

ceribell®

Corporate Presentation

August 2026

Disclaimers

This presentation contains forward-looking statements, within the meaning of the Private Securities Litigation Reform Act of 1995, about the Company and its industry. These statements and the outcomes of the events they describe involve substantial risks, uncertainties and potentially inaccurate assumptions, some of which cannot be predicted or quantified. All statements other than statements of historical facts contained in this presentation, including statements regarding the Company's strategy, financial guidance and projections, future results of operations and financial performance, future operations, projected costs, prospects, plans, objectives, expected products, market size, market expansion, growth opportunities and competitive position, as well as assumptions relating to the foregoing, are forward-looking statements. The Company cautions you that the foregoing list does not contain all of the forward-looking statements made in this presentation. In some cases, you can identify forward-looking statements by words such as "may," "will," "shall," "should," "expects," "plans," "anticipates," "could," "intends," "target," "projects," "contemplates," "believes," "estimates," "predicts," "potential," "goal," "objective," "seeks," "aims," "forecasts," "guidance" or the negative of these words or other similar terms or expressions that concern the Company's expectations, strategy, plans, or intentions.

You should not rely upon forward-looking statements as predictions of future events, future financial performance or future trends. The Company has based the forward-looking statements contained in this presentation primarily on its current expectations, estimates, forecasts, and projections about future events and trends that it believes may affect its business, financial condition, results of operations, and prospects. The Company cannot guarantee or ensure that the future results, performance, events, circumstances or any other outcome expressed, anticipated or reflected in the forward-looking statements will be achieved or occur in whole or in part. Actual results, events, or circumstances could differ materially from those described or assumed in the forward-looking statements. Moreover, the Company operates in a very competitive and rapidly changing environment. New risks and uncertainties emerge from time to time, and it is not possible for the Company to predict all risks and uncertainties that could have an impact on the forward-looking statements in this presentation. This presentation contains estimates, projections, and other information concerning the Company's industry and its business, as well as data regarding market research, estimates, and forecasts prepared by its management or third parties. Information that is based on estimates, forecasts, projections, market research, or similar methodologies is inherently subject to uncertainties. These data and the industry in which the Company operates are subject to a high degree of uncertainty and risk due to a variety of factors which could cause results to differ materially from those expressed in these estimates, publications, and reports made by third parties or the Company.

Among the factors that could cause actual results to differ materially from past results and future plans and projected future results are the following: risks related to our limited operating history and history of net losses; our ability to successfully achieve substantial market acceptance and adoption of our products; competitive pressures; our ability to adapt our manufacturing and production capacities to evolving patterns of demand and customer trends; the manufacturing of a substantial number of our product components and their assembly in China and Vietnam; product defects or complaints and related liability; the complexity, timing, expense, and outcomes of clinical studies; the timing and success of our product development, regulatory clearances and commercial launches; our ability to obtain and maintain adequate coverage and reimbursement levels for our products, including the temporary and conditional nature of new technology add-on payments and our ability to obtain renewal of any such designation; developments in rulemakings, including proposed and final rules affecting payment rates, coverage policies and new technology add-on payment eligibility; the performance of our artificial intelligence-powered algorithms and evolving regulatory treatment of such technologies; cybersecurity incidents and data privacy obligations; our ability to comply with changing laws and regulatory requirements and resulting costs; our dependence on a limited number of suppliers; tariffs, trade policy developments, geopolitical conflicts and related supply chain disruptions; the outcome, timing and costs of intellectual property enforcement and infringement proceedings, including challenges to the validity, enforceability or scope of our patents in litigation or in proceedings before the U.S. Patent and Trademark Office; our ability to comply with the terms of our indebtedness; delays in regulatory, litigation or other matters affecting our business, and other risks and uncertainties, including those described under the heading "Risk Factors" in our Annual Report on Form 10-K, our subsequent Quarterly Reports on Form 10-Q and other reports filed with the U.S. Securities and Exchange Commission ("SEC"). These filings are available on the Investor Relations section of our website at <https://investors.ceribell.com/> and on the SEC's website at <https://sec.gov/>.

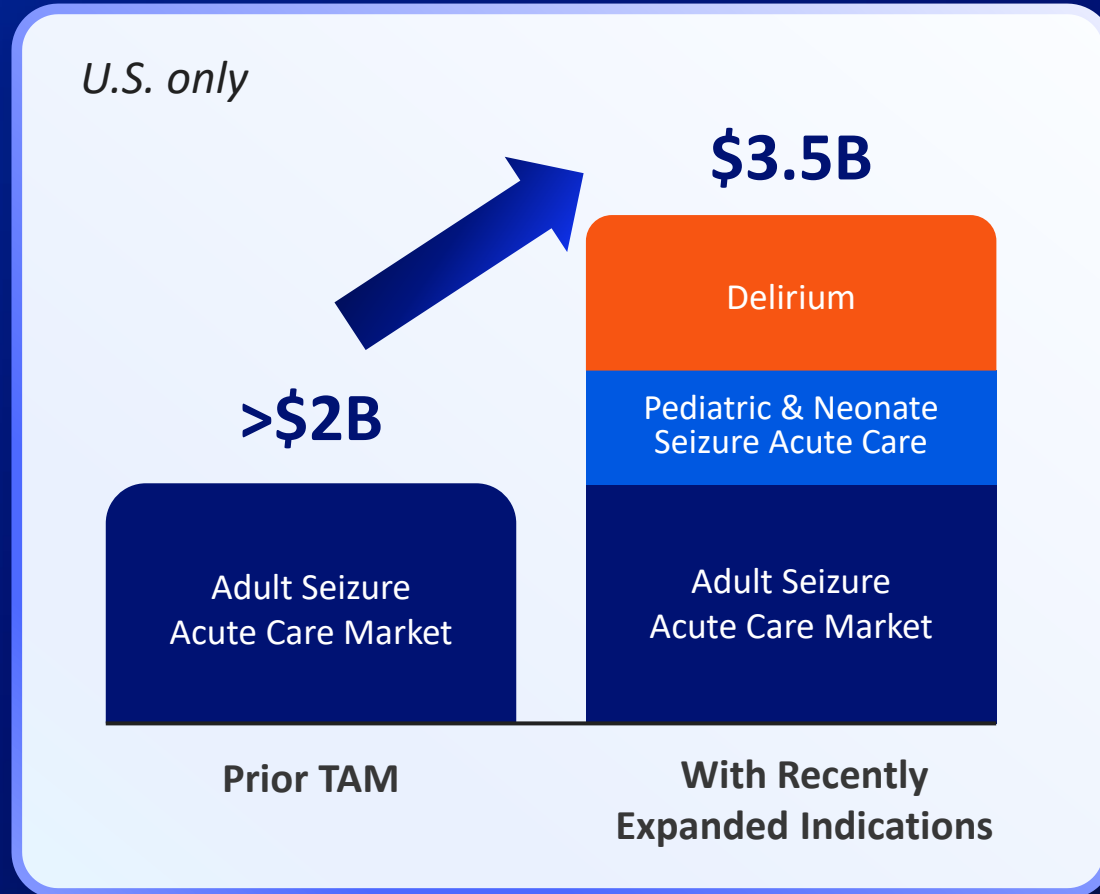
The forward-looking statements made in this presentation relate only to events as of the date on which the statements are made. The Company undertakes no obligation to update any forward-looking statements made in this presentation to reflect events or circumstances after the date of this presentation or to reflect new information or the occurrence of unanticipated events, except as required by law. The Company's forward-looking statements do not reflect the potential impact of any future acquisitions, mergers, dispositions, joint ventures, or investments it may make. In addition, the results of studies referenced in this presentation concerning the Clarity algorithm apply only to the algorithm version that was in use at the time of the analysis and do not reflect subsequent algorithm updates. Most studies referenced in this presentation were conducted with small sample sizes and were not powered for statistical significance, did not control for other clinical variables, or have other design limitations (e.g., the studies may be retrospective and are not randomized controlled trials). In addition, some of the studies were sponsored, funded or supported by the Company or involved employees or consultants of the Company.

