



HEMATOLOGY, TRANSFUSION AND CELL THERAPY

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Letter to the Editor

National multiple myeloma cohort: Gaps and opportunities for research in Brazil

1 Dear Editors,

2 Multiple myeloma (MM) is a relatively uncommon hemato-
3 logic malignancy. It accounts for approximately 1 % of all can-
4 cers and is the second most frequent hematologic
5 malignancy [1] However, national data on the epidemiological
6 profile, access to innovative therapies, and clinical outcomes
7 in Brazil remain scarce [2]

8 The study “Overall survival in multiple myeloma in Brazil:
9 A cohort of 16 years”, published in Hematology, Transfusion
10 and Cell Therapy, provides a comprehensive overview of sur-
11 vival among MM patients in the country. This study evaluated
12 25,370 patients treated within the Brazilian Unified Health
13 System (SUS) between 2000 and 2015, constituting one of the
14 largest national MM cohorts ever published. It allows for
15 important reflections on gaps and opportunities in research
16 and health policy [3]

17 The epidemiological data of the study align with the
18 national literature. Key characteristics included: a median
19 age of patients was 62 years, a predominance of males,
20 regional differences (highest concentration in the southeast-
21 ern region), and overall survival (OS) of 37 months. These
22 findings are consistent with other Brazilian analyses, [4] but
23 contrast with countries such as the USA and France, where
24 the median age at diagnosis is higher (66–74 years) and sur-
25 vival is longer [5,6] This underscores differences in demo-
26 graphics, epidemiology, and access to therapy.

27 The study also evaluated the impact of different therapeu-
28 tic regimens. Although bortezomib (Bortezomib) was only for-
29 mally incorporated into the SUS in 2020, patients treated with
30 this agent had an OS of 67 months, while those on thalido-
31 mide-based regimens reached 54 months. Notably, patients
32 undergoing hematopoietic stem cell transplantation (HSCT)
33 exhibited an even higher survival, with a median of 87
34 months [3] These results reinforce the clinical benefits of
35 innovative therapies and transplantation, highlighting the
36 importance of early access to effective treatments.

37 However, despite the robustness of the administrative
38 database used by the authors (DATASUS), some gaps remain.

There are no data on treatment adherence, toxicity, cyto- 39
genetic stratification, or responses to specific lines of therapy. 40
Additionally, the absence of information on time to diagnosis, 41
functional status, and associated comorbidities limits individ- 42
ualized interpretation and prevents understanding the deter- 43
minants of survival. Furthermore, the methodology adopted 44
—classifying patients according to therapy exposure at any 45
point rather than only the first line regimen—reflects real- 46
world complexity, although it may introduce selection bias. 47

The difference in survival of individuals undergoing HSCT 48
underscores its relevance and efficacy as a standard of care. 49
However, it is noteworthy that HSCT was performed in only 50
26.9 % of the population, despite its established efficacy as a 51
standard therapeutic cornerstone for eligible patients [3] In 52
the Brazilian context, this low rate reflects structural barriers, 53
including an insufficient number of specialized centers, 54
regional disparities in healthcare access, long waiting times, 55
and socioeconomic limitations. Thus, concerns remain 56
regarding the underutilization of transplantation in Brazil [7] 57
Additionally, the late incorporation of new drugs, such as bor- 58
tezomib (Bortezomib), demonstrates the need for more agile 59
health technology assessment processes without compromis- 60
ing safety and efficacy [8] Consequently, it is plausible that 61
many eligible patients never undergo HSCT, limiting the 62
potential gains shown by the national cohort. 63

Importantly, this study offers key lessons for Brazil. In 64
summary, three points should be considered: there is room 65
for more detailed investigations that can translate evidence 66
into concrete improvements in clinical practice; it is essential 67
to discuss strategies that expand equitable access to HSCT, 68
such as increasing the network of specialized centers, cou- 69
pled with rapid incorporation policies and adequate funding. 70
Robust national data can support health policy, guide thera- 71
peutic protocols, and reduce reliance on evidence from other 72
countries, which often have distinct population profiles. 73

In conclusion, the 16-year national cohort analysis high- 74
lights the impact of innovative drugs, the importance of long- 75
term studies for MM in Brazil, exposes failures in access to 76
innovative therapies and HSCT, and underscores the urgency 77

of public policies capable of democratizing this access. Encouraging the collection of population-based data, expanding access to medications, and increasing the transplantation network are essential steps to improve care for patients with MM in Brazil.

Conflicts of interest

The authors declare no competing financial interests.

REFERENCES

- Centers for Disease Control and Prevention (CDC). Myeloma basics [internet]. Myeloma; Available at <https://www.cdc.gov/myeloma/about/index.html> [cited 21 Oct 2025].
- De Oliveira Santos M. Estimate/2020 – Cancer incidence in Brazil. Rev Bras Cancerol. 2020;66(1):e00927. Mar 20 Available at <https://rbc.inca.gov.br/index.php/revista/article/view/927>.
- Marta D, Menezes C, Prata WM, Laper IZ, Azevedo PS, Sabino P, et al. Overall survival in multiple myeloma in Brazil: a 16-year cohort. Hematol. Transfus. Cell Ther. 2025;47(4):103962. <https://doi.org/10.1016/j.htct.2025.103962>.
- Garcia CS, Righes CS, Muller KTC, Ricas SMMC, Almeida EBC. Epidemiological profile of patients diagnosed with multiple myeloma in a referral hospital for malignant hematologic neoplasms. Rev Bras Anal Clin. 2020;52(3):200–5. Available at <https://rbac.org.br/artigos/perfil-epidemiologico-de-pacientes-diagnosticados-com-mieloma-multiplo-em-hospital-de-referencia-para-neoplasias-malignas-hematologicas/>.
- Arias E, Xu J, Kochanek K. United States life tables, 2021. Natl. Vital. Stat. Rep. 2023;72(12):1–64. Available at <https://www.cdc.gov/nchs/data/nvsr/nvsr72/nvsr72-12.pdf>.

- Hungria VT. Multiple myeloma in Brazil: clinical and demographic aspects and validation of the International Staging System (ISS) in Brazilian patients. Rev. Bras. Hematol. Hemoter. 2007;29(1):10–3. Available at <https://pesquisa.bvsalud.org/portal/resource/pt/lil-537336>.
- Instituto Oncoguia. Stem Cell Transplantation For Multiple Myeloma [Internet]. São Paulo: Oncoguia; 2015. Available at <https://www.oncoguia.org.br/conteudo/transplante-de-celulas-tronco-para-mieloma-multiplo/2009/397/> [cited 21 Oct 2025].
- National Committee for Health Technology Incorporation (CONITEC). Recommendation Report No. 557: Bortezomib [Internet]. Brasília: Ministry of Health; 2020. Available at <https://www.gov.br/conitec/pt-br/assuntos/avaliacao-de-tecnologias-em-saude/recomendacoes-da-conitec/2021> [cited 21 Oct 2025].

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