

PREFERENCES OF PATIENTS WITH MULTIPLE SCLEROSIS FOR ATTRIBUTES OF INJECTABLE MULTIPLE SCLEROSIS TREATMENTS IN THE UNITED KINGDOM AND FRANCE

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Objectives: Adherence to injectable disease-modifying treatments in patients with multiple sclerosis (MS) impacts outcomes and can be influenced by perceptions of treatment efficacy, side effects, injection frequency, and the duration of injection. This study aimed to quantify preferences for selected attributes of injectable treatments among individuals with MS in the United Kingdom and France.

Methods: Respondents with a self-reported diagnosis of MS completed an online discrete-choice-experiment survey, consisting of a series of treatment-choice questions. Each choice question presented two hypothetical treatments, each with six attributes (years until disability progression, relapses in the next 4 years, injection time, injection frequency, flu-like symptoms (FLS), and injection-site reactions), each with various levels. Mixed-logit regression analysis was used to estimate preference weights for attribute levels and to calculate the relative importance of changes in treatment attributes (vertical distance between preference weights). Minimum acceptable efficacy estimates indicate improvement in efficacy that respondents would require in exchange for worsening injection frequency and FLS.

Results: In both countries, 100 respondents completed the survey. In the United Kingdom and France, respectively, improving the time until disability progression from 2 to 4 years, reducing injection frequency from “daily” to “every 2 weeks”, and reducing FLS from 3 days after every injection to none had a relative importance of 2.9 and 2.6, 3.0 and 3.5, and 2.5 and 3.1. Given the ranges included in the study, changes in these attributes were more important than most changes in other attributes assessed.

Conclusions: Reductions in the injection frequency of MS treatments and FLS can be as important to patients as improvements in treatment efficacy.

Keywords: multiple sclerosis, stated preference, discrete-choice experiment, injectable treatments, adherence

Injectable disease-modifying treatments (DMTs) are the standard therapy for multiple sclerosis (MS), and their efficacy and safety profiles are well-established (1). Dosing mode and frequency varies across injectable DMTs: Interferon beta-1b is administered subcutaneously (SC) every other day, interferon beta-1a is administered as a once-weekly intramuscular (IM) injection or a three-times-weekly SC injection, peginterferon beta-1a is a SC injection administered once every 2 weeks, and glatiramer acetate is an SC injection administered three times a week or once per day, depending on the dose. Numerous studies have demonstrated that adherence to DMT is suboptimal (2;3). Injection fatigue, injection fear, and injection-related side effects are common reasons for non-adherence (2–4), and adherence has also been demonstrated to be lower when dosing frequency is higher (2;3;5–8).

The most commonly reported adverse events (AEs) associated with injectable DMTs include injection-site reactions (ISRs; particularly with SC formulations) (9), and flu-like symptoms (FLS; associated with interferon therapies in particular) (4;10;11). Perceptions of and experiences with treatment efficacy, side effects, dosing frequency, and dosing complexity influence patient adherence to injectable DMTs, which impacts clinical outcomes (5–7). The literature on adherence to injectable DMTs for patients with MS suggests that improvements in administration, including reduced dosing frequency and/or increased ease of administration by means of the use of auto-injectors, could improve adherence to treatment and, hence, outcomes (2;12;13). Examination of patients’ preferences for features of injectable DMTs can provide quantitative measures of the tradeoffs patients are willing to make between the benefits of treatment, the AEs associated with treatment, and other dosing features (14). Minimum acceptable efficacy (MAE) is one of several ways to quantify preferences (14).

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Several recent papers have reported preferences for treatment attributes of DMTs. One quantified preferences for features or attributes of quality of life but not for features of treatment (15). Others quantified preferences for features of injection devices (16) or outcomes (13) for DMTs, but did not include treatment side effects or dosing attributes. Wicks et al. (17) quantified preferences for oral DMTs only including efficacy, side effects, and dosing frequency, and found that side effects were weighted more heavily than efficacy.

Conversely, Utz et al. (18) quantified preferences for oral and injectable DMTs and found that route of administration and treatment frequency are important to patients and can be more important than mild side effects. However, in this study, the mild side effects were not specified, so preference weights for specific side effects could not be estimated. One study from a single university-based center in the United States (U.S.) (19;20) quantified preferences for features of oral, injectable, and intravenously administered DMTs including efficacy, side effects, and dosing characteristics among U.S. patients to estimate risk preferences. These studies found that severe side effects, MS symptoms, and progression were most important over the ranges examined. However, the study could not dissociate effects of dosing frequency from mode of administration on patient preference.

