

Real-World Healthcare Resource Utilization and Costs Associated With Elranatamab Initiation in Multiple Myeloma: The ALTITUDE-1 Study

Objectives



To determine the changes in healthcare resource utilization (HCRU) and costs of patients with RRMM who initiate elranatamab

Conclusions



- Among the elranatamab-treated patient population with RRMM included in the index period:
 - Nearly half of the patients were penta-drug exposed
 - History of infection and peripheral neuropathy was common
- In patients with RRMM, HCRU and costs per-patient-per-month (PPPM) evolved after initiating elranatamab
 - All-cause inpatient visits increased while all-cause outpatient visits decreased
 - All-cause outpatient costs remained stable while inpatient and pharmacy costs decreased
 - Overall, all-cause medical and pharmacy costs remained stable
- Results presented here reflect preliminary data of patients receiving elranatamab, which may change with longer follow-up



Electronic Poster

Please scan this Quick Response (QR) code with your smartphone app to view this poster. If you do not have a smartphone, access the poster via the internet at: <https://scientificpubs.congressposter.com/p/239fq4qqlzmgc89j>
Copies of this poster obtained through QR Code are for personal use only and may not be reproduced without permission from the author of this poster.

References: 1. Elrexfio (elranatamab-bcmm). Prescribing information. Pfizer; 2023. 2. Costa LJ, et al. Blood 2024;144:2401-2402.

Acknowledgments: The study was sponsored by Pfizer. Medical writing and/or editorial services provided by Robyn Roth, PhD, from Nucleus Global were funded by Pfizer.

Disclosures: **RB:** reports consultancy for Adaptive Biotech, BMS, Caribou Biosciences, Genentech, Janssen, Karyopharm, Legend Biotech, Pfizer, and SparkCures; research for Novartis and Peak Health. **MM:** reports advisory board or consulting fees from Sanofi SA, BMS, Celgene, Pfizer, Janssen, and Legend Biotech; research funding from Sanofi SA, BMS, and Celgene. **BS:** reports advisory board or consulting fees from Pfizer. **PP:** reports no conflicts. **NG:** reports stock/equity ownership in Aetion, Inc. **BC and EB:** report consulting fees from Moderna. **DH, GN, PH, RS, BL, AM, CHK, IPC, MS and MD:** report current employment and stock ownership at Pfizer Inc.

Contact: Rahul Banerjee, rahulban@uw.edu

Copyright © 2025

Rahul Banerjee,¹ Meera Mohan,² Bhavesh Shah,³ Patricia Prince,⁴ Nileesa Gautam,⁴ Brian Conroy,⁴ Elisha Beebe,⁴ David Hughes,⁵ Guido Nador,⁶ Patrick Hlavacek,⁷ Rickard Sandin,⁸ Benjamin Li,⁷ Aster Meche,⁷ Chai Hyun Kim,⁷ Isabel Perez Cruz,⁷ Mohsena Sumaya,⁷ Marco DiBonaventura⁷

Background

- Elranatamab-bcmm is a bispecific antibody approved in the US for the treatment of adult patients with relapsed/refractory multiple myeloma (RRMM) who have been treated with ≥1 proteasome inhibitor, ≥1 immunomodulatory agent, and ≥1 anti-CD38 monoclonal antibody¹
- Prior studies have shown that elranatamab is associated with improved clinical outcomes compared with standard regimens in real-world practice² but its impact on HCRU and costs are unclear

Results

PATIENTS AND TREATMENT

- As of November 26, 2024, 59 patients treated with elranatamab were included in the study (**Table 1**)
 - Almost half (45.8%) were penta-drug exposed and 25.4% had received prior commercial BCMA-directed therapy (chimeric antigen receptor T cell and/or antibody drug conjugate therapies)
 - Common (≥25%) relevant disease history included infection (91.5%), hypertension (91.5%), and peripheral neuropathy (81.4%)
 - Common (≥25%) comorbidities included renal disease (30.5%), congestive heart failure (27.1%), and metastatic solid tumor (25.4%)
- Median follow up was 4.7 months (IQR 2.5-7.0)

