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Economic review of point-of-care EEG

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ABSTRACT

Aims: Point-of-care electroencephalogram (POC-EEG) is an acute care bedside screening tool for the identification of nonconvulsive seizures (NCS) and nonconvulsive status epilepticus (NCSE). The objective of this narrative review is to describe the economic themes related to POC-EEG in the United States (US).

Materials and methods: We examined peer-reviewed, published manuscripts on the economic findings of POC-EEG for bedside use in US hospitals, which included those found through targeted searches on PubMed and Google Scholar. Conference abstracts, gray literature offerings, frank advertisements, white papers, and studies conducted outside the US were excluded.

Results: Twelve manuscripts were identified and reviewed; results were then grouped into four categories of economic evidence. First, POC-EEG usage was associated with clinical management amendments and antiseizure medication reductions. Second, POC-EEG was correlated with fewer unnecessary transfers to other facilities for monitoring and reduced hospital length of stay (LOS). Third, when identifying NCS or NCSE onsite, POC-EEG was associated with greater reimbursement in Medical Severity-Diagnosis Related Group coding. Fourth, POC-EEG may lower labor costs *via* decreasing after-hours requests to EEG technologists for conventional EEG (convEEG).

Limitations: We conducted a narrative review, not a systematic review. The studies were observational and utilized one rapid circumferential headband system, which limited generalizability of the findings and indicated publication bias. Some sample sizes were small and hospital characteristics may not represent all US hospitals. POC-EEG studies in pediatric populations were also lacking. Ultimately, further research is justified.

Conclusions: POC-EEG is a rapid screening tool for NCS and NCSE in critical care and emergency medicine with potential financial benefits through refining clinical management, reducing unnecessary patient transfers and hospital LOS, improving reimbursement, and mitigating burdens on healthcare staff and hospitals. Since POC-EEG has limitations (i.e. no video component and reduced montage), the studies asserted that it did not replace convEEG.

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Introduction

From a clinical perspective, status epilepticus (SE) is a medical emergency comprised of prolonged seizure activity that can induce irreversible neuronal damage or death^{1–5}. Estimates of the annual incidence of SE in the United States (US) are variable at 18⁶ to 41⁷ patients per 100,000 population. Median daily hospitalization costs for SE inpatient admissions in the US are approximated at \$2,366 to \$3,359, depending on SE refractoriness⁸.

The American Clinical Neurophysiology Society³ provides specific terminology defining SE in their guidelines. Electroclinical SE refers to SE involving both electroencephalogram (EEG) findings and clinical manifestations of seizures, which may include prominent motor activity leading to the

classification of convulsive SE. Electrographic SE, in contrast, refers to SE that primarily involves EEG data without requiring clinical manifestations of seizures³. Nevertheless, since SE must be promptly addressed¹, a physician's decision to treat a patient for SE may be empirical. This is particularly relevant for nonconvulsive seizures (NCS) and nonconvulsive SE (NCSE), as NCSE does not have the prominent motor activity characteristic of convulsive SE^{3,9}. Accordingly, clinical symptoms in patients with NCSE can be difficult to detect or even absent, obfuscating empirical treatment decisions^{2,3,10–12}. Underlying causes of NCSE include brain injury^{13,14}, undifferentiated comatose states¹⁵, and systemic medical conditions, such as sepsis¹⁶; therefore, physicians often require EEG to confirm NCSE in patients with impaired consciousness.

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Clinical acumen combined with EEG assessment is ultimately needed to corroborate an NCSE diagnosis^{3,10}.

NCSE is a common medical condition in hospitalized patients, spurring clinical and financial burdens. In an observational cohort study across three years at a tertiary care center, nonconvulsive and subtle SE (with minimal clinical symptoms) accounted for 81% of SE episodes¹⁷. Laccheo et al.¹¹ reported that the median length of stay (LOS) in the intensive care unit (ICU) was 3 days longer for neurological patients with than those without NCS or NCSE. The former group also had more than double the mortality rate of the latter¹¹. In the US, the median cost of SE-related admissions among adults is \$15,595¹⁸.

Conventional EEG for NCS and NCSE detection

Due to the challenges in the clinical identification of NCS and NCSE, together with the medical^{11,12} and financial repercussions^{8,18}, accurate and objective means of facilitating NCSE diagnosis are imperative. Conventional 16-channel EEG (convEEG) using the International 10–20 electrode system remains the standard modality in detecting NCSE, as well as other forms of electroclinical and electrographic SE, and for informing treatment decisions^{3,10,19–21}. The Neurocritical Care Society² states that continuous EEG initiation should occur within 1 h of clinical suspicion of SE, including NCSE. Based on these guidelines², the American Clinical Neurophysiology Society²² recommends that “critical care continuous EEG should be initiated as soon as possible when NCS are suspected, since prolonged NCS and NCSE are associated with higher morbidity and mortality and treatment is likely to be more effective earlier in the course.”

Despite the urgency intrinsic to detecting and treating NCS and NCSE, implementation of convEEG is resource-heavy and time-consuming; in academic hospitals, the median delay for convEEG is nearly 4 h long²³. The reasons are manifold. First, EEG technologists must operate convEEG, but staffing and EEG equipment availability can be expensive and scarce. Since it is unknown when or where SE will occur in a hospital, both EEG technologists and convEEG may already be occupied. This complicates triaging, for when EEG supply is needed to assess the severity of a patient’s condition, it may be unprocurable. Further, inpatient EEG requests may stack such that convEEG in use for active monitoring must be disconnected, packed, and transported to patients in other departments. When the EEG arrives at bedside, additional time is expended in setup and troubleshooting by the EEG technologist for proper signal acquisition. Finally, only once a neurologist or neurology subspecialist interprets the EEG data, can the study be resulted^{24,25}.