Poulos et al. (2016) quantified preferences for features of injectable DMTs in a sample of patients from across the United States (21) and replicated the study in Germany (22). In both the U.S. and German studies (21;22), delaying disease progression and preventing FLS were key drivers of preferences, and some changes in dosing frequency were as important as changes in efficacy and FLS. These studies used a discrete choice experiment, a stated preference method, to examine preferences for treatment features characterizing treatments that were both currently available and possibly available in future treatments. Riñon et al. (22) show that the percentage of patients initiating DMTs or interrupting or discontinuing treatment vary considerably across different countries, suggesting that preferences vary across countries.

To further explore preferences in different countries, a study similar to the Poulos et al. (2016) study (21) was carried out among individuals living with MS in the United Kingdom (U.K.) and France to quantify preferences for attributes of injectable DMTs and assess the relative importance of changes in these attributes needed to offset changes in other treatment attributes. We are aware of no previous studies that have been carried out in these regions.

METHODS

The study replicated a study design previously administered in the United States and Germany (21;23) The study was approved by the Institutional Review Board at the Office of

Research Protection at RTI International. The study involved minimal risk; no personally identifiable information was collected from respondents. It was determined that local ethics review was not required in the United Kingdom due to the fact that respondents were not recruited through the National Health Service. Local ethics review in France was determined to be unnecessary due to the fact that the study was noninterventive.

Discrete-Choice Experiment

The study used a discrete-choice experiment (DCE) (24;25) to elicit preferences using a series of paired comparison questions in which respondents were asked to choose which of two hypothetical treatments they preferred. Each hypothetical treatment had six attributes (years until disability progression, number of relapses in the next 4 years, injection time, injection frequency, FLS, and ISRs) and each attribute had various levels (Supplementary Table 1). DCE studies postulate that the benefit or utility of a treatment is a weighted sum of its attributes, where the estimated weights reflect respondents' perceptions about the strength of preference for each attribute level and can be used to calculate patients' willingness to trade-off among attribute levels. The weights for each attribute level, indicating strength of preference for each attribute level, were estimated by means of statistical analysis of the series of treatment choices.

The draft survey instrument was pretested before the U.S. study using in-depth, in-person interviews with fifteen adult patients with a self-reported physician diagnosis of MS. The interviews were used to evaluate and improve the understandability of the survey instrument, appropriateness of descriptive information, level of difficulty of the choice questions, and ranges of attribute levels. The pretests indicated that the information in the survey and the survey questions were understood and accepted by participants and the survey was neither too long nor too complex. The survey was adapted for the United Kingdom and France, translated (for France), and then refined based on five semi-structured in-person interviews in each country.

Experimental Design

The treatments and treatment pairs were determined using a best-practice experimental design with statistical properties that optimize the statistical information generated from a given sample size (26). The U.K. and French studies used the experimental design developed for the U.S. and German studies (21;23).

Hypothetical treatments were constructed using the information in Supplementary Table 1. The treatment attributes and levels of the hypothetical treatments were informed by consultation with experts and characteristics of currently available injectable DMTs as well as levels that may characterize

treatments that are currently unavailable, but may be possible in the future. DCE (and other stated preference methods) permit the exploration of preferences for treatment features that lie outside the range of currently available treatments. This can help us understand the importance of treatment features over a broader range than presently observed, but one that could be realized in the future.

Each respondent was asked nine choice questions, including one repeated choice question. Each choice question included two hypothetical injectable MS treatments and respondents were asked to indicate which treatment they would choose if these were the only two available. Example treatment-choice questions from the U.K. and France surveys are shown in Figure 1. Patients were provided with a hypothetical reference scenario (progressing from mild to moderate symptoms, as described using the Hohol, or Disease Steps, scale) (27) to ensure that each patient was considering the same initial disability level and the same change in disability when answering questions. The Hohol scale was selected because of the simplicity of the scale compared with the Expanded Disability Status Scale.

