

Designing Multi-arm Multistage Adaptive Trials for Neuroprotection in Progressive Multiple Sclerosis

Vivien Li, MBBS, FRACP,* Baptiste Leurent, PhD,* Frederik Barkhof, MD, PhD, Marie Braisher, MSc, Fay Cafferty, PhD, Olga Ciccarelli, MD, PhD, Arman Eshaghi, MD, PhD, Emma Gray, PhD, Jennifer M. Nicholas, PhD, Mahesh Parmar, PhD, Guy Peryer, PhD, Jenny Robertson, MPsych(Health), Nigel Stallard, PhD, James Wason, PhD, and Jeremy Chataway, MD, PhD

Correspondence

Dr. Li
vivien.li@unimelb.edu.au

Neurology® 2022;98:754-764. doi:10.1212/WNL.0000000000200604

Abstract

There are few treatments shown to slow disability progression in progressive multiple sclerosis (PMS). One challenge has been efficiently testing the pipeline of candidate therapies from preclinical studies in clinical trials. Multi-arm multistage (MAMS) platform trials may accelerate evaluation of new therapies compared to traditional sequential clinical trials. We describe a MAMS design in PMS focusing on selection of interim and final outcome measures, sample size, and statistical considerations. The UK MS Society Expert Consortium for Progression in MS Clinical Trials reviewed recent phase II and III PMS trials to inform interim and final outcome selection and design measures. Simulations were performed to evaluate trial operating characteristics under different treatment effect, recruitment rate, and sample size assumptions. People with MS formed a patient and public involvement group and contributed to the trial design, ensuring it would meet the needs of the MS community. The proposed design evaluates 3 experimental arms compared to a common standard of care arm in 2 stages. Stage 1 (interim) outcome will be whole brain atrophy on MRI at 18 months, assessed for 123 participants per arm. Treatments with sufficient evidence for slowing brain atrophy will continue to the second stage. The stage 2 (final) outcome will be time to 6-month confirmed disability progression, based on a composite clinical score comprising the Expanded Disability Status Scale, Timed 25-Foot Walk test, and 9-Hole Peg Test. To detect a hazard ratio of 0.75 for this primary final outcome with 90% power, 600 participants per arm are required. Assuming one treatment progresses to stage 2, the trial will recruit $\approx 1,900$ participants and last ≈ 6 years. This is approximately two-thirds the size and half the time of separate 2-arm phase II and III trials. The proposed MAMS trial design will substantially reduce duration and sample size compared to traditional clinical trials, accelerating discovery of effective treatments for PMS. The design was well-received by people with multiple sclerosis. The practical and statistical principles of MAMS trial design may be applicable to other neurodegenerative conditions to facilitate efficient testing of new therapies.

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*These authors contributed equally to this work as co-first authors.

From the Florey Institute of Neuroscience and Mental Health (V.L.), University of Melbourne; Department of Neurology (V.L.), Royal Melbourne Hospital, Australia; Department of Medical Statistics (B.L., J.M.N.) and International Statistics and Epidemiology Group (B.L.), London School of Hygiene and Tropical Medicine, UK; Department of Radiology and Nuclear Medicine, Neuroscience Campus Amsterdam (F.B.), VU University Medical Center, Amsterdam, the Netherlands; Queen Square Institute of Neurology and Centre for Medical Image Computing (F.B.), Department of Neuroinflammation, Queen Square Multiple Sclerosis Centre (M.B., O.C.), and NMR Unit, Department of Neuroinflammation (A.E.), Faculty of Brain Sciences, UCL Queen Square Institute of Neurology; MRC Clinical Trials Unit at UCL, Institute of Clinical Trials and Methodology (F.C., M.P., J.C.), and Department of Computer Science, Centre for Medical Image Computing (A.E.), University College London; National Institute for Health Research (F.B., O.C., J.C.), University College London Hospitals Biomedical Research Centre; UK Multiple Sclerosis Society (E.G., G.P., J.R.), London; Faculty of Medicine and Health Sciences (G.P.), University of East Anglia, Norwich; Statistics and Epidemiology, Division of Health Sciences (N.S.), Warwick Medical School, University of Warwick, Coventry; and Population Health Sciences Institute (J.W.), Newcastle University, UK.

Go to [Neurology.org/N](https://www.neurology.org/N) for full disclosures. Funding information and disclosures deemed relevant by the authors, if any, are provided at the end of the article.

The Article Processing Charge was funded by Medical Research Council.

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Glossary

9HPT = 9-Hole Peg Test; **EDSS** = Expanded Disability Status Scale; **MAMS** = multi-arm multistage; **MS** = multiple sclerosis; **PBVC** = percentage of brain volume change; **PMS** = progressive multiple sclerosis; **PPI** = patient and public involvement; **PPMS** = primary progressive multiple sclerosis; **RRMS** = relapsing-remitting multiple sclerosis; **SPMS** = secondary progressive multiple sclerosis; **T25FW** = Timed 25-Foot Walk.

Progressive multiple sclerosis (PMS) is a significant health problem worldwide¹ and has considerable financial costs for health care systems, patients, and their caregivers, with costs increasing at higher levels of disability.²⁻⁴ Despite extensive efforts, there are few proven therapies for PMS. Compared to the predominantly inflammatory pathology in relapsing multiple sclerosis (MS) targeted by current treatments, the neurodegenerative processes driving progression in PMS are complex and less well-defined.^{5,6} There is a pipeline of candidate therapies from pre-clinical studies, but the challenge is testing them efficiently in clinical trials with appropriate outcome measures to determine whether they can successfully slow disability progression.

