Suzetrigine for Acute Pain: Effectiveness and Value

Public Meeting — February 28, 2025

Meeting materials available at: <u>https://icer.org/assessment/acute-pain-2025</u>





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Participating Members of the Midwest CEPAC

Jill Johnson, PharmD, Chair

Professor, College of Pharmacy, University of Arkansas for Medical Sciences

- Eric Armbrecht, PhD, Professor, Saint Louis University
- Alan J. Balch, PhD, CEO, Patient Advocate Foundation
- **Bijan Borah, PhD,** Professor of Health Services Research, Mayo Clinic College of Medicine and Science
- **Donald Casey, MD, MPH, MBA, MACP, FAHA**, Associate Professor of Internal Medicine, Rush Medical College
- **Gregory Curfman, MD,** Executive Editor, JAMA, American Medical Association
- Sneha Dave, Executive Director, Generation Patient
- Yngve Falck-Ytter, MD, AGAF, Professor, Case Western Reserve University

- Heather Guidone, BCPA, Program Director, Center for Endometriosis Care
- Jayani Jayawardhana, Ph.D, Associate Professor, University of Kentucky
- David Kim, PhD, Assistant Professor, University of Chicago
- **Timothy McBride, PhD,** Washington University in St. Louis
- Jimi Olaghere, Patient Advocate
- Stuart Winston, DO, Patient Experience Lead Consultant, Trinity Health IHA Medical Group

Patient Experts

Nicole Hemmenway, CEO, US Pain Foundation

 The U.S. Pain Foundation receives over 25% of its funding from a diverse range of sources, including healthcare companies, industry groups such as Vertex Pharmaceuticals, family foundations, and individual donors. Additionally, the organization previously engaged in a fee-for-service collaboration with Humana Neighborhood Centers providing educational content about pain. All educational materials are developed independently, without any input or review from funders, and intended solely for educational purposes.

Gabriel Smith, Patient

• No conflicts to disclose.

Clinical Experts

Benjamin Friedman, MD, MS, Professor of Emergency Medicine, Montefiore Einstein

• Dr. Friedman is an attending physician at Montefiore Einstein and has no conflicts to disclose.

Andrew Kolodny, MD, Medical Director, Opioid Policy Research, Heller School, Brandeis University

• Dr. Kolodny has served as an expert witness in litigation involving the opioid industry.



ICER Speakers



Sarah K. Emond, MPP President & CEO



Brett McQueen, PhD Lead Modeler & Associate Professor, Department of Clinical Pharmacy, University of Colorado



Dan Ollendorf, PhD, MPH *Chief Scientific Officer and Director of HTA Methods and Engagement*



David Rind, MD, MSc *Evidence Author & Chief Medical Officer*

Why are we here today?

"I don't have an MD at the end of my name, but you are not in my body. You can't tell me what I am feeling. The pain I have is real."

"I lost my identity and my career. I wasn't able to hike, run or play with my kids like I wanted to. There's so much I wish I could have done."

Real patient stories from the InvisibleProject.Org

Why Are We Here Today?

- What happens the day these treatments receive FDA approval?
- Questions about:
 - What are the risks and benefits?
 - How do new treatments fit into the evolving landscape?
 - What are reasonable prices and costs to patients, the health system, and the government?
 - What lessons are being learned to guide our actions in the future?



The Impact on Rising Health Care Costs for Everyone

DIAGNOSIS: DEBT

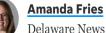
100 Million People in America Are Saddled With Health Care Debt

By Noam N. Levey JUNE 16, 2022



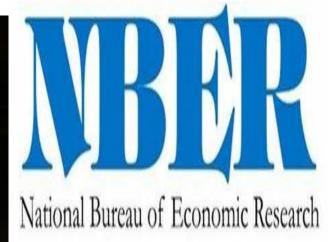


Why Delaware is eying a 27% premium hike on state employees' health insurance



Delaware News Journal

Published 4:35 a.m. ET Feb. 1, 2024 | Updated 9:29 p.m. ET Feb. 6, 2024



WHO PAYS FOR RISING **HEALTH CARE PRICES?**



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100 Million People in America Are Saddled With Health Care Debt (KFF Health News)

Why Delaware is eyeing a 27% premium hike on state employees' health insurance (Delaware Online)





Organizational Overview







Funding 2025

ICER Analytics Manufacturer **Subscribers** Philanthropy/Other Contributions 9% 1% Health Plans and 13% **Provider Group** Contributions 8% Nonprofit Foundations 69%

ICER Policy Summit and non-report activities only



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How Was the ICER Report Developed?