Study Sample

Nielsen Healthcare (formerly Harris Interactive) invited Web panelists with MS in the United Kingdom and France to participate in an online survey by means of email invitations with a link to the survey. All respondents were aged 18 years or older and had a self-reported physician diagnosis of MS. Data collection was terminated when the target sample of 100 respondents was achieved for each country. The 25-min online survey was administered in the United Kingdom in April 2014 and in France in July and August of 2014. Patients were given Harris Interactive Loyalty Points if they completed at least one treatment-choice question in the survey. These points may be accumulated by panel members and exchanged for cash or gifts. The study was approved by one of RTI International's Institutional Review Boards (Federalwide Assurance #3331). All patients provided informed consent before their inclusion in the study.

Model and Analysis

These studies replicated the statistical analysis plan completed in the U.S. and German studies (21;23).

Treatment-choice data were analyzed using a mixed logit (or random parameter logit [RPL]) model, which controls for heterogeneity in preferences and the panel nature of data (28;29). The dependent variable was the treatment (Medicine A or Medicine B) selected in each DCE question for each respondent. Independent variables included all attribute levels in Supplementary Table 1. All independent variables were effect coded with normally distributed random parameters.

RPL regression parameters are preference weights, or utilities, of attribute levels. The vertical distance, or difference, between preference weights associated with different levels of an attribute measures the relative importance, or utility differences, of the corresponding change in each attribute (30).

The estimated preference weights were used to calculate MAE estimates for various changes in treatment-related features and outcomes. MAE is a measure of the relative importance of efficacy and is higher when changes have a greater perceived importance. It is calculated as the delay in disability progression (in years) or the reduction in the number of relapses that would exactly offset the perceived importance of a given worsening in dosing frequency or a given worsening in FLS. The MAE is the mean minimum level of change in efficacy that respondents consider to be equivalent to a given change in treatment-related attributes or outcomes. The mean MAE is calculated as the ratio of the relative importance of a given change in a treatment attribute divided by the relative importance of a unit change in either the delay in disability or the number of relapses.

For example, the MAE in terms of delay in disability progression for an increase in dosing frequency from 1 to 12 times per month is calculated as the ratio of the relative importance of the change in dosing frequency (i.e., the vertical distance between the preference weights corresponding to 12 and 1 doses per month) to the relative importance of a 1-year delay in disability progression. To calculate the denominator, the relative importance of a 1-unit change in efficacy was calculated using linear interpolation between the effect coded parameter estimates over the utility range defined in the ratio's numerator.

RESULTS

Sample

In both the United Kingdom and France, 100 respondents completed the survey and provided sufficient data for analysis. Nielsen Healthcare invited 2,111 individuals in France to be screened for participation in the survey. Of those invited, 521 (25 percent) responded to the invitation. Of those who responded, 127 (24 percent) were eligible to participate. Of those who were eligible to participate, 123 (97 percent) consented to participate. Of those who were eligible and consented to participate, 100 (81 percent) completed the survey. In the United Kingdom, Nielsen Healthcare invited 1,246 patients in the United Kingdom to participate in the survey. Of those invited, 455 individuals (37 percent) responded to the invitation. Of those who responded, 141 (31 percent) were eligible to participate. Of those who were eligible to participate, 129 (91 percent) consented to participate. Of those who were eligible and consented to participate, 100 (78 percent) completed the survey.

Panel A: From UK study
 Considering all features below, please tell us whether you would prefer Medicine A or Medicine B, and which medicine you would be more likely to miss.

Medicine Feature	Medicine A	Medicine B			
Number of years until MS symptoms get worse					
Number of flare-ups in the next 4 years	3 flare-ups in the next 4 years	4 flare-ups in the next 4 years			
Injection time	10 seconds	3 seconds			
Frequency of injections	Weekly (4 times each month)	Monthly (1 time each month)			
Flu-like symptoms	For 3 days after <u>some</u> injections	For 1 day after <u>some</u> injections			
Injection-site reactions	Mild	None			
Which would you choose?	<input type="radio"/>	<input type="radio"/>			
How likely is it that you would miss doses with these medicines?	<input type="radio"/> Much more likely to miss doses with Medicine A	<input type="radio"/> A little more likely to miss doses with Medicine A	<input type="radio"/> Equally likely to miss doses with Medicine A and Medicine B	<input type="radio"/> A little more likely to miss doses with Medicine B	<input type="radio"/> Much more likely to miss doses with Medicine B

Note: This is a screenshot from the final online survey in the UK.

