

Clinical Outcomes of Migraine Patients Receiving Calcitonin Gene-Related Peptide Antagonists from an Integrated Health System Specialty Pharmacy

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BACKGROUND

- Migraine is a highly prevalent neurological disease affecting over 1 billion people worldwide. The Calcitonin Gene-Related Peptide (CGRP) antagonists block CGRP, a neuropeptide implicated in migraines. Clinical trials demonstrate that CGRP antagonists robustly reduce monthly migraine days by 33 to 43% compared to placebo in episodic and chronic migraine.
- Limited data exist to demonstrate the real-world clinical outcomes of migraine patients on CGRPs managed within integrated health system specialty pharmacies (HSSPs).
- The purpose of this analysis is to evaluate the percent reduction in patient-reported monthly migraine days of patients receiving CGRPs from an integrated health system specialty pharmacy (HSSP).

METHODS

Study Design: Retrospective observational analysis of adult patients with episodic or chronic migraine receiving a CGRP from NYU Langone Health Specialty Pharmacy between January 21, 2023 and December 31, 2023.

- Inclusion Criteria: Patients enrolled in the patient management service for ≥ 4 months with both a baseline and follow up number of monthly migraine days.
- Exclusion Criteria : Patients receiving a CGRP with ICD-10 codes unrelated to episodic migraine, chronic migraine, migraine with aura, and migraine without aura

Primary Outcome: Reduction in patient reported monthly migraine days from baseline

Data identification: Data were collected through the electronic medical record or specialty pharmacy management system. Treatment duration was defined as number of days between the baseline and follow up assessment; medication adherence was measured by proportion of days covered (PDC).

Analysis: An ordinal logistic regression model was utilized to evaluate the impact of various patient factors on the change in month migraine days (improved, no change, declined).

RESULTS

Age (years)¹

Sex (n. %)

Medication (n, %) Aimovig (erenumab)

Ajovy (fremanezumab)

Insurance Type (n, %)

Government

Commerical

Diagnosis Chronic Migraine

Days¹

(OOPC)1

PDC¹

Unknown/Other

Episodic Migraine

Other Migraine

Baseline Migraine

Treatment Duration¹

Medication Status

New to medication

Out-of-Pocket Cost

Treatment experienced

Emgality (galcanezumab)

Μ

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Within the study period, 643 patients meeting the inclusion criteria were identified for analysis. **Table 1** outlines the baseline patient characteristics and mean PDC of the cohort. An average reduction in monthly migraine days of 35.9% was observed in the entire cohort. Patients new to the CGRP medication demonstrated a 44.7% reduction in monthly migraine days (baseline 14.8 days) compared to a 21.3% reduction in existing to medication patients (baseline 8.9 days). A higher number of baseline migraine days and longer treatment duration were associated with a reduction in number of migraine days. Patients with episodic migraines were more likely to experience a reduction in migraine days compared to patients with chronic migraines.

Table 1: Patient Characteristics

p-value

0.240

0.230

0.930

0.460

0.620

0.040

0.050

< 0.001

0.020

< 0.001

0.260

0.690

¹Mean ²Medi

45

112 (17%)

531 (83%)

208 (32%)

316 (49%)

119 (19%)

157 (24.4%)

407 (63.3%)

79 (12.3%)

232 (36.1%)

315 (49.0%)

96 (14.9%)

11.86

144.2

321 (49.9%)

322 (50.1%)

\$9.60

Clinical Outcomes

94.6%

Figure 1: Change in Average Monthly Migraine Days

OR ³	All Patients	Total		-4.26			
1.503	Age	Age ≤ 65		-4.12			
		Age > 65		-5.61			
- 0.739	Sex	M F		-4.82 -4.14	•		
-	Medication	Aimovig (erenumab) Ajovy (fremanezumab) Emgality (galcanezumab)		-3.70 -4.02 -4.72	•		
0.974 1.177	Insurance Type	Government Commercial Unknown		-4.37 -4.06 -5.06	•		
- 1.116	Diagnosis	Chronic Episodic Other		-3.98 -4.20 -5.14	•		
- 1.498 1.744	Baseline Migraine Days	1-5 6-10 11-15 16-20 21-25 26-30	4.06	-2.46	-9.73	-12.90	
1.061		1-40 41-80		-2.70 -4.46			
1.002	Treatment Duration (Days)	81-120 121-160		-4.57			
3.001		161-200 201-240 New to Medication		-4.56			
-	Medication Status	Treatment Experienced		-1.90			
1.266	OOPC	Copay = 0 Copay > 0		-4.33			
	PDC	PDC ≤ 90% PDC > 90%		-5.43 -4.26	•		
1.062		C	0 2 4	6 8 10 12 A	14 16 18 verage Migraine Da	20 22 24 ays	26 28 30
ian ³ Odds ratio				Baseline	+	Followup	٠

CONCLUSION

- This real-world analysis demonstrated clinically meaningful reductions in monthly migraine days for migraine patients receiving CGRP antagonists managed within an integrated HSSP, consistent with clinical trial efficacy.
- Patients new to CGRP therapy demonstrated a more robust reduction in monthly migraine days compared to medication experienced patients. Higher baseline
 migraine frequency, longer treatment duration, and new user status were associated with greater likelihood of reduction.
- Overall, the findings provide real-world evidence supporting CGRP antagonists' effectiveness when managed within a HSSP care model.