One potential avenue is improving efficiency of trials by incorporating adaptive elements in a multi-arm multistage (MAMS) platform design. MAMS trials aim to evaluate multiple experimental arms and seamlessly integrate traditional phase II and III evaluations into a single trial. They have been successful in accelerating evaluation of therapies and changing practice in other disease settings, such as cancer⁷ and infectious diseases.⁸ They are also increasingly being considered for neurologic conditions such as Alzheimer disease,⁹ Parkinson disease¹⁰ and motor neuron disease.^{11,12} These neurodegenerative conditions share commonalities with PMS, where there is a marked translational gap between the relative abundance of early phase trials stemming from increased understanding of disease pathobiology and lack of positive phase III trials leading to disease-modifying treatments.

Adaptive MAMS platform designs offer flexible features that can provide efficiencies at various levels¹³ (Table 1). These include simultaneous evaluation of multiple treatments against a common standard of care, reducing both time and numbers of patients required; the ability to add new treatments as they become relevant, avoiding lengthy setup times for multiple trials; dropping treatments that are not showing sufficient promise, allowing redirection of resources; and incorporation of the traditional separate phase II and III evaluations within a single protocol with seamless transitions.

With the aim of designing a MAMS trial in PMS, the UK MS Society Expert Consortium for Progression in MS Clinical Trials set up 4 working groups on outcome measures, trial design, treatment selection, and trial infrastructure. Each group included members with relevant expertise and worked closely with the patient and public involvement (PPI) group throughout the development process. The work of the treatment selection working group on identifying and

shortlisting candidate treatments, focusing on licensed drugs that can be repurposed, has been reported elsewhere.¹⁴

This article describes the work of the trial design and outcome measures working groups. We discuss key elements of the MAMS trial design based on evaluation of 3 candidate treatments against standard of care in 2 analysis stages, including selection of the primary interim and final outcomes, sample size, and other statistical considerations.

Methods

Outcome Measures

The outcome measures working group comprised individuals with expertise in MS trials, imaging, and biomarkers, as well as people with MS with lived experience of the condition. The group reviewed the literature to determine outcome measures relevant to a MAMS trial evaluating predominantly neuroprotective treatments. Individual members submitted proposed outcome measures based on their expertise with final prioritization of outcomes determined in consensus meetings.

The MAMS design allows distinct interim (stage 1) and final (stage 2) outcomes. The final outcome should be clinically derived and relevant to patients and regulators. The interim outcome serves as an early indicator of whether a treatment is likely to be effective and hence should be continued into the second stage of the trial while minimizing the likelihood of ceasing truly effective treatments. It should reflect the underlying association between the treatment and the clinical outcome. The absence of effect on the interim outcome should be indicative of the absence of effect on the final outcome, although the converse may not necessarily hold.¹⁵

Trial Design

The trial design working group, comprising experts in design and implementation of MAMS trials, statisticians, and MS clinicians, was tasked with generating design options for running an efficient, scalable, and flexible clinical trial by exploring different scenarios to determine the best design type. The group reviewed data from phase II and III randomized controlled PMS trials from January 1, 2009, to January 1, 2019, to inform key design measures for both stage 1 and 2 analyses, such as effect size, and the relationship between the interim and final outcomes.

To assess the statistical operating characteristics of the trial (e.g., type I and type II error rates), we simulated multiple

Table 1 Glossary of Terms

Adaptive trial: A trial with scheduled interim analyses to evaluate observed data during the course of the trial, which inform and allow prespecified changes to be made while maintaining the overall integrity of the trial

Platform trial: A long-term trial that evaluates multiple hypotheses, defined around core elements but allowing for flexibility, such as adding new treatment arms or changing the standard of care

Multi-arm, multistage trial: A trial that tests several interventions simultaneously against a shared contemporaneous control group, with potentially seamless transition from phase II to phase III evaluations

Multi-arm multistage platform and adaptive trial: All the above elements combined in the same trial

Phase II trials: Trials typically involving up to a few hundred participants aiming to demonstrate that the treatment is sufficiently safe and promising, usually demonstrating effect on an intermediary (biological) outcome

Phase III trials: Trials typically involving some hundreds or thousands of participants aiming to provide definitive clinical evidence of treatment efficacy

Stage 1 or interim analysis: Analyses performed during the course of the trial at a prespecified timepoint to assess for early evidence of activity or futility of a treatment

Stage 2 or final analysis: Analyses performed at the end of a trial to assess the efficacy of promising treatments based on interim analysis, usually based on relevant clinical outcomes

Type I error or alpha: Rejecting a true null hypothesis (false-positive finding)

Type II error: Accepting a false null hypothesis (false-negative finding)

Power: The probability that a statistical test will reject a false null hypothesis, i.e., probability of detecting a specific difference when it truly exists; equal to 1 – type II error rate

trials with different correlation structures for the treatment effects on stage 1 and stage 2 outcomes, under different treatment effect assumptions. We also modeled the expected trial progress over time based on different assumptions (such as recruitment rates) and design parameters (such as sample size and treatment stopping rules) to the second stage. Further details of the simulation methods are reported in eAppendix 1 and 2 (links.lww.com/WNL/B900). To support the design of the stage 1 analysis, we analyzed brain atrophy data from the MS-STAT1¹⁶ and ASCEND¹⁷ clinical trials (for full Methods, see eAppendix 1, links.lww.com/WNL/B900). Modeling was conducted in Stata version 15 (StataCorp) and Microsoft Excel 2016.