Panel B: From French study
 En tenant compte de toutes les caractéristiques ci-dessous, indiquez si vous préférez le Médicament A ou le Médicament B et lequel de ces médicaments vous risquez le plus d'oublier de prendre.

Caractéristiques	Médicament A	Médicament B			
Nombre d'années jusqu'à la dégradation des symptômes					
Nombre de poussées au cours des 4 prochaines années	1 poussée dans les 4 prochaines années	3 poussées dans les 4 prochaines années			
Durée de l'injection	10 secondes	3 secondes			
Fréquence des injections	2 fois par mois (bimensuel)	1 fois par mois (mensuel)			
Symptômes grippaux	Pendant 3 jours suite à <u>chaque</u> injection	Pendant 3 jours suite à <u>certaines</u> injections			
Réactions au site d'injection	Leger	Aucun			
Lequel choisiriez-vous?	<input type="radio"/>	<input type="radio"/>			
Quelle est la probabilité d'oublier de prendre vos prises avec ces médicaments?	<input type="radio"/> Il est bien plus probable que j'oublie de prendre mes prises avec le Médicament A	<input type="radio"/> Il est légèrement plus probable que j'oublie de prendre mes prises avec le Médicament A	<input type="radio"/> Il est aussi probable que j'oublie de prendre mes prises avec le Médicament A que le Médicament B	<input type="radio"/> Il est légèrement plus probable que j'oublie de prendre mes prises avec le Médicament B	<input type="radio"/> Il est bien plus probable que j'oublie de prendre mes prises avec le Médicament B

Note: This is a screenshot from the final online French survey.

Figure 1. Example treatment-choice questions.

Sample characteristics are reported in Table 1. In both samples, the mean age was 46 years and the mean time since receiving a diagnosis of MS was 10 years. Both samples were weighted toward female respondents (59 percent and 71 percent in the United Kingdom and France, respectively) with relapsing-remitting MS (45 percent and 51

percent, respectively). Twenty-four percent of respondents in the United Kingdom were treatment-naïve and 11 percent of respondents in France were treatment-naïve. A total of 50 percent of the respondents in the U.K. sample were injection-naïve, while only 24 percent were injection-naïve in the French sample.

Table 1. Characteristics of survey respondents

Characteristic	U.K. (N = 100)	France (N = 100)
Age, mean (SD)	46.4 (12.4)	46.2 (10.2)
Female, n (%)	59 (59.0%)	71 (71.0%)
Employed full-time, n (%)	30 (30.0%)	30 (30.0%)
Type of MS, n (%)		
Relapsing-remitting	45 (45.0%)	51 (51.0%)
Secondary-progressive	30 (30.0%)	21 (21.0%)
Primary-progressive	15 (15.0%)	15 (15.0%)
Progressive-relapsing	4 (4.0%)	4 (4.0%)
Number of years since MS diagnosis, mean (SD)	10 (7)	10 (6)
Number of relapses in past 2 years, mean (SD)	3.6 (4.3)	2.9 (4.2)
Has never been prescribed a medicine to take on a regular basis to manage MS, n (%)	24 (24.0%)	11 (11.0%)
Injection-naïve, n (%) ^a	50 (50.0%)	24 (24.0%)
MS symptoms, on most days, n (%) ^b		
No limitations	9 (9.0%)	13 (13.0%)
Mild symptoms	24 (24.0%)	29 (29.0%)
Moderate symptoms	21 (21.0%)	18 (18.0%)
Need cane for long distances	11 (11.0%)	12 (12.0%)
Need cane for short and long distances	18 (18.0%)	18 (18.0%)
Need bilateral support	7 (7.0%)	2 (2.0%)
Need wheelchair or scooter	10 (10.0%)	8 (8.0%)
Seconds it takes for medicine to come out of injection device, mean (SD)		
Among those currently receiving regular injections	8.3 (3.0) (n = 19)	8.5 (5.0) (n = 41)
Among those who have received regular injections in the past	8.3 (3.1) (n = 22)	9.1 (6.4) (n = 25)
Ever had flu-like symptoms caused by your MS medicines, n (%)	31 (31.0%)	53 (53.0%)
Among those who have ever received regular injections: has had a mild injection-site reaction, n (%)	35 (71.4%) (n = 49)	48 (64.0%) (n = 75)

MS, multiple sclerosis; SD, standard deviation; U.K., United Kingdom.

^a Has never received injections on a regular basis to treat MS.