Trade names, trademarks and service marks of other companies appearing in this presentation are the property of their respective owners. Solely for convenience, the trademarks and trade names referred to in this presentation appear without the ® and ™ symbols. This does not indicate, in any way, that the Company will not assert its rights, and the rights of any applicable licensor, to these marks and tradenames, to the fullest extent under applicable law. This presentation is not a substitute for the Company's filings with the SEC, and recipients should conduct their own analysis of the market, the Company's market position and the potential future performance of its business.

AI-Powered Point-of-Care EEG Platform Targeting Serious Neurological Conditions in the Acute Care Setting



Large & Expanding Potential Market Opportunity¹



Financial Highlights

\$114M – \$117M

FY 2026 Guidance²

33%

Q2 2026 YoY Revenue Growth³

90%

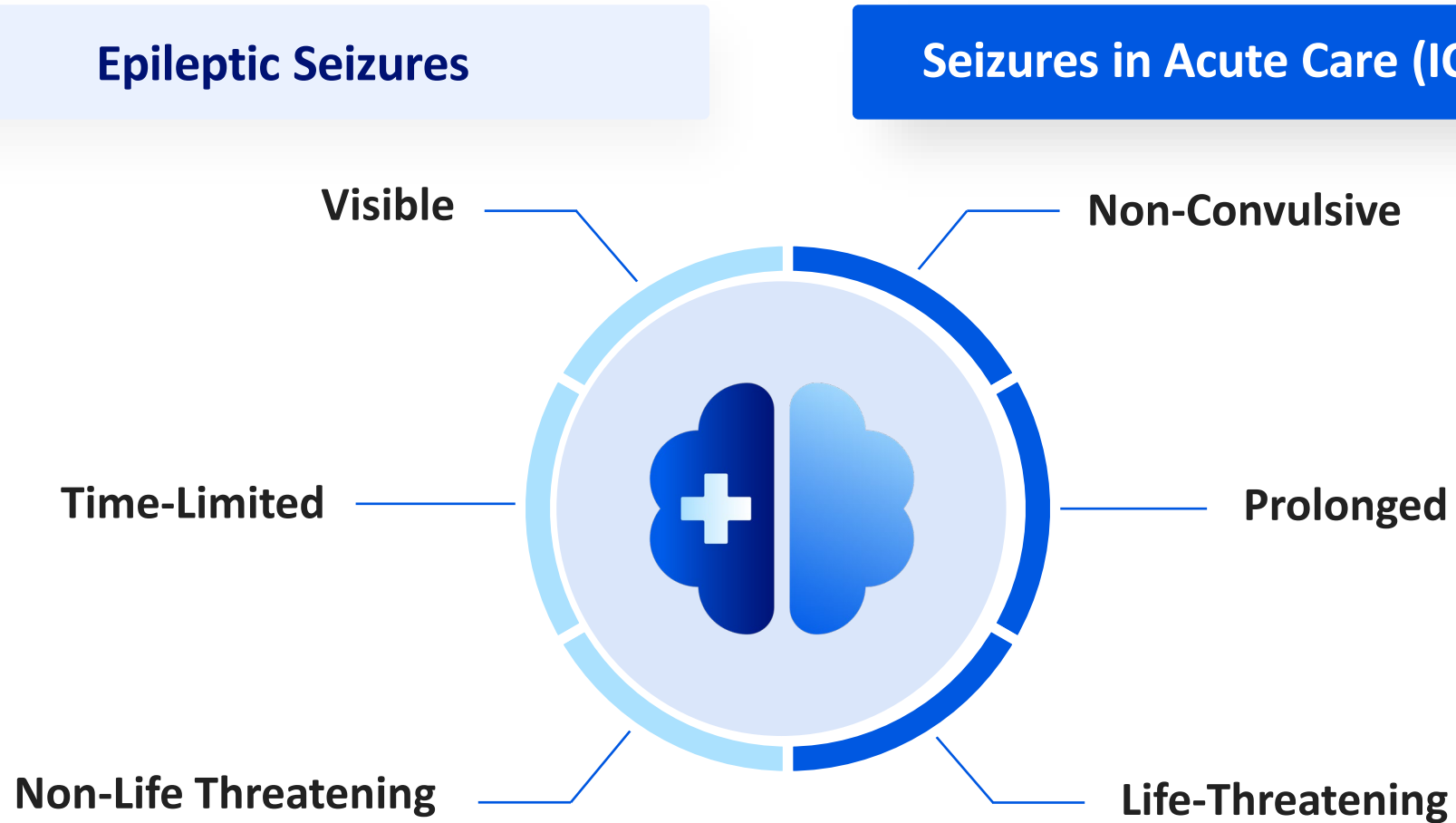
YTD 2026 Gross Margin^{3,4}

1. TAM based on management estimates. Actual results may differ materially.
2. Revenue guidance issued August 10, 2026.
3. As of June 30, 2026.
4. 2Q 2026 gross margin includes impact of \$1.0 million in tariff refunded in COGS.

