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# Daratumumab-Based Maintenance After ASCT in Newly Diagnosed Multiple Myeloma: Real-World Outcomes From a Single-Center Cohort

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## INTRODUCTION

- The integration of daratumumab into frontline therapy for newly diagnosed multiple myeloma (NDMM) has led to significant improvements in response depth and progression-free survival (PFS). Phase III trials such as CASSIOPEIA¹ and PERSEUS² demonstrated that daratumumab-based induction regimens, followed by maintenance with daratumumab alone or in combination with lenalidomide, provide superior disease control compared to conventional therapies. Based on these results, daratumumab-based maintenance has become a new standard of care post-autologous stem cell transplantation (ASCT).
- We present a preliminary real-world analysis of daratumumab-based maintenance—either as monotherapy or in combination with lenalidomide—in a group of NDMM patients undergoing ASCT, treated during a period when this approach was being progressively adopted into clinical practice based on emerging trial evidence.

## **RESULTS**

#### **Baseline Characteristics of NDMM Patients (n = 19)**

A total of 19 patients with newly diagnosed multiple myeloma (NDMM) received daratumumab-based maintenance following autologous stem cell transplantation (ASCT).

The median age at diagnosis was 59 years (range 44–69), and 7 patients (36.8%) were female.

n	(%)
7	(36.8%)
3	(15.8%)
4	(21.1%)
1	(5.3%)
2	(10.5%)
2	(10.5%)
	7 3 4 1 2

133 Stage	LI LI	( /0)
I	6	(31.6%)
ll l	10	(52.6%)
III	3	(15.8%)
R-ISS Stage	n	(%)
R-ISS Stage	<b>n</b> 6	(%)
R-ISS Stage I		
1	6	(31.6%)
	6 10	(31.6%) (52.6%)

High-risk cytogenetics were present in 2 patients (10.5%), one with del(17p) and one with t(4;14). Gain of 1q was detected in 4 patients (21.1%), 2 of whom also presented a complex karyotype.

## **Treatment and Response Outcomes**

#### **Treatment Overview**

All patients received induction with daratumumab-based regimens:

- D-VTd: 14 patients (73.7%)
  D-VRd: 4 patients (21.1%)
- **DVMP**: 1 patient (5.3%)

All patients underwent **ASCT** with conditioning using melphalan.

- 17 patients (89.5%) received melphalan at 200 mg/m<sup>2</sup>
- 2 patients (10.5%) received reduced-intensity melphalan

**Post-transplant consolidation** was administered to 18 patients (94.7%). **Maintenance therapy**:

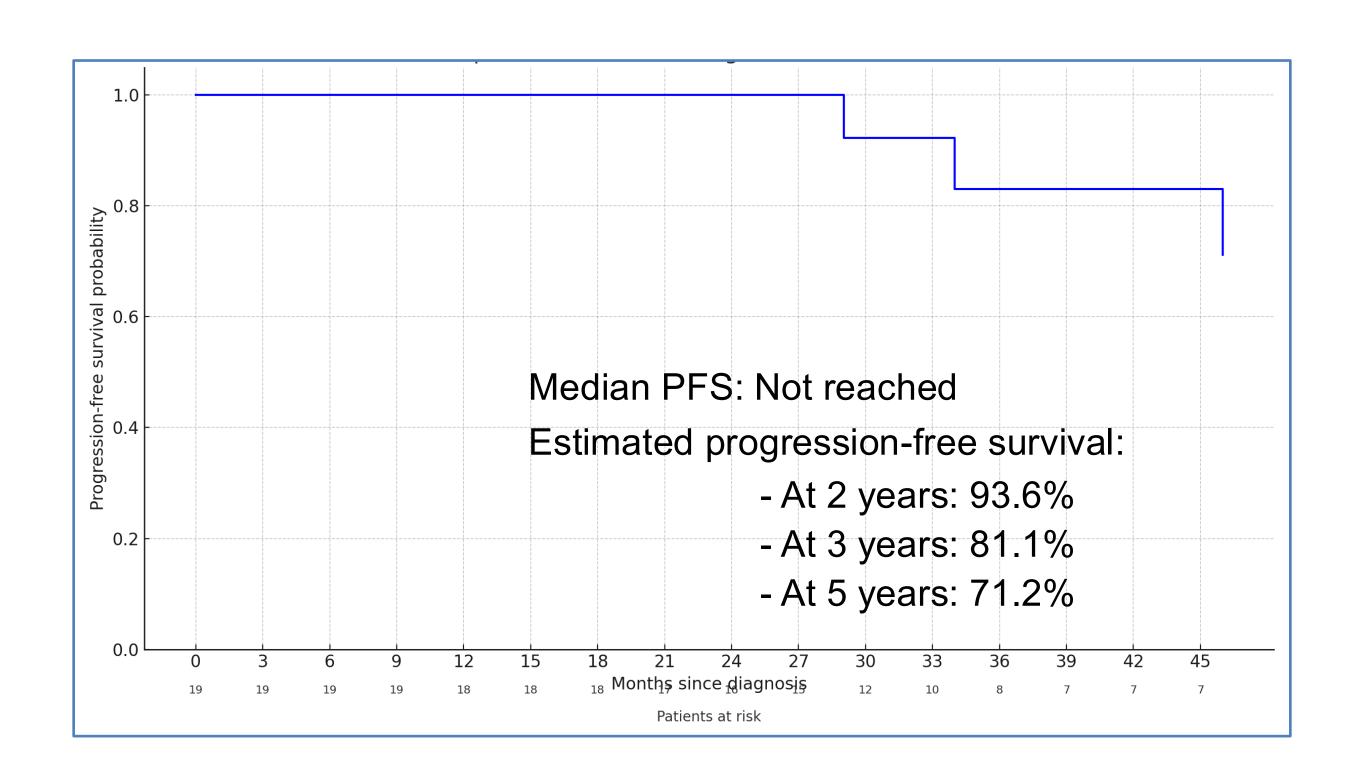
- •Daratumumab monotherapy: 12 patients (63.2%) → 4 discontinued without relapse
- •Daratumumab + lenalidomide (D-R): 7 patients  $(36.8\%) \rightarrow 1$  discontinued daratumumab without relapse

## **Response by Phase**

Treatment Phase	sCR	CR	VGPR	PR
Post-induction (n=19)	2 (10.5%)	4 (21.1%)	10 (52.6%)	3 (15.8%)
Post-ASCT (n=19)	8 (42.1%)	4 (21.1%)	6 (31.6%)	1 (5.3%)
Post-consolidation (n=18)	10 (55.6%)	3 (16.7%)	5 (27.8%)	_
Best response during maintenance (n=19)	14 (73.7%)	3 (15.8%)	2 (10.5%)	_

## **Estimated Progressión Free Survival**

Two patients relapsed (1 on Dara, 1 on D-R); 1 additional progression without strict criteria managed by adding lenalidomide to Dara.



#### **CONCLUSIONS**

- In this real-world preliminary cohort of NDMM patients, daratumumab-based maintenance after ASCT—either as monotherapy or in combination with lenalidomide—was well tolerated and led to deep, sustained responses.
- The majority of patients achieved stringent complete response during maintenance, and estimated 3-year progression-free survival remained above 80%. These results are consistent with those reported in pivotal trials such as PERSEUS, supporting the use of daratumumab-based maintenance as a standard post-transplant strategy in clinical practice.

#### **REFERENCES**

<sup>1</sup>Moreau P et al. *Lancet*, 2019; 394(10192):29-38.

<sup>2</sup>Sonneveld P et al. *N Engl J Med*. 2023;389(24):2284–2297.

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