Patient and Public Involvement

A PPI strategy group was involved since the earliest conception of the project, including members of each of the expert consortium working groups. The PPI strategy group included 4 members of the MS Society Research Network (patients with MS) and the MS Society Public Involvement Officer. They contributed to discussions as the project developed and focused on ensuring that the research would meet the needs of the MS community. Additional workshops attended by a total of 34 people with MS held in 3 UK locations brought in further expertise of people with MS on topics including relevance, feasibility, and acceptability of all aspects of the trial design as well as recruitment and engagement strategies.

Data Availability

Data not published in this article will be made available by request from any qualified investigator.

Review of Previous Trials

Our review identified 15 eligible phase II (n = 8) and phase III (n = 7) randomized trials in PMS (Table 2). The median trial

size was 374 participants (range 54–1,651) and median follow-up duration was 2 years (range 1–4.5 years). Trials included secondary progressive MS (SPMS) (n = 6), primary progressive MS (PPMS) (n = 6), and mixed PMS (n = 3). Confirmed disability progression on the Expanded Disability Status Scale (EDSS) was reported in 9 trials at different time intervals ranging from 3 to 6 months and a composite outcome was reported in 4. Two (29%) of the phase III trials (siponimod, ocrelizumab) and 4 (50%) of the phase II trials (ibudilast, lipoic acid, biotin, simvastatin) were positive for their primary endpoints.

Proposed MAMS Trial Design

Overview

We propose the design of a 2-stage MAMS trial in PMS: 1 interim analysis to examine early evidence of treatment effect (stage 1) and 1 final confirmatory analysis of efficacy (stage 2). The trial would include 4 arms in stage 1: 1 standard of care (control) arm and 3 experimental arms. Any treatment that is sufficiently promising at the interim analysis will progress to stage 2, which will continue until the required number of events is reached, as represented in Figure 1.

It is expected that in the initial phase of the trial the standard of care arm for most participants will comprise best supportive care. Whereas ocrelizumab and siponimod have been approved for PMS, these treatments are not currently available to or suitable for all patients, in particular nonambulatory patients, who would be eligible for this proposed trial. If an efficacious therapy is subsequently found, this would then become the standard of care for future participants entering the platform.

The number of experimental arms was informed by feasibility constraints and the treatment selection group's work on

Table 2 Description of PMS Randomized Controlled Trials Included in the Review

Trial (drug assessed)	Participants	PPMS/SPMS	Trial duration, mo	Data on confirmed progression ^a	Data on whole brain atrophy ^b	Reference
Phase III						
EXPAND (siponimod)	1,651	SPMS	36	EDSS 3 mo, EDSS 6 mo	Direct change	33
ASCEND (natalizumab)	887	SPMS	36	Composite 6 mo	Direct change	17
ORATORIO (ocrelizumab)	732	PPMS	Up to 50	EDSS 3 mo, EDSS 6 mo, composite 3 mo, composite 6 mo	Direct change	42
INFORMS (fingolimod)	823	PPMS	Up to 60	EDSS 3 mo, composite 3 mo	Direct change	22
PROMESS (cyclophosphamide)	138	SPMS	24	EDSS 4 mo	None	43
CUPID (dronabinol)	498	PPMS/SPMS	36	EDSS 6 mo	Direct change	44
OLYMPUS (rituximab)	439	PPMS	24	EDSS 3 mo	Indirect change	45
Phase II						
Lamotrigine	120	PPMS	24	None	Direct change	46
SPRINT-MS (ibudilast)	255	PPMS/SPMS	24	EDSS 5 mo	Brain parenchymal fraction	25
Lipoic acid	54	SPMS	24	None	Direct change	47
MS-SPI (biotin)	154	PPMS/SPMS	12	EDSS 3 mo	None	48
MS-STAT1 (simvastatin)	140	SPMS	24	EDSS unconfirmed	Direct change	16
IPPoMS (idebenone)	85	PPMS	24	Composite 6 mo	Unclear	49
MS-SMART (amiloride, riluzole, and fluoxetine)	440	SPMS	24	None	Direct change	24
ARPEGGIO (laquinimod)	374	PPMS	12	EDSS 3 mo	Unclear	50

Abbreviations: EDSS = Expanded Disability Status Scale; PMS = progressive multiple sclerosis; PPMS = primary progressive multiple sclerosis; SPMS = secondary progressive multiple sclerosis.

^a Disability progression confirmed after X months.

^b Direct change: registration-based techniques, such as boundary shift integral and SIENA (structural image evaluation, using normalisation, of atrophy).

number of repurposed therapies ready for clinical testing.¹⁴ Participants will be randomized with an equal probability between each of the 4 arms (1:1:1:1 ratio). In a standard multi-arm trial with n experimental arms, the optimal allocation ratio would be $\sqrt{n} : 1$ in favor of the control arm. This is because the control participants contribute to each of the pairwise comparisons. However, for a MAMS trial, this depends on the number of arms continuing into stage 2, which is unknown.¹⁸ Unequal allocation would also make the trial less attractive to people with MS, as it results in a lower likelihood of being randomized to an experimental arm.