^b As described in the Hohol scale.²⁶

Preference Weights

The results of the RPL estimation (Figure 2) show that respondent preferences were generally consistent with the expectation that better outcomes (e.g., better efficacy, fewer side effects, less frequent administration) would be preferred to worse outcomes in the United Kingdom and France. The U.K. results suggest that four relapses are preferred to three relapses and a longer injection (10 sec) is preferred to a shorter injection (3 sec), but in both cases the differences between the pairs of preference weights are not statistically significantly different, suggesting that respondents did not distinguish between these

levels when making choices in the survey. Regression results are shown in Supplementary Tables 2 and 3.

In the United Kingdom, improving the time until disability progression from 2 to 4 years and reducing injection frequency from “daily” to “every 2 weeks” had a relative importance (vertical distance between preference weights for attribute levels) of 2.9 and 2.6, respectively (Figure 2a). The importance of these changes was similar to the importance of reducing the frequency of FLS from 3 days after every dose to none (2.5), and between 3.5 and 3 times more important, respectively, than reducing the number of relapses in the next year from 4 to 1 (0.8).

In France, improving the time until disability progression from 2 to 4 years and reducing injection frequency from “daily” to “every 2 weeks” had a relative importance of 3.0 and 3.5, respectively (Figure 2b). The importance of this change in dosing frequency was similar to the importance of reducing FLS from 3 days after every dose to none (3.4), and more than 2.5 times as important as reducing the number of relapses in the next year from 4 to 1 (1.3). In both countries, these changes in disability progression, dosing frequency, and FLS were more important than changes in the other attributes assessed, and the relative importance of improvements in injection time and ISRs was much smaller than changes among the other attributes.

Minimum Acceptable Efficacy

The MAEs (in terms of delay in disability progression and decrease in relapses over 5 years) are shown for different changes in dosing frequency and FLS in Table 2 for both countries. The MAEs were higher for the U.K. sample than for the French sample, although the differences are not statistically significant. On average, respondents in the United Kingdom and France, respectively, would trade an increase in injection frequency from 2 to 30 times per month for 9.1 and 8.2 fewer relapses in the next 5 years or for 2.4 and 2.5 years of delay in disability progression. As measured by MAE, these changes in dosing frequency from 2 to 30 times per month are similar to the changes in FLS from none to 3 days after each injection.

DISCUSSION

In this study, both in the United Kingdom and France, lengthening the time until disability progression, reducing injection frequency, and reducing FLS were key drivers of patient preferences. These results suggest that people with MS might be willing to trade off efficacy for improvements in dosing frequency and improvements in FLS, and the MAE estimates indicate the tradeoffs that might, on average, be acceptable. Most of the mean MAE estimates were statistically similar for the United Kingdom and France.

