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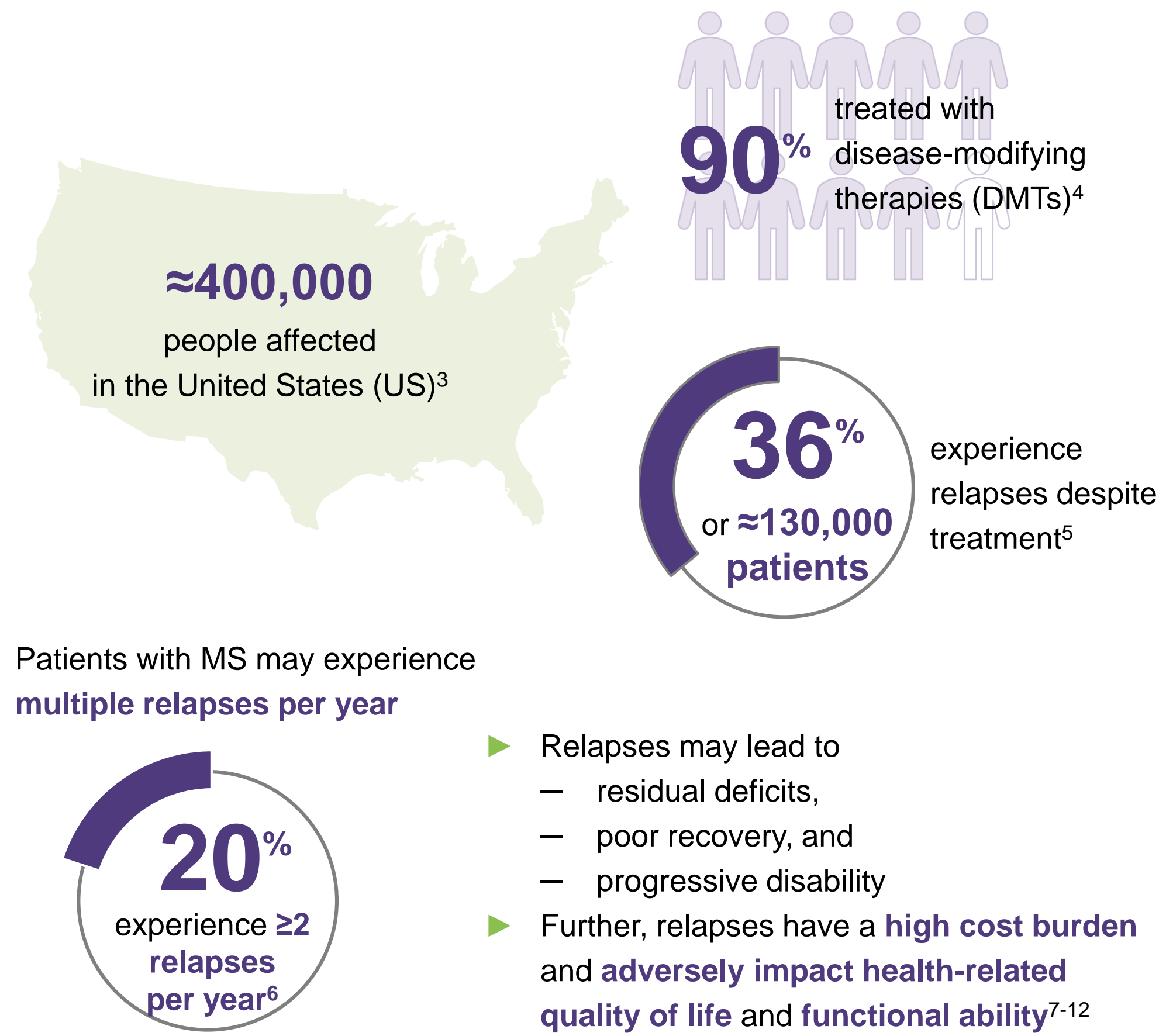
Cost Per Response Analysis of Repository Corticotropin Injection versus Other Late-Line Treatments for Multiple Sclerosis Relapses in Adults