Scoping	Evidence Synthes and Model Development		Review	Draft Report	Public Comment and Revision	Evidence Report
Guidance fro patients, clini experts, manufacturer and other stakeholders	n Evidence ana cal ICER team a cost-effective	ysis by id iess • vith v of	Medicine, S University Benjamin I Medicine, M Kit Simpso Leadership South Caro Nicole Hen Pain Found	nmenway, Chief Exe	alth Sciences essor of Emergency of Health Care Medical University of ecutive Officer, US	Structured to support Midwes CEPAC voting and policy discussion



Value Assessment Framework: Long-Term Value for Money

Special Social/Ethical Priorities

Benefits Beyond "Health"

Total Cost Overall Including Cost Offsets

Health Benefits: Return of Function, Fewer Side Effects

> Health Benefits: Longer Life



Agenda (Central Time)

10:00 AM	Meeting Convened and Opening Remarks				
10:20 AM	Presentation of the Clinical Evidence				
11:00 AM	Presentation of the Economic Model				
11:40 AM	Public Comments and Discussion				
12:00 PM	Lunch Break				
12:50 PM	Midwest CEPAC Deliberation and Vote				
1:50 PM	Break				
2:00 PM	Policy Roundtable Discussion				
3:30 PM	Reflections from Midwest CEPAC				
4:00 PM	Meeting Adjourned				

Presentation of the Clinical Evidence

David Rind, MD, MSc

Chief Medical Officer

Institute for Clinical and Economic Review



Key Collaborators

Team Role	Assigned Team Member
Research Lead	Dmitriy Nikitin, MSPH
Research Assistants	Finn Raymond, BS, Sol Sanchez, BA

Disclosures

David Rind, Dmitriy Nikitin, Finn Raymond, and Sol Sanchez are employees of the Institute for Clinical and Economic Review (ICER) and have no conflicts to disclose.



Acute Pain

- Acute pain is ubiquitous
 - Definitions vary, but time-limited
- At least 80 million in the US annually receive prescription medications for acute pain
- Nearly all systemic treatments include NSAIDs, acetaminophen, or opioids
- Today we are talking about acute pain, not chronic pain



Patient Impact

- Most patients'/patient groups' focus is around chronic pain.
- As with sickle cell disease, concerns around:
 - Undertreatment of pain
 - Overuse of opioids
 - Stigma
- Inadequate treatment leads to ER care
- Lack of access to multimodal pain management
- OUD and the opioid epidemic



Opioids

- Being prescribed less frequently and in smaller amounts
- Risks include sedation, confusion, falls, GI side effects, respiratory depression
- Rate of opioid use disorder (OUD) after short treatment of acute pain is uncertain
- Risk is clearly high in people with prior OUD



- Readily available for minor to moderate pain
- Risks include GI bleeding, acute kidney injury, CV events
- Risks with short-term treatment uncertain
- Some pain experts feel these are being underutilized





Oral medication for moderate to severe acute pain in adults



Administered every 12 hours



Inhibitor of Na_v1.8 (new mechanism of action)



FDA approved on January 30th, 2025 ("Journavx®")



Insights from Discussions with Patients

- Need for education around appropriate use of opioid pain medication
- Different types of pain elicit different pain responses
- Openness to alternative treatments for acute pain without risk of addiction
- Patient-important outcomes include quality of life, physical functioning and interference in activities of daily living, development of chronic pain, use of rescue medication, and opioid avoidance





- How does suzetrigine efficacy compare to other options?
- How does suzetrigine safety compare to other options?



Clinical Evidence

Pivotal Trials: NAVIGATE-1 and -2

Study Design

- Post-operative abdominoplasty and bunionectomy with ≥4 on the NPRS
- Phase III double-blind trials
- Randomized 2:2:1 to suzetrigine, hydrocodone 5 mg/APAP 325 mg, or placebo
- Rescue with ibuprofen 400 mg
- No peer-reviewed publication

Baseline Characteristics

- 2191 participants across both trials
- 92% Female
- About 1/3 severe pain; 2/3
 - moderate pain at baseline
 - Higher pain in abdominoplasty

