

# Treatment patterns in the management of multiple myeloma in a Colombian HMO during 2015-2023: a Real-World Data study

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## BACKGROUND

- Multiple myeloma (MM), characterized by clonal plasma cell proliferation in the bone marrow, is the second most common hematologic malignancy<sup>1</sup>.
- The objectives in managing MM focus on symptomatic relief, eliminating of myeloma cells from the bone marrow, reducing the amount of M protein, improving quality of life, generating the longest possible period of response, and enhancing overall survival rates<sup>2</sup>.
- The main treatments of MM are composed by immunomodulatory drugs (such as thalidomide, lenalidomide, and pomalidomide), proteasome inhibitors (bortezomib, carfilzomib, and ixazomib), monoclonal antibodies (elotuzumab, daratumumab and isatuximab), and bispecific antibodies (tedastamab, talquetamab, and elranatamab) which depend on transplant eligibility<sup>3</sup>.
- MM is still considered an incurable disease. Although there are several therapies that have led to longer remission periods, patients develop relapse and/or become refractory to treatment needing new approaches<sup>4,5</sup>.
- In Colombia, there are no available studies on the treatment patterns used to manage MM.

## OBJECTIVE

- To identify treatment patterns in the management of multiple myeloma (MM) in a Colombian Health Maintenance Organization (HMO) between 2015-2023.

## METHODS

- A retrospective, descriptive cohort study of MM patients from 2015 to 2023 was conducted.
- Patients were identified based on the C90.0 code for MM.
- The index date was when MM was diagnosed, and patients were followed until death, insurance discontinuation, or December 31<sup>st</sup>, 2023, whichever occurred first.
- Demographics, treatment, and medical care were assessed at the index and during follow-up using available databases, including but not limited to: EHR, claims records, laboratory, diagnostic imaging records, and clinical records.
- According to International Myeloma Working Group, refractory myeloma is a disease that is non-response to therapy or progresses within 60 days of the last line, and relapsed myeloma is previously treated myeloma that has progressed after prior therapy and requires new therapy<sup>6</sup>. In this study, a patient that failed the first line of treatment was considered relapsed or refractory MM.
- Patients who achieved strict complete response, complete response, very good partial response or partial response were classified as responders.
- The frequency of treatments was presented by line of treatment as well as the percentage of responders.

## RESULTS

- N=700 patients were included. At diagnosis, approximately 50.5% of these patients presented arterial hypertension, with a median age of 64 years (IQR 16, Q1-Q3: 55-71) and more than half (51.3%) of the patients were at Stage III (A or B) of the Durie-Salmon classification (Table 1).
- At the last contact, 65% were alive, 51.3% had registered a complete response, and 23.1% had progressed.

## RESULTS (cont)

- The most common first-line (1L) treatment was cyclophosphamide/bortezomib/dexamethasone (44.8%), where 85.0% were responders and 13.4% progressed. The second most common treatment was bortezomib/lenalidomide/dexamethasone (16.7%), with 91.4% of these patients responding (Figure 1).
- At 2L (513 patients), autologous transplant was the preferred option (n=229, 44.6%) where majority of patients (94.8%) were responders and 5.2% progressed. The following treatments most frequent were Bortezomib/lenalidomide/dexamethasone treatment scheme (8.6%), bortezomib/dexamethasone (6.2%), and daratumumab/lenalidomide/ dexamethasone (6.0%) were with progression in 18.2%, 12.5%, and 16.13%, respectively.
- At 3L (271 patients), autologous transplant (18.5%), bortezomib/lenalidomide/dexamethasone (12.5%), and daratumumab/bortezomib/dexamethasone (9.9%) were the most common treatment strategies.
- The proportion of patients with later line data was small (4L = 18.4%, 5L = 8%, 6L = 2.7%, and 7L = 0.6%).
- The regimen bortezomib/lenalidomide/dexamethasone was frequently used in 1L, 2L, and 3L. Daratumumab increased its used after 2L.
- 297 patients (42.4%) were treated with autologous transplant, mainly in 2L (77% of patients) (Figure 1).
- Although there was a 66% response rate at 4L, this rate was decreasing with each subsequent line of treatment (Figure 2).
- Bortezomib/cyclophosphamide/dexamethasone (n=170) and bortezomib/lenalidomide/dexamethasone (n=44), and bortezomib/dexamethasone (n=37) were the most common regimens at 1L for autologous transplant.
- 127 patients (18.14%) had triple-class refractory MM. The most frequent treatment after triple refractory were daratumumab/bortezomib/dexamethasone (14.2%), daratumumab/carfilzomib/dexamethasone (12.6%), daratumumab/dexamethasone (11.0%), and daratumumab/lenalidomide/dexamethasone (10.2%) (Table 4).

