Phase 3, Randomized, Open-Label Study of Pembrolizumab Versus Brentuximab Vedotin for Treatment of Relapsed or Refractory Classical Hodgkin Lymphoma: KEYNOTE-204

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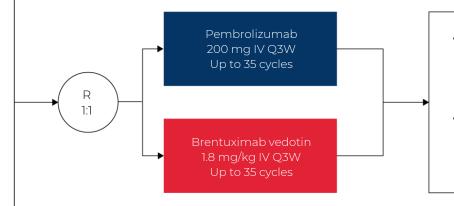




KEYNOTE-204 Study Design (NCT02684292)

Key Eligibility Criteria

- · Relapsed or refractory cHL
- Relapse after auto-SCT or ineligible for auto-SCT and failed 1 prior line of therapy
- Measurable disease per IWG 2007 criteria¹
- ECOG PS 0-1
- BV-naive and BV-exposed patients eligible



- Response assessed Q12W per IWG 2007 Revised Response Criteria for Malignant Lymphoma¹
- AEs evaluated Q3W throughout the trial period, and Q12W during follow-up

Stratification Factors

- Prior auto-SCT (yes vs no)
- Status after 1L therapy (primary refractory vs relapsed <12 months vs relapsed ≥12 months after end of 1L therapy)

Primary End Point: PFS per blinded independent central review (BICR) by IWG 2007 criteria including clinical and imaging data following auto-SCT or allogeneic stem cell transplantation (allo-SCT); OS

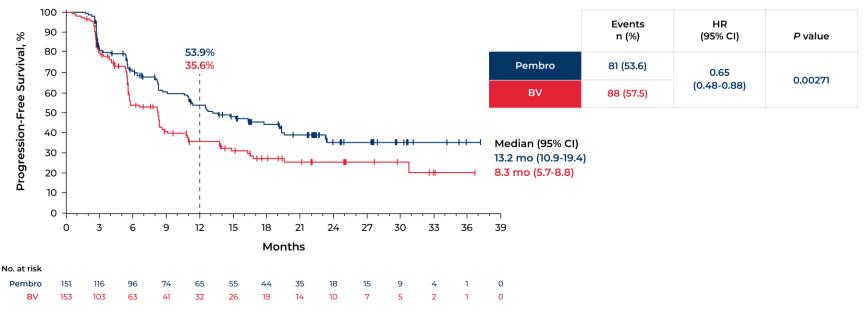
Secondary End Points: PFS per BICR by IWG 2007 criteria excluding clinical and imaging data following auto-SCT or allo-SCT; ORR by BICR per IWG 2007; PFS per investigator review; DOR, safety

Cheson BD et al. J Clin Oncol. 2007;25:579-586.



Primary End Point: Progression-Free Survival per Blinded Independent Central Review

Including clinical and imaging data following auto-SCT or allo-SCT

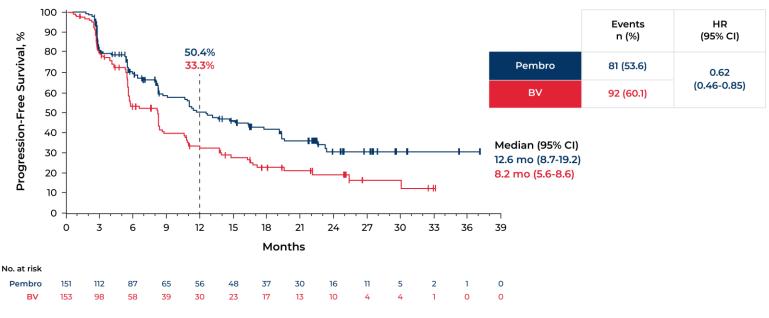


Data cutoff: January 16, 2020



Progression-Free Survival per Blinded Independent Central Review

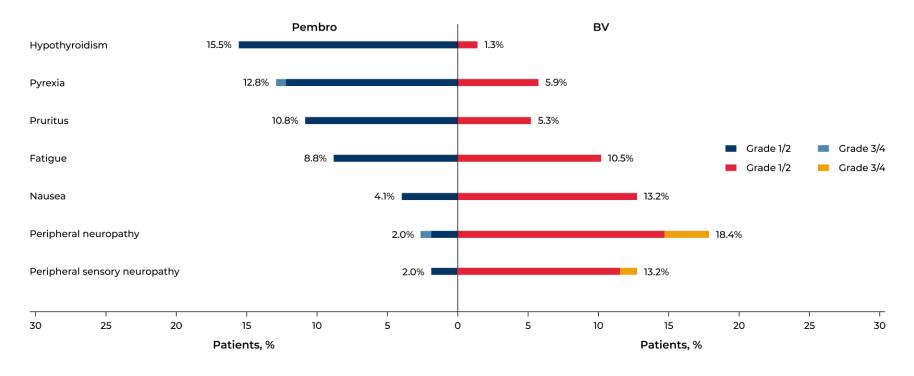
Excluding clinical and imaging data following auto-SCT or allo-SCT



Data cutoff: January 16, 2020



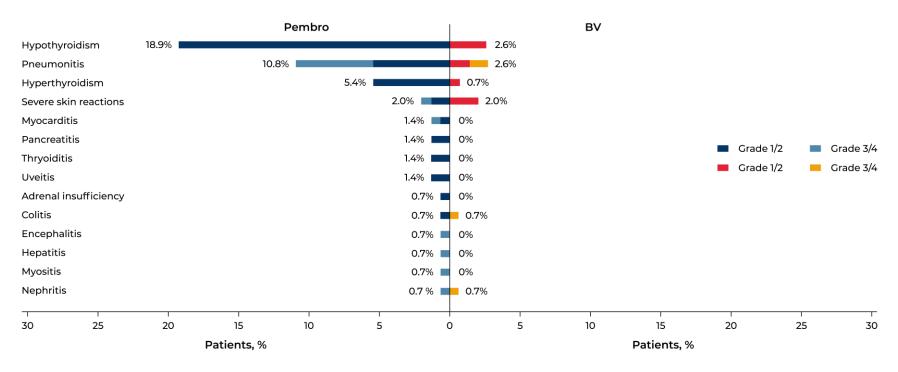
Treatment-Related AEs (≥10% Either Arm)



Data cutoff: January 16, 2020



Immune-Mediated AEs



Based on a list of terms specified by the sponsor and included by the investigator regardless of atribution to study treatment or immune relatedness. Data cutoff: January 16, 2020