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BACKGROUND

- Multiple Sclerosis (MS) is an autoimmune, inflammatory, demyelinating disease of the central nervous system, characterized by **relapsing-remitting disease**^{1,2}



- Patients with MS may experience **multiple relapses per year**
- Treatment guidelines suggest corticosteroids as first-line agents for managing MS relapses¹³
 - However, not all patients **tolerate or respond effectively to corticosteroids**
 - Other late-line MS relapse treatment options include Acthar® Gel (repository corticotropin injection [RCI]), plasmapheresis (PMP), or intravenous immunoglobulin (IVIg)¹⁴
- There is a **lack of evidence comparing the cost per response of these late-line treatments** for the resolution of MS relapses

RESULTS

Base case results (Figure 2)

- The base case annual cost per RCI response (\$141,970) was **lower** than that with PMP/IVIg (\$253,331)

Sensitivity analysis results (Figure 3)

- The cost per response for RCI was sensitive to the response rate and annual cost of care
 - Cost per response**, range: \$126,879 - \$157,019 for change in response rate from 96.9% to 78.3%, respectively
 - Cost per response**, range: \$124,090 - \$161,450 for change in average annual cost of care from \$107,462 to \$139,816, respectively
- RCI had a lower cost per response compared with PMP/IVIg with changes in annual cost of care and response rate