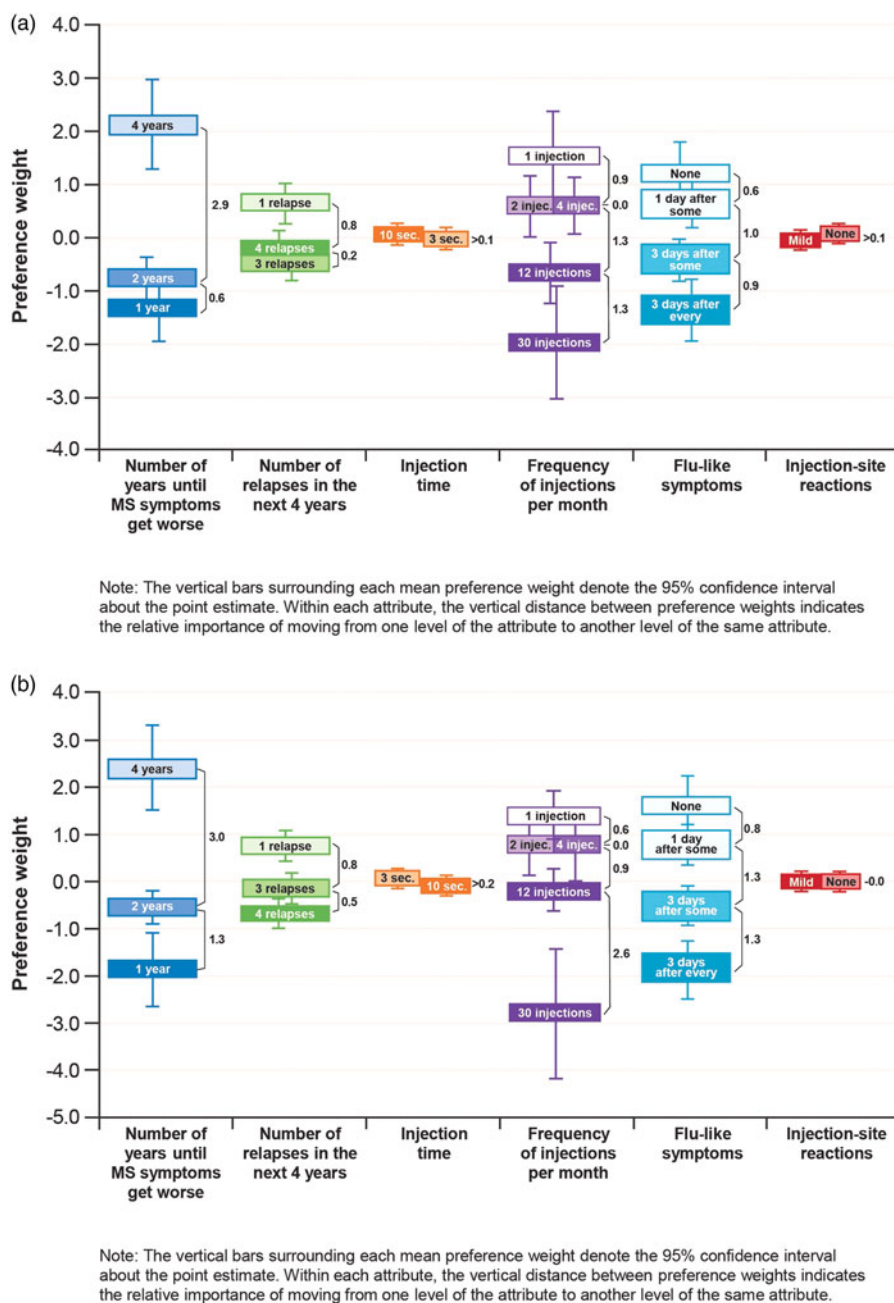


Figure 2. Preference weights for patients surveyed in the United Kingdom (a) and France (b); (N = 100 each).

These results are similar to the study findings in the United States and Germany. Among 192 respondents from the United States, we found that similar levels of importance were placed on reduction in injection frequency or FLS as were placed on treatment efficacy (21). The German study, in 189 patients, also showed similar levels of importance placed on improvements in the frequency of dosing and delay in disability progression (23).

The present study was the first study examining patient preferences for DMT attributes in the United Kingdom and France, and is the only study to specify the mild side effects separately and evaluate tradeoffs among efficacy and different side effects.

This study is focused only on injectable treatments. Wilson (19;20) is a comprehensive study, but carried out only in the United States with a geographically limited sample. These results, combined with results from previous studies on patient preferences for MS therapies, suggest that some changes in injection frequency of MS injectable treatments can be as important to patients as some changes in efficacy or side effect attributes of those treatments. These studies highlight the perceived benefits of therapies with lower dosing frequencies, as a reduction in dosing complexity and injection-related side effects may lead to improvements in patient adherence.

Table 2. MAE estimates (95% CI)

Change in treatment attribute or outcome	MAE in terms of the number of years of delay until MS symptoms worsen (95% CI)		MAE in terms of decreases in the number of relapses in the next 5 years (95% CI)	
	U.K. sample (N = 100)	French sample (N = 100)	U.K. sample (N = 100)	French sample (N = 100)
Attribute	To	French sample (N = 100)	U.K. sample (N = 100)	French sample (N = 100)
Frequency of injections per month	From			
	1	2.1 (1.3, 2.7)	1.2 (0.6, 1.9)	7.8 (3.1, 23.1) ^a
	1	3.0 (2.1, 3.8) ^a	2.9 (2.1, 3.9)	12.4 (5.7, 38.0) ^a
	2	1.5 (0.4, 2.1)	0.7 (0.0, 1.5)	4.4 (0.9, 12.7) ^a
	2	2.4 (1.5, 3.1)	2.5 (1.2, 3.6)	9.1 (3.9, 26.4) ^a
Flu-like symptoms	None	1.0 (-0.3, 1.7)	0.6 (0.1, 1.3)	2.1 (-0.5, 8.9)
	None	1.7 (1.1, 2.3)	1.6 (0.9, 2.3)	5.7 (2.4, 18.8) ^a
	None	2.3 (1.7, 3.1)	2.5 (1.8, 3.4)	9.0 (4.4, 29.8) ^a
	1 day after some injections	1.3 (0.5, 1.9)	1.0 (0.4, 1.6)	3.6 (1.1, 10.4) ^a
	1 day after some injections	1.9 (1.4, 2.6)	1.9 (1.2, 2.8)	7.0 (3.6, 22.0) ^a
	3 days after some injections	1.2 (0.5, 1.8)	1.0 (0.5, 1.7)	3.3 (0.8, 12.4) ^a
	1 day after some injections			
	3 days after some injections			
	3 days after each injection			
	3 days after each injection			
	3 days after each injection			