Each of these issues compounds the time to diagnosis and delays proper care of NCSE. Standard business hours²⁶ of EEG coverage from 8:00am–4:00pm or 9:00am–5:00pm, Monday through Friday in some hospitals only comprises about 24% of the entire week, yet again, seizure occurrence is not confined to weekday workday hours. The demand for inpatient EEG is unpredictable and on-call services may in turn be constrained. A NCSE patient could experience several

seizures before receiving convEEG evaluation given lengthy delays, which exist even among academic hospitals with onsite EEG technologists during and outside business hours^{23–25,27}. Postponement of convEEG assessment and SE treatment to hours later, the next workday, or even longer may jeopardize patient health^{23,27,28}. Although continuous EEG has been linked with more seizure detection and alterations to antiseizure treatment, some evidence has indicated that it may not benefit patient outcomes as compared with routine EEG²⁹. Hospitalization charges may also rise with the use of continuous EEG³⁰.

In the absence of timely EEG assessment, healthcare providers relying on incomplete information (i.e. clinical evaluation only) are at risk for overtreating in some cases and administering inadequate treatment in others. Delays in convEEG for a suspected actively-seizing patient may precipitate empirical treatment; thus, some patients may be erroneously treated for NCSE based on clinical manifestations alone^{25,31}. Benzodiazepines, which are first-line antiseizure medications (ASMs) to abort seizures, can have substantial side effects² if prescribed to patients without NCSE or other seizures. Spatola et al.³² found that excess benzodiazepine administration was associated with an increased use of orotracheal intubation, and subsequently, extended hospital LOS. In a clinical trial (ESETT) of rapid treatment of seizure without the benefit of EEG monitoring, researchers similarly noted that the lack of improvement in some patients may have been due to the sedative nature of ASMs prescribed to those who might not have been in SE. These patients therefore may have incorrectly received anti-epileptic treatment³¹.

Moreover, an analysis of data from ESETT and from another clinical trial on prehospital benzodiazepine administration (RAMPART)³³ revealed that about 8% of patients had prolonged psychogenic nonepileptic seizures, even though they had been treated for SE. Over 20% of these inappropriately treated patients required an ICU stay and almost 30% experienced adverse events³⁴. High rates of benzodiazepine administration to patients with dissociative (i.e. nonepileptic) seizures have been observed in a different study at a university hospital as well³⁵. With the accuracy of diagnosing video recordings of epileptic versus psychogenic nonepileptic seizures ranging from about 44% to 58% among non-neurology healthcare staff³⁶, the importance of EEG monitoring to minimize overmedication of benzodiazepines is clear. Lastly, in addition to the detrimental impact on patient health, overtreatment in general drains resources, with approximately \$75.7 to \$101.2 billion wasted per year from overtreatment or low-value care in the overall US healthcare system³⁷.

Without prompt confirmation of seizures *via* EEG, patients could conversely receive insufficient treatment²⁵. Benzodiazepine underdosing is prevalent and may lower seizure recovery rates^{38,39}. ASMs are less effective when prescribed late into prolonged seizure and protracted SE is linked to worse health outcomes²⁸; early detection of SE is again vital. Delays may be particularly inimical for patients who are then prescribed suboptimal benzodiazepine doses³⁸, though these findings on treatment dosing and timing may be less suited for electrographic seizures, focal SE, and NCSE. Moreover, inpatient

hospital admission costs are inversely proportional to treatment success. Compared with the costs associated with non-refractory SE, those of refractory SE and super-refractory SE are two and five times greater, respectively¹⁸. Like overtreatment, inadequate treatment of SE may have financial consequences as well.

Point-of-care EEG for NCS and NCSE detection

Over the past years, new high-quality EEG devices have been developed and introduced to hospitals for patients at risk for SE, especially NCSE^{21,23,24,40,41}. The purpose of these technologies is to facilitate faster bedside detection without mandating technologist involvement. This narrative review defines such devices as point-of-care EEGs (POC-EEGs). With fewer electrodes than convEEG, POC-EEG serves as a screening test for NCS and NCSE²¹. The speed of POC-EEG encapsulates its value; its setup time has been reported as about 5 to 6 min^{23,42,43}, with an approximate time from order and time from arrival at the emergency department (ED) or ICU, respectively, to initiation of 23⁴³ and 96⁴¹ minutes. The median duration of POC-EEG monitoring is about 2 h⁴⁴, which is distinct from that of routine EEG, with its recording time of around 0.3–0.5 h⁴⁵.

Non-neurologist physicians, nurses, and other allied health professionals can deploy POC-EEG, as its operation is independent of EEG technologists and only basic training is necessary for application^{25,42}. Although POC-EEG application is non-specialized, neurologists or neurology subspecialists are still responsible for the interpretation of recordings. Since the incorporation of POC-EEG in various hospital settings, its impact on workflow and outcomes has been illustrated^{44,46,47}. POC-EEG, however, neither antiquates nor supplants convEEG^{48–50}, nor does it replace the role of EEG technologists or neurologists. ConvEEG underpins patient diagnosis and treatment strategy^{10,19,20}. The utility of POC-EEG is in time-sensitive ED and ICU cases, including ruling out NCSE^{48–50}. Continuous EEG, when available, should ideally be acquired for patients with NCS or NCSE detected on POC-EEG to ensure ongoing evaluation and long-term treatment planning.