population

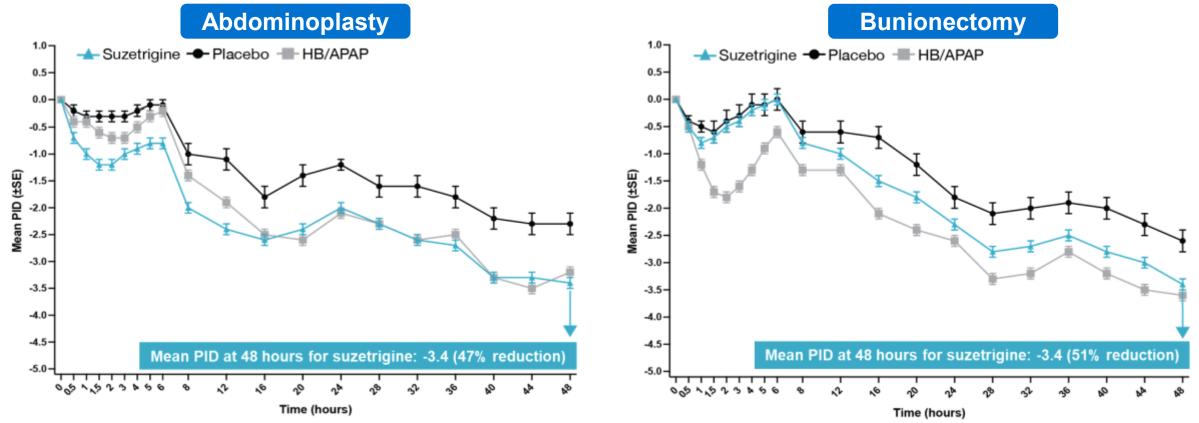




- Primary outcome: Time-weighted pain intensity over 48 hours
- Other outcomes:
 - Time to ≥2-point reduction in pain intensity on the Numeric Pain Rating Scale from baseline



Clinically Meaningful Pain Relief at 48 Hours Observed With Suzetrigine After Procedures



Note: Figures include participants who were randomized and received at least one dose of study drug. Participants were analyzed according to their randomized treatment. HB/APAP: hydrocodone bitartrate/acetaminophen, PID: pain intensity difference, SE: standard error, SPID48: time-weighted sum of the pain intensity difference as recorded on the NPRS from 0 to 48 hours

inical and Economic Review

Reprinted from "Randomized, Placebo-Controlled, Phase 3 Trials of Suzetrigine, a Non-

Opioid, Pain Signal Inhibitor for Treatment of Acute Pain After Abdominoplasty or 28 Bunionectomy," by Todd Bertoch, 2024, presented at The Anesthesiology Annual Meeting. Copyright (2024), with permission from Vertex Pharmaceuticals.

What Don't We Know About Efficacy?

- No head-to-head trials with NSAIDs
- Opioid dose was low
- Rescue medication use (new data)
- Network meta-analysis to examine NSAIDs and higher-dose opioids



Network Meta-Analysis Overview

- Population: Adults undergoing bunionectomy and abdominoplasty procedures with moderate to severe acute pain
- Interventions: Suzetrigine vs NSAIDs vs low- and high-dose opioids vs placebo
- Outcome: Time-weighted pain intensity over 48 hours



Network Meta-Analysis Results

• Wide confidence intervals around results

High-Dose Opioid				
0.14 (-0.13, 0.41)	Suzetrigine			
0.17 (-0.08, 0.47)	0.04 (-0.14, 0.24)	Low-Dose Opioid		_
0.22 (-0.04, 0.5)	0.08 (-0.18, 0.35)	0.05 (-0.24, 0.3)	NSAID	
0.56 (0.37, 0.76)	0.42 (0.24, 0.61)	0.39 (0.18, 0.56)	0.34 (0.15, 0.52)	Placebo

NSAID: nonsteroidal anti-inflammatory drugs

Standardized mean differences greater than 0 favor the column-defining treatment. Significant results are in bold.

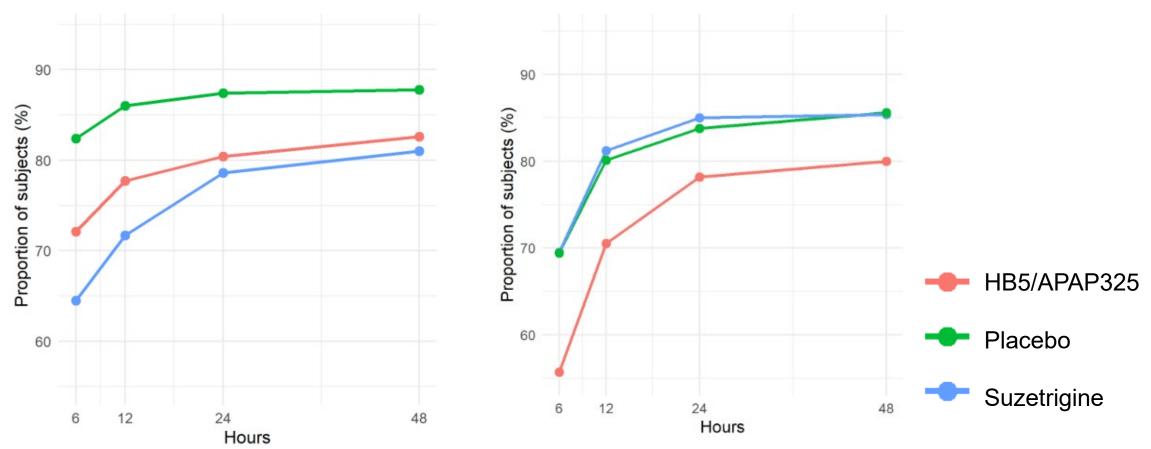


Rescue Medication

- Rescue with ibuprofen 400 mg
- Rates of rescue medication not initially reported
- Implications of rescue medication
- Likely effects:
 - Versus placebo
 - Versus hydrocodone/APAP