Figure 2: Base case cost per response

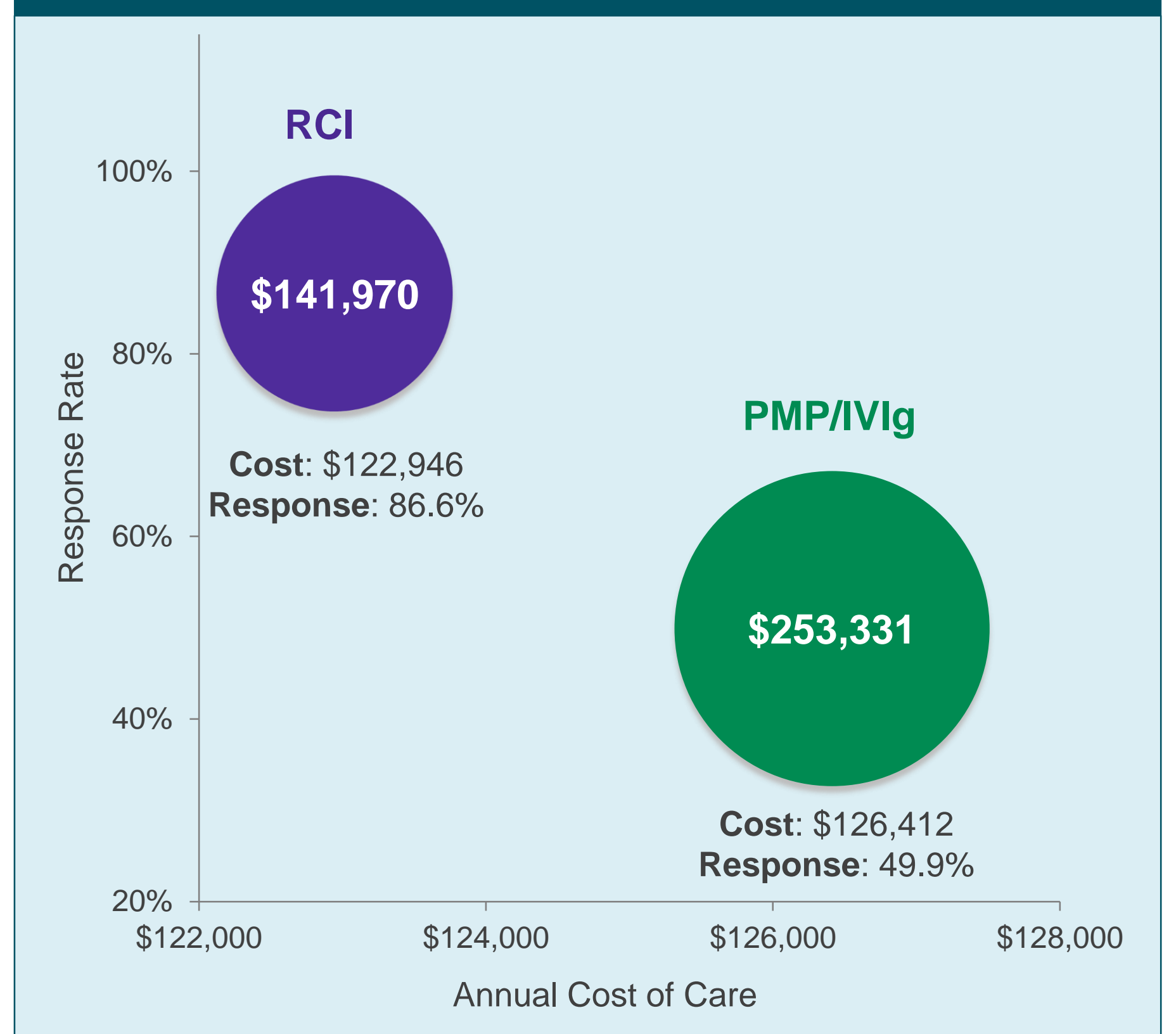
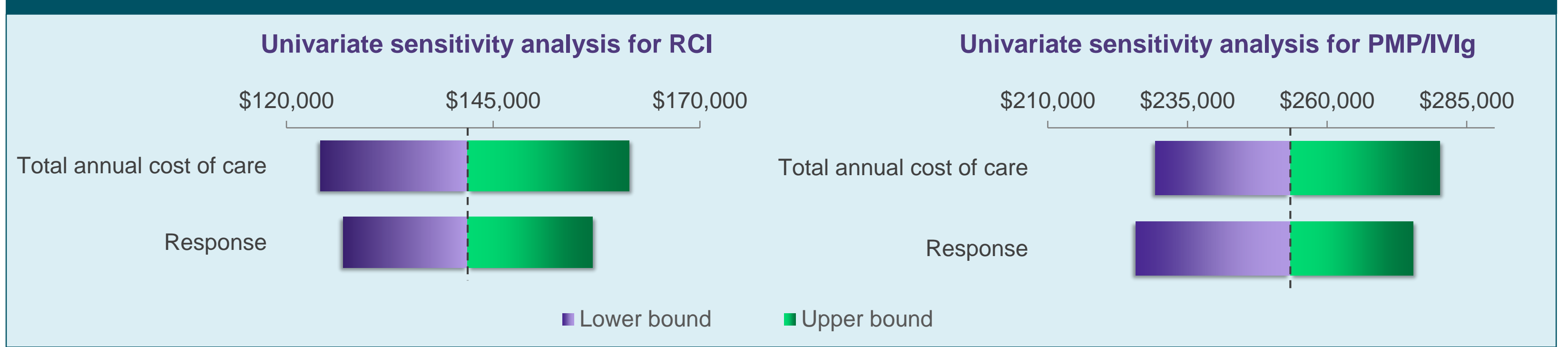


Figure 3: Univariate sensitivity analysis of cost per response



OBJECTIVE

To estimate the cost per response of MS relapse treatment with RCI versus PMP/IVIg among patients with acute exacerbations of MS (≥1 relapse per year) from a payer perspective

METHODS

- Sample:** Patients who experienced ≥2 MS exacerbations
- Treatments:** RCI and PMP/IVIg
- Average annual cost of care: (Figure 1)**
 - Data source:** Truven Health Analytics MarketScan® Commercial Claims and Encounters Databases between July 1, 2007 and December 31, 2012¹⁵
 - Costs included MS-related inpatient, outpatient, and medication costs
 - Adjusted for number of relapses prior to index date, days between exacerbations, comorbid diabetes without complications, year of index exacerbation, and number of outpatient services, hospitalizations, and medications in the 6 months prior to the index exacerbation
 - Costs were inflated to 2019 USD using medical consumer price index
- Response rate: (Table 1)**
 - Data sources:** Humana Comprehensive Health Insights Database® (January 1, 2008 through July 31, 2015)⁶ and HealthCore Integrated Research Database™ (January 1, 2006 through November 30, 2016)¹⁶
- Response:** No additional relapse treatments or procedures within 30 days
 - Relapse:** defined using established claims-based methodology which included an inpatient or outpatient claim with a diagnosis of MS followed by receipt of a relapse treatment or procedure (RCI, PMP, or IVIg)¹⁷
- Cost per response:** Designed to compare annual cost of care per patient achieving MS relapse resolution