Since POC-EEG utilization has grown in the last few years among large tertiary and academic hospitals, as well as in community hospitals, device performance has been investigated^{48,51,52}. Research on the economic impacts of POC-EEG, nonetheless, is relatively sparse. The aim of this narrative review is to summarize the literature on the economic themes related to POC-EEG use for the bedside detection of NCS and NCSE in US hospital patients.

Methods

Research related to economic themes for POC-EEG evaluation of hospital patients in the US was identified. The evidence base consisted of articles provided by the study funder and those found through targeted searches on PubMed and Google Scholar. The relevant search terms are listed in [Table S1](#). As this was a narrative review, Preferred Reporting Items

for Systematic Reviews and Meta-Analyses (PRISMA) or other formal systematic review protocols were not utilized.

Studies were restricted to peer-reviewed publications involving US hospitals, written in English, and conducted with hospital patients at bedside; these studies had to feature economic or potential economic themes associated with POC-EEG. Economic or potential economic themes considered were budgetary implications, cost-benefits, cost-savings, healthcare resource utilization (including staffing), reimbursement, and other monetary-related data (such as LOS), all of which could be actual, possible, or theoretical. Studies about clinical management modifications associated with POC-EEG utilization that could lead or relate to any of these themes were also considered. Adhering to these same guidelines, pertinent references within relevant publications were assessed for applicability.

Conference abstracts, gray literature offerings, frank advertisements, white papers, and studies conducted outside the US were ineligible for this narrative review. Further excluded studies were those describing POC-EEG accuracy, feasibility, performance, or use that did not feature any implicit or explicit economic findings, such as research that tested the sensitivity and/or specificity of POC-EEG in seizure detection. [Figure 1](#) summarizes study selection.

POC-EEGs were defined as reduced-montage, rapid-EEG systems utilized at bedside in US hospitals. Ambulatory, mobile, or video EEG devices, as well as EEG devices in the outpatient setting, were not regarded as POC-EEGs for this narrative review, nor were post-hoc full-montage to reduced-montage reconfigurations. These systems and others have been examined elsewhere, along with POC-EEGs in other countries^{48,51–53}. Economic studies on convEEG or continuous/video EEG were additionally excluded.

Results

Twelve publications on the economic themes associated with POC-EEG were identified and cited in the results of this narrative review. These studies were published from 2018 to 2023; most were single center. All investigated a single rapid circumferential headband system. [Table 1](#) provides a more detailed overview of the literature presented. After examination of the economic themes presented in this included research, the study results were organized into four categories of patient care-driven economic evidence for POC-EEG: refining clinical management, reducing unnecessary patient transfers and hospital LOS, improving reimbursement, and mitigating burdens on healthcare staff and hospitals.

Refining clinical management

Findings in eight studies indicated that POC-EEG could help fine-tune clinical management, which has possible economic implications. Overall, the trend was that POC-EEG supported physicians' ability to rule out seizures, prompting decreased administration or planned administration of unnecessary ASMs.