Choice of Stage 2 (Final) Primary Outcome

The classic measurement tool and regulatory standard has been the EDSS,¹⁹ used to determine the time to disability progression. Its strengths and limitations are well-documented²⁰ and numerous attempts have been made to evolve it, including using a composite measure based on progression in 1 or more of 3 endpoints: (1) increase in EDSS (of ≥ 1 point if baseline EDSS was < 5.5 or ≥ 0.5 points if baseline EDSS was ≥ 5.5), (2) $\geq 20\%$ increase in 9-Hole Peg

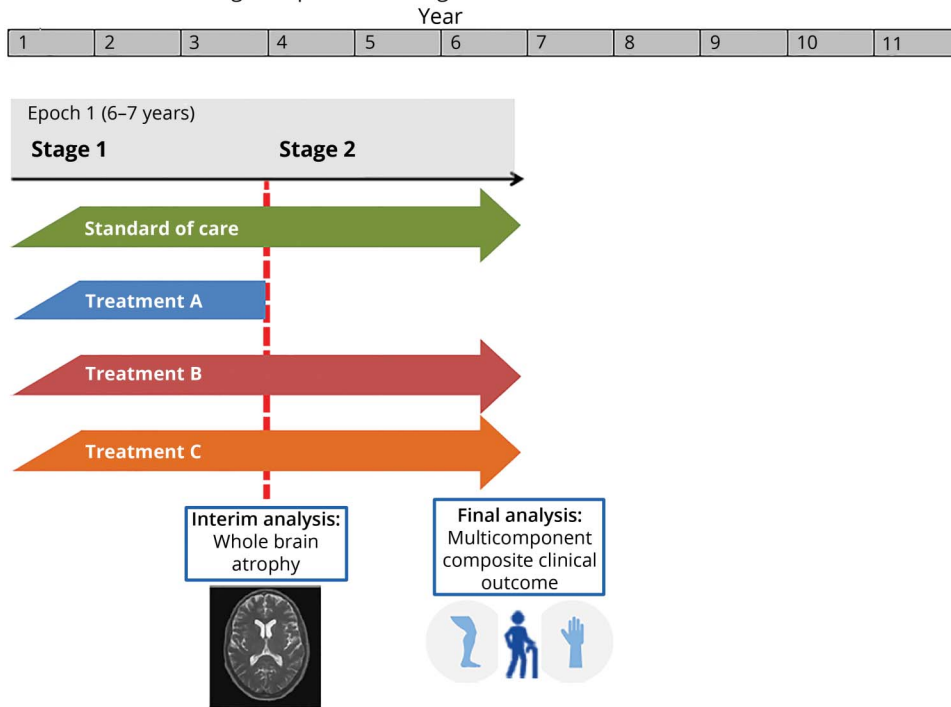
Test (9HPT), or (3) $\geq 20\%$ increase in Timed 25-Foot Walk (T25FW) (if ambulant).²¹

Composite measures achieve higher event rates than single measures (eAppendix 3, links.lww.com/WNL/B900), which can reduce trial duration and sample size. For example, in the INFORMS phase III trial of fingolimod in PPMS, $>70\%$ of participants had reached progression on the 3-month confirmed disability composite outcome by 3 years, as opposed to 50% based on EDSS alone.²² Inclusion of a measure of upper limb function also addresses the PPI group's interest in expanding the traditionally narrow EDSS inclusion criteria to include patients with higher levels of disability, to whom arm function is critical and measures of ambulation less relevant.²³

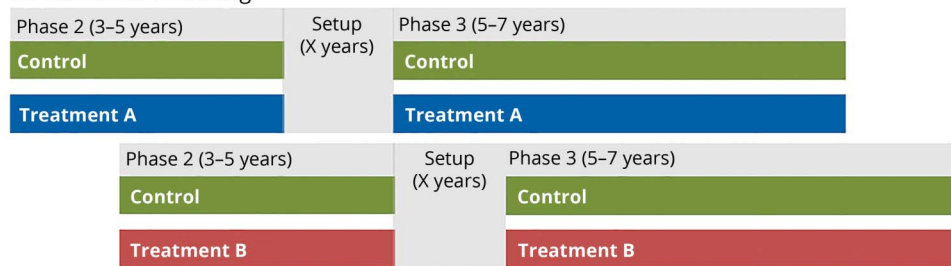
Based on these considerations, we selected time to 6-month confirmed composite disability progression as the primary outcome for the final (stage 2) analysis. The composite outcome will be measured at baseline and every 6 months until the end of the follow-up. The time to progression will be from randomization until date of the initial disability progression (if subsequently

Figure 1 Schematic Representation of the MAMS Trial Comparing 3 Experimental Arms With Standard of Care in 2 Stages and Traditional 2-Arm Phase II and III Clinical Trials

A. Multi-arm, multistage adaptive trial design



B. Traditional trial design



(A) Multi-arm multistage (MAMS) adaptive trial design. (B) Traditional trial design.

confirmed). Based on earlier trials (eAppendix 3, links.lww.com/WNL/B900), we expect the rate of 6-month confirmed composite disability progression to be around 50% at 3 years.

Choice of Stage 1 (Interim) Primary Outcome

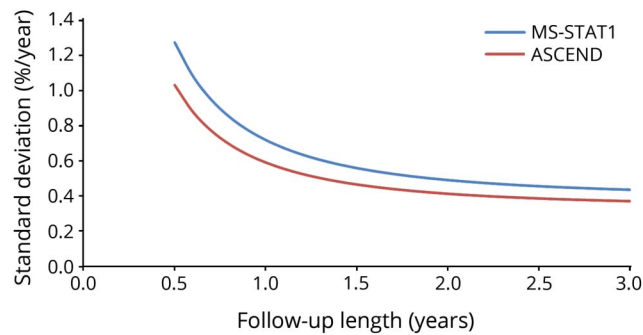
Whole brain atrophy on MRI, measured as annualized percentage of brain volume change (PBVC), was selected as the primary interim outcome, based on the initial candidate drugs having primarily neuroprotective mechanisms of action. Brain atrophy reflects underlying neuroaxonal loss, which contributes to accrual of disability in PMS, and has been successfully used as a primary outcome in phase II trials, including MS-STAT1,¹⁶ MS-SMART,²⁴ and SPRINT-MS.²⁵ Importantly for a multistage trial, the treatment effect size on atrophy has been found to correlate with the clinical disability endpoint in a meta-analysis of relapsing-remitting MS (RRMS) trials.²⁴

Methods of measuring PBVC include registration-based techniques (such as boundary shift integral and SIENA

[structural image evaluation, using normalisation, of atrophy]) or brain parenchymal fraction, which quantify the amount of brain tissue contained within a contour surrounding the entire brain including CSF.²⁶ Some therapies, particularly those with anti-inflammatory effects, can excessively reduce brain volume in the first months (pseudoa-trophy),²⁷ so it was recommended to assess PBVC also after at least 6 months on treatment.