Table 1. Baseline and treatment characteristics	
	N=59
Age, mean (SD), years	70.5 (10.4)
Sex, n (%)	
Female, n (%)	28 (47.5)
Male, n (%)	26 (44.1)
Unknown/Missing, n (%)	5 (8.5)
Race, n (%)	
African American	14 (23.7)
Asian/Pacific Islander	1 (1.7)
Caucasian	32 (54.2)
Hispanic/Latino	5 (8.5)
Other	3 (5.1)
Unknown/Missing	4 (6.8)
Region on index date ^a , n (%)	
South	23 (39.0)
Northeast	19 (32.2)
West	11 (18.6)
Midwest	6 (10.2)
Time from initial MM diagnosis to index date, median (IQR), months	77.5 (50.2-97.1)
Prior treatment history ^b , n (%)	
Penta-drug exposed ^c	27 (45.8)
BCMA-directed therapy	15 (25.4)
CAR-T	9 (15.3)
Talquetamab	2 (3.4)
Relevant disease history ^b , n (%)	
Any infection	54 (91.5)
Hypertension	54 (91.5)
Peripheral neuropathy	48 (81.4)
Neutropenia	40 (67.8)
Use of IV anti-infective ^d	38 (64.4)
Hypercalcemia	20 (33.9)
Amyloidosis	9 (15.3)
Extramedullary disease	9 (15.3)
CCI score ^e , mean (SD)	3.2 (3.1)
Categorical CCI score ^e , n (%)	
0 (no comorbidities)	15 (25.4)
1 to 2 (mild)	16 (27.1)
3 to 4 (moderate)	10 (16.9)
≥5 (severe)	18 (30.5)
CCI comorbidities (≥5%) ^e , n (%)	
Renal disease	18 (30.5)
Congestive heart failure	16 (27.1)
Metastatic solid tumor	15 (25.4)
Diabetes	14 (23.7)
Chronic pulmonary disease	12 (20.3)
Cerebrovascular disease	9 (15.3)
Perivascular disease	8 (13.6)
Myocardial infarction	3 (5.1)

BCMA, B cell maturation agent; CAR-T, chimeric antigen receptor T cell therapy; CCI, Charlson Comorbidity Index; IQR, interquartile range
^aAssessed from any time prior to index date to index date; ^bAssessed from the initial MM diagnosis date to one day prior to index date; ^cExposed to 2 unique proteasome inhibitors, 2 unique immunomodulatory agents, and CD38 monoclonal antibodies; ^dAssessed from 14 days prior to index date to index date; ^eAssessed from 180 days prior to index date to index date

¹Fred Hutchinson Cancer Center, Seattle, WA, USA; ²Medical College of Wisconsin, Milwaukee, WI, USA; ³Boston Medical Center, Boston, MA, USA; ⁴Aetion, New York, NY, USA; ⁵Pfizer Inc, Cambridge, MA, USA; ⁶Pfizer Ltd, Surrey, UK; ⁷Pfizer Inc, New York, NY, USA; ⁸Pfizer AB, Stockholm, Sweden

Methods

- ALTITUDE-1 (EUPAS1000000229) is an ongoing, non-interventional database study on RW treatment patterns, HCRU, and costs (among other outcomes) of patients with RRMM treated with elranatamab
 - The study analyzed de-identified data from the Komodo US claims dataset
 - The study cohort included patients with RRMM who initiated elranatamab between August 14, 2023 (US approval date) and November 26, 2024

HEALTHCARE RESOURCE UTILIZATION

- The median number of all-cause inpatient visits increased from pre-index (0.67 PPPM; IQR 0.25-1.75) to post-index (1.14 PPPM; IQR 0.28-3.46) (**Figure 1**)
 - The number of patients with an inpatient visit was 39 (66.1%) pre-index and 34 (57.6%) post-index
- All-cause outpatient visits decreased from pre-index (2.83 PPPM; IQR 1.83-4.00) to post-index (2.33 PPPM; IQR 1.67-3.96)
 - The number of patients with an outpatient visit was 59 (100%) pre-index and 57 (96.6%) post-index
- The median number of all-cause emergency room visits were 0 PPPM [IQR 0.00-0.00] pre- and post-index
 - The number of patients with an emergency room visit was 12 (20.3%) pre-index and 8 (13.6%) post-index

ALL-CAUSE PATIENT COSTS

- Total median all-cause medical and pharmacy costs remained stable from pre-index (\$15,323 PPPM; IQR \$5800-\$33,846) to post-index (\$14,722; IQR \$5658-\$40,242) (**Figure 2**)
 - Median all-cause inpatient costs numerically decreased from \$1892 PPPM (IQR \$28-\$4395) pre-index to \$1071 PPPM (IQR \$32-\$5429) post-index
 - Median all-cause outpatient costs also remained stable at \$6477 PPPM (IQR \$2770-\$16,064) pre-index and \$6353 PPPM (IQR \$1,505-\$27,599) post-index
 - Median all-cause pharmacy costs also decreased from \$584 (IQR \$89-\$7767) pre-index to \$201 (IQR \$43-\$607) post-index
 - Median all-cause emergency department costs were \$0 pre- and post-index