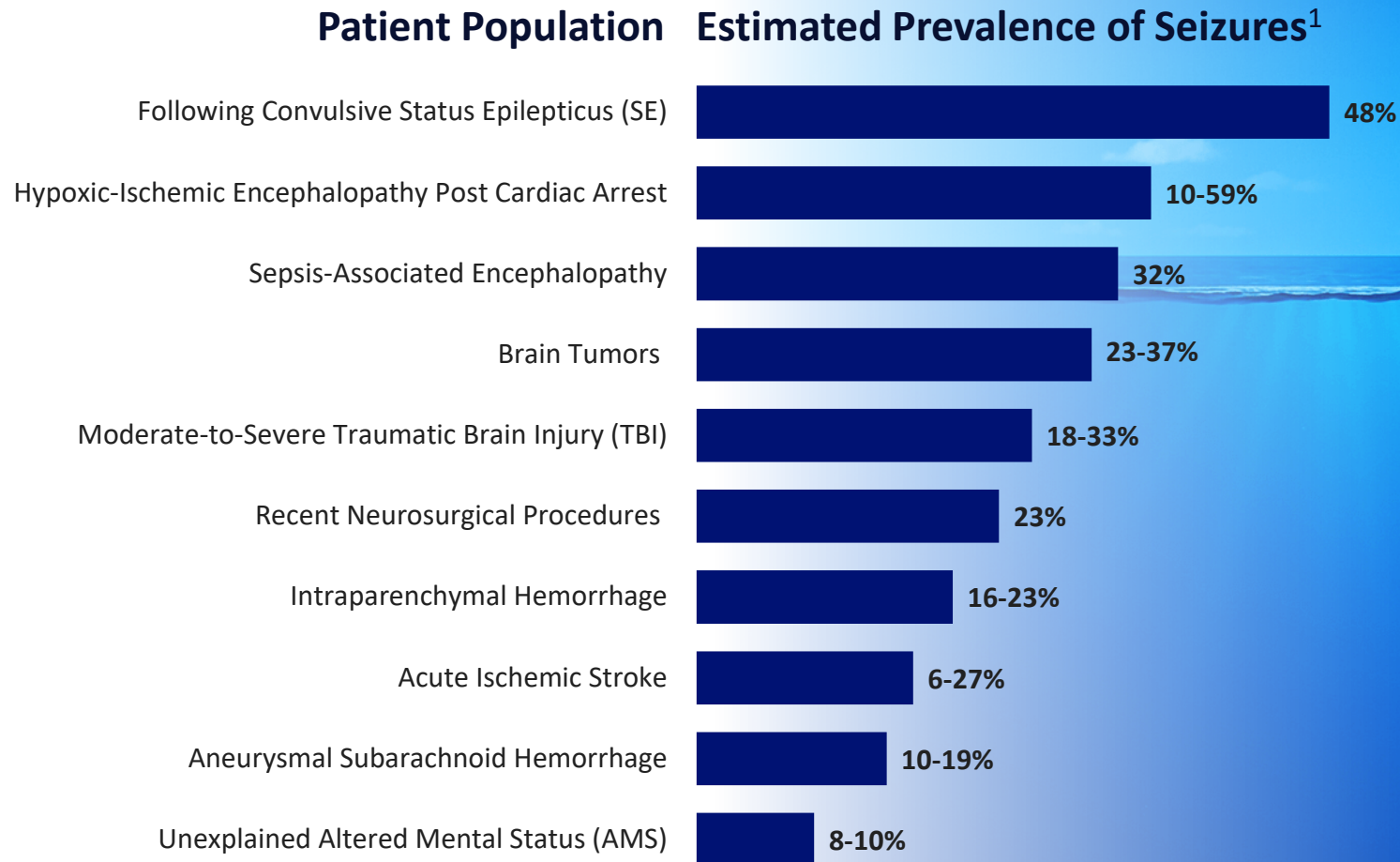
Ceribell's Three Growth Horizons



Targeting Seizures in the Acute Care Setting



Seizures Are Highly Prevalent in Critically-Ill Patients and Often Go Undiagnosed



up to **92%**
of seizures in the ICU
are non-convulsive^{2,3}

EEG required for diagnosis

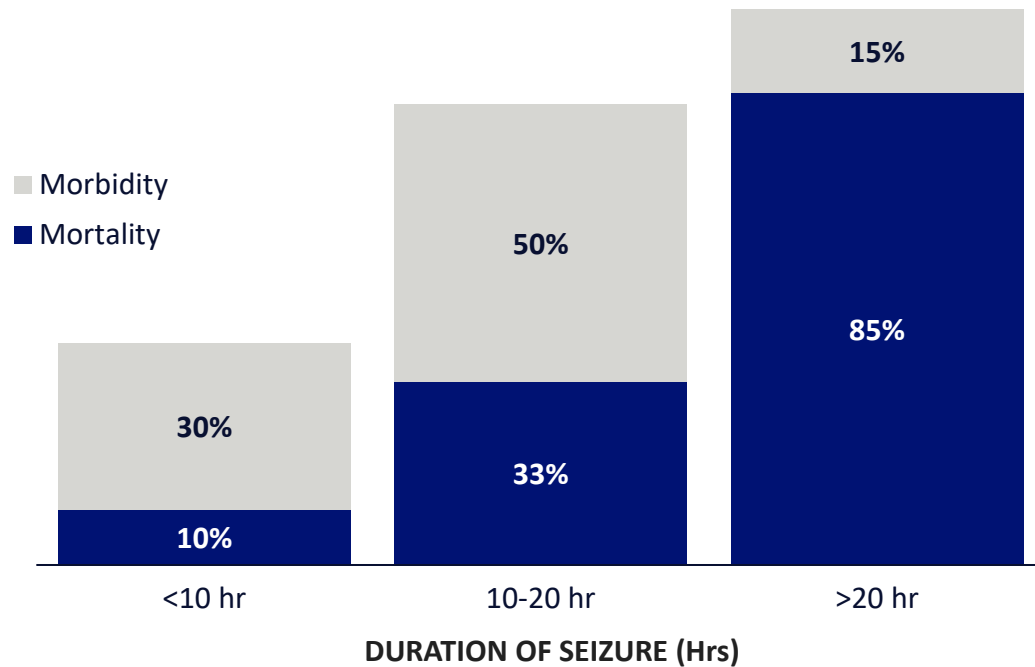
1. Herman, S.T., et al. (2015) *J Clin Neurophysiol.* 32(2):87-95

2. Claassen, J., et al. (2004). *Neurology.* 62(10):1743-1748

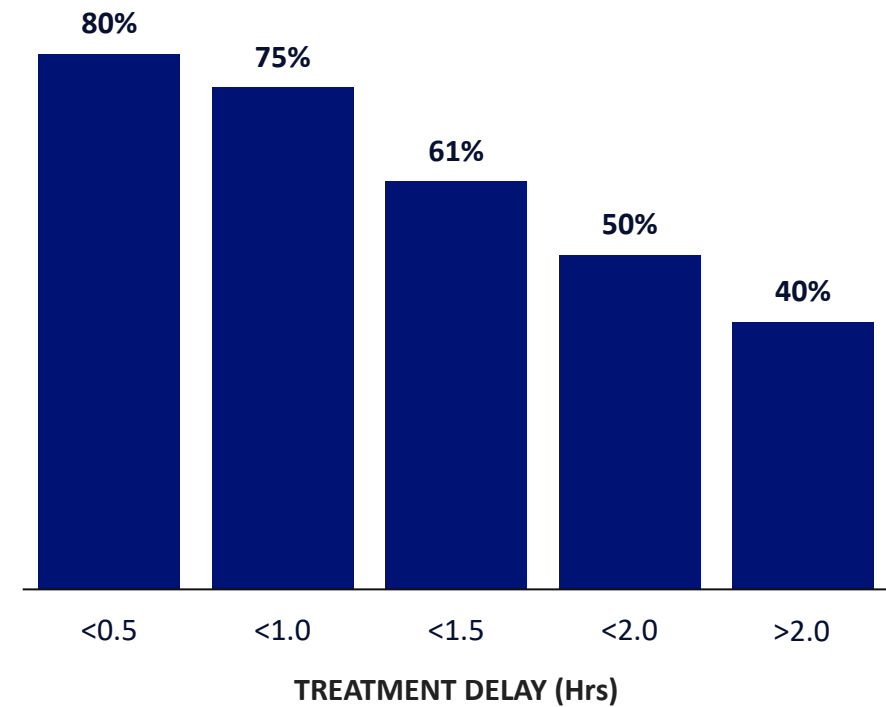
3. Rudin, D., et al. (2011) *Epilepsy Res.* 96(1-2):140-50

"Time is Brain"

STATUS EPILEPTICUS ALL-CAUSE MORBIDITY & MORTALITY RATE¹



PATIENT RESPONSE RATE TO FIRST-LINE TREATMENT²



1. Young, G.B., et al. (1996). *Neurology*, 47(1):83-89
2. Lowenstein, D.H., et al. (1993) *Neurology*, 43(3 Pt 1):483-488

Guidelines and Recommendations Include Timely EEG to Detect and Manage Seizures Across Different Disease States

2012

NEUROCRITICAL
CARE SOCIETY

EEG should be initiated **within 15-60 minutes** to “evaluate for NCSE if [patient is] not waking up after clinically obvious seizures cease.”¹

2020

American **Heart** Association

“Recommend **promptly performing and interpreting EEG** for the diagnosis of seizures in **all comatose patients** after the return of spontaneous circulation (ROSC)” from **cardiac arrests**.²