Rescue Medication Use

Abdominoplasty



Bunionectomy

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Figures reprinted from FDA CDER Integrated Review of JOURNAVX Reference ID: 5521299

What Don't We Know Around Safety/Harms?

- Pain medications need to be very safe
- Information from clinical trials and biologic plausibility
- Unknown risks
- Comparator risks



Suzetrigine Harms from Phase III Trials

- Well tolerated
- Side effects similar to placebo
- Lower incidence of nausea and vomiting compared with hydrocodone/APAP



Uncertainty Around Harms

- Renal harms
 - Six of 55 patients with diabetic neuropathy treated for 12 weeks had decreased renal function
- Brugada Syndrome
 - Na_v1.8 is encoded by the gene SCN10A
- Unknown harms
 - Rofecoxib
 - Addiction risk

Comparator Harms

- Widely varying estimates of OUD risk with one week of treatment
- NSAID short-term risks (cardiac, GI, renal) also uncertain



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Overall Uncertainties

- Dose of hydrocodone/APAP
- Time to benefit
- Rescue medication
- Comparison with NSAIDs
- Known and unknown risks



Benefits Beyond Health and Special Ethical Priorities

- Unmet need: Many patients not good candidates for existing pain medications
- Populations underserved by the healthcare system may both be undertreated for pain and overtreated with opioids
- Caregiver benefits if OUD is reduced
- New mechanism of action

Public Comments Received

- Manufacturer notes that HC 5 mg/APAP 325 mg is the most commonly prescribed dose for acute pain
- Patient group expressed hope that suzetrigine would decrease progression to chronic regional pain syndrome
- Patient input: many stories of impacts of pain and OUD



Summary

- Efficacy:
 - Greater than placebo
 - Likely similar to low-dose opioids; bunionectomy versus abdominoplasty?
 - Uncertain comparisons with NSAIDs and high-dose opioids
- Harms:
 - Possible renal harms; less likely cardiac harms
 - Concerns around unknown harms
 - Will require time on the market to be sure; sequencing of patient selection

ICER Evidence Ratings

Treatment	Comparator	Evidence Rating
	No systemic therapy	P/I
Suzetrigine	Opioid analgesics	P/I
	Nonsteroidal anti-inflammatory drugs	P/I

P/I: "Promising but Inconclusive" - Moderate certainty of a small or substantial net health benefit, small (but nonzero) likelihood of a negative net health benefit





Presentation of the Economic Model

Brett McQueen, PhD

Associate Professor

University of Colorado, Anschutz Medical Campus



Key Review Team Members

Team Role	Assigned Team Member
Modelers	Brett McQueen, PhD, Michael DiStefano, PhD, MBE, Antal Zemplenyi, PhD
Economics Lead	Woojung Lee PharmD, PhD

Disclosures

WL is an employee of the Institute for Clinical and Economic Review (ICER) and has no conflicts to disclose.

BM, MD, and AZ have no conflicts to disclose defined as more than \$10,000 in healthcare company stock or more than \$5,000 in honoraria or consultancies relevant to this report during the previous year from health care manufacturers or insurers.



Objective

Acute Pain

To evaluate the lifetime cost-effectiveness of suzetrigine compared to hydrocodone bitartrate/acetaminophen (HB/APAP) for the treatment of moderate-to-severe acute pain.



Unmet Need

Condition	Absolute evLY Shortfall	Proportional evLY Shortfall		
Acute Pain	0.24	0.8%		
Other Example Conditions				
High Cholesterol	1.7	11%		
Multiple Sclerosis	18.9	52%		
Osteoporosis	2.6	19%		