We considered other imaging-based measures, including spinal cord atrophy, which contributes to MS disability progression and occurs at a faster rate than brain atrophy,²⁸ neurite indices derived from diffusion MRI, which reflect the microstructural changes of axons and dendrites, and magnetization transfer imaging, which reflects demyelination and axonal loss.¹⁹ However, technical challenges limit widespread implementation and standardization across multiple centers.²⁷ Although biofluid markers such as neurofilament light chain are associated in high concentrations with disability and brain atrophy,²⁹ there are

Figure 2 Predicted SD of Atrophy Rate for Varying Follow-up Length, Based on Modeling of MS-STAT1 and ASCEND Trial Data^{20,35}



mixed findings on whether they are sensitive to treatment and they are not ready to be used as primary outcome measures until a validated, standardized, and widely accessible assay is available, with normative values of neurofilaments across age groups. Moreover, there is divergence of their utility in relapsing MS compared to PMS.^{30,31}

Predicted Brain Atrophy Rate

Nine of the reviewed trials reported a direct measure of change in whole brain volume (eAppendix 1, links.lww.com/WNL/B900). Brain atrophy rate varied between 0.4%/year and 0.7%/year in control arms. There was no clear pattern of differences between trials in PPMS or SPMS or by follow-up length. The SD for atrophy rate decreased with increasing follow-up length, ranging from 0.59%/year to 0.78%/year over 1 year, and 0.37%/year to 0.60%/year over 2 years.

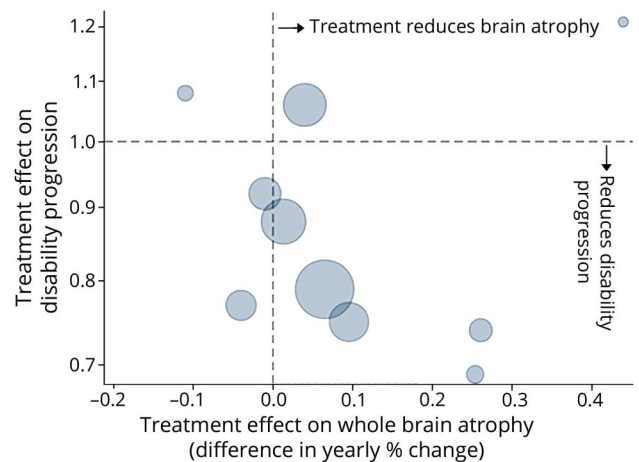
The predicted SDs based on applying our statistical model (eAppendix 1, links.lww.com/WNL/B900) to the data from MS-STAT1 and ASCEND are shown in Figure 2. The SD is expected to decrease rapidly with increasing length of follow-up, especially in the first 12 months. After 18–24 months, the reduction in SD becomes much smaller.

Timing of the interim analysis is an important consideration of adaptive designs. It should occur after accruing sufficient participant data to make a reliable decision on continuing or dropping treatment arms, but early enough relative to total trial recruitment to have value in informing adaptation of the trial design.³² Based on these considerations, PBVC after 18 months' follow-up will be assessed at interim analysis. This choice achieves a balance between reducing variance of the measure and ensuring that the interim analysis was sufficiently timely to make it worthwhile (see below). The SD at this point is predicted to be around 0.55%/year.

Treatment Effects on Brain Atrophy and Clinical Progression

A key criterion for the stage 1 outcome is the ability to identify treatments expected to be ineffective and also potentially

Figure 3 Association Between Treatment Effect on Brain Atrophy and Disability Progression in Progressive Multiple Sclerosis Trials



The size of each circle is proportional to the trial size.

effective in terms of the final (stage 2) outcome. We reviewed trials reporting treatment effect on both brain atrophy rate and clinical progression. Trial results are reported in eAppendix 2 (links.lww.com/WNL/B900) and summarized in Figure 3. There is a negative correlation, indicating that drugs with a stronger effect on reducing brain atrophy in PMS were more effective on clinical outcomes, confirming findings in RRMS.²⁶

Our trial targets a treatment effect of 25% relative reduction in the 6-month confirmed disability progression rate, that is, a hazard ratio of 0.75. This is a clinically important effect in slowing progression in ambulation, upper limb function, or disability, which has been achieved in previous trials.³³ Assuming 50% of patients experience a disability progression by 3 years in the control arm, a 25% relative reduction would equate to a 12.5% absolute difference (50% control vs 37.5% active treatment).

Based on the review of previous trials, we assumed an effective treatment would reduce the rate of whole brain atrophy by around 0.15%/year, from 0.55%/year to 0.40%/year.