2021

American **Stroke** Association

“**EEG** [is recommended] for a *change in mental status* or depressed mental status out of proportion to the **[ischemic] stroke**.”³

2023

American **Stroke** Association

“Monitoring with continuous EEG can detect nonconvulsive seizures, especially in **[aneurysmal subarachnoid hemorrhage] patients** with depressed consciousness or fluctuating neurological examination.”⁴

1. Brophy, G., et al. (2012) Neurocrit Care. 17(1):3-23

2. Panchal, A.R., et al. (2020) Circulation. 142(suppl 2):S366-S468

3. Perman S.M., et al. (2024) Circulation 149(5):e254-e273

4. Green, T.L., et al. (2021) Stroke 52(5):e179-e197

Conventional EEG Has Significant Limitations in the Acute Care Setting

Overview of EEG

An EEG is a non-invasive tool used to **measure** and **display** electrical activity in the brain



*Designed for use in the **outpatient setting**, primarily for managing epilepsy patients*

Conventional EEG systems were not designed for the acute care setting

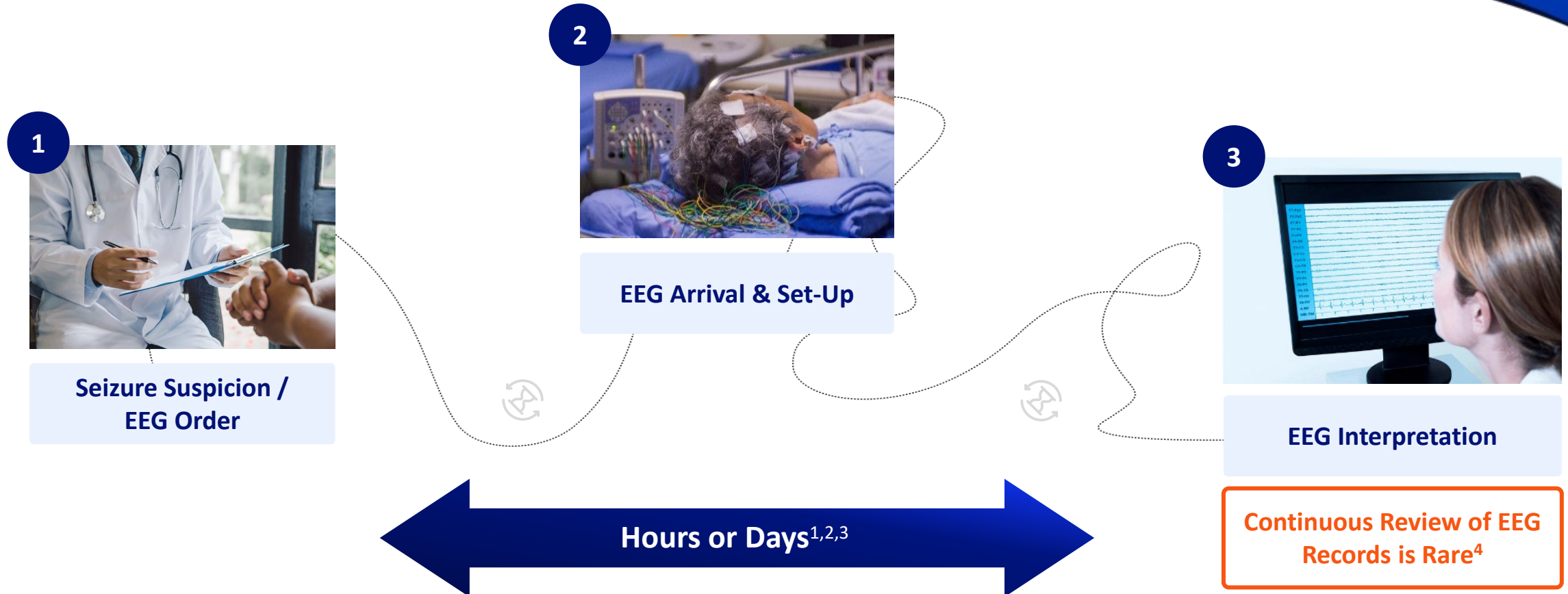
Hardware & Access Challenges

- ❌ Requires EEG Technician (Typically 9-5 Monday – Friday)
- ❌ Long Set-Up Process

Interpretation Challenges

- ❌ Requires Interpretation by a Specially-Trained Neurologist
- ❌ Continuous Monitoring Rarely Performed in Practice

Clinical Reality: Conventional EEG is Not Suited for the Acute Care Setting and Leads to Long Delays

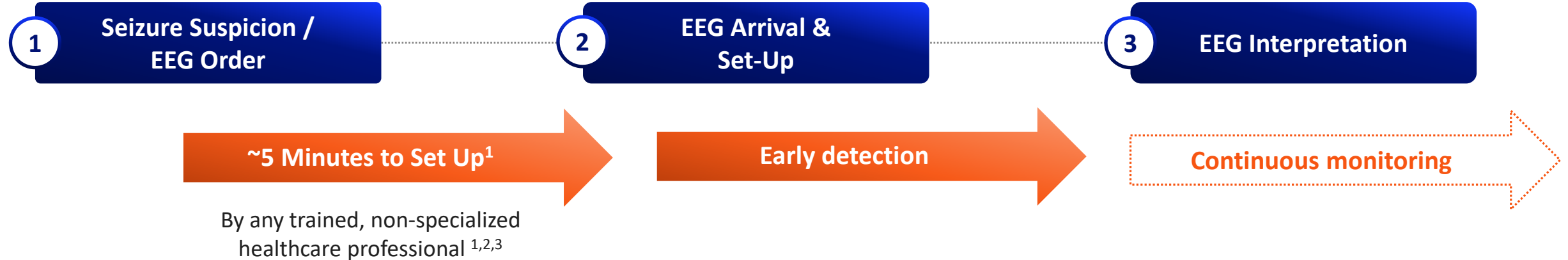


1. Gururangan, K., et al. (2016) *Clinical Neurophysiology*. 127(10):3335-3340. Maximum time from EEG order to arrival and set-up
2. Vespa, P., et al. (2020) *Crit Care Med*. 48(9):1249-1257. Median time from EEG order to arrival and set-up

3. Quigg, M. et al. (2001) *J Clin Neurophysiol*. 18(2):162-165. Range of time from request to interpretation
4. Gavvala, J., et al. (2014) *Epilepsia*. 55(11):1864-1871

Ceribell EEG System: Suspicion to Diagnosis in Minutes, Enabling Earlier & More Accurate Treatment

The **ceribell**[®] System

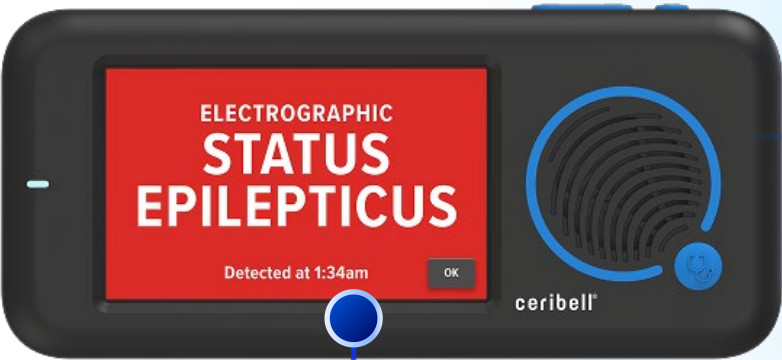


1. Yazbeck et al. (2019) *Journal of Neuroscience Nursing*
2. Hobbs et al. (2018) *Neurocritical Care*
3. Eberhard et al. (2023) *Clinical Nursing Focus*

