

Cost-effectiveness of Fingolimod, Teriflunomide, Dimethyl Fumarate and Interferon Beta-1a in Relapsing-remitting Multiple Sclerosis

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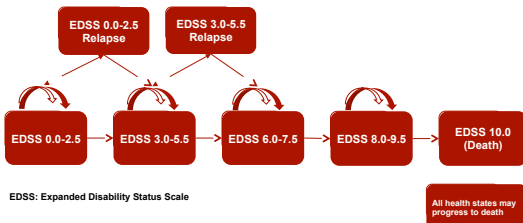
Objective

To compare the cost-effectiveness of fingolimod, teriflunomide, dimethyl fumarate and IM IFN β -1a as first-line therapies in treatment of patients with relapsing-remitting multiple sclerosis (RRMS).

Methods

- A Markov model was developed to simulate the disease progression and evaluate the cost-effectiveness of disease-modifying drugs from a US societal perspective. Time horizon in base case was 5 years [Figure 1].
- Model parameters were obtained from randomized controlled trials, natural history studies of multiple sclerosis, cross-sectional surveys and federal supply schedule drug prices.
- Outcomes included quality-adjusted life years (QALYs), incremental net monetary benefit (INMB) and incremental cost-effectiveness ratio (ICER). The societal willingness-to-pay (WTP) threshold was assumed to be \$150,000 per QALY.
- Costs were reported in 2012 US dollars and both costs and outcomes were discounted at 3% annual rate in base case.
- One-way sensitivity analyses and probabilistic sensitivity analyses were conducted to test the robustness of the model results.

Figure 1. Markov model for disease progression of multiple sclerosis



Base case

- The 5 years' total costs per patient were estimated at \$322,694, \$339,457, \$324,512 and \$298,875 for IM IFN β -1a, fingolimod, teriflunomide, and dimethyl fumarate, respectively. The accumulated QALYs associated with each drug were 3.34, 3.69, 3.68 and 3.72, respectively [Table 1].
- Compared with IM IFN β -1a, at the WTP of \$150,000, INMBs were estimated at \$36,146, \$50,441, and \$80,970 for fingolimod, teriflunomide, and dimethyl fumarate, respectively. Compared with IM IFN β -1a, ICERs were \$47,523 and \$5,218 for fingolimod and teriflunomide, respectively [Table 1].
- Dimethyl fumarate dominated all other drugs over the range of WTPs from \$0 to \$180,000 [Figure 2].

Sensitivity analysis

- One-way sensitivity analyses found model results were robust to most parameter variations. The only exception is when the monthly cost of fingolimod was beyond \$5,121, fingolimod would no longer be cost-effective compared with IM IFN β -1a [Figure 3]. Other one-way sensitivity analysis comparison figures are available on request. For all of the three oral therapies versus IM IFN beta-1a, INMB increased as the time horizon became longer [Figure 4].
- Probabilistic sensitivity analysis showed that for more than 90% of the simulations, dimethyl fumarate was the optimal therapy across all willingness-to-pay values [Figure 5].

Table 1. Base case results, compared with IM IFN β -1a (WTP=\$150,000)

	Cost	QALY	NMB	INMB vs. IM IFN β -1a	CER	ICER vs. IM IFN β -1a
IM IFN β -1a	\$322,694	3.34	\$177,656		\$96,741	
Fingolimod	\$339,457	3.69	\$213,802	\$36,146	\$92,034	\$47,523
Teriflunomide	\$324,512	3.68	\$228,097	\$50,441	\$88,085	\$5,218
Dimethyl fumarate	\$298,875	3.72	\$258,626	\$80,970	\$80,415	Dominant

Results

Figure 2. INMB of oral drugs vs IM IFN β -1a

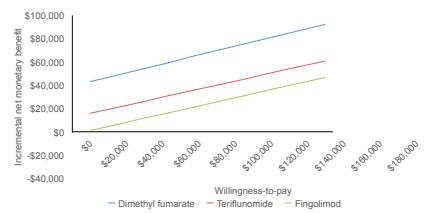


Figure 3. One-way sensitivity analysis

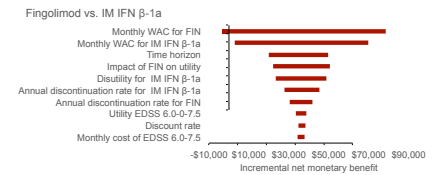


Figure 4. INMB vs IM IFN β -1a over time

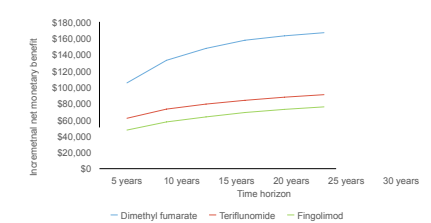
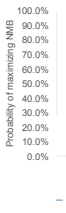


Figure 5. P...



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