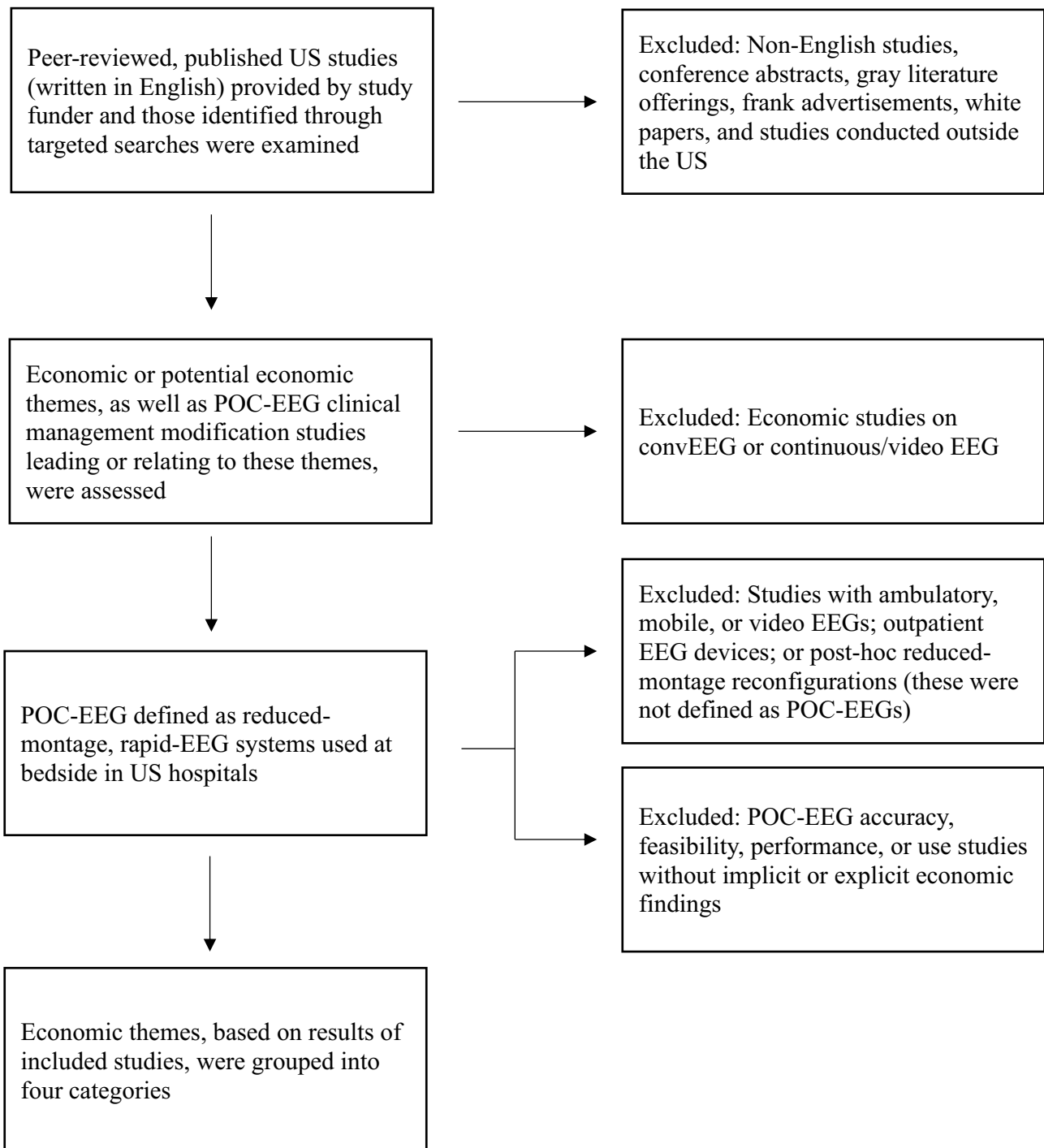


Figure 1. Overview of study design.

ConvEEG, Conventional electroencephalogram; EEG, Electroencephalogram; POC-EEG, Point-of-care electroencephalogram; US, United States.

The multicenter prospective observational DECIDE trial was the pivotal study that demonstrated POC-EEG's utility in influencing clinical management among physicians. It was conducted at five academic hospitals across the US. Study researchers distributed the same questionnaire to physicians before and after POC-EEG use to quantify treatment plan changes for patients with altered mental status. Due to evaluation with POC-EEG, physicians modified their diagnostic suspicion for seizures and adjusted treatment decisions in approximately 40% ($n=72/179$) and 20% ($n=36/179$) of

cases, respectively. They were also less likely to intensify treatment with ASMs following utilization of POC-EEG in almost 13% ($n=23/179$) of patients²³.

Findings similar to those in the DECIDE trial²³ have been repeated in other prospective studies in various US hospitals, further corroborating that POC-EEG can play a role in improving clinical management. In Stanford University Hospital (SUH) ICUs, Hobbs et al.⁴³ showed that POC-EEG usage after initial assessment spurred physicians to amend treatment in 40% ($n=14/35$) of suspected NCSE cases,

Table 1. Summary of included studies.

Main Economic or Potentially Economic Findings Cited for POC-EEG on:							
Author(s)	Journal and Publication Year	Study Type	Study Population (Patients)	Refining Clinical Management	Reducing Unnecessary Patient Transfers and Hospital LOS	Improving Reimbursement	Mitigating Burdens on Healthcare Staff and Hospitals
Eberhard and Beckerman ⁴⁷	<i>J Neurosci Nurs</i> (2023)	Retrospective cohort	102	–	LOS decreased by 3 days; almost \$740,000 in potential annual savings	10 additional DRGs “with MCC/CC” billed; nearly \$146,000 in potential annual gain	–
Hobbs et al. ⁴³	<i>Neurocritic Care</i> (2018)	Prospective cohort	34	Clinical management amended in 40% of suspected NCSE cases; fewer ASMs prescribed ($p = 0.01$)	–	–	POC-EEG user-friendliness
Kamoussi et al. ⁴⁰	<i>Neurocritic Care</i> (2021)	Retrospective cohort	353	–	–	Potential additional DRG “with MCC/CC” billing	–
Kozak et al. ⁴⁴	<i>J Am Coll Emerg Physicians Open</i> (2023)	Retrospective cohort	157	Clinical management amendments correlated with POC-EEG findings ($p < 0.001$)	ED ($p = 0.04$) and hospital ($p < 0.001$) LOS associated with door-to-EEG time ^a	–	Extended EEG coverage with POC-EEG
Kurup et al. ⁵⁵	<i>Epileptic Disord</i> (2022)	Retrospective chart review	100	Decreased probability of initiation/continuation of ASMs ($X^2 = 7.086, p = 0.0078$)	–	–	–
LaMonte ⁵⁰	<i>Epilepsia Open</i> (2021)	Prospective cohort with retrospective controls	50	Decreased overtreatment for 90% of patients concerning for SE	–	–	Diminished after-hours EEG technologist labor ($p = 0.02$); healthcare staff positively commented on POC-EEG impact
Madill et al. ⁵⁷	<i>Epileptic Disord</i> (2022)	Retrospective cohort	74	–	After POC-EEG implementation, only 6% of patients were transferred offsite for EEG; >\$39,000 in ground transportation savings	Potential to decrease reimbursement lost to other hospitals	Extended EEG coverage with POC-EEG
Vespa et al. ²³	<i>Crit Care Med</i> (2020)	Prospective multicenter nonrandomized observational	181 ^b	Adjusted treatment decision in 20% of cases; projected 51% decrease in parental ASM treatments and intubations ^c	ICU and hospital LOS decreased by 0.4 and 1.2 days, respectively; \$3,971/POC-EEG use saved ^c	–	POC-EEG user-friendliness
Ward et al. ⁴⁶	<i>Front Digit Health</i> (2023)	Prospective cohort with retrospective controls	88 ^d	–	Reduced transfer rate to 1.1 transfers/month; >\$37,000 in potential annual savings and potential gain of nearly \$14,000/non-transferred patient	Potential to decrease reimbursement lost to other hospitals; greater amount collected (\$11,161) by hospitals for non-transferred patients assessed with POC-EEG	–
Wright et al. ⁵⁴	<i>Emerg Med J</i> (2021)	Prospective cohort	38	Impacted clinical management in 53% of suspected NCSE cases	–	–	–
Yazbeck et al. ⁴²	<i>J Neurosci Nurs</i> (2019)	Prospective cohort	10	De-escalated antiseizure treatment in 40% of patients	–	–	POC-EEG user-friendliness

