessarily represent the views of the US Food and Drug Administration or the United States.

The copyright line for this article was changed on 5 August 2019 after original online publication.

Abstract

Background

Despite the rapid adoption of immunotherapies in advanced non–small cell lung cancer (advNSCLC), knowledge gaps remain about their real-world (rw) performance.

Methods

This retrospective, observational, multicenter analysis used the Flatiron Health deidentified electronic health record-derived database of rw patients with advNSCLC who received treatment with PD-1 and/or PD-L1 (PD-[L]1) inhibitors before July 1, 2017 (N = 5257) and had ≥6 months of follow-up. The authors investigated PD-(L)1 line of treatment and PD-L1 testing rates and the relationship between overall survival (OS) and rw intermediate endpoints: progression-free survival (rwPFS), rw time to

JCO® Clinical Cancer Informatics An American Society of Clinical Oncology Journal

Characterizing the Feasibility and Performance of Real-World Tumor Progression End Points and Their Association With Overall Survival in a Large Advanced Non–Small-Cell Lung Cancer Data Set

Sandra D. Griffith, PhD¹; Rebecca A. Miksad, MD, MPH¹ Geoff Calkins¹; Paul You, MPH¹; Nicole G. Lipitz¹; Ariel B. Bourla, MD, PhD¹; ...

ABSTRACT

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Article

Using Electronic Health Records to Derive Control Arms for Early Phase Single-Arm Lung Cancer Trials: Proof-of-Concept in Randomized Controlled Trials

First published: 27 July 2019 | https://doi.org/10.1002/cpt.1586

This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1002/cpt.1586

Abstract

Oncology drug development increasingly relies on single-arm clinical trials. External controls (EC) derived from electronic health record (EHR) databases may provide additional context. Patients from a US based oncology EHR database were aligned with patients from randomized controlled trials (RCTs) and trial specific eligibility criteria were applied to the EHR dataset. Overall survival (OS) in the EC derived control arm was compared to OS in the RCT experimental arm. The primary outcome was OS, defined as time from randomization or treatment initiation (EHR) to death. Cox regression models were used to obtain effect estimates using EHR data. EC-derived hazard ratio (HR) estimates aligned closely with those from the corresponding RCT with one exception. Comparing log HRs between all RCT and EC results gave a Pearson correlation coefficient of 0.86. Properly selected control arms from contemporaneous EHR data could be used to put single-arm trials of OS in advanced non-small cell lung cancer (aNSCLC) into context.

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2019 World Conference on Lung Cancer

MA14.09 REAL-WORLD SURVIVAL OF RELAPSED

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BMC Part of Springer Nature

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JCO° Clinical Cancer Informatics

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Characterizing the Feasibility and Performance of Real-**World Tumor Progression End Points and Their Association With Overall Survival in a Large Advanced** Non-Small-Cell Lung Cancer Data Set

Sandra D. Griffith, PhD¹; Rebecca A. Miksad, MD, MPH^{1 🔀}; Geoff Calkins¹; Paul You, MPH¹; Nicole G. Lipitz¹; Ariel B. Bourla, MD, PhD¹; ...



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