

A cost-of-illness analysis of relapsed or refractory acute myeloid leukemia (R/R AML) with *FLT3* mutations in Chile: a public payer perspective

Juan Guillermo Ariza, MD, MBA, MS¹; Kenny Arciniegas, MD, CPS, MS²; Giovanni Montoya, MS³; Joel Castellano, MBA³; Camilo Tamayo, MS⁴; Alexandra Gonzalez, MS, MD⁴

¹Astellas Pharma International Markets, Bogotá, Colombia; ²Astellas Farma South Cone - International Markets, Bogotá, Colombia; ³Astellas Farma Colombia S.A.S., Bogotá, Colombia; ³IQVIA LatAM, Santiago, Chile; ⁴IQVIA LatAM, Bogotá, Colombia

INTRODUCTION AND OBJECTIVE

- Most newly diagnosed patients with acute myeloid leukemia (AML) will relapse or develop refractory (R/R) disease, despite complete response rates as high as 80% to first-line chemotherapy and targeted therapies¹
- Patients with R/R *FMS-like tyrosine kinase 3* mutation-positive (*FLT3*^{mut+}) AML have particularly poor clinical outcomes with standard salvage chemotherapy, characterized by low response rates and dismal survival²
- A substantial proportion (40%) of patients with R/R *FLT3*^{mut+} AML receive best supportive care services, incurring a heavy burden on the healthcare system^{3,4}
- We estimated the direct costs associated with the treatment of patients with R/R *FLT3*^{mut+} AML in the Chilean public health system

METHODS

- The target population was estimated, using cost-of-illness analysis model, based on the Chilean population in 2023⁵
 - AML prevalence was derived from the International Agency for Research on Cancer database 2020⁶
 - Proportion of patients with AML who received *FLT3* testing and proportion of patients with R/R *FLT3*^{mut+} AML were estimated from published literature^{7,8}
- Health resource utilization, frequency of usage, and quantities were estimated from literature review, clinical practice guidelines, and semi-structured interviews with five Chilean clinical hematology experts
- Cost inputs were based on the CENABAST public purchasing system, public hospital, and Chilean diagnosis-related group fees for relevant clinical events⁹
 - Costs were adjusted by inflation and converted to United States Dollars (USD)
 - The average November 2023 exchange rate was used, with 1 USD = 886.61 Chilean Peso
- Costs were calculated per patient per year for 2023 by using an “incident” approach and estimated for the overall target population

CONCLUSIONS

- In the Chilean public health system, this cost-of-illness analysis demonstrated the substantial healthcare resource utilization and economic burden among patients with R/R *FLT3*^{mut+} AML
- Treatment strategies that reduce drug administration and hospitalization costs among patients with R/R *FLT3*^{mut+} AML may alleviate some of the economic burden associated with AML

RESULTS

Estimation of target population

- A cohort of 15,492,606 (77.6%) adults (≥ 18 years) was estimated from the overall Chilean population⁵ (Table 1)

Treatment distribution

- According to Chilean expert opinion, the most frequently used treatments in the target population were off-label venetoclax combination regimens (37%), followed by low-intensity chemotherapy (29%), targeted therapies (22%), and high-intensity chemotherapy (12%)