ASM, Antiseizure medication; DRG, Diagnosis related group; ED, Emergency department; EEG, Electroencephalogram; ICU, Intensive care unit; LOS, Length of stay; MCC/CC, Major complication or comorbidity/complication or comorbidity; NCSE, Nonconvulsive status epilepticus; POC-EEG, Point-of-care electroencephalogram; SE, Status epilepticus.

^aDoor-to-EEG time was defined as time from patients’ first ED evaluation to the initiation of POC-EEG monitoring.⁴⁴

^bThere were missing data for two patients for the refining clinical management analysis.²³

^cProjected treatment decreases, LOS results, and monetary savings were quantified by Ney et al.²⁵

^dThis was the number of patients assessed with POC-EEG; the number of patients assessed with convEEG (control group) was not provided.⁴⁶

precipitating treatment with significantly fewer ASMs ($p=0.01$). Additionally, POC-EEG evaluation may have averted intubation of a patient scheduled for further medical care⁴³. At a California community hospital ICU, POC-EEG for suspected seizures helped enable de-escalation of antiseizure treatment in 40% ($n=4/10$) of patients⁴². A study at a Maryland teaching hospital likewise found that negative POC-EEG evaluations decreased overtreatment with ASMs in 90% ($n=9/10$) of patients concerning for SE⁵⁰. Furthermore, at the EDs of Stanford Health Care (SHC) in California and Episcopal Hospital (EH) in Pennsylvania, POC-EEG impacted clinical management in over half (53%, $n=20/38$) of suspected NCSE cases at the two hospitals. Compared with treating only on clinical suspicion, EH emergency physicians, who were auditorily interpreting sonified POC-EEG data, prescribed fewer ASMs in 43% ($n=6/14$) of their cases after POC-EEG monitoring. (On-call neurologist visual inspection verified the diagnoses⁵⁴.)

Two retrospective studies of POC-EEG also emphasized this economic theme of refining clinical management. Kurup et al.⁵⁵ performed a chart review of 100 patients assessed with POC-EEG for any reason at SUH ED and ICU. For the 19 (19%) patients presenting with highly epileptiform or generalized/focal seizures on POC-EEG, 84% ($n=16/19$) received or continued to receive ASMs, in contrast to 51% ($n=41/81$) of those presenting with normal or nonepileptic encephalopathic patterns (NL/SL) on POC-EEG. ASM initiation or continuation was significantly less probable for patients with NL/SL after POC-EEG, compared with patients for whom seizures were not ruled out after POC-EEG ($X^2=7.086$, $p=0.0078$)⁵⁵. Kozak et al.⁴⁴ analyzed POC-EEG data over the span of one year from 157 ED patients at Providence Mission Hospital Mission Viejo, a community hospital in California. POC-EEG evaluation resulted in discontinuation of ASMs in 49 (53%) of the 93 patients who had received ASMs prior to POC-EEG. Of the 88 patients treated with ASMs after POC-EEG assessment, 44 (50%) patients had received ASMs based on POC-EEG findings. Altogether, clinical management amendments were significantly correlated with POC-EEG findings ($p < 0.001$)⁴⁴.

To determine financial implications from POC-EEG's influence on clinical management, Ney et al.²⁵ developed a decision-analytic model from extracted data of the DECIDE trial²³ and other sources, such as the 2017 National Inpatient Sample dataset⁵⁶. The model examined the costs and benefits of treating patients with suspected seizures based on POC-EEG versus treating based only on clinical assessment. For each POC-EEG use, the authors determined an average savings of \$3,971, given the projected 51% decrease in parenteral ASM treatments and intubations, along with the shorter ICU and hospital LOS. These conclusions, including the monetary savings, were theoretical, since they were derived from a model, not from a randomized clinical trial or prospective study²⁵.

Reducing unnecessary patient transfers and hospital LOS

Five studies illustrated that POC-EEG can reduce both unneeded patient transfers to other sites for EEG assessment

and extended hospital LOS. In conjunction with refining clinical management, decreasing unnecessary patient transfers and LOS may retrench hospital expenses.

