Engaging multiple stakeholder groups in the development of standards for multiple sclerosis care: a modified Delphi process

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Background

- Although broad guidelines exist for multiple sclerosis (MS) care,^{1,2} there is no benchmark for timely care.
 - The need for prompt diagnosis and early treatment of MS was highlighted by the widely endorsed policy report Brain health: time matters in multiple sclerosis.³
- The current study aimed to engage multiple stakeholder groups in defining standards for the timing of key steps in the MS care pathway.
- These standards will inform the content of tools to help MS clinics strive for the highest level of care.

Developing standards for MS care

Methods

- A modified Delphi process was conducted, including both a core Delphi Consensus Panel and an additional Reviewing Group (Figure 1).
- Responses were collected via online surveys; the Panel remained anonymous to analysts and Chairs throughout.

Participants

- Four Chairs directed the process; they represented neurology, patient-reported outcomes, nursing/policy and the patient perspective.
- Participants were invited from regions where MS prevalence is high:⁴ North America