- The index date was the date of the first prescription or medical claim for elranatamab
- Patients were required to be ≥18 years on the index date with ≥180 days of continuous closed-claims medical and pharmacy enrollment prior to index, with no prior anti-BCMA directed BsAbs treatment
- This interim analysis reported the changes in HCRU and costs from pre-index (180 days prior) to post-index (while treated with elranatamab) on a per-patient-per-month (PPPM) basis (ie, HCRU and costs were rescaled to reflect a monthly average for each patient)
- Results were reported descriptively with a focus on median values

Figure 1. Median all-cause inpatient and outpatient visits pre- and post-index date among patients receiving elranatamab

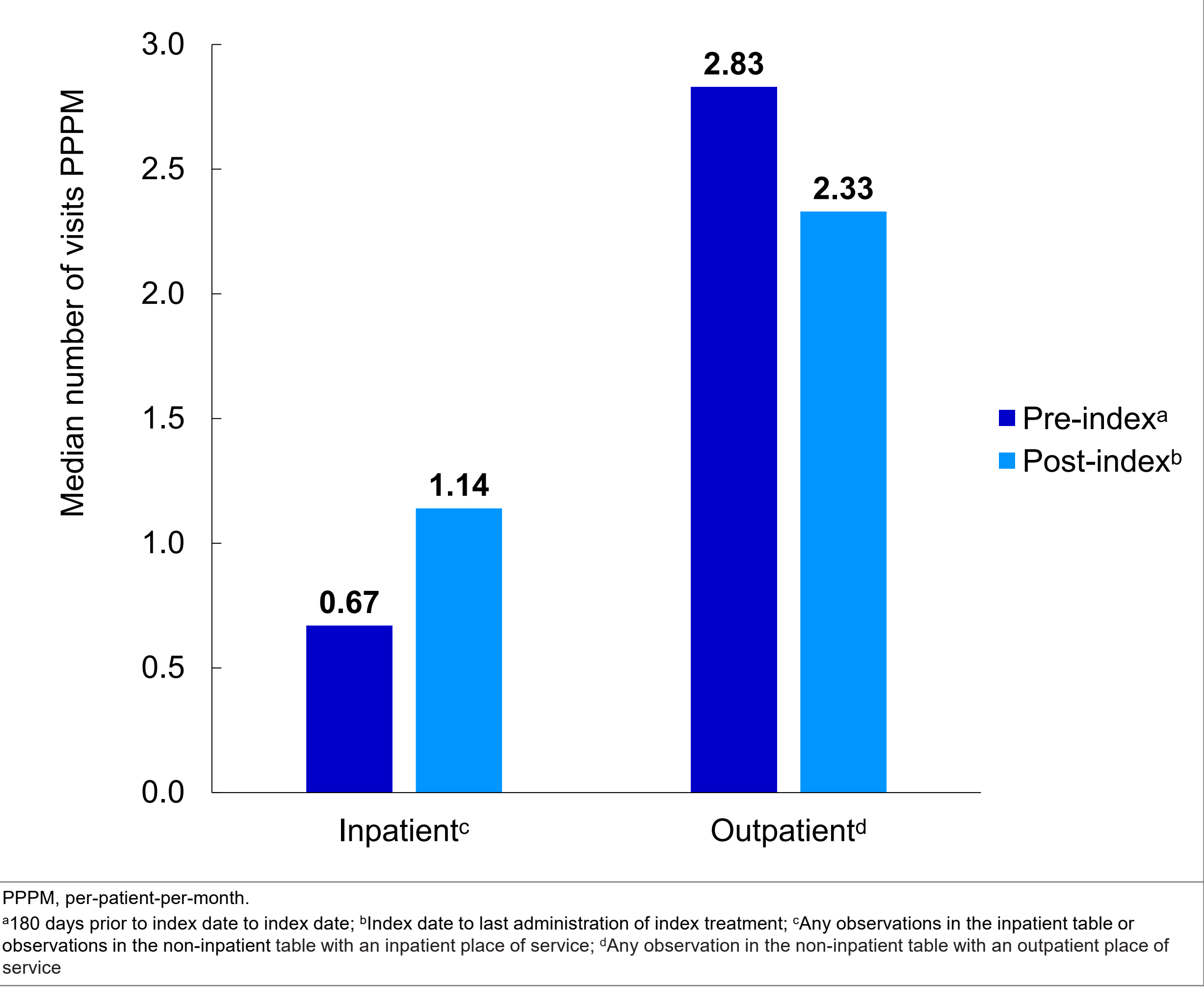


Figure 2. Median all-cause patient costs (PPPM) pre- and post-index date among patients receiving elranatamab^{a,b}