Madill et al.⁵⁷ examined the introduction of POC-EEG at SHC ValleyCare, a community hospital in California. Before the introduction of POC-EEG, patients arriving outside business hours with suspected NCS were transferred to SUH for convEEG or were evaluated during the next business day. The authors found 33 cases of suspected seizure in sixteen months that would have met criteria for transfer before POC-EEG implementation. With POC-EEG, 2 (6%, $n=2/33$) of these patients were transferred to SUH for EEG. Based on an average cost of \$1,274 per patient for inter-facility transfer *via* ambulance⁵⁸, the researchers calculated savings of over \$39,000 just for ground transportation during the study period from December 2018 to the end of March 2020⁵⁷.

In a prospective cohort study of POC-EEG in a New Jersey community hospital setting, a mean of 2 ($n=22$ across 11 months) patients each month were transferred offsite for emergent EEG prior to POC-EEG introduction. Over the ten months since using POC-EEG, these two hospitals experienced a 45% reduction in mean transfer rate (1.1 transfers per month) for emergent EEG. Ward et al.⁴⁶ calculated a net loss of about \$3,463 for each historically transferred patient (\$4,037, the transfer reimbursement, subtracted from \$7,500, the transfer cost). For each non-transferred patient evaluated with POC-EEG, POC-EEG was attributed with a total gain of almost \$14,000, according to the approximate avoided transfer cost (\$3,463) combined with the mean amount collected (\$11,161) and acknowledging the headband cost (\$688). Ultimately, the savings from POC-EEG from reducing unnecessary transfers were projected at over \$37,000 for the year ($[\$3,463.11/\text{patient transfer} \times \text{mean of 2 patient transfers/month} \times 12 \text{ months}] - [\$3,463.11/\text{patient transfer} \times \text{mean of 1.1 patient transfers/month} \times 12 \text{ months}] = \$37,401.59$). Further cost offsets could occur if interfacility transfers were by air ambulance, due to remoteness or medical urgency, in which averted transfers could save over \$40,000 in transportation costs per patient⁴⁶.

Despite these potential financial benefits, the price of POC-EEG must also be considered. The authors of the previous research elaborated that POC-EEG would have to preclude about 0.7 transfers per month, or 8.6 patients annually, to offset the multi-hospital system cost, including disposables, of the POC-EEG (\$119,700 per year) in their study⁴⁶.

Along with fewer transfers, some of the research demonstrated that POC-EEG usage and abbreviated length of hospitalization were linked. Kozak et al.⁴⁴, in their retrospective study at a California community hospital ED, found a positive association between ED ($p=0.04$) and hospital LOS ($p < 0.001$) and the time from patients' first ED evaluation to the initiation of POC-EEG. In another retrospective study at a different community hospital in northern California, hospital days were also fewer for patients receiving POC-EEG. Among 62 patients assessed with POC-EEG, Eberhard and Beckerman⁴⁷ found a median LOS decrease of 3 days (4 versus 7 days before and after POC-EEG introduction at the hospital, respectively), yet this was not significant ($p=0.058$).

Ney et al.²⁵ reported that shorter LOS (0.4 and 1.2 days fewer for ICU and hospital LOS, respectively) contributed to the estimated average savings (\$3,971 per patient) with POC-EEG utilization in the decision-analytic model based on DECIDE trial findings²³. LOS for false-positive suspected seizure was the main factor in total cost variance, since these patients received ASMs, intubations, and ventilations that could have been avoided with more informed treatment decisions²⁵. Similarly, Eberhard and Beckerman⁴⁷ noted potential cost reductions due to decreases in LOS with POC-EEG usage in their aforementioned study at the northern California community hospital. From an extrapolation of their six-month findings across one year, they postulated that fewer hospitalization days and ED discharges with POC-EEG implementation would be linked with nearly \$740,000 in annual savings. The costs of equipment were incorporated in the financial analysis⁴⁷. Again, however, these two models by Ney et al.²⁵ and Eberhard and Beckerman⁴⁷ were derived from calculations of real-world data, not the results of a *priori* clinical trial analyses.

Improving reimbursement

Combined with the other categories of economic impact, securing additional reimbursement for NCS or NCSE patients *via* POC-EEG may support hospital finances, as shown in four of the studies identified for this narrative review. Medical Severity-Diagnosis Related Group (MS-DRG) with baseline DRG, as well as elevated reimbursement categories relating to complication or comorbidity (CC) or major complication or comorbidity (MCC) assignment, plays a role in optimizing hospital reimbursement with POC-EEG.

As Madill et al.⁵⁷ explained, transferring hospitals may forfeit a proportion of reimbursement within the Medicare Part A MS-DRG to the hospitals that receive and treat the transferred patient⁵⁹. In the 2023 study examining patient transfers at the New Jersey community hospital setting, the approximate mean amount collected (\$11,161) for a non-transferred patient assessed with POC-EEG was greater than that of a historically transferred patient (\$4,037) in the control group. As described above, POC-EEG would have to preclude the transfer of 8.6 patients annually to offset its multi-hospital system cost (\$119,700/year ÷ \$13,936.44/patient = 8.59 patients/year), which accounted for the reimbursement differences for transferred versus non-transferred patients⁴⁶.

Regarding MS-DRG appropriate medical severity coding, a retrospective cohort study analyzed over 350 POC-EEG recordings from patients at six US academic and community hospitals. In total, epileptiform abnormalities or seizures were detected in 33% ($n = 141/353$) of cases⁴⁰. If the primary diagnosis for these cases were non-seizure, then POC-EEG could have enabled hospitals to code for the additional CC/MCC DRGs, thus bolstering reimbursement.