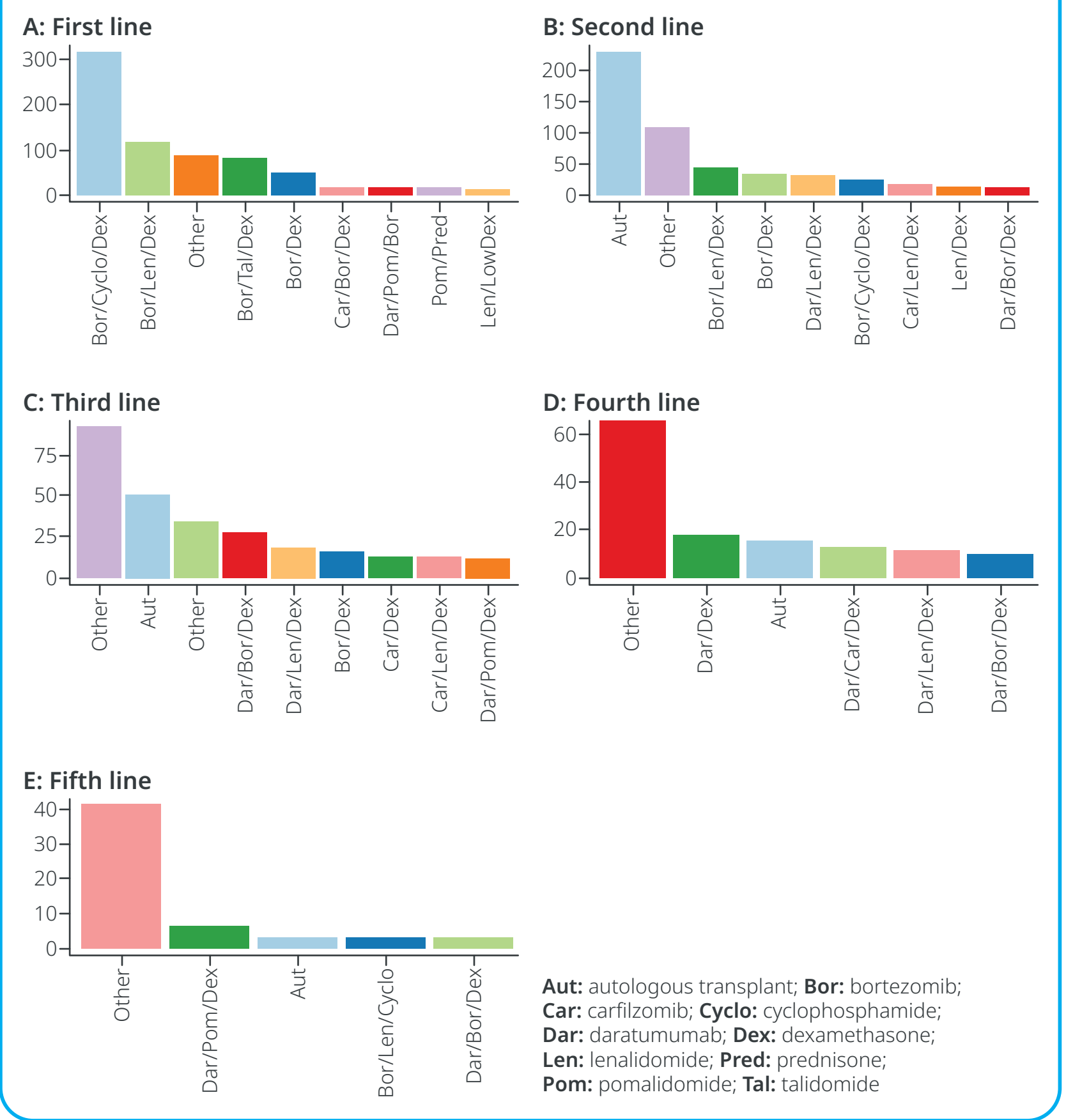
**Table 1.** Clinical and demographic characteristics of patients included in the study

