

Cost per Responder for Teclistamab-Daratumumab versus Daratumumab, Pomalidomide, Dexamethasone (DPd), and Daratumumab, Bortezomib, Dexamethasone (DVd) in Relapsed Refractory Multiple Myeloma Patients

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Key Takeaway



This analysis demonstrates that TecDara provides greater economic value than DVd/DPd, driven by deeper responses, thus supporting its use in earlier lines of therapy

Conclusion



The cost per CR+ was lower for TecDara versus DVd and DPd despite the availability of generic pomalidomide and bortezomib



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Introduction

- The MajesTEC-3 trial of teclistamab in combination with daratumumab (TecDara) showed markedly superior efficacy over standard of care regimens daratumumab combined with dexamethasone and bortezomib or pomalidomide (DVd/DPd) in patients with relapsed refractory multiple myeloma (RRMM) with 1-3 prior lines of therapy.¹
- At a median follow-up of 34.5 months, 81.8% of patients in the TecDara cohort and 32.1% of patients in the DVd/DPd cohort had a complete response or better (CR+) (P<0.001).¹
- Progression-free survival (PFS) rates at 36 months were significantly higher in the TecDara cohort than the DVd/DPd cohort (HR: 0.17; 95% CI: 0.12-0.23; P<0.001), and overall survival at 36 months was significantly higher in the TecDara cohort (83.3% vs 65.0%; P<0.0001).¹
- With the availability of bortezomib and pomalidomide generics, it is important to evaluate the economic value of TecDara compared with DVd/DPd.
- Given the association between deep responses and improved clinical outcomes, cost per CR+ was used as a clinically meaningful and readily interpretable measure of treatment value.

Table 1: Key model inputs and assumptions

1. Model time horizon of 36 months	
2. Drug costs	
i. All unit drug costs are based on WAC (April 2026). WAC for tocilizumab is based on the WAC for Actemra® and pomalidomide and bortezomib are based on generic pricing.	vi. Cost of per day IP stay from Tan et al, ⁴ inflated to costs in 2026.
ii. Dosing schedule for TecDara, including SUD for Tec, is as described in the United States Prescribing Information (USPI). ² For DVd/DPd, drug dosing schedules as described in the MajesTEC-3 trial protocol ¹	3. OP administration and supportive care costs
iii. For TecDara, assumes 28.5% of patients switch from 3 mg/kg to 1.5mg/kg at week 29. ¹	i. Drug administration cost for sub-cutaneous drugs is based on the CMS Schedule. ³
3. SUD costs	ii. Supportive care costs included monthly recurring cost of treatment monitoring and non-myeloma related healthcare costs. ⁵
i. Applicable for TecDara only.	4. AE post-SUD hospitalization costs
ii. Patients receive the first and second teclistamab step-up doses (SUDs) in one hospitalization on days 1 and 3 and (discharged on day 4; length of stay = 4 days), in accordance with the USPI. ² The first daratumumab dose is assumed to be administered OP on day 1 (prior to the hospitalization), while the second daratumumab dose is assumed to be administered on day 7 prior to the third teclistamab SUD (first full treatment dose) on day 7 in the OP setting.	i. Cost of AE management for each treatment was calculated as sum-product of risks and per-episode costs of all-cause grade 3-5 AEs occurring with a frequency of at least 5% for at least one of the treatment regimens.
iii. Patients who experience CRS or ICANS of grade 3+ are assumed to be hospitalized and finish SUD in IP settings.	ii. AE rates are based on MajesTEC-3 for TecDara and DVd/DPd ¹
iv. Costs for radiology exam, neurological consult, and a course with Anakinra are assumed for patients with Grade 2+ ICANS. Costs for treatment with tocilizumab is assumed for patients with Grade 2+ CRS. Costs for radiology exam and neurological consult are based on CMS schedule. ³	iii. Costs of each AE type per episode were obtained from the Healthcare Cost and Utilization Project (HCUP) National Inpatient Sample 2022.
v. The rates of CRS and ICANS are based on MajesTEC-3 results. ¹	5. Subsequent therapy costs
	i. The PFS Kaplan-Meier curve associated with each treatment from the MajesTEC-3 trial were used as a proxy for the proportion of patients remaining on treatment over time.
	ii. Subsequent treatment costs were assumed for 56.7% of patients who discontinue initial regimens.
	iii. Applied as a monthly cost based on a real-world evidence study reporting on all-cause healthcare cost ⁵ and applied from progression to death or after 36 months of treatment.

AE, adverse event; CRS, cytokine release syndrome; Dara, daratumumab; DVd/DPd, daratumumab and dexamethasone with either bortezomib (V) or pomalidomide (P); ICANS, immune effector cell-associated neurotoxicity syndrome; IP, inpatient; OP, outpatient; PFS, progression-free survival; SUD, step-up dose; Tec, teclistamab; USPI, United States Prescribing Information; WAC, wholesale acquisition cost.

Results

- The cost of care per patient for the first 36-months of treatment was \$1,393,647 for TecDara, \$635,099 for DVd, and \$704,441 for DPd (Figure 1).
- Despite higher total cost of care, TecDara demonstrated the lowest cost per CR+ (\$1,703,998) vs DVd (\$1,978,833) and DPd (\$2,194,890), representing savings of \$274,836 and \$490,892 for each cost per CR+, respectively (Figure 2).

