

Epilepsy-related healthcare resource use and costs in commercially insured patients before and after initiating brivaracetam: a retrospective claims database analysis

Silky Beaty¹
Adina Estrin²
Edward Lee¹
Kelly Guntrum¹
Andrea Loewendorf¹
Michelle Skornicki²

1. UCB Pharma, Smyrna, GA, USA
2. Aetion, Inc., New York, NY, USA

EE86

Background

- Brivaracetam, an antiseizure medication (ASM) that can be started without titration, is approved in the United States as monotherapy or adjunctive therapy for focal (partial-onset) seizures in patients 1 month of age and older.¹
- Newer ASMs that allow for therapeutic doses at treatment initiation may affect healthcare resource use (HRU) and costs.

Objective

- To understand HRU and costs for commercially insured patients with epilepsy in the United States treated with brivaracetam in the 12 months pre- and post-treatment initiation.

Methods

- Retrospective cohort analysis using IBM MarketScan Commercial Claims and Encounters Database.
- Commercially insured patients ≥18 years of age with baseline epilepsy/seizure diagnosis and continuous medical and pharmacy benefit for 12 months pre- and post-brivaracetam initiation between March 1, 2016 and September 30, 2018 were included.
- Epilepsy-related costs (inpatient, outpatient, and pharmacy), epilepsy-related medical costs (inpatient and outpatient), and inpatient and outpatient HRU were defined as all claims with a diagnosis of epilepsy (*International Classification of Diseases, Ninth Revision, Clinical Modification [ICD-9-CM] codes: 345.X, 780.3X, 333.2; International Classification of Diseases, Tenth Revision, Clinical Modification [ICD-10-CM] codes: G40X, R56X, G25.3*) in any position or procedure code for ASM (C9254, J2560, C9238, J1953, J1165).
 - Total costs (payments made by commercial payers + out-of-pocket costs) were adjusted to 2019 United States dollars (USD).
- Patients with brivaracetam use during baseline period were excluded.
- The follow-up period was 12 months of continuous medical and pharmacy benefit post-index date.
- Baseline characteristics, ASMs used in the 12 months before starting brivaracetam, and epilepsy-related HRU and costs in the 12 months pre- and post-brivaracetam initiation were assessed.

Results

PATIENT DISPOSITION AND BASELINE DEMOGRAPHICS

- A total of 479 commercially insured patients treated with brivaracetam were identified.
- During the baseline period, 67.4% of patients received at least two ASMs.

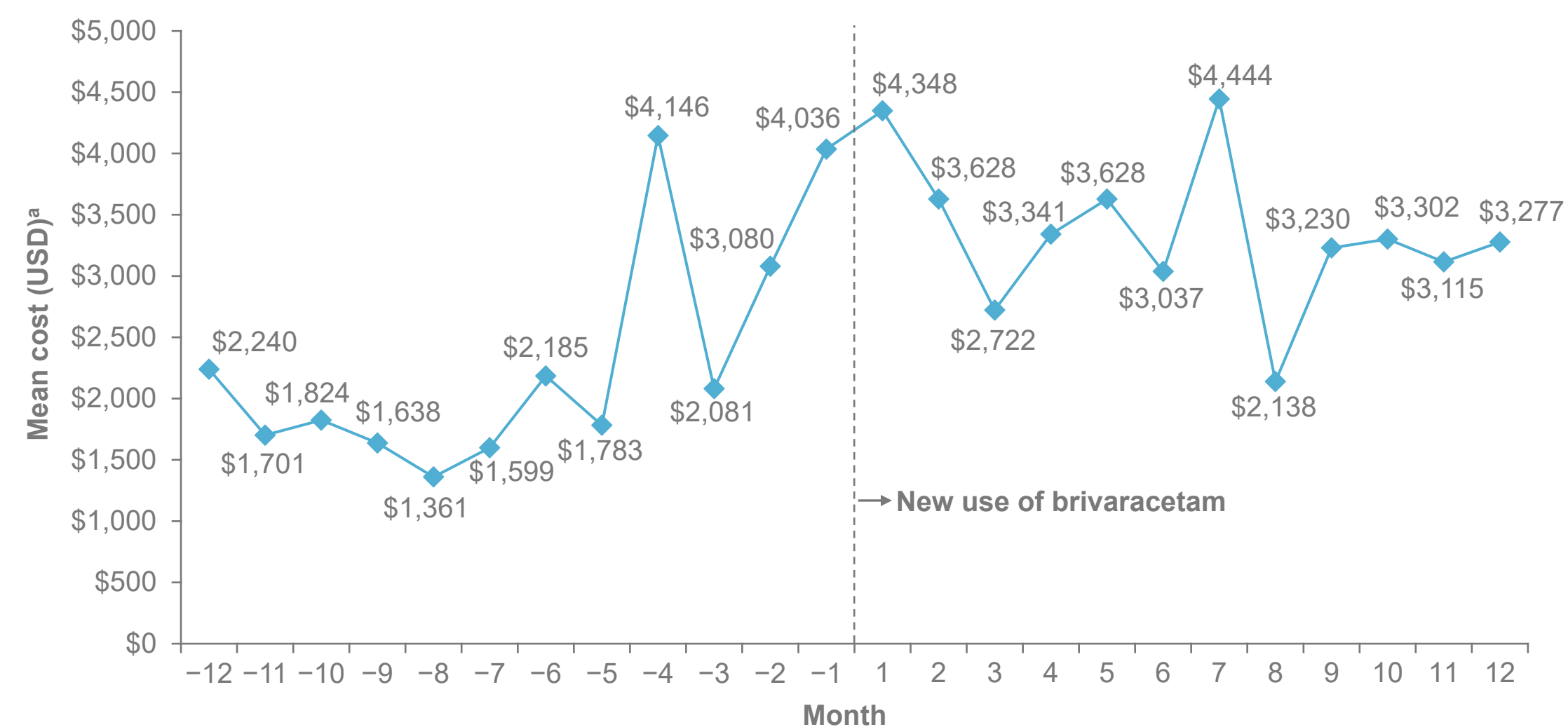
Baseline demographics and epilepsy characteristics