| Characteristics                                 | (n=700, %)  |
|---|-------------|
| Women (%)                                       | 367 (52.4)  |
| Age ( mean (SD))                                | 62.9 (11.3) |
| <b>Raze</b>                                     |             |
| African american                                | 12 (1.7)    |
| White   | 65 (9.3)    |
| Indigenous                                      | 1 (0.1)     |
| Mestizoes                                       | 466 (66.6)  |
| Mulatto   | 1 (0.1)     |
| Raizales  | 1 (0.1)     |
| Zambaigo  | 1 (0.1)     |
| Non information                                 | 153 (21.9)  |
| <b>International Staging System (ISS) Stage</b> |             |
| I   | 98 (14.0)   |
| II  | 257 (36.7)  |
| III   | 345 (49.3)  |
| <b>Durie Salmon Stage</b>                       |             |
| IA  | 78 (11.1)   |
| IB  | 7 (1.0)     |
| IIA   | 193 (27.6)  |
| IIB   | 63 (9.0)    |
| IIIA  | 248 (35.4)  |
| IIIB  | 111 (15.9)  |

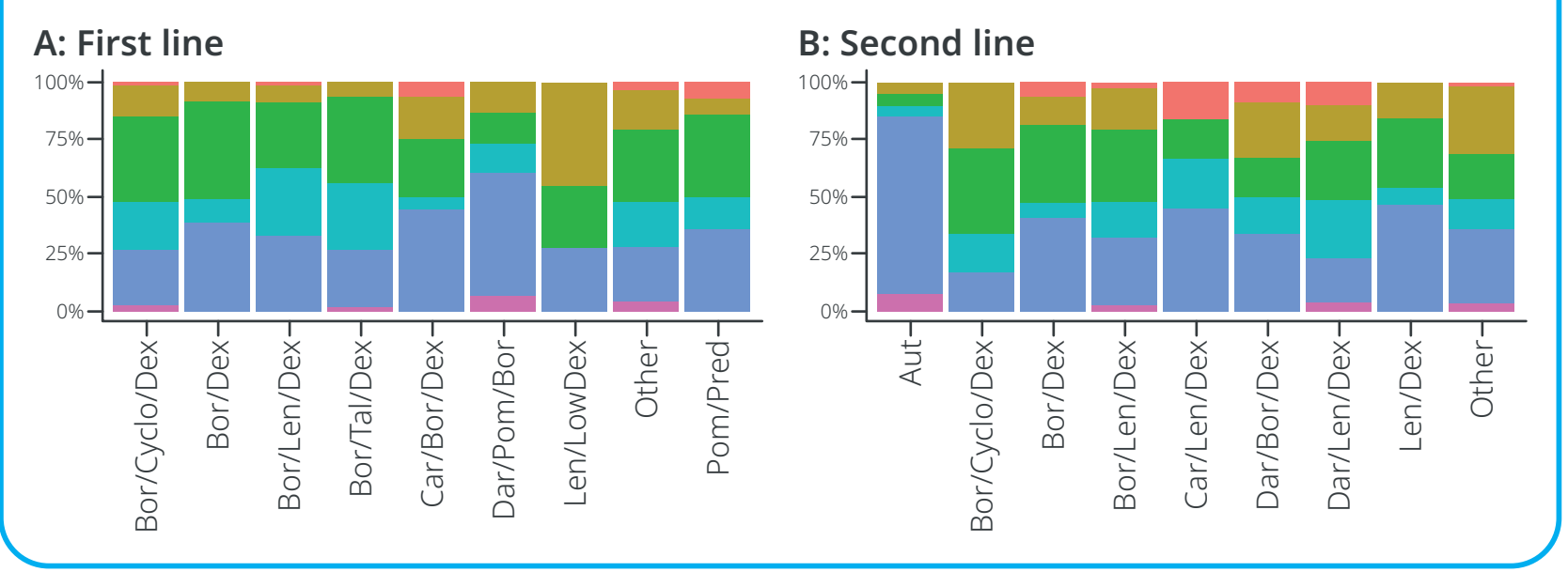
## RESULTS (cont)