Cost of treatment

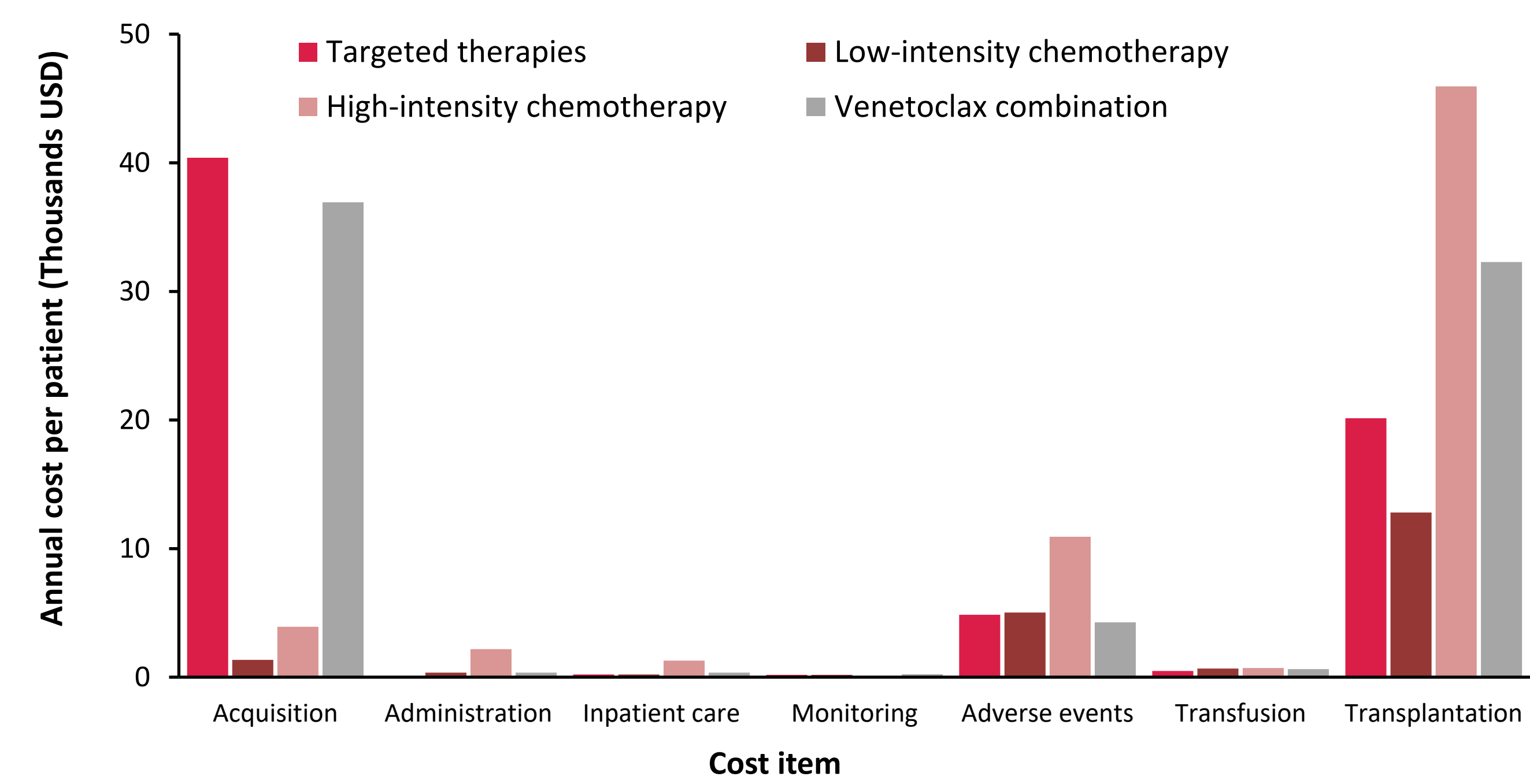
- Treatment-associated costs per patient varied across the four treatment categories analysed (Figure 1)
 - The costs associated with targeted therapies and venetoclax combination treatment were highest for drug acquisition, followed by hematopoietic stem cell transplantation (HSCT); whereas the costs associated with chemotherapy were highest for transplantation, followed by adverse events costs
 - Critical care hospitalization was the main driver for the post-progression AML care costs (Figure 2)
- The estimated annual per capita cost for a new patient with R/R *FLT3*^{mut+} AML was USD 67,761.07 with an overall cost for the target population of USD 7,724,761.70 (Table 2)

Table 1. Estimation of target population

Eligible patients	Input value	Population flow
Chilean population in the year 2023	19,960,889	
Adults ≥18 years	77.6%	15,492,606
Adults with AML	0.0205%	3,175
Adult patients with AML receiving <i>FLT3</i> testing	45.0%	1,429
Adult patients with <i>FLT3</i> ^{mut+} AML	20.0%	286
Adult patients with R/R <i>FLT3</i> ^{mut+} AML	40.0%	114

