## **MM-431**

## A Comparison Between Daratumumab and Non-Daratumumab-Based Salvage Regimens Used at First Relapse Post-Lenalidomide Maintenance in Multiple Myeloma

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Context: Lenalidomide maintenance therapy after autologous stem cell transplantation (ASCT) is standard of care for patients with multiple myeloma (MM). However, most patients relapse, and the optimal choice of therapy at first-relapse post-lenalidomide maintenance is not established. Objective: To compare the efficacy of salvage therapies at first relapse post-lenalidomide maintenance. Design: Retrospective study of consecutive patients with MM seen at Mayo Clinic, Rochester between 1/1/2005-12/31/2016. Setting: Tertiary referral center. Patients or Other Participants: Two hundred thirteen patients treated with lenalidomide/lenalidomidedexamethasone maintenance post-ASCT were screened; 136 patients experienced a first relapse. One hundred eighteen patients receiving salvage therapy were included in subsequent analyses. Interventions: N/A Main Outcomes Measures: The main outcome measure was second progression-free survival (2nd PFS), which was calculated from the start of therapy at first relapse after maintenance until therapy discontinuation. Results: The median follow-up was 5.4 (95% CI: 4.9, 5.9) years from diagnosis. The median 2nd PFS was significantly longer in patients treated with daratumumab-based regimens [18.4 (95% CI: 10.9, 25.9) months; n=32] versus patients without daratumumab [8.9 (95% CI: 5.5, 12.3) months; n=86; p=0.006]. Adjusting for age, ISS stage 3, cytogenetic risk, Rd maintenance, and lenalidomide refractoriness at salvage, daratumumab-based therapy was associated with a significantly improved PFS [hazard ratio 0.31 (95% CI: 0.16, 0.61; p=0.001)]. The median 2nd PFS was superior in daratumumab + IMiD (n=16) compared to daratumumab + PI (n=15) [NR versus 1 yr (95% CI: 0.5, 1.5), respectively; p=0.004]. In patients not receiving daratumumab, median 2nd PFS was comparable between PI-based combinations [9.2 (95% CI: 6.6, 11.7) months; n=44], IMiD-based combinations [6.7 (95% CI: 0.82, 12.6) months; n=18; p=0.7], or PI + IMiD-based combinations [11.2 (95% CI: 0, 28.4 months; n=24; p=0.17]. Without daratumumab, there was also no significant difference in median 2nd PFS between patients who received lenalidomide-based combinations [6.7 (95% CI: 0, 15.4) months] compared with pomalidomide-based regimens [20.1 (95% CI: 0, 41.4) months; p=0.5]. Conclusions: Daratumumab-based therapies at relapse are associated with improvement in 2nd PFS, and daratumumab-IMiD combination was superior to daratumumab-bortezomib combination. Without daratumumab, there was no significant difference between doublet *versus* triplet therapies, IMiD-versus PI-based regimens or lenalidomide versus pomalidomide-based combinations. Keywords: MM, multiple myeloma, lenalidomide, maintenance, daratumumab

## **MM-435**

Propensity Score Matching
Analysis to Evaluate Bortezomib/
Cyclophosphamide /Dexamethasone
and Bortezomib/Thalidomide/
Dexamethasone from Real-World Data
in Patients with Newly Diagnosed
Multiple Myeloma, on Behalf of the
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Context: The most frequently used induction treatments for newly diagnosed multiple myeloma in Colombia are VTD and CyBorD for transplant-eligible patients. Objective: At present, there are no clinical studies in our region that compare the effectiveness of these two regimens. Design: To reduce the bias between the groups, we performed a 1:1 propensity score matching (PSM) technique for analysis. Setting: The Registro Epidemiológico de Neoplasias Hematológicas en Colombia (RENEHOC) collected data electronically on Colombian patients with multiple myeloma between 2010 and 2018. Patients or Other Participants: After