	All patients (N=479)
Age, mean (SD), years	37.5 (13.8)
Female, n (%)	288 (60.1)
ESCI score, mean (SD)	1.2 (1.8)
Seizure type at index, n (%)	
Focal	175 (36.5)
Generalized	45 (9.4)
Unspecified	71 (14.8)
Seizure/convulsion	76 (15.9)
Other	29 (6.1)
Mixed type	83 (17.3)
Most common comorbid conditions (≥15% of patients), n (%)	
Anxiety	152 (31.7)
Hypertension	121 (25.3)
Headache	119 (24.8)
Depression	118 (24.6)
Hyperlipidemia	108 (22.5)
Cardiac arrhythmias	81 (16.9)
Number of ASMs used during the baseline period^a, n (%)	
0	20 (4.2)
1	136 (28.4)
2	159 (33.2)
3	110 (23.0)
≥4	54 (11.3)

^aASMs used at any time during 12 months before initiation of brivaracetam treatment. ASM, antiseizure medication; ESCI, St Germaine-Smith epilepsy-specific comorbidity index.²

EPILEPSY-RELATED COSTS AND RESOURCE UTILIZATION

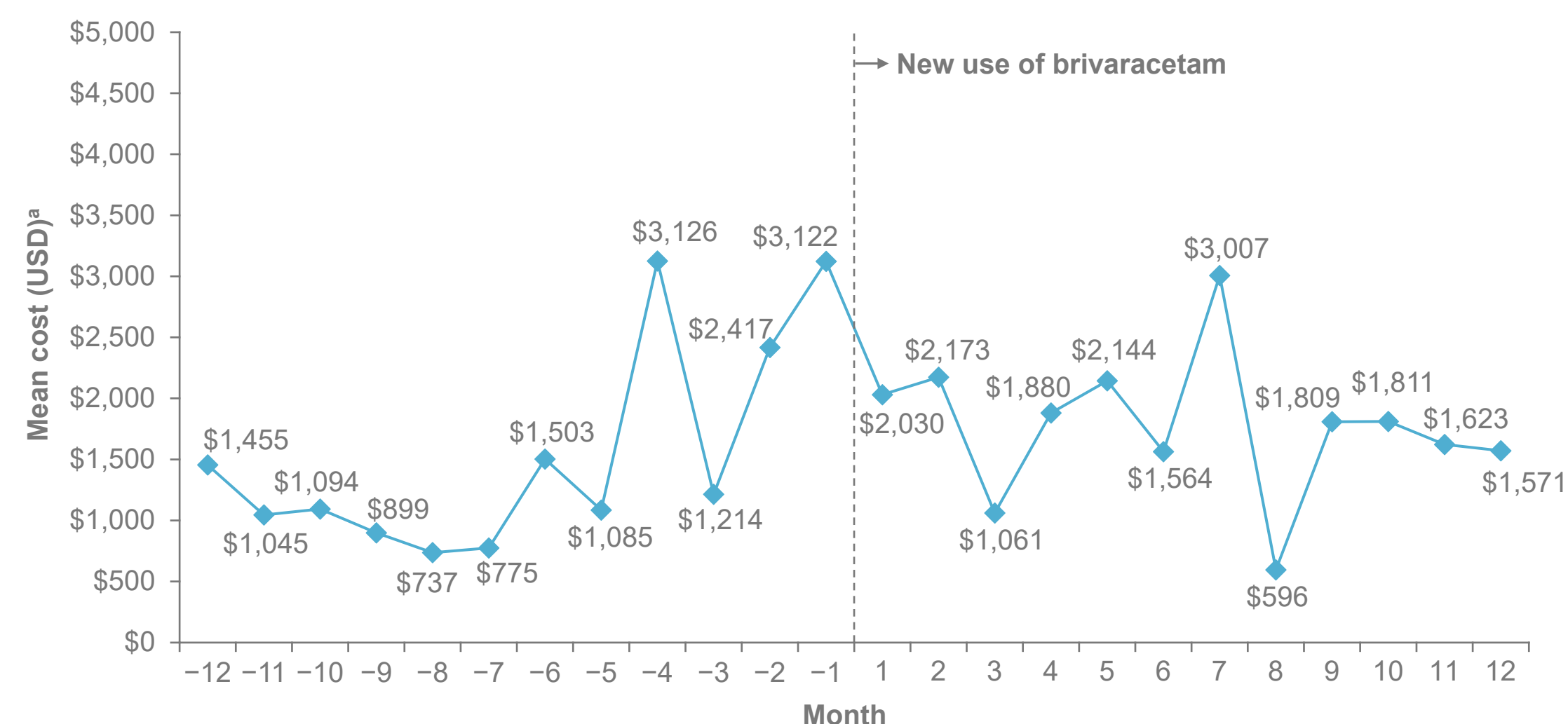
Mean monthly epilepsy-related costs



^aHealthcare costs were expressed in 2019 constant USD, adjusted using the medical care components of the consumer price index.

- Mean total epilepsy-related costs increased 45% during the 12-month follow-up period vs the 12-month baseline period (\$40,212 vs \$27,671, respectively), which was mainly driven by pharmacy costs.