evLY: equal value of life years

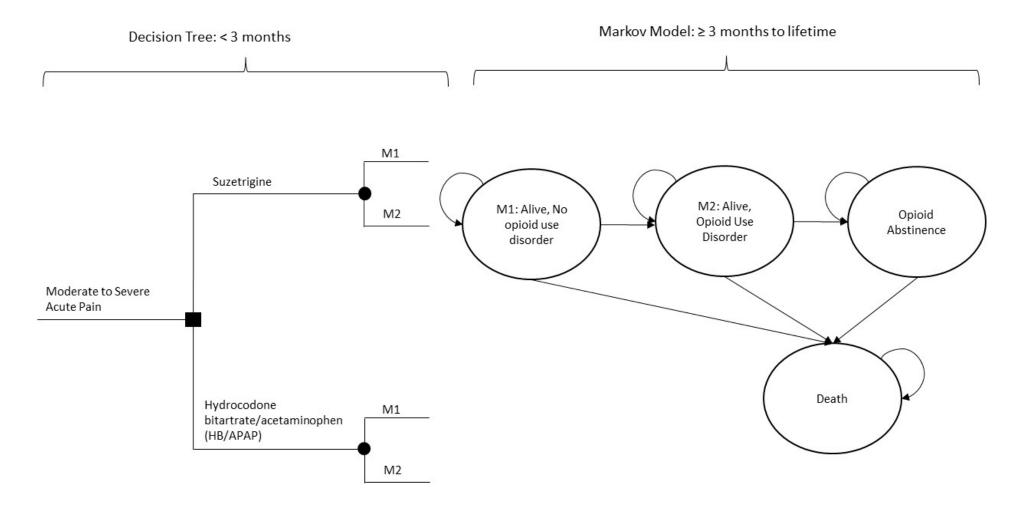
Methods in Brief

Methods Overview

Domain	Approach		
Model	Decision Tree and Markov Model		
Setting	United States		
Perspective	Health Care Sector Perspective and Modified Societal Perspective		
Time Horizon	Lifetime		
Discount Rate	3% per year (costs and outcomes)		
Cycle Length	Annual cycle after first three months		
Primary Outcome	Cost per quality-adjusted life year (QALY) gained, evLY gained, life years gained, cost per opioid use disorder (OUD) case averted		



Model Schematic



Model Characteristics

- Target Population
 - Mean age (years): 45.3
 - Percent male: 45.5



Key Assumptions

Assumption #1

 The model focused on acute pain requiring up to one week of prescription pain medication, such as surgery or other acute events causing pain, and did not include treatment for sub-acute or chronic pain.

Assumption #2

• The proportion of patients allocated to OUD from the suzetrigine arm equaled 0%.

Assumption #3

• There is no further transition to OUD after three years.



Key Assumptions

Assumption #4

 Consistent with long-term evidence on OUD (sustained five-year abstinence), a proportion of patients transitioned to opioid abstinence without the chance of moving back to the OUD health state.

Assumption #5

 A weighted average of quality of life for OUD was estimated for those seeking OUD treatment and those not seeking OUD treatment.

Assumption #6

• Adverse effects from opioid use were modeled over a lifetime.



Key Model Inputs: Treatment-Related Efficacy

Characteristic	Value	Source
3-year Incidence of OUD in Suzetrigine Arm	0%	Assumption
3-year Incidence of OUD in HB/APAP Arm	0.43%	Schoenfeld et al., 2024
5-year Proportion of Patients Achieving Abstinence from OUD	0.052	Dowell et al., 2024, Zhu et al. 2018, Authors' calculation

HB/APAP: hydrocodone bitartrate/acetaminophen, OUD: Opioid Use Disorder



Key Model Inputs: Mortality

Characteristic	Value [95% Cl]	Source
All-Cause Mortality from Extramedical Opioid Use (Standardized Mortality Ratio)	5.02 [4.21, 5.98]	Larney et al. 2020
All-Cause Mortality Among those who are Abstinent versus those with Untreated OUD (Rate Ratio)	0.40 [0.34-0.46]	Santo et al. 2021
All-Cause Mortality	Varies by age and sex	US Life Tables

CI: confidence interval, OUD: Opioid Use Disorder, US: United States



Key Model Inputs: Treatment Costs

Costs	Value	Source
Suzetrigine, 7-day prescription (Journavx [®])	\$232.50	Vertex, Authors' calculation
HB/APAP, 7-day prescription (Multiple Brand Names)	\$10.64	US Redbook

HB/APAP: hydrocodone bitartrate/acetaminophen, US: United States



Key Model Inputs: Related and Unrelated Health Care Costs

Costs	Value [95% Cl]	Source
Annual Mean Excess Health Care Costs for People with OUD	\$17,370	Davenport et al., 2019
Annual Cost of MAT	\$7,676 [6,928-8,463]	Fairley et al., 2021; Authors' calculation
Future Unrelated Health Care Costs (Background Health Care Costs)	Varies by age and sex	Jiao et al., 2021

CI: confidence interval; MAT: medication-assisted therapy; OUD: Opioid Use Disorder



Key Model Inputs: Utilities

Health State	Value	Source
NPRS Levels	Varies	Dixon et al., 2011; Authors' calculation
OUD State (Disutility)	0.231	Wittenberg et al, 2016, Wu et al. 2016, Dowell et al. 2024, Authors' calculation
Abstinence (Disutility)	0.081	Wittenberg et al., 2016, Zhu et al., 2018, Authors' calculation