|                                   |            |
|-----------------------------------|------------|
| <b>ECOG</b>                       |            |
| 0                                 | 74 (10.6)  |
| 1                                 | 356 (50.9) |
| 2                                 | 210 (30.0) |
| 3                                 | 54 (7.7)   |
| 4                                 | 6 (0.9)    |
| <b>Skeletal-related events</b>    |            |
| Bone fractures                    | 342 (48.8) |
| Hypercalcemia                     | 183 (26.2) |
| Spinal cord compression           | 135 (19.3) |
| Radiation therapy                 | 82 (11.7)  |
| Dialysis at diagnosis             | 66 (9.4)   |
| Plasmapheresis                    | 13 (1.90)  |
| Spinal cord decompression surgery | 48 (6.9)   |

**Figure 1.** Treatment patterns of Colombian patients with MM diagnosed between 2015 to 2023

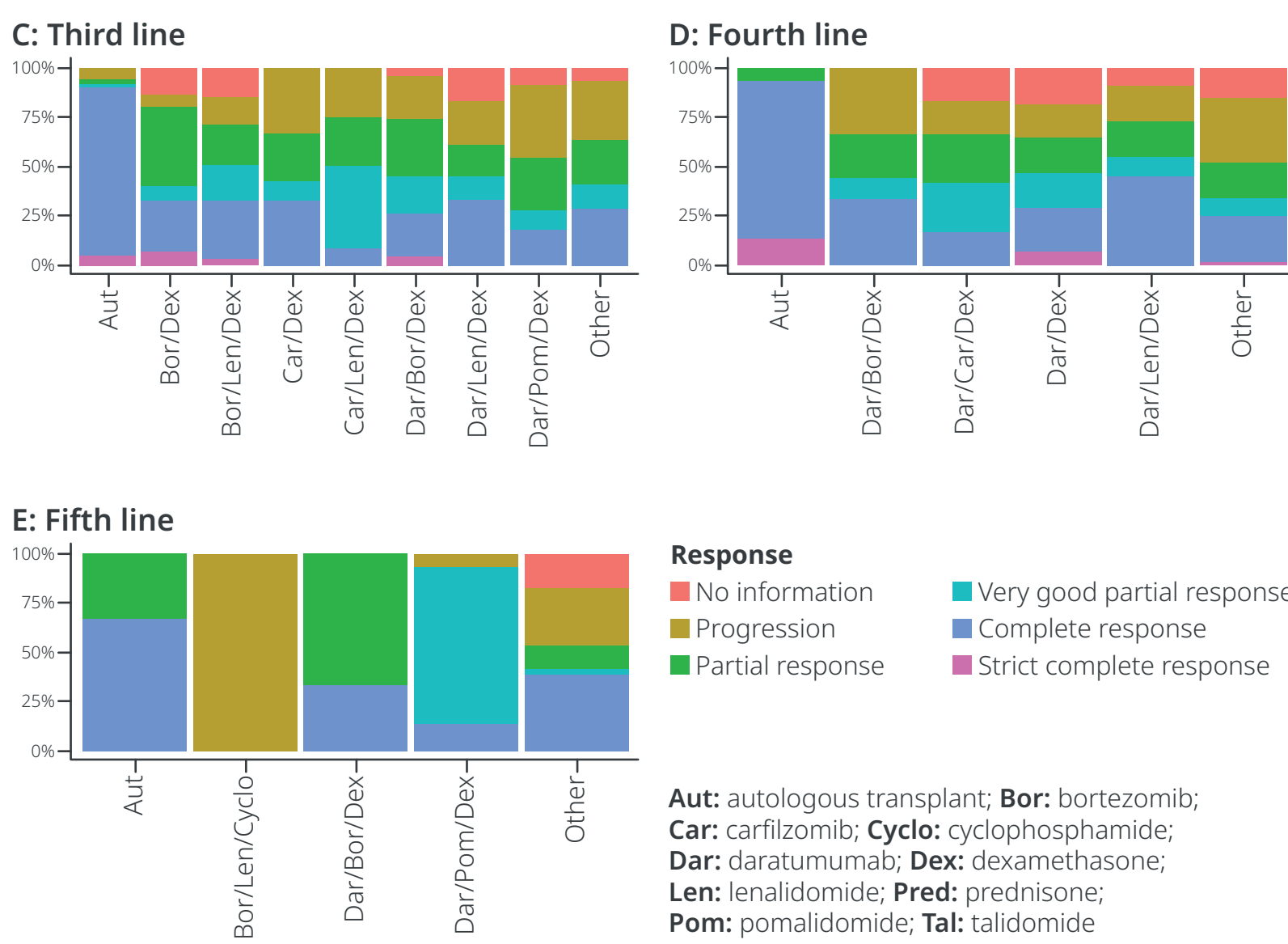


**Figure 2.** Response by treatment Line (per IMWG Criteria) for patients initiating treatment in 2015 to 2023

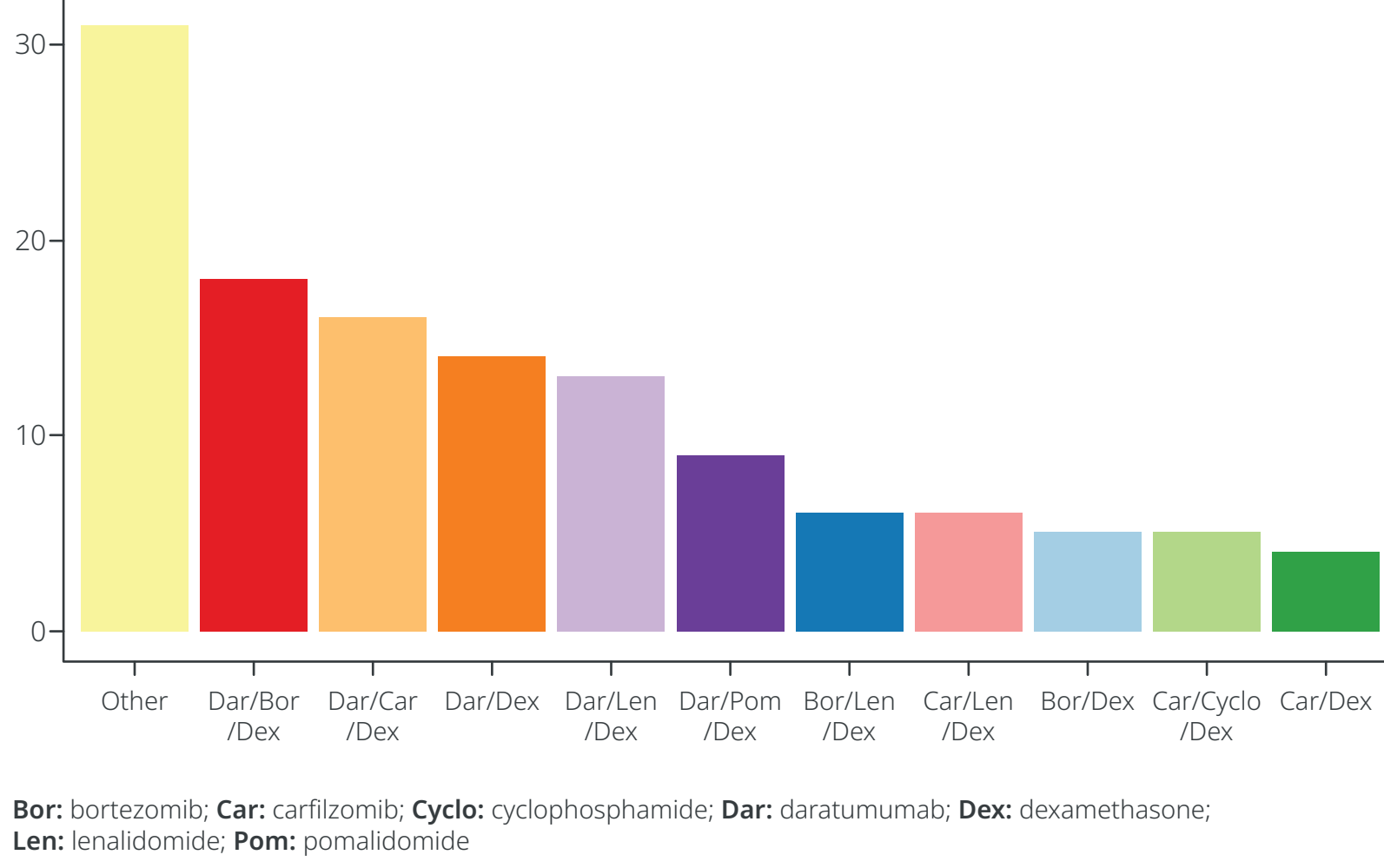


## RESULTS (cont)

**Figure 2. Cont.**



**Figure 3.** Treatment patterns of Colombian patients with triple refractory MM diagnosed between 2015 to 2023



## CONCLUSION

- Patients were in later stages of the disease according to Durie-Salmon, most of them achieved complete response at final contact, and cyclophosphamide/bortezomib/dexamethasone was the preferred treatment for first line and autologous transplant for the second and third lines.

## REFERENCE

- Palumbo A, Anderson K. Multiple myeloma. N Engl J Med. 2011 Mar 17;364(11):1046-60
- Kyle RA, Rajkumar SV. Multiple myeloma. N Engl J Med 2004;351:1860-73
- Cowan AJ, Green DJ, Kwok M, et al. Diagnosis and Management of Multiple Myeloma: A Review. JAMA. 2022;327(5):464-477.
- Sonneveld P, De Wit T, Moreau P (2017) How have evolutions in strategies for the treatment of relapsed/refractory multiple myeloma translated into improved outcomes for patients? Crit Rev Oncol Hematol 112:153–170
- National comprehensive cancer network. Multiple Myeloma. Version 1. 2025. URL: [https://www.nccn.org/professionals/physician\\_gls/pdf/myeloma.pdf](https://www.nccn.org/professionals/physician_gls/pdf/myeloma.pdf). Date access: 09/04/2025
- Rajkumar SV, Dimopoulos MA, Palumbo A, Blade J, Merlini G, Mateos MV, et al. International Myeloma Working Group updated criteria for the diagnosis of multiple myeloma. Lancet Oncol. 2014;15(12):e538-48.