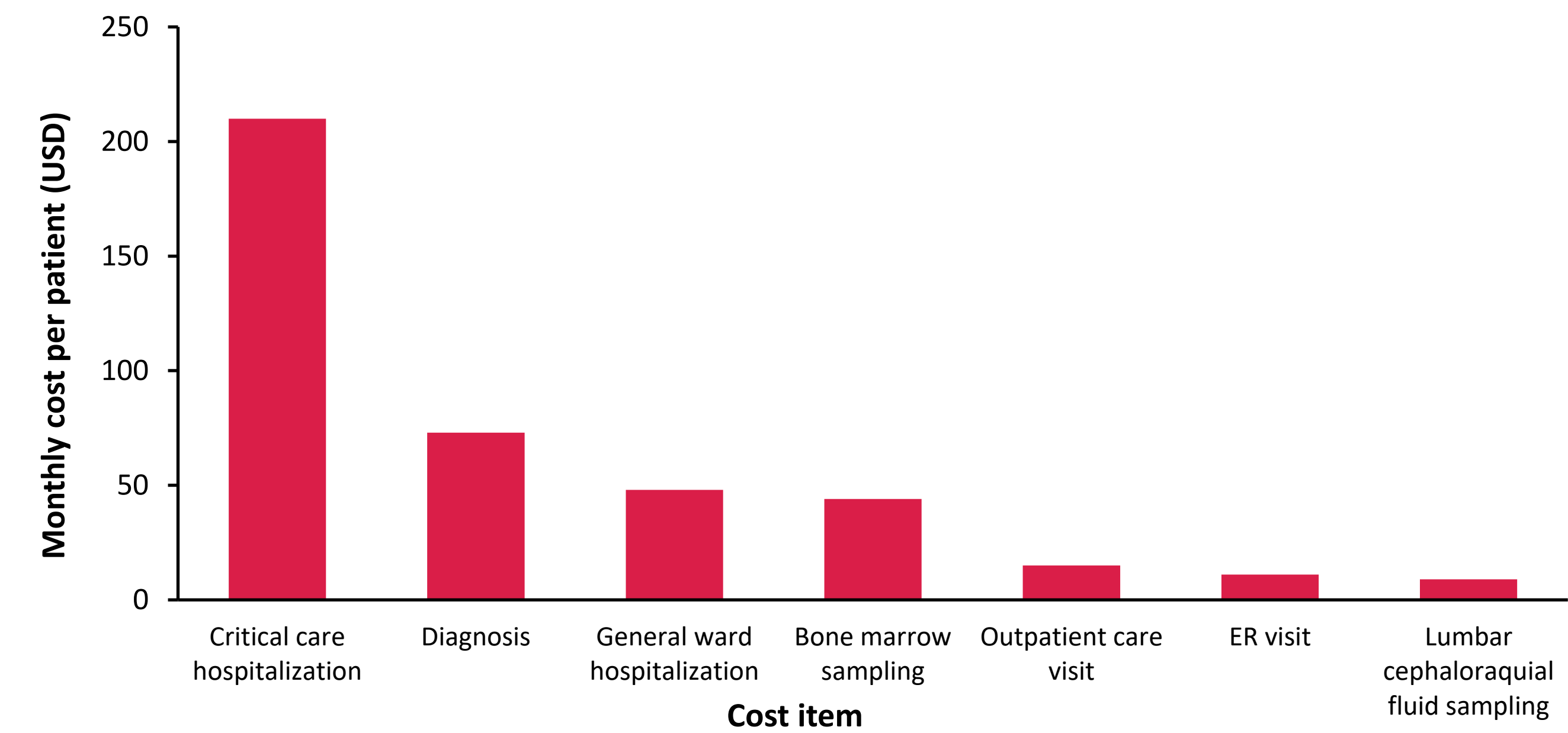
AML, acute myeloid leukemia; *FLT3*^{mut+}, *FMS-like tyrosine kinase 3* mutation-positive; R/R, relapse or refractory

Figure 1. Treatment-associated costs per patient with R/R *FLT3*^{mut+} AML by treatment category



AML, acute myeloid leukemia; *FLT3*^{mut+}, *FMS-like tyrosine kinase 3* mutation-positive; R/R, relapse or refractory

Figure 2. Post-progression monthly costs of treating patients with R/R *FLT3*^{mut+} AML



AML, acute myeloid leukemia; *FLT3*^{mut+}, *FMS-like tyrosine kinase 3* mutation-positive; R/R, relapse or refractory

Table 2. Overall treatment costs of treating patients with R/R *FLT3*^{mut+} AML*

Cost item	Cost per patient per year, USD	Cost for the target population per year, USD (N = 114)	Percentage of overall cost, %
Drug administration and hospitalization	36,238.14	4,131,148.29	53
Monitoring	1,027.90	117,180.40	2
Adverse event	5,007.50	570,855.40	7
Blood and platelet transfusion	634.74	72,360.38	1
Subsequent HSCT	20,079.78	2,289,094.58	30
Post-progression	4,773.01	544,122.65	7
Overall	67,761.07	7,724,761.70	100

*Results in the accepted abstract have been updated here based on revised assumptions for transplantation and some medication costs.
AML, acute myeloid leukemia; *FLT3*^{mut+}, *FMS-like tyrosine kinase 3* mutation-positive; HSCT, hematopoietic stem cell transplantation; R/R, relapse or refractory; USD, United States Dollar

Acknowledgments

This study was funded by Astellas Pharma Global Development Inc. Medical writing support was provided by Oana Coban, PhD, and Eden Godfrey-Shaw, BSc, of Lumanity, funded by Astellas Pharma Global Development Inc.

Author disclosures

JGA, KA, and GM are employees of Astellas Pharma International Markets (Colombia). JC, CT and AG are employees of IQVIA, a health economics and outcomes research vendor contracted to generate data on behalf of Astellas Pharmaceuticals.

References

- Ramos N, et al. *J Clin Med*. 2015;4(4):665-695; 2. Mangan JK, Luger SM. *Ther Adv Hematol*. 2011;2(2):73-82; 3. Griffin JD et al. *Eur J Haematol*. 2019;102(4):341-350; 4. Cannas G et al. *Transfus Clin Biol*. 2015 0;22(5-6):341-7; 5. The National Institute of Statistics. Available at: https://www.inec.gov.cl/docs/default-source/proyecciones-de-poblacion/publicaciones-y-anuarios/base-2017/inec_estimaciones-y-proyecciones-de-poblacion-1992-2050_base-2017_s%C3%ADntesis.pdf?sfvrsn=c623983e_6 Accessed March 2024; 6. The Global Cancer Observatory. Available at: <https://gco.iarc.fr/> Accessed March 2024; 7. Nagel et al. *Ann Hematol*. 2017;96(12):1993-2003; 8. Dombret H, et al. *Blood*. 2015;126(3):291-9; 9. Cenabast. Available at <https://www.cenabast.cl/compras-cenabast/> Accessed March 2024