The
ceribell[®] System

Combining highly portable, simple-to-use and rapidly deployable hardware with AI-powered algorithms

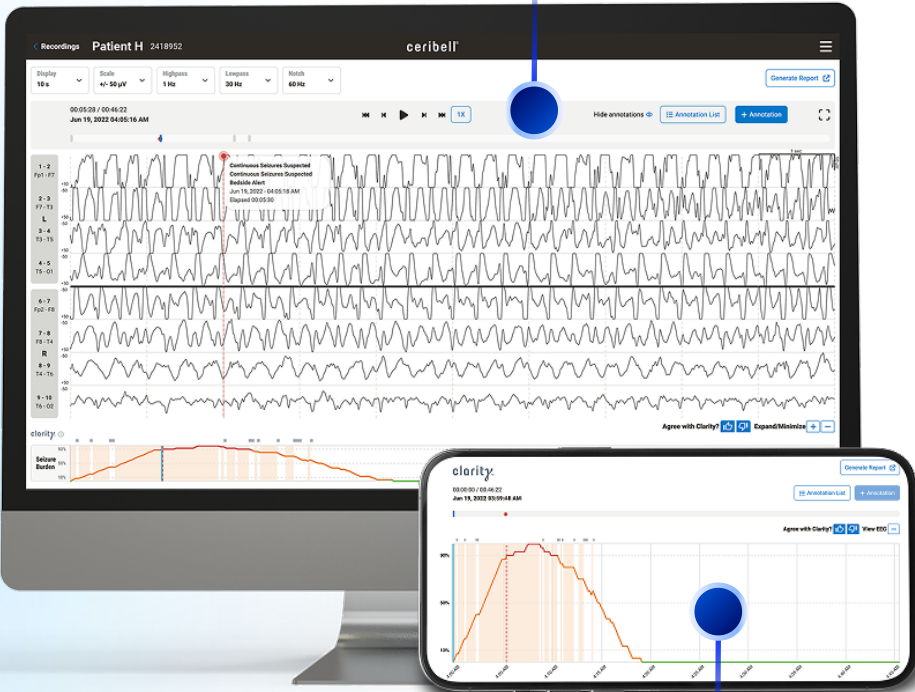
Ceribell EEG Headband



Ceribell EEG Recorder



Ceribell EEG Portal



AI-powered Seizure Detection Algorithm

clarity[®]

Clarity: Our Proprietary AI-Powered Seizure Detection Algorithm

clarity®



✓ Provides seizure burden to facilitate EEG reading for neurologists



✓ Bedside Alert

✓ Real-time feedback on response to medication

Ceribell Supports Precise Patient Care for Status Epilepticus: Expediting Diagnosis and Continuously Monitoring



CAUTION: Device does not substitute for EEG review by a qualified clinician. Before use, review the manual for indications, contraindications, warnings, precautions, potential adverse events and Instructions for Use. Sale requires the order of a physician.

Validated Cost Savings

51 Peer-Reviewed Publications & 107 Abstracts*

Reduced
Over-administration of
Anti-Seizure Medication (ASM)¹



40%

changed diagnostic
suspicion and **20%** changed
treatment decisions⁵

43%

of patients with reduced ASM⁷ +
51% potential reduction in
intubation and parenteral ASM³

53%

changed clinical management
and expedited disposition for
21% of patients⁶

Reduced
Length of Stay (LOS)
in the ICU and Hospital



4.1 days

ICU LOS
reduction⁷

Trend of
3 days

hospital LOS
reduction²

Potential
0.4 days

ICU LOS reduction³

1.2 days

hospital LOS reduction³

Reduced
Patient Transfers⁴



94%

of transfers that would
have been made avoided⁸

100%

of non-seizure
patients retained⁹

* As of June 30, 2026

- 1. Wright, N., et al. (2021) EMJ. 38(12):923-926
- 2. Eberhard, E., et al. (2023) J Neurosci Nurs. 55(5):157-163
- 3. Ney, J.P., et al. (2021) J Med Econ. 24(1):318-327

- 4. Madill et al. (2022) Epileptic Disord 24 (3): 507-516
- 5. Vespa, P., et al. (2020) Crit Care Med. 48(9):1249-1257
- 6. Wright, N., et al. (2021). EMJ. 38(12):923-926

- 7. Desai et al. (2024). Critical Care Medicine. 52(1):p S268, 589
- 8. Madill, E.S., et al. (2022) Epileptic Disorders. 24(3):507-516
- 9. Ward, J., et al. (2023) Front. Digit. Health. 5(1)

Evidence-Based Clinical & Economic Benefits



Study Findings

Outcome	convEEG (N = 62)	Ceribell (N = 62) ²	Δ Delta	P- Value
Median door-to-EEG time (hours)	25.3	5.9	19.4 hours faster door-to-EEG time	p < 0.0001
Median ICU LOS	8.0 Days	3.9 days	4.1 days shorter ICU LOS	P = 0.003
mRS ³ greater than or equal to 4 at discharge	76%	58%	18% better clinical outcomes ⁴	p = 0.047

SAFER Study Overview

The Seizure Assessment and Forecasting with Efficient Rapid-EEG (SAFER-EEG) study is a multisite retrospective study of adult patients who received EEG during hospital stay. Most centers had 24/7 conventional EEG with technician onsite or on-call.

Study Sites :

- Yale University
- Mass General
- University of New Mexico

1. Desai, M., et al. (2024) Neurocrit Care.

2. The cohorts were matched 1:1 with propensity scores to have equivalent age, admission scores, diagnosis group and seizure suspicion.

3. The Modified Rankin Scale (mRS) is a 6-point disability scale with possible scores ranging from 0 to 5. 0 is healthy and 5 is severe disability. A separate category of 6 is usually added for patients who expire.

4. Using mRS greater than or equal to 4 at discharge as an indicator of functional disability. Results with Ceribell vs. conventional EEG.

New for 2026: Pediatric & Neonate Indication Expansion & Commercial Launch



- **Received FDA clearances for Clarity algorithms for age 1+ and neonate**, including pre-term in 2025

- Unlocked **incremental \$400M market opportunity**¹, including ~280 children's hospitals targets



- **Initiated commercial launch** in Q1 2026

1. TAM based on management estimate . Actual results may differ materially

Seizures in Critically Ill Neonates are Common, Requiring EEG for Accurate Diagnosis