NPRS: Numeric Pain Rating Scale; OUD: Opioid Use Disorder



Results

Base-Case Results

Drug	Cost	QALYs	evLYs	Life Years	OUD Cases (per 100,000)
Suzetrigine	\$197,500	18.65	18.65	21.92	0
HB/APAP	\$197,900	18.61	18.61	21.89	429

evLYs: equal value of life years, HB/APAP: hydrocodone bitartrate/acetaminophen, OUD: Opioid Use Disorder, QALYs: quality-adjusted life years



Base-Case Incremental Results

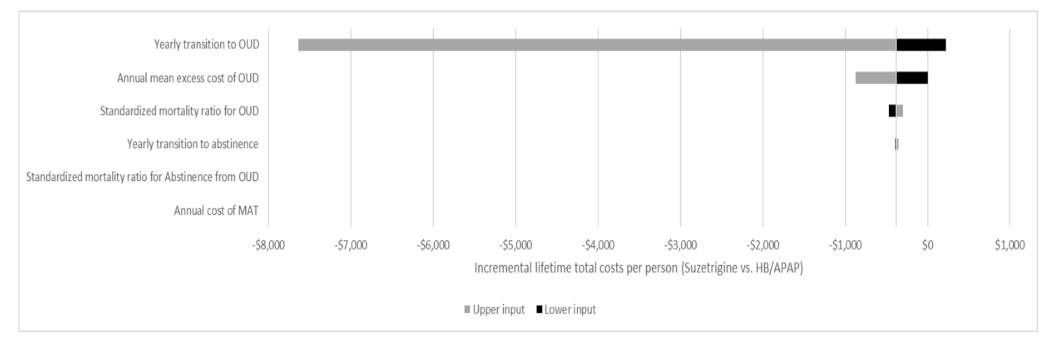
Drug	Comparator	Cost per QALY Gained	Cost per evLY Gained	Cost per Life Year Gained	Cost per OUD Case Averted
Suzetrigine	HB/APAP	Less costly, more effective	Less costly, more effective	Less costly, more effective	Less costly, more effective

evLYs: equal value of life years, HB/APAP: hydrocodone bitartrate/acetaminophen, OUD: Opioid Use Disorder, QALYs: quality-adjusted life years



One Way Sensitivity Analyses

Tornado Diagram for Incremental Lifetime Costs

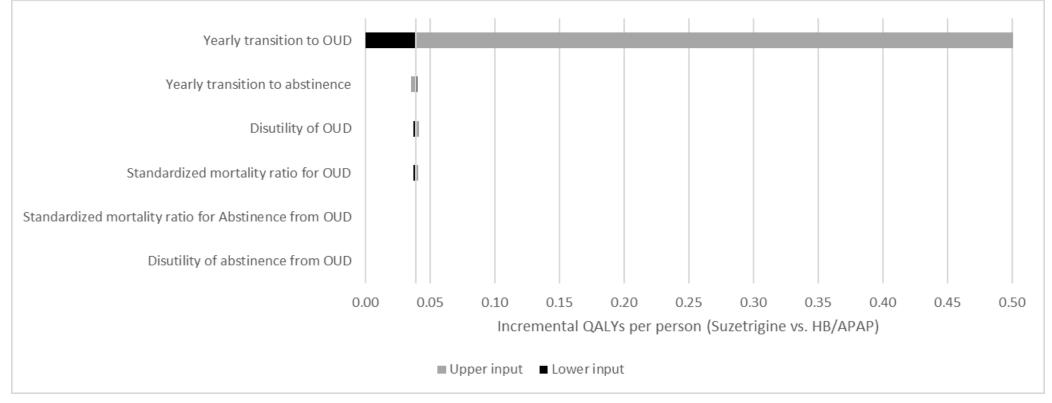


HB/APAP: hydrocodone bitartrate/acetaminophen, MAT: medication-assisted therapy, OUD: Opioid Use Disorder



One Way Sensitivity Analyses

Tornado Diagram for Incremental Quality-Adjusted Life Year Gains



HB/APAP: hydrocodone bitartrate/acetaminophen, OUD: Opioid Use Disorder, QALYs: quality-adjusted life years



Probabilistic Sensitivity Analysis

Drug	Cost-Effective at	Cost-Effective at	Cost-Effective at
	\$50,000 per	\$100,000 per	\$150,000 per
	QALY/evLY	QALY/evLY	QALY/evLY
Suzetrigine	92%	94%	96%

evLY: equal value of life years, QALY: quality-adjusted life years



Scenario Analyses

Analysis 1: Modified societal perspective with components such as productivity losses, criminal justice and incarceration, and caregiver disutilities applied to the OUD health state.