Indeed, Eberhard and Beckerman⁴⁷ evinced that use of POC-EEG was associated with higher MS-DRG coding ("with CC/MCC" DRGs) in their retrospective study on POC-EEG implementation at the northern California hospital. Within

the six-month period after POC-EEG was introduced, POC-EEG detected seizures in 15 patients, 10 (67%) of whom had a primary diagnosis of non-seizure. Subsequently, 1 (10%, $n = 1/10$) patient had an MS-DRG "with MCC," while the other 9 (90%, $n = 9/10$) patients coded for MS-DRG "with CC." The authors projected a total gain of almost \$146,000 for their hospital from MS-DRG reimbursements for one year of POC-EEG usage. This value, combined with the nearly \$740,000 in annual savings from fewer hospitalization days and ED discharges, and including the additional equipment costs (over \$200,000), would lead to a positive yearly income (about \$683,000) for this hospital ($[\$145,580 + \$737,818] - \$200,240 = \$683,158$)⁴⁷. When considering the impact of both transfer savings^{46,57} and the reimbursements associated with POC-EEG⁴⁷, the overall annual financial impact of POC-EEG may be greater, even as operating costs are applied.

Mitigating burdens on healthcare staff and hospitals

Lastly, several studies suggested that the adoption of POC-EEG could mitigate burdens on healthcare staff and hospitals. For 38% ($n = 15/40$) of control cases with convEEG assessment from the earlier study at the Maryland teaching hospital by LaMonte⁵⁰, EEG technologists were requested after normal business hours. However, none of the patients evaluated with POC-EEG required EEG technologist support outside workday hours, representing a marked diminution in after-hours EEG technologist labor ($p = 0.02$)⁵⁰.

Further, of 118 POC-EEG cases cited by Madill et al.⁵⁷, only 31% ($n = 37/118$) overlapped with SHC ValleyCare's typical business hours; this included 1 (10%, $n = 1/10$) seizure case. Kozak et al.⁴⁴ noted that just under half (45%, $n = 71/157$) of POC-EEG evaluations were conducted during business hours in the community hospital ED setting, consisting of 14 (64%) of 22 seizure cases. In these studies, the capacity for after-hours convEEG was either limited⁴⁴ or nonexistent⁵⁷, which may encumber EEG technologists and incur overtime or other additional costs when occurring⁴⁴.

Healthcare staff, including during COVID-19 isolation, have also positively commented on the impact of POC-EEG⁵⁰, such as its user-friendliness^{23,42,43}, and this is crucial for non-specialist operation. Although more causal proof is necessary, POC-EEG may alleviate scheduling and work burdens where both labor (EEG technologists) and equipment (convEEG) are scarce.

Discussion

POC-EEGs are rapid screening tests for NCS and NCSE in critical care and emergency medicine^{21,48}; they can confirm or rule out NCS or NCSE diagnoses when convEEG technology is constrained. This narrative review examined the economic value of POC-EEG, a resource that has been implemented in hospitals across the US in recent years to expedite patient care in the detection of NCS and NCSE^{48,51,52}. Key patient care-driven economic categories were refining clinical management, reducing unnecessary patient transfers and hospital

LOS, improving reimbursement, and mitigating burdens on healthcare staff and hospitals.

Utilization of unnecessary anti-epileptic treatments and procedures is costly and can lengthen hospitalization stay for patients who must recover from excess benzodiazepines and ventilations^{25,32}. POC-EEG can fine-tune clinical management *via* quickly screening for NCS and NCSE, particularly when convEEG and EEG technologists are not readily available^{23,25,42–44,50,54,55}. This in turn could precipitate cost-savings²⁵. Moreover, POC-EEG has shown economic value in eliminating the need for patient transfers, which can be expensive, through enabling physicians to assess for NCS and NCSE on-site^{46,57}. Transfers can additionally generate anxiety for patients and their family members, squander healthcare resources, and encumber society through inflating Medicare and/or Medicaid costs for federally insured patients^{60,61}.

Prolonged LOS can be harmful for patients, including those with NCS or NCSE and those incorrectly treated for it, as well as to the healthcare system³². Since POC-EEG assessment can affect clinical management, it may curb overtreatment and shorten hospitalization stay^{25,44,47}, thereby paring expenses and improving clinical outcomes. POC-EEG has the capacity to bolster reimbursement through avoiding unnecessary transfers and triggering higher Medical Severity DRG-related billing codes^{46,47,57}. The latter is notable because the average additional reimbursement for an MS-DRG “with MCC” is about \$7,000 for conditions linked frequently with NCSE^{22,62}. Finally, POC-EEG adoption could lead to savings in EEG technologist labor expenditures for staffing convEEG requests outside business hours^{44,50,57}. Overtime and lengthy working hours have been correlated with burnout across various fields of medicine, in conjunction with health risks, such as ischemic heart disease, motor vehicle accidents, stroke, workplace injury, and death^{63–65}. Of course, neurologist or neurology subspecialists must still be present outside weekday workday hours, including overnight shifts, to interpret POC-EEG recordings.