Diagnostic challenges can lead to under- and over-treatment



up to
90% of neonatal seizures are non-convulsive^{1,2}

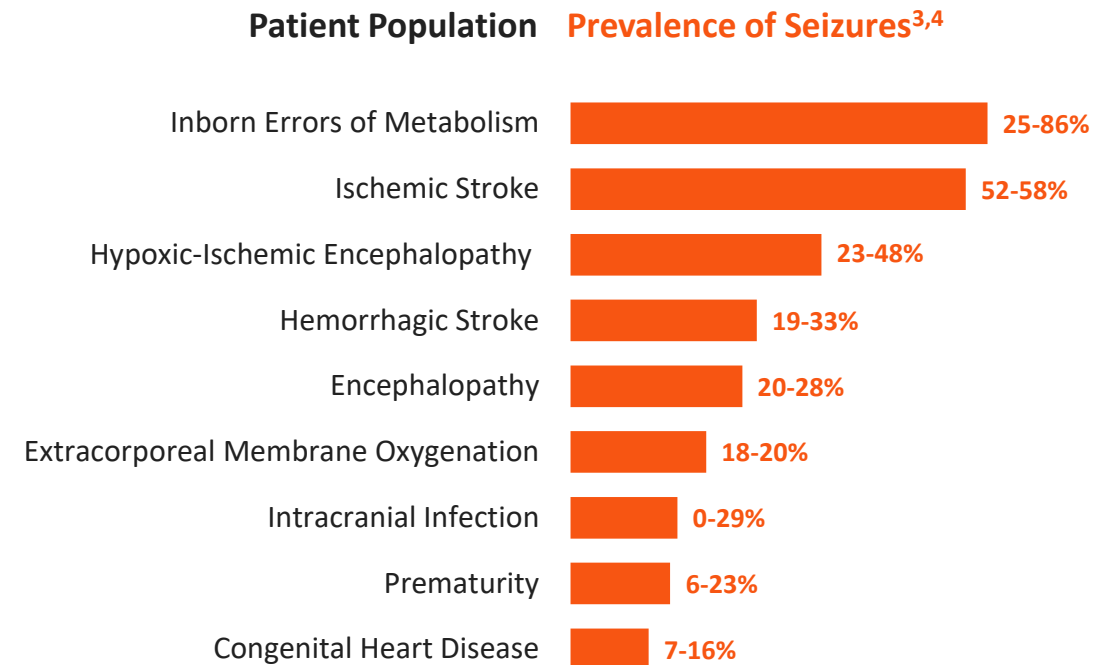


up to
73% of clinically observed “seizures” are not seizures³

“We suggest cEEG use to monitor neonates at risk for seizure **in the absence of clinically evident seizures.**”

— American clinical Neurophysiology Society

Seizures are the most common neurological emergency in newborns



1. Murray, D.M, et al. (2008). Arch Dis Child Fetal Neonatal Ed. 93:F187-F191

2. Massey, S, et al. (2018). Seminars in Fetal & Neonatal Medicine. 23(2018):168-174

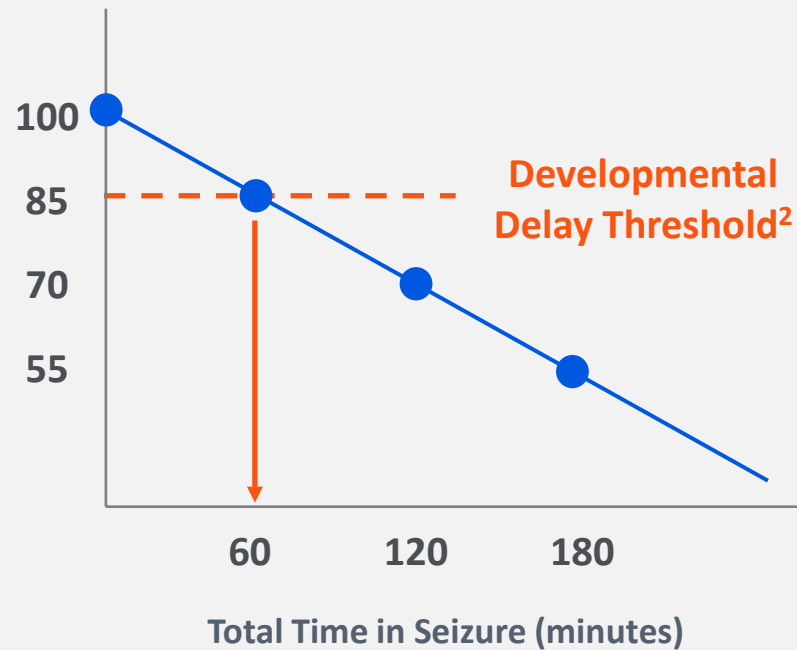
3. Sheth, R., et al. (1999). J Perinatol. 19(1):40-3

4. Yan, K., et al. (2023). Jama Network Open. 6(7):e2326301

Neonatal Seizures Require Urgent Diagnosis and Management

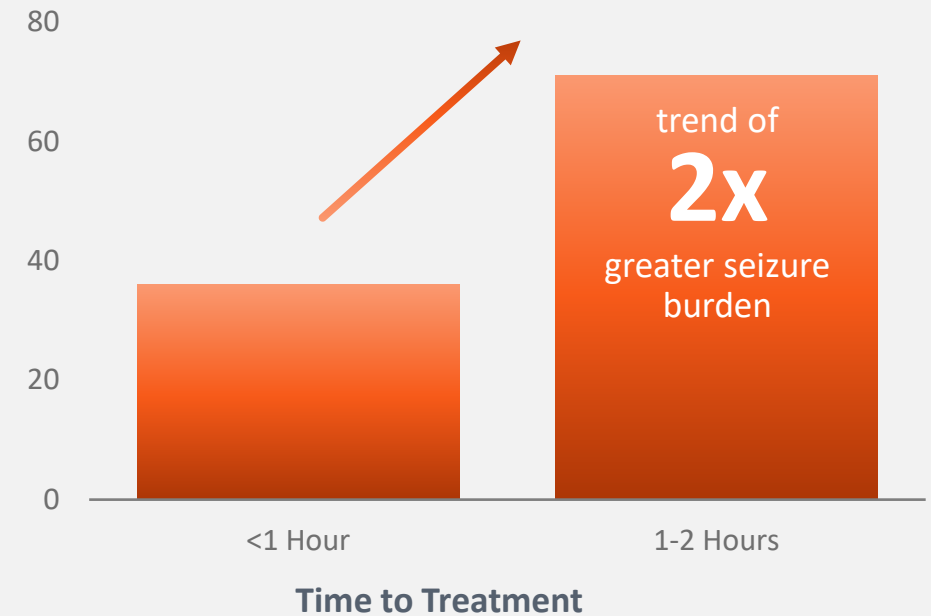
1 hour in seizure is associated with **language impairment and cognitive deterioration**¹

Predicted Language Score at 18 Months (Bayley-III)



Just **1 hour** delay to treat seizure may lead to **higher seizure burden**³

Median Seizure Burden



1. Alharbi, H.M., et al. (2022). *Neurology*. 100:e1976-e1984

2. Johnson, S, et al. (2014). *Pediatr Res* 75, 670-674

3. Pavel, A.M., et al (2021) *J Pediatr*. 243:61-68.e2

Business Model: Two Sources of Recurring Revenue

~25% Subscription

(SaaS + loaned capital)

clarity®

+



+



AI Algorithm
SaaS

Recorder
Capital

Portal
SaaS



~75% Product

(single-patient disposable)