Stage 1 Sample Size

The sample size for stage 1 analysis was based on pairwise comparisons of whole brain atrophy rate at 18 months between each intervention arm and standard of care. A 1-sided test is used for the interim analysis, with a treatment continuing to the second stage if there is evidence in favor of a lower atrophy rate compared to standard of care. We chose 95% power because a priority of the interim analysis is to minimize the chance of stopping an arm when the treatment is genuinely active in slowing brain atrophy (i.e., avoid false-negatives). Stage 1 alpha (type I error rate) captures the

Table 3 Trial Operating Characteristics According to Number of Effective Treatments Entering the Trial

	Number of (truly) effective treatments at start of trial, %			
	0	1	2	3
Number of experimental arms continuing to 2nd stage				
0	40.5	4.3	1.0	0.4
1	26.8	45.8	6.4	2.3
2	19.3	30.7	58.8	9.2
3	13.4	19.2	33.8	88.1
Detected at least 1 truly effective treatment (power)	—	87.3	96.2	98.3
At least 1 ineffective treatment found significant (type I error)	3.7	2.6	1.3	—

Results are column %, based on 10,000 simulations for each of the 4 scenarios (number of effective treatments at start).

probability of an ineffective treatment to continue to the second stage. It should be chosen to balance minimizing this risk while ensuring the timeliness of the interim analysis. Designs were considered with stage 1 alpha between 20% and 50% with the final choice of 35% representing an achievable sample size and timely interim analysis (see trial timeline). This is in line with other MAMS trials,¹⁵ but differs from the 5% commonly used in confirmatory analysis, as the objectives here are different. Assuming an SD of 0.55%/year (see above), 111 observations per arm will allow 95% power to detect a 0.15%/year difference at a 1-sided significance level (α) of 35%. Allowing for 10% drop-out, 123 participants are needed per arm.

Therefore, we recommended that stage 1 analysis be conducted once 18 months' brain atrophy data are available for 111 participants per arm, with pairwise comparison for each experimental arm compared to the control arm. If the 1-sided p value is below 0.35, then the treatment arm is continued into stage 2.

Stage 2 Sample Size

The sample size for the stage 2 analysis was based on comparing the time to confirmed disability progression between each intervention arm to standard of care. For each pairwise comparison, to have 90% stage 2 power to detect a hazard ratio of 0.75 at the 2-sided stage 2 significance level of 5% (or equivalently a 2.5% 1-sided significance level), 281 progression events are required in the control arm and 600 participants per arm are needed.

The stage 2 significance level was set at a 2-sided 5% level, as in standard confirmatory trials, corresponding to a 1-sided level of 0.025. The question of multiplicity (adjusting significance level due to multiple comparisons) has been discussed before in

MAMS.¹⁸ We aimed to select drugs with different mechanisms, which might be viewed as independent evaluations, similar to multiple trials being conducted,³⁴ and therefore did not apply any correction for multiple comparisons. If drugs of similar action are selected (e.g., different doses of the same drug), an appropriate correction (e.g., Dunnett³⁵) should probably be applied. The statistical power in a time-to-event analysis is determined by the number of events. Recruiting 600 participants per arm should be sufficient to observe the required 281 progression events in the control arm in a timely manner. This number of events is anticipated to occur around 18 months after the last participant has been enrolled, assuming a 10% drop-out rate and 50% disability progression rate by 3 years and recruitment rate, as described in eAppendix 4 (links.lww.com/WNL/B900) (see trial timeline).

Trial Operating Characteristics

We conducted simulations to assess the operating characteristics of the proposed trial design under different scenarios (see eAppendix 5, links.lww.com/WNL/B900, for Methods and full Results). Table 3 shows the overall trial characteristics, depending on the number of truly effective treatments at the start of the trial. In all scenarios, the probability to wrongly conclude that 1 or more treatments is effective (false-positive) is below 4%. The chance of correctly concluding that at least 1 treatment is effective (power) if a single effective drug enters the trial is around 87%, but this increases to above 96% if more than 1 effective drug enters the trial.

Trial Timeline

An important consideration in adaptive trials is to anticipate the possible dynamics of the trial over time, including the relative timing of the interim and final analyses. eAppendix 4 (links.lww.com/WNL/B900) describes the assumptions made and how the timeline was modeled. Results are summarized in Table 4. Under a base case scenario of 40–50 participants recruited per month and 1 experimental arm continuing into stage 2, we expect the interim analysis to be conducted after around 3.4 years, and the final analysis after 6.1 years (ranging between 5.7 and 6.6 years depending on different scenarios modeled).

Discussion

MAMS trials have considerable potential in PMS, where there are many candidate therapies, as well as relevant interim outcome measures that have appropriate relationship to final clinical outcomes. We propose a MAMS trial design that could potentially accelerate the evaluation of new treatments in PMS.

Advantages

The proposed MAMS design leads to efficiencies in both sample size and trial duration compared to traditional separate phase II and III trials of single treatments. A single control arm is used to assess multiple experimental arms and participants recruited in stage 1 seamlessly continue to be included in the stage 2 analysis without additional setup time in between. This

Table 4 Expected Trial Duration Under Different Assumptions

Scenario	Expected time of stage 1 analysis, y	Expected time of stage 2 analysis, y	Percent of total participants by stage 1 analysis ^a
Base case ^b	3.4	6.1	58
Recruitment 20% slower	3.7	6.5	51
Recruitment 20% faster	3.3	5.7	66
2 experimental arms in stage 2	3.4	6.2	58
3 experimental arms in stage 2	3.4	6.3	58
45% progression rate at 3 years	3.4	6.6	58
55% progression rate at 3 years	3.4	5.7	58

^a Number of participants recruited in trial at the time of the interim analysis/total trial size for arms continuing to stage 2.

^b Assumed recruitment rate of 40 participants per month during stage 1 and 50 per month during stage 2, 50% disability progression rate at 3 years, and 1 experimental arm continuing into stage 2.

trial is expected to last 6–7 years with 1,900 participants, encompassing the stage 1 and 2 evaluations of 3 initial candidate treatments (Figure 1 and eAppendix 4, links.lww.com/WNL/B900).