POC-EEG tradeoffs

Although convEEG systems have their own costs, both upfront and annualized⁶⁶, they differ from those of POC-EEG and depend on specific hospital infrastructure. It however must be reiterated that POC-EEG does not replace convEEG, the work of EEG technologists, or the neurology review of EEG recordings, nor does it render them obsolete^{48–50}. POC-EEG indeed has limitations in comparison with convEEG. Overall, convEEG *via* the International 10–20 electrode system provides a more comprehensive evaluation of electrographic activity in patients than does POC-EEG with its fewer electrodes^{21,49,50}. The rapid circumferential headband system mentioned in this narrative review, for example, does not place electrodes in the midline or parasagittal regions of the brain. Consequently, it may not detect midline or parasagittal seizures, which have a prevalence of about 1%, or other epileptic abnormalities in these areas^{43,49}.

Further, convEEG has the potential to be used with concomitant video, as endorsed by the American Clinical Neurophysiology Society²² to aid physicians in patient care, whereas POC-EEG does not. Signal artifacts have in turn been more frequent in POC-EEG than convEEG evaluation; this may obfuscate interpretation of electrographic data⁴¹. POC-EEG also has a lower sampling rate (250 Hz) than convEEG (200–1,000 Hz)⁴⁶, thus limiting analysis in the high frequency domain. Ease of POC-EEG application by a range of healthcare providers^{23,42,43} for rapid assessment of patients, as previously described, may counter some of these limitations. Similarly, when integrated with an automated algorithm, POC-EEG can be used to continuously monitor and alert bedside providers of suspected NCS or NCSE within minutes, whereas convEEG requires human monitoring at an infrequent rate⁴⁰. Ultimately, when utilized as a screening tool and monitor for NCS or NCSE in critical care and emergency medicine, POC-EEG complements but does not supplant convEEG^{48–50}.

Limitations

This narrative review was based on targeted searches, as depicted in Table S1 and Figure 1. It was not a systematic review conducted under PRISMA guidelines and should not be held to the same reporting standards. The studies identified in this narrative review were largely local or regional in geographic scope. As such, the findings may not be generalizable to all hospitals in the US or inpatient medical centers in non-US settings, like those in low-income countries. Additionally, it is unknown whether the financial gains of POC-EEG would be applicable to pediatric patients because the research in this narrative review was almost exclusively based on adult patients.

Sample sizes in some of the included studies were also modest, with two having only 10 patients receiving POC-EEG^{42,50}. In the study by Kozak et al.⁴⁴, neither ED nor hospital LOS was significantly associated with the time to first ASM treatment after POC-EEG utilization. Further, Ney et al.²⁵ primarily sourced clinical parameters from the DECIDE trial²³ for their model; its generalizability outside academic centers is uncertain²⁵. The twelve included studies were observational, not randomized controlled trials, which exposes them to systemic biases in how the data were collected and reported⁶⁷, particularly as the findings were in favor of POC-EEG.

All studies presented here investigated a single rapid circumferential headband system; the evidence on this device may not translate to other POC-EEGs. In the absence of a systematic review or meta-analytic framework, formal testing for publication bias, such as Begg’s⁶⁸ and Egger’s tests⁶⁹, are not available. On average, quality of the cohort studies cited within the results was fair (Table S2) according to the Newcastle-Ottawa scale⁷⁰; in general, nonetheless, assessment of these publications was not well suited to this scale given their aims and design. Examining the disclosure statements of the twelve cited publications also revealed that the majority of study authors received payment or support from

the manufacturer of that POC-EEG, who funded this narrative review. Similarly, negative economic outcomes linked with this particular POC-EEG were not identified.

Future directions

We have endeavored to survey pertinent available evidence for POC-EEG, yet a systematic review would likely expand this narrative review. The research on clinical management was the most robust, but overall, more studies analyzing this topic and these evidence categories are imperative, especially those with greater sample sizes, more geographic variety, and the resulting outcomes of clinical decisions. Assessing the long-term financial consequences of NCS and NCSE accrued over time, along with the influence of POC-EEG on reimbursement through Medicare Part A MS-DRGs⁵⁹, would be valuable as well. Since POC-EEGs have been implemented in other countries^{48,53}, it would furthermore be interesting to analyze whether POC-EEG utilization is more widespread around the globe and which unique challenges have arisen from it.

More broadly, point-of-care testing expands past POC-EEG utilization in the detection of NCS or NCSE in US hospital patients. Medical point-of-care devices, such as cardiac monitors⁷¹, point-of-care ultrasound⁷², and pupillometers⁷³, have been developed for multiple indications. Some devices, like cardiac monitors, are more widespread than POC-EEG, for individuals can use them at their own discretion⁷¹. Others must be actuated by qualified personnel and are restricted to inpatient units or hospitals^{72,73}. The goal of all these point-of-care technologies, nevertheless, mirrors that of POC-EEG; they aim to quickly and specifically assess certain aspects of patient health so that precise treatment is initiated sooner⁷⁴. Like with POC-EEG, the utility and finances of these devices compared with conventional devices should continue to be explored, both in and outside the US.

Conclusions

POC-EEG can refine clinical management of hospitalized patients with suspected seizures, reduce unnecessary patient transfers and hospital LOS, improve reimbursement, and mitigate burdens on healthcare staff and hospitals, all of which are accompanied with potential economic benefits. As an adjunct to convEEG, POC-EEG is an expeditious screening device for identifying NCS or NCSE in critical care and emergency medicine^{48–50} with the promise of financial advantages over standard care.

Transparency

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