$$\frac{\text{Average annual cost of care}}{\text{Percent patients with resolved relapse}}$$
- Sensitivity analysis:** A one-way deterministic sensitivity analysis was also performed to assess the impact of model inputs on the results for cost per response
- Model assumptions:**
 - Population across the studies were assumed to be homogeneous for the diagnosis of MS
 - Annual cost of care assumes that patients are treated with intravenous methylprednisolone for the initial relapse and subsequently treated with RCI or PMP/IVIg for the next relapse

Figure 1: Annual cost of care for RCI versus PMP/IVIg

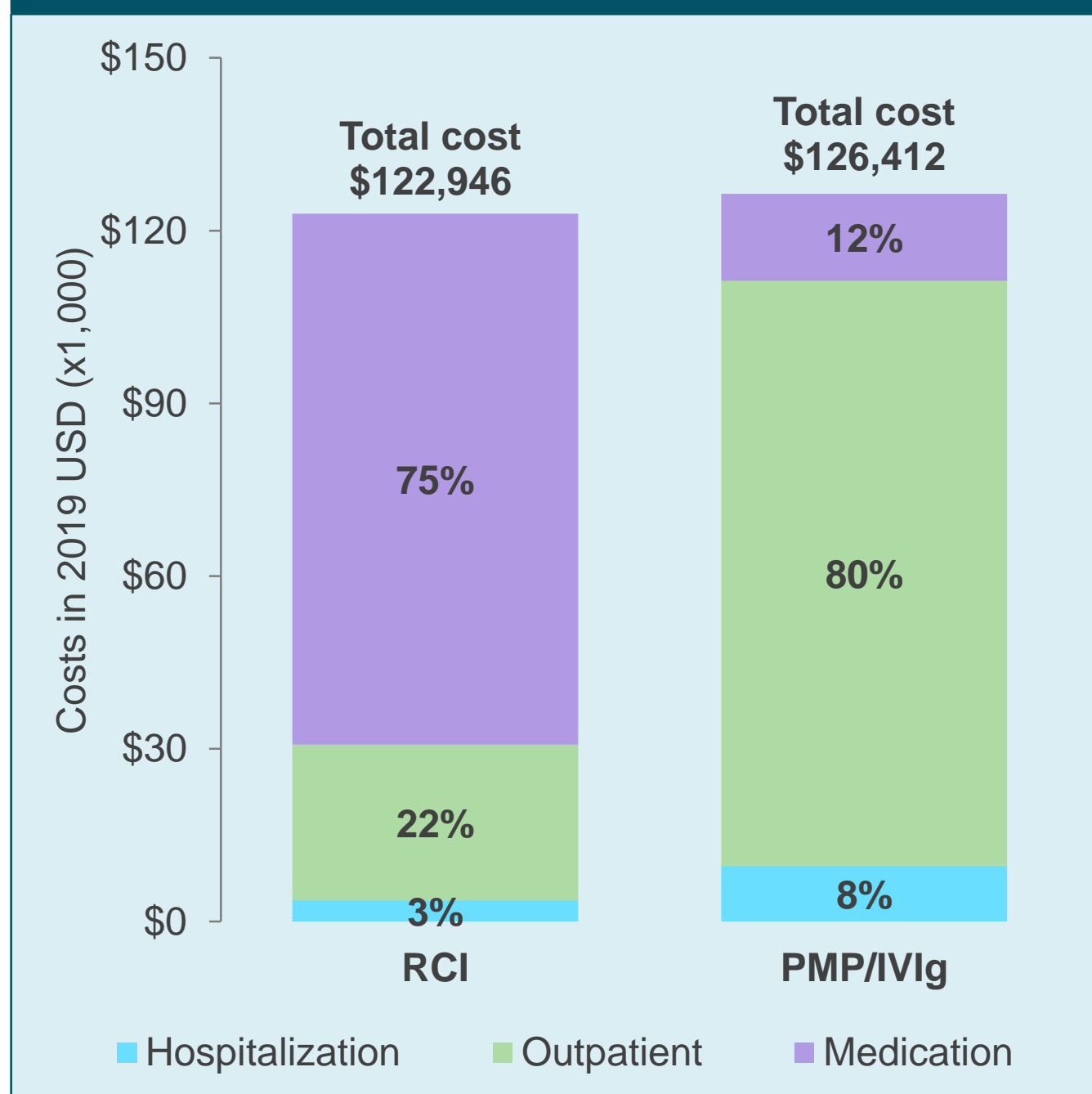


Table 1. Weighted response rates for patients treated with RCI and PMP/IVIg

Data source	Response rate	
	RCI	PMP/IVIg
Humana Comprehensive Health Insights Database ⁶	96.9% (189 / 195)	45.9% (112 / 244)
HealthCore Integrated Research Database ¹⁶	78.3% (188 / 240)	56.0% (89 / 159)
Average response rates	86.6% (377 / 435)	49.9% (201 / 403)

ACKNOWLEDGMENTS

Falcon Research Group (North Potomac, MD, USA) provided medical writing and editorial assistance

DISCLOSURES

George Wan and John Niewoehner are employees of Mallinckrodt; Ishveen Chopra was a paid consultant at Mallinckrodt Pharmaceuticals

LIMITATIONS

- Relapses were identified based on treatment-seeking behavior across two databases using an established claims-based algorithm; treatment received outside a healthcare visit was not addressed
- Unrestricted enrollment could underestimate unresolved relapses. PMP and IVIg may be administered as courses of therapy, which would also lead to an underestimation
- The total annual cost of care did not account for treatment convenience and compliance and the safety profile associated with each therapy