In comparison, under the traditional approach, around 630 patients would be required in each of 3 phase II studies to have 90% power with 5% type I error under the same assumptions. If one of these treatments was found to be effective and proceeded to phase III, 1,200 additional participants would be required, totaling 3,090 participants. Separate phase II and III trials of a single treatment would be expected to take more than 10 years, with 3–5 years for phase II, 5–7 years for phase III, and additional setup time between the two (Figure 1). For example, evaluation of high-dose simvastatin is following a more conventional path with separate phase II (MS-STAT)¹⁶ and phase III (MS-STAT2)³⁶ trials. Recruitment to MS-STAT started in 2011 and MS-STAT2 is expected to be completed by 2025, which corresponds to 14 years overall.

Challenges

Planning and setting up a MAMS adaptive platform trial is considerably more complex than standard phase II and III trials and may take up to 12 to 18 months. In particular, statistical simulations examining different design options, scenarios, and parameters are essential to optimize efficiency and select appropriate trial operating characteristics while preserving the overall integrity of the trial. The initial modest investment in time and resources will be further offset by shorter subsequent setup times for further treatments added to the platform.

As adaptive platform designs are relatively novel in neurodegenerative diseases, there is a perception that regulatory agencies may not immediately accept them as equal to more conventional phase III studies. However, a precedent has been set for regulatory approval of MAMS platform trials in settings such as oncology⁷ and infectious diseases⁸ and our experience in these other disease areas suggests that regulators are becoming more open to, and knowledgeable and welcoming about, such designs.

Patient and Public Involvement

The PPI group actively participated in the entire trial design process, as well as treatment and outcome measure selection, to ensure the needs of people with MS were being met. For example, it was important, particularly for nonambulatory people with MS, to include an assessment of upper limb function in the primary efficacy endpoint and proposed secondary outcomes included patient-reported outcome measures of key symptoms such as fatigue. Feedback indicated the trial design was well-received and acceptable, despite being more complex. Perceived advantages included the ability to evaluate multiple candidate treatments and the relatively lower likelihood of randomization to placebo. If a participants' treatment arm is discontinued after interim analysis, there is the potential opportunity to re-enter the trial in a continuing arm or future trials in the platform. Some participants expressed concern about the total trial duration, but this was offset by the favorable consensus overall regarding potential acceleration of treatment discovery.

Scope and Future Developments

This article is based on work conducted by the trial design working group and in many senses is an evolution from our work carried out a decade ago.³⁷ A program grant proposal based on activity of all working groups of the UK MS Society's Expert Consortium for Progression in MS was submitted to the UK MS Society in November 2019 and received favorable international peer and lay review. Funding has been awarded to develop the protocol and deliver the first active arms plus standard of care in the MAMS trial platform, with recruitment expected to commence in 2022. Whereas this article focuses on evaluation of only the first 3 candidate therapies, the adaptive MAMS platform will allow addition of new treatment arms⁷ and re-randomization of participants from discontinued arms in the future.³⁸ Drugs with predominantly remyelinating potential will likely require additional and alternative endpoints at the interim analysis stage. Further aspects of the trial protocol, for example secondary and exploratory outcomes and recruitment infrastructure, are beyond the scope of this article.

Adaptive Platform Trials in Other Neurologic Disorders

Like PMS, conditions such as Parkinson disease, Alzheimer disease, and motor neuron disease are increasing in prevalence, have significant impact on patients, carers, and health care systems, and have no or few therapies that slow or prevent

progression. An improved understanding of disease pathophysiology in recent years has led to a growing pipeline of potential therapeutics. For example, a 2020 review identified 121 agents in 136 phase I to III clinical trials for Alzheimer disease, with an increasing number of disease modification treatment candidates over the past 5 years.³⁹ However, these conditions face similar challenges of efficiently translating candidate drugs into effective treatments with many disappointing phase III clinical trial results to date. Various reasons for this have been proposed, including the need to improve trial design.^{10,12,40}

MAMS designs are particularly relevant when there are multiple candidate therapies to be trialed and when a reliable early marker of clinical efficacy is available. MAMS adaptive platform trials have been planned and initiated to accelerate successful drug discovery in these disorders. The Motor Neuron Disease–Systematic Multi-arm Adaptive Randomization Trial (MND-SMART) will initially test 2 repurposed drugs against a common placebo.¹¹ The Dominantly Inherited Alzheimer's Network Trials Unit (DIAN-TU) platform trial established in 2012 was a multi-arm trial of 2 anti-amyloid monoclonal antibodies. Although neither drug met the primary cognitive endpoint,⁴¹ lessons learned, including refinements in participant and outcome measure selection and trial duration, have led to several emerging platform trials, such as the AHEAD study evaluating different doses of an anti-A β monoclonal antibody in 2 phase III clinical trials that respectively use amyloid PET and cognitive testing as the primary outcome measures.⁹

The principles of designing a PMS MAMS trial as outlined in this article are relevant to other neurodegenerative conditions, but each condition will present unique considerations and challenges, including selection of biologically and clinically relevant, sensitive, and timely interim and final outcome measures, determination of the most appropriate patient population for inclusion, and trial duration required to detect a meaningful effect.

We propose a design for a MAMS trial in PMS for evaluation of 3 repurposed neuroprotective drugs compared to standard of care. Although more complex in design, efficiencies in participant numbers and trial duration, as well as the ability to incorporate adaptive elements and continually test newly identified treatments through an ongoing platform, make this approach more likely to succeed in finding effective therapies that target disability progression in PMS in a timely manner.

Acknowledgment

Susan Scott, Research Network member of the UK Multiple Sclerosis Society, provided advice during the trial design process and reviewed the manuscript.