Figure 1: Total cost of care per patient of TecDara versus DVd/DPd during the first 36 months of treatment

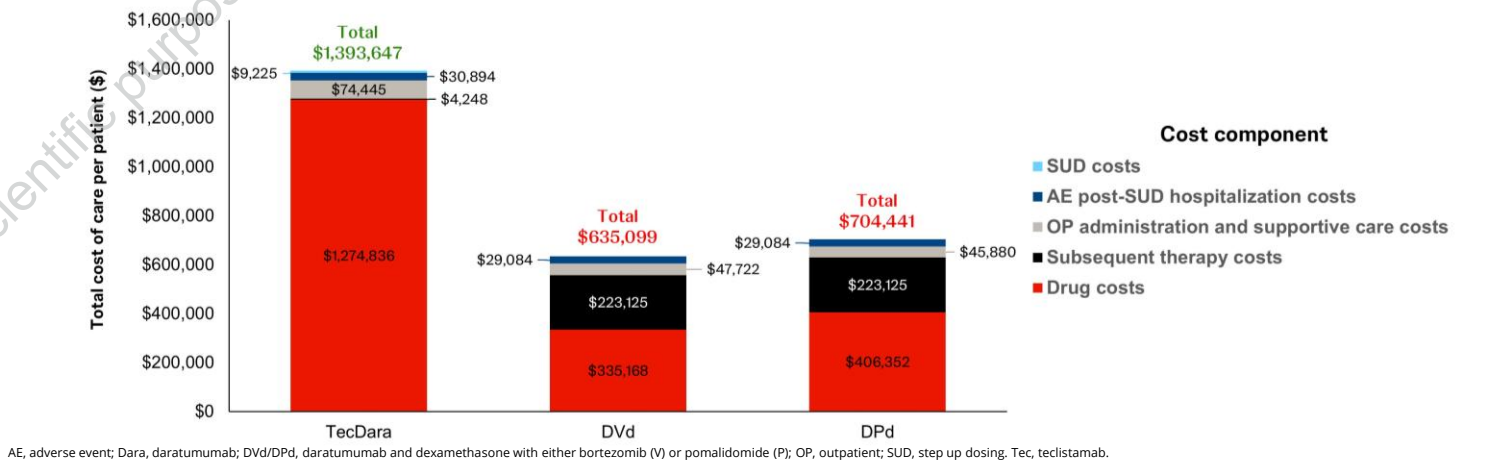
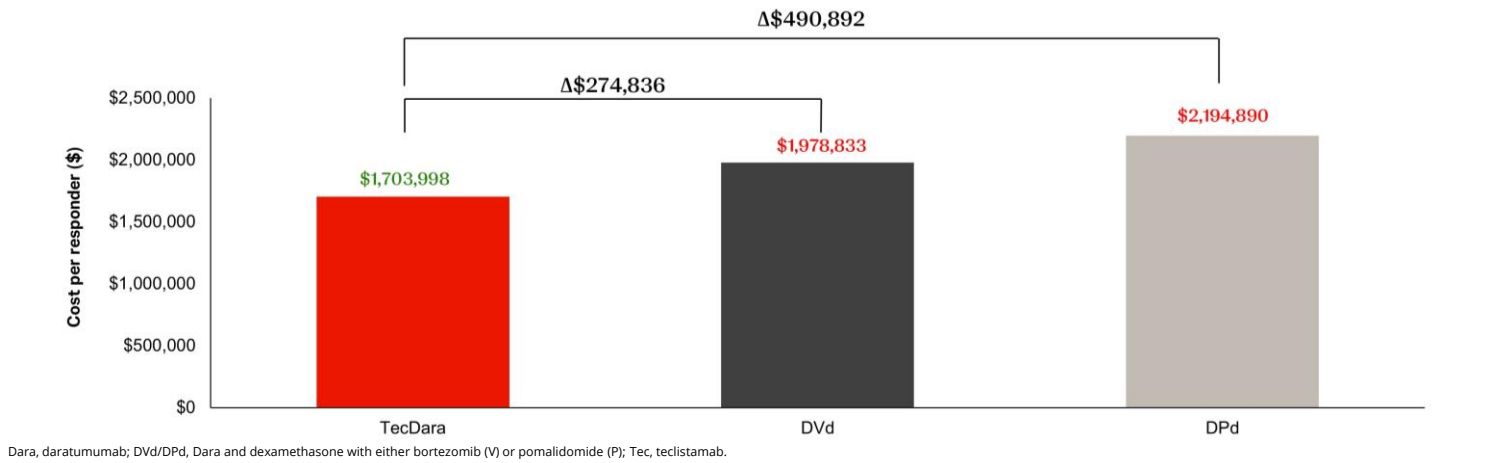


Figure 2: Cost per responder (CR+) of TecDara versus DVd/DPd during the first 36 months of treatment



Dara, daratumumab; DVd/DPd, Dara and dexamethasone with either bortezomib (V) or pomalidomide (P); Tec, teclistamab.

Study limitations

- Clinical efficacy and safety inputs for both DVd and DPd were based on the MajesTEC-3 comparator arm where most patients received DPd, and therefore, the clinical outcomes may be more representative of DPd than DVd.
- As dose reduction timing was highly variable from cycle 7 onwards in MajesTEC-3, the model assumes all dose reductions occurred at Week 29, which may overestimate the duration of dose reduction.

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