Mean monthly epilepsy-related medical costs

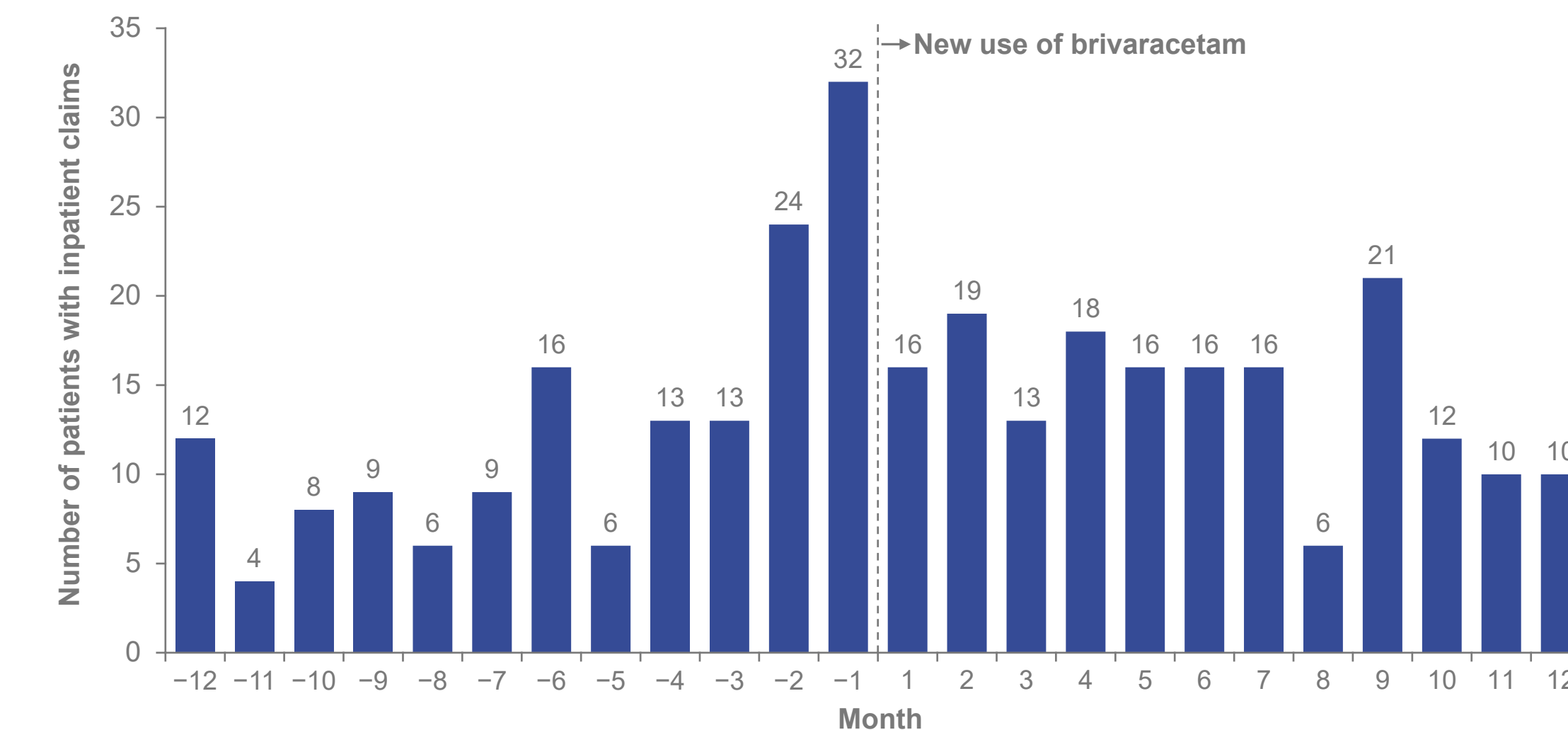


^aHealthcare costs were expressed in 2019 constant USD, adjusted using the medical care components of the consumer price index.

- Within 1 month post-initiation of brivaracetam treatment, mean total epilepsy-related medical costs decreased 35% vs the month pre-initiation of brivaracetam treatment (\$2,030 vs \$3,122).
- Mean total epilepsy-related medical costs increased 15% during the 12-month follow-up period vs the 12-month baseline period (\$21,269 vs \$18,472, respectively).
- Mean monthly epilepsy-related medical costs were lower during the 12-month follow-up period vs 6 months before initiation of brivaracetam treatment (\$1,772 vs \$2,078, respectively).

UCB Pharma-sponsored. UCB Pharma was involved in the design of the study, analysis and interpretation of data, and review of the poster. The authors acknowledge Kathleen Richards, PhD (UCB Pharma, Smyrna, GA, USA) for managing the development of the poster, and Emma Budd, PhD (Evidence Scientific Solutions, Horsham, UK) for writing assistance, which was funded by UCB Pharma. Author disclosures: S Beaty, E Lee, K Guntrum, and A Loewendorf are employees of UCB Pharma. A Estrin and M Skornicki are employees of Aetion, Inc.

Monthly epilepsy-related inpatient healthcare resource use



- From 6 months to 1 month pre-initiation of brivaracetam treatment, the number of patients with epilepsy-related inpatient or outpatient HRU increased (16 to 32 and 158 to 363, respectively), suggesting uncontrolled epilepsy.
- Within 1 month of initiation of brivaracetam treatment, there was a 50% (32 vs 16) and 37.5% (363 vs 227) reduction in the proportion of patients with evidence of an epilepsy-related inpatient or outpatient event, respectively, compared with the month before initiation of brivaracetam treatment, although utilization varied in later months post-initiation of brivaracetam treatment.