Study Funding

This work was supported by UK Multiple Sclerosis Society project grants 101 and 110; an award from the International Progressive MS Alliance (reference number PA-1412-02,420); and the ACORD collaboration: A Collaboration of Groups Developing, Running and Reporting Multi-arm, Multi-stage

Trials in Neurodegenerative Diseases (mrcctu.ucl.ac.uk/studies/all-studies/a/acord/).

Disclosure

V. Li and B. Leurent report no disclosures relevant to the manuscript. F. Barkhof acts as a consultant for Bayer-Schering, Biogen-Idec, IXICO, Merck-Serono, Novartis, Janssen, and Roche; has received grants from IMI for the Amyloid Imaging to Prevent Alzheimer's Disease (AMYPAD) initiative, the Dutch MS Society, ECTRIMS-MAGNIMS, the Dutch Research Council (NWO), the UK MS Society, Biogen, Merck, and GE Healthcare; is supported by the UCLH Biomedical Research Centre through the National Institute for Health Research (NIHR); is co-founder and a stockholder of Queen Square Analytics Ltd.; and is on the editorial boards of *Radiology*, *Neuroradiology*, *Multiple Sclerosis Journal*, and *Neurology*[®]. M. Braisher and F. Cafferty report no disclosures relevant to the manuscript. O. Ciccarelli is Deputy Editor of *Neurology* and serves on the editorial board of *Multiple Sclerosis Journal*; and receives research support from the NIHR UCLH/UCL Biomedical Research Centre, NIHR, MS Society of Great Britain and Northern Ireland, National MS Society, and Rosetrees Trust. A. Eshaghi has received speaker's honoraria from Biogen and At The Limits educational programme; travel support from the National Multiple Sclerosis Society; honoraria from the *Journal of Neurology*, *Neurosurgery and Psychiatry* for editorial commentaries; research grants from Biogen and Roche through his institutions; holds equity stake in Queen Square Analytics; and serves on the editorial board of *Neurology*. E. Gray, J. Nicholas, M. Parmar, G. Peryer, J. Robertson, N. Stallard, and J. Wason report no disclosures relevant to the manuscript. J. Chataway has received support from the Efficacy and Evaluation Programme, a Medical Research Council and NIHR partnership, the Health Technology Assessment Programme (NIHR), the UK MS Society, the US National MS Society, and the Rosetrees Trust in the past 3 years; and is supported in part by the National Institute for Health Research, University College London Hospitals, Biomedical Research Centre, London, UK. Go to Neurology.org/N for full disclosures.

Publication History

Received by *Neurology* October 26, 2021. Accepted in final form March 10, 2022. Submitted and externally peer reviewed. The handling editors were Elizabeth Silbermann, MD, and Brad Worrall, MD, MSc, FAAN.

Appendix Authors

Name	Location	Contribution
Vivien Li, MBBS, FRACP	Florey Institute of Neuroscience and Mental Health, University of Melbourne; Department of Neurology, Royal Melbourne Hospital, Melbourne	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data

Appendix (continued)

Name	Location	Contribution
Baptiste Laurent, PhD	Department of Medical Statistics, London School of Hygiene and Tropical Medicine; International Statistics and Epidemiology Group, London School of Hygiene and Tropical Medicine, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data
Frederik Barkhof, MD, PhD	Department of Radiology and Nuclear Medicine, Neuroscience Campus Amsterdam, VU University Medical Center, the Netherlands; Queen Square Institute of Neurology and Centre for Medical Image Computing, University College London; National Institute for Health Research, University College London Hospitals Biomedical Research Centre, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data
Marie Braisher, MSc	Queen Square Multiple Sclerosis Centre, Department of Neuroinflammation, UCL Queen Square Institute of Neurology, Faculty of Brain Sciences, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design
Fay Cafferty, PhD	MRC Clinical Trials Unit, Institute of Clinical Trials and Methodology, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data
Olga Ciccarelli, MD, PhD	National Institute for Health Research, University College London Hospitals Biomedical Research Centre; Queen Square Multiple Sclerosis Centre, Department of Neuroinflammation, UCL Queen Square Institute of Neurology, Faculty of Brain Sciences, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design
Arman Eshaghi, MD, PhD	NMR Unit, Department of Neuroinflammation, UCL Queen Square Institute of Neurology, Faculty of Brain Sciences, University College London; Centre for Medical Image Computing, Department of Computer Science, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data
Emma Gray, PhD	Multiple Sclerosis Society, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design

Appendix (continued)

Name	Location	Contribution
Jennifer M. Nicholas, PhD	Department of Medical Statistics, London School of Hygiene and Tropical Medicine, London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data
Mahesh Parmar, PhD	MRC Clinical Trials Unit, Institute of Clinical Trials and Methodology, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data
Guy Peryer, PhD	Multiple Sclerosis Society; Faculty of Medicine and Health Sciences, University of East Anglia, Norwich, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design
Jenny Robertson, MPsych(Health)	Multiple Sclerosis Society, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design
Nigel Stallard, PhD	Statistics and Epidemiology, Division of Health Sciences, Warwick Medical School, University of Warwick	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data
James Wason, PhD	Population Health Sciences Institute, Newcastle University, UK	Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data
Jeremy Chataway, MD, PhD	National Institute for Health Research, University College London Hospitals Biomedical Research Centre; MRC Clinical Trials Unit, Institute of Clinical Trials and Methodology, University College London, UK	Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data

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Vivien Li, Baptiste Leurent, Frederik Barkhof, et al.

Neurology 2022;98;754-764 Published Online before print March 23, 2022

DOI 10.1212/WNL.0000000000200604

This information is current as of March 23, 2022

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