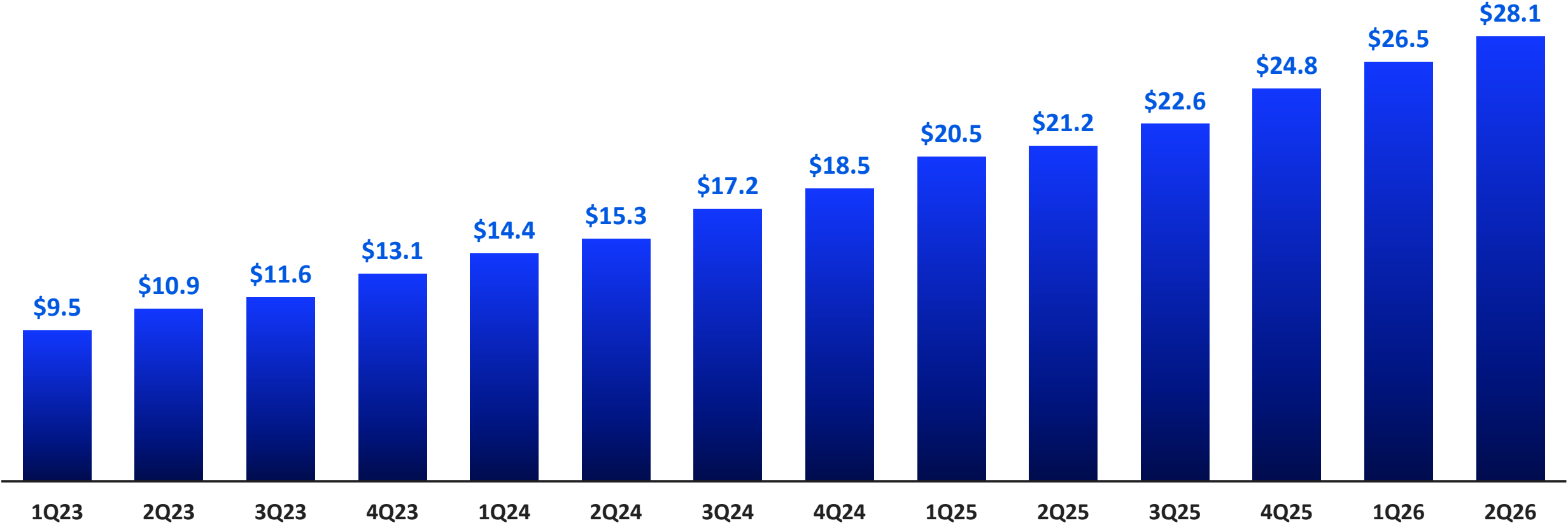
Single-Use Headband
Disposable



Rapid Revenue Growth & Projectable Business Model

Quarterly Revenue

(\$ in million)



Significant Opportunity For Continued Growth Within Core Seizure Market

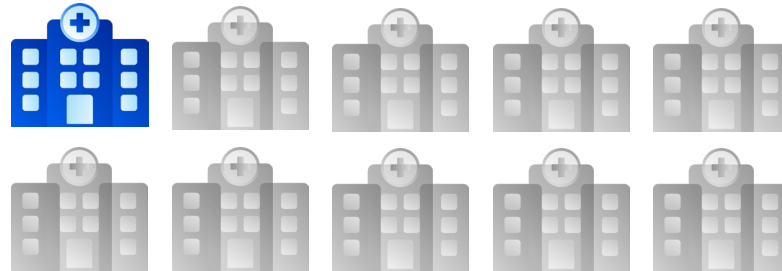
<4%

of addressable
Market¹



12%

hospital penetration

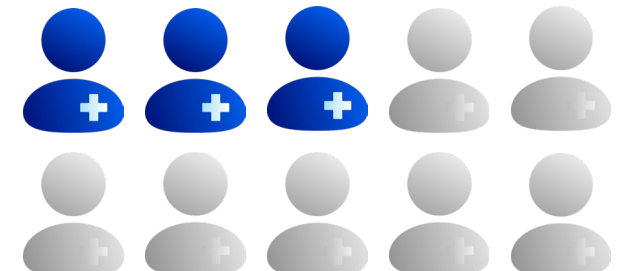


Ceribell is currently active in only
712 out of roughly 6,000 hospitals
providing acute care services

X

30%

within existing accounts



Ceribell's top customers use
approximately **3x the number of
devices** as average customers,
when controlling for hospital size

2026 Commercial Growth Drivers in Core Seizure Market



Account Acquisition

- Execute on proven account acquisition strategy
- Drive productivity of recently expanded commercial infrastructure
- Expand in VA hospitals
- Build out health system infrastructure and playbook
- Expand to children's hospitals



Drive Utilization in Existing Accounts

- Expand to new departments, EDs, ICUs
- Train more providers in all shifts
- Integrate Ceribell into various patient population protocols based on established guidelines
- Expand to NICU, PICU, and Ped ED

Two-Pronged Platform Expansion Strategy



Algorithm Development

- Develop novel algorithms to support clinical expansion to new patient populations, and enhance seizure management offering

- ✓ FDA 510(k) clearance for artifact reduction algorithm
- ✓ FDA 510(k) clearance for epileptiform abnormality detection algorithm



Hardware Enhancements

- Provide comprehensiveness of conventional EEG and speed of point-of-care EEG
- Launching new product platform in 2027

- ✓ FDA 510(k) clearances for new recorder with video, ECG, and long-term monitoring
- ✓ FDA 510(k) clearances for two new headband designs

Ceribell's Three Growth Horizons



Ceribell's Goal: Make EEG A New Vital Sign



ceribell®



Stroke



Seizure



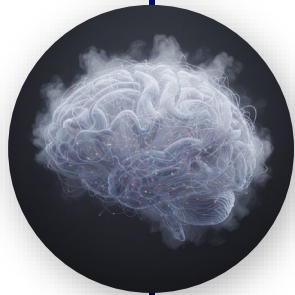
Delirium



Other

Moving Beyond Seizure in the Acute Care Setting

Delirium



Received FDA Clearance:
Delirium Algorithm



Activated First Pilot Sites



Targeting commercial launch
by year-end 2026

Received NTAP Favorable
Final Rule from CMS

in August 2026 for delirium monitoring
solution

Up to \$2,171 per patient²

Effective October 1, 2026

Unlocks >\$1B potential market opportunity¹

Stroke



Received FDA Breakthrough Designation:

LVO stroke detection algorithm in inpatient setting

1. U.S. acute care delirium market opportunity estimated by applying preliminary pricing assumptions to estimated patients at risk of delirium, less overlapping seizure patients. Actual results may differ materially.
2. Reflects NTAP per eligible Medicare patient case.

Delirium Represents a Significant Unmet Need

Delirium, defined as an acute change in attention and awareness, is often called “**acute brain failure**”

High Prevalence

- **3+ million** patients in the US¹⁻⁴
- **~30%** of Intensive Care Unit patients⁵
- up to **80%** of mechanically ventilated patients⁶

Poor Clinical Outcomes

- 1 ICU delirium day associated with **10% mortality risk increase**⁷
- **60% increase** in likelihood of developing **dementia** after surviving delirium in the ICU⁸

Unmet Need

- Current diagnosis tool (CAM-ICU) is dependent on nurse training, **binary**, and typically only administered **once or twice per day**