CONCLUSIONS

- Although average annual cost of late-line treatments is similar, the cost per response of RCI is lower than other late-line treatments
- Sensitivity analysis shows that RCI was still cost-saving compared to other late-line treatments
- These findings shed light on the importance of late-line treatment selection strategies for patients with acute exacerbations of MS
- Robust management of MS relapse should reflect timely resolution with appropriate treatment to minimize patient burden

REFERENCES

- National Institute of Neurological Disorders and Stroke: www.ninds.nih.gov/disorders/multiple_sclerosis/detail_multiple_sclerosis.htm
- Cook S.D. et al International CMSC Consensus Conference, 2012, Int. J. MS Care. 14, 105-104
- Hersh CM, Fox RJ. Multiple sclerosis http://www.clevelandclinicmeded.com/medicalpubs/diseasemanagement/neurology/multiple_sclerosis/
- Data on file: MS HCP Tracking Study Wave 7 Final Report – May 2017
- Data calculations on file: Estimated ARR and percent patients relapsing from DMT PIs. Mallinckrodt Pharmaceuticals.
- Nazareth T et al. 7th JointECTRIMS – ACTRIMS, Paris, France, October 25-28, 2017
- Lublin FD et al. Neurology. 2003;61(11):1528-1532
- Hirst C et al. J Neurol. 2008;255(2):280
- Leray E et al. Brain. 2010;133(pt 7):1900-1913
- Scott TF et al. J Neurol Sci. 2010;292(1-2):52-56
- Mowry EM et al. Neurology. 2009;27(7):602-608
- Ross AP et al. Int J MS Care. 2012;14(3):148-159
- National Multiple Sclerosis Society, treating MS, managing relapses
- Nickerson M et al BMC Neurology. 2013;13:119
- Gold LS et al. Adv Ther. 2016;33(8):1279-92
- Nazareth T et al. 7th JointECTRIMS, Berlin, Germany, October 10-12, 2018
- Chastek et al J Med Econ. 2010;13(4):618-25