Analysis 2: The proportion of patients with OUD in the opioid comparator arm that result in scenarios for suzetrigine that meet commonly cited cost-effectiveness thresholds.

Analysis 3: Exclusion of unrelated health care and death costs.

Drug	Analysis 1	Analysis 2	Analysis 3
Suzetrigine	Less costly, more effective	0.02% with OUD by three years in the opioid arm to meet \$100,000 per QALY and evLY thresholds	Less costly, more effective

evLY: equal value of life years, OUD: Opioid Use Disorder, QALY: quality-adjusted life years



Health Benefit Price Benchmark (HBPB)

Intervention	Weekly Price (WAC)	Weekly Price at \$100,000/QALY Threshold	Weekly Price at \$150,000/evLY Threshold	Discount from WAC to Reach Threshold Prices
Suzetrigine	\$232.50	\$4,500	\$6,500	N/A

evLY: equal value of life years, N/A: not applicable, QALY: quality-adjusted life years, WAC: wholesale acquisition cost



Limitations

Top Limitations

- The cost-effectiveness of suzetrigine for acute pain compared to HB/APAP depends greatly on the incidence of OUD from a short course of HB/APAP.
- The societal perspective estimates may underestimate the economic burden of OUD.



Comments Received

- Updated wholesale acquisition cost for suzetrigine following FDA approval.
- Additional text added to the Uncertainty and Controversies section of the Evidence Report indicating potential for underestimate of societal costs from avoiding OUD.



Conclusions

 Suzetrigine for treating moderate-to-severe acute pain is slightly cost-saving due to averting OUD compared to HB/APAP, using the wholesale acquisition cost of \$232.50 for a one-week prescription to treat acute pain.



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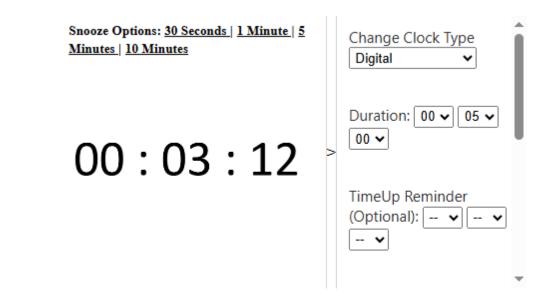


Manufacturer Public Comment and Discussion

Paul Negulescu, Ph.D Senior Vice President and Disease Area Executive, Vertex Pharmaceuticals

Conflicts of Interest:

• Dr. Negulescu is a full-time employee at Vertex Pharmaceuticals.



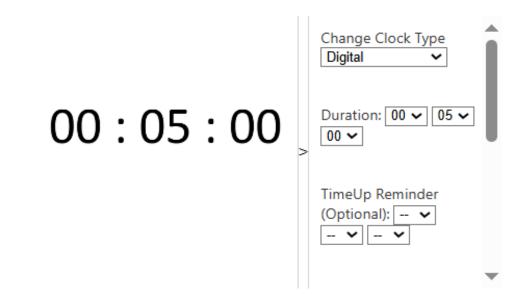


Public Comment and Discussion

Kathy Sapp CEO, American Chronic Pain Association

Conflicts of Interest:

• No conflicts to disclose.

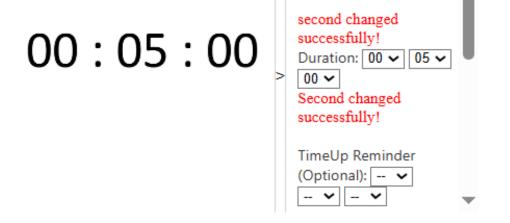


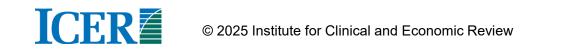


Marvell Adams CEO, Caregiver Action Network

Conflicts of Interest:

- No conflicts to disclose. ٠
- Marvell Adams has collaborated with Forbes Tate Partners to compose this ٠ Change Clock Type public comment. Digital



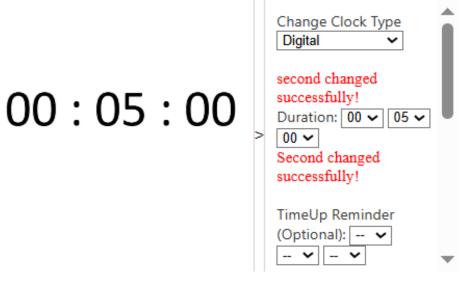


~

Cindy Steinberg, BA, BS Director of Policy and Advocacy, US Pain Foundation

Conflicts of Interest:

• US Pain Foundation receives funding from pharmaceutical companies including Vertex Pharmaceuticals, as well as grants from foundations and individuals.