1. Oh E.S., et al. JAMA. 2017 Sep 26;318(12):1161–1174

2. Lindroth H., et al. J Acad Consult Liaison Psychiatry, 65 (5) (2024), pp. 417-430

3. Watt J., et al. Journal of general internal medicine 33, no. 4 (2018): 500-509

4. American Delirium Society, <https://www.americandeliriumsociety.org/What-Is-Delirium>

5. Krewulak, K.D., et al. (2018) Crit Care Med 46(12):p 2029-2035

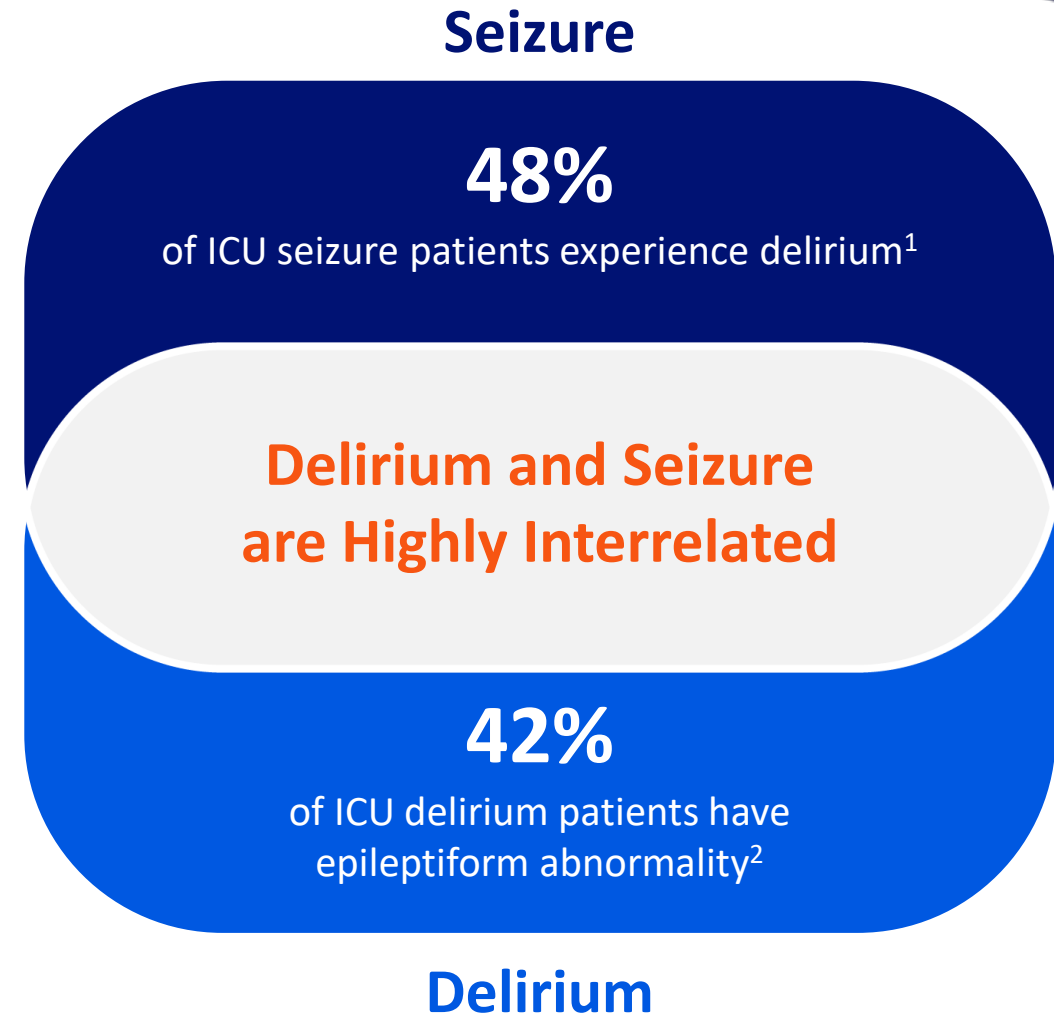
6. Girard, T.D., et al. (2008). Crit Care 12 Suppl 3(Suppl 3):S3

7. Ely EW., et al. Delirium as a predictor of mortality in mechanically ventilated patients in the intensive care unit. JAMA 291(14):1753–62 (2004)

8. Wang, S. et al. (2024) Alzheimer's & Dementia, 20(1), 278-287.

Delirium & Seizure: A Hidden Clinical Overlap

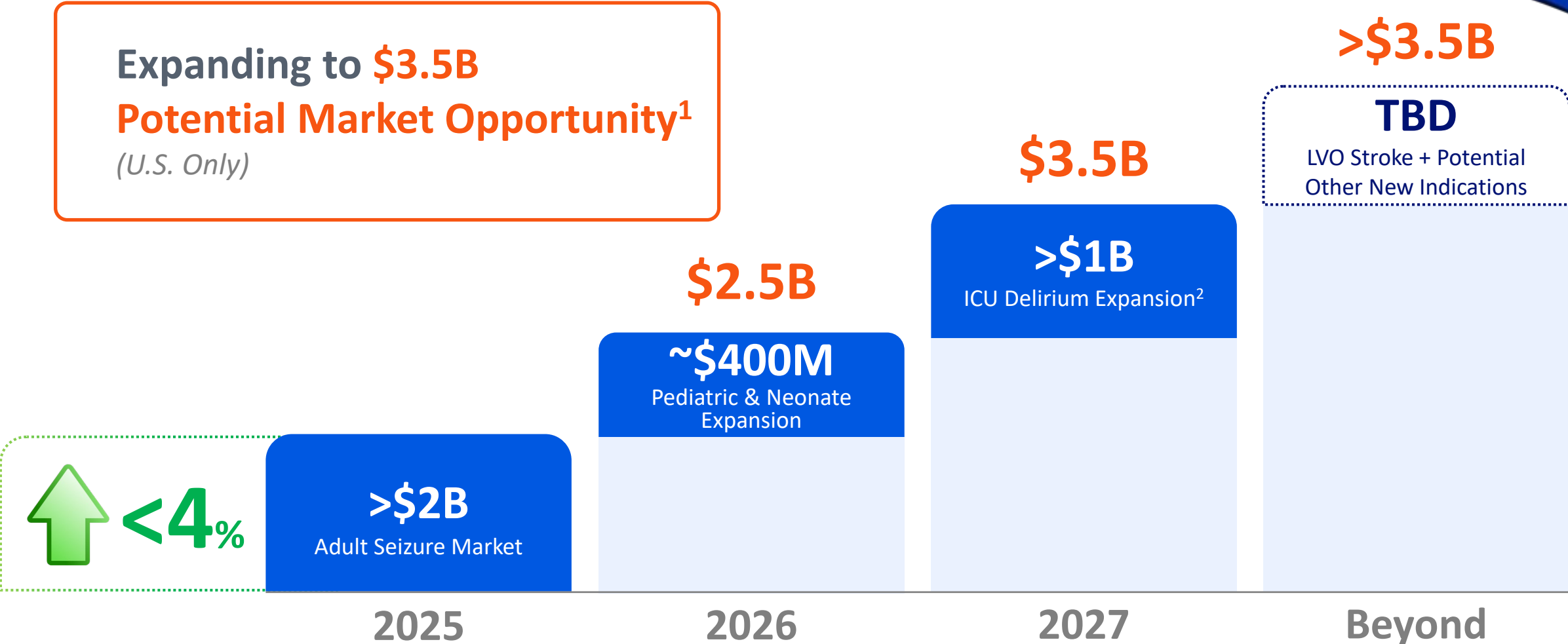
- Similar presentation...
- Very different treatment approaches



1. Frei A.I., et al. (2023) J Neurol. 271(1):231–240

2. Sambin S., et al. (2019) Front Neurol. 10:263

Pipeline Programs Significantly Expand Ceribell's TAM



1. Chart is not to scale. TAM based on management estimate. Actual results may differ materially.
2. Acute care delirium market estimated by applying preliminary pricing assumptions to estimated patients at risk of delirium, less overlapping seizure patients.

ceribell[®]

The First and Only

Status Epilepticus

FDA Breakthrough and CMS NTAP

Seizure Detection Algorithm

for Preterm Neonates

Seizure Detection Algorithm

for Ages 1+

Epileptiform Abnormality Pattern Detection Algorithm

FDA Cleared

FedRAMP[®] High

Authorization

Delirium Detection Algorithm

FDA Breakthrough and Cleared

LVO Monitoring Algorithm

Breakthrough Designation

ceribell®

Clarity When It's Critical