CI, confidence interval; MAE, minimum acceptable efficacy; MS, multiple sclerosis; U.K., United Kingdom.

^a For this change, the MAE was calculated by extrapolating preferences beyond the range for the levels included in the study.

One potential limitation of this study is that a substantial number of patients in both samples were injection-naïve, which might be expected to limit their preferences for DMT treatment attributes. However, we found no evidence that preferences among patients with and without injection experience were statistically different from one another (data available upon request).

General limitations of this type of assessment include the evaluation of hypothetical treatment profiles as opposed to actual treatments, and the omission of possible important contextual factors, such as cost, which may influence actual treatment choices. In particular, because hypothetical treatment profiles were constructed, the selection of attributes and levels used in this study may have influenced the results observed. However, the study design was informed by the reviews of medical experts to ensure that the attributes selected were clinically meaningful and reflective of the treatment attributes that, based on their experience, influence treatment choice.

The survey was also pretested with patients to explore whether there were salient treatment attributes that were not included in the survey. The pretests and consultations with experts suggested that the selection of attributes and levels was clinically appropriate and did not noticeably influence survey responses. In addition, there is always the possibility that small samples in surveys may not be representative of the overall patient populations in the areas studied. Furthermore, the lack of data on the populations of MS patients limit our ability to evaluate the representativeness of the sample. However, the majority of study respondents in the United Kingdom and France were women (59 percent and 71 percent, respectively), which is consistent with the results of epidemiological studies showing that MS occurs more often in women than men (31–34).

The physician diagnoses of MS were self-reported, which is challenging because not all patients will know, but the design attempted to minimize the possibility of misidentification by screening respondents for study eligibility without revealing the subject of the survey. The screening question involved selecting health conditions that had been diagnosed by a health care provider from a prespecified list that included MS. While this will not eliminate the chance of misidentification, it is likely to reduce the risk of strategic bias in self-reporting. There are also acknowledged limitations of the use of an online panel to recruit patients, including potential sample bias and mode effects. However, research has shown that results from online DCEs are, in general, not statistically different from those elicited through face-to-face or telephone interviews (35;36).

Strengths of this study include the design of the survey in collaboration with clinical experts and pretesting using in-depth interviews with MS patients. The treatment-choice data were analyzed using RPL methods that avoid estimation bias

from unobserved variation across sample and within-sample correlation for each patient. In this study, unlike previous studies, the importance of changes in selected attributes are calculated in terms of MAE, which allows more direct comparison between two countries. To our knowledge, these measures have not been used previously to explore the relative importance of DMT treatment features.

There are no direct implications of this research for public policy. However, the results may be of importance among physicians and payers who influence the selection of treatments to add to formulary, reimburse, or offer to patients.

In conclusion, the results of these surveys suggest that some changes in the injection frequency and side effects (FLS) of injectable treatments for MS can be as important to some patients as improvements in treatment efficacy. This information may be useful to providers when discussing treatment choices with their patients.

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SUPPLEMENTARY MATERIAL

Supplementary Table 1:

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Supplementary Table 2:

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Supplementary Table 3:

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CONFLICTS OF INTEREST

C. Poulos is an employee of RTI Health Solutions, an independent scientific research organization. J. Posner was an employee of RTI Health Solutions at the time of the study. The study that is the subject of this manuscript was conducted by RTI Health Solutions and funded by Biogen. E. Kinter, J. van Beek and K. Christensen are employees of Biogen. The authors have full control of all primary data and agree to allow the journal to review the data if requested. *Ethical standards:* The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2008.

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