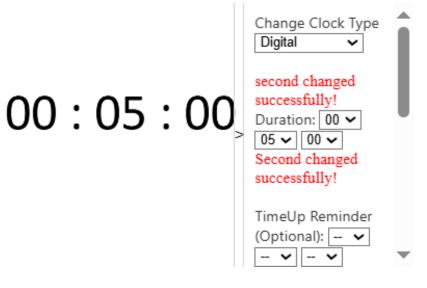
Margaret French Senior Director, Federal Affairs, Voices for Non-Opioid Choices

Conflicts of Interest:

- Margaret is a former employee of America's Essential Hospitals.
- Voices for Non-Opioid Choices has received funding

from Vertex Pharmaceuticals and Pacira Biosciences

that is equal to or greater than 25% of their overall budget. ${f U}$





Lunch

Meeting will resume at 12:50PM CT



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Voting Questions

Patient Population for all questions: Adult patients with acute pain that is not adequately controlled with non-systemic therapies (e.g. heat therapy, local anesthetic).

Clinical Evidence



1. For patients with acute pain, is the current evidence adequate to demonstrate that the net health benefit of suzetrigine in addition to nonsystemic therapies (e.g. heat therapy, local anesthetic) is greater than that of non-systemic therapies alone?



2. For patients with acute pain, is the current evidence adequate to distinguish the net health benefit of suzetrigine from that of oral opioid analgesics (with or without acetaminophen), each in addition to non-systemic therapies?



2a. If "Yes", which has a greater net health benefit?



3. For patients with acute pain, is the current evidence adequate to distinguish the net health benefit of suzetrigine from that of oral NSAIDs, each in addition to non-systemic therapies?



3a. If "Yes", which has a greater net health benefit?

Benefits Beyond Health and Special Ethical Priorities

To help inform judgments of overall long-term value for money, please indicate your level of agreement with the following statements:



4. There is substantial unmet need despite currently available treatments.



5. This condition is of substantial relevance for people from a racial/ethnic group that have not been equitably served by the healthcare system.

To help inform judgments of overall long-term value for money, please indicate your level of agreement with the following statements based on the relative effects of suzetrigine versus non-systemic therapies (e.g. heat therapy, local anesthetic), non-opioid analgesics including NSAIDs, acetaminophen, opioid analgesics, and combination with acetaminophen:



6. The treatment is likely to produce substantial improvement in caregivers' quality of life and/or ability to pursue their own education, work, and family life.



7. The treatment offers a substantial opportunity to improve access to effective treatment by means of its mechanism of action or method of delivery.

Long-Term Value for Money

8. Given the available evidence on comparative clinical effectiveness and incremental cost effectiveness, and considering benefits beyond health and special ethical priorities, what is the long-term value for money of suzetrigine compared to oral opioid analgesics (with or without acetaminophen) at current pricing?



8. What is the long-term value for money of suzetrigine compared to oral opioid analgesics (with or without acetaminophen) at current pricing?

Break

Meeting will resume at 2:00PM CT



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Policy Roundtable

Policy Roundtable

Participant	Conflict of Interest
Vicky Brown, PharmD, BCOP, Associate Vice President Clinical Drug Strategy, Humana	Dr. Brown is a full-time employee at Humana.
Jaime Rubin Cahill, MA, MPH, Vice President, Health Economics and Outcomes Research, Vertex Pharmaceuticals	Jaime Rubin Cahill is a full-time employee at Vertex Pharmaceuticals.
David Dohan, MD, MHCM , Medical Director of Pharmacy, Point32 Health	Dr. Dohan is a full-time employee at Point32 Health.
Benjamin Friedman, MD, MS, Professor of Emergency Medicine, Montefiore Einstein	No conflicts to disclose.
Nicole Hemmenway , Chief Executive Officer, US Pain Foundation	Nicole Hemmenway is a full-time employee of the US Pain Foundation. The US Pain Foundation receives greater than 25% of funding from health care companies, industry groups, family foundations, and individual donors.
Andrew Kolodny, MD , Medical Director, Opioid Policy Research, Heller School for Social Policy and Management, Brandeis University	Dr. Kolodny has served as an expert witness in litigation involving the opioid industry.
Gabriel Smith, Patient, Arlington Chronic Pain Support Group	No conflicts to disclose.



Midwest CEPAC Reflections

Next Steps

- Meeting recording posted to ICER website next week
- Final Report published on or around March 27th, 2025
 - Includes description of Midwest CEPAC votes, deliberation, policy roundtable discussion
- Materials available at: <u>https://icer.org/assessment/acute-pain-</u>
 <u>2025/</u>







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