Limitations

- This study was not designed to detect statistically significant differences pre- and post-initiation of brivaracetam treatment.
- Findings from this study relied on the accuracy of diagnosis, medication, and procedure codes contained in the claims data.
- As with all claims-based analyses, study results may not be generalizable to the overall population or patients without health insurance.
- Patients were required to meet enrollment criteria, and results from this study may not be generalizable to patients with shorter coverage.
- Eligible patients remained in the cohort throughout this analysis regardless of any switches or discontinuation.
- This analysis did not consider the duration of epilepsy, or severity of disease, which impact HRU and cost outcomes. Patient epilepsy characteristics may have changed between the 12-month baseline period and the 12-month follow-up period.

Conclusions

- In a commercially insured seizure/epilepsy patient population newly starting on brivaracetam treatment, annual epilepsy-related costs increased but epilepsy-related HRU and medical costs were reduced in the period immediately post-initiation of brivaracetam treatment.
- To our knowledge, this was the first retrospective study of detailed payer-specific costs incurred in the periods pre- and post-initiation of brivaracetam treatment.

References

1. Briviact® (brivaracetam) US Prescribing Information. UCB Inc. 2021. <https://www.briviact.com/briviact-PI.pdf> Accessed March 21, 2022.
2. St Germaine-Smith C, et al. *Epilepsia* 2011;52(12):2161-2167.

For a copy of this poster, use your smartphone to scan the QR code, download from the website below, or contact UCBCares®

Website: [UCBposters.com/ISPOR2022](https://ucbposters.com/ISPOR2022); Poster ID: EE86
Phone: 844-599-CARE (2273)
Email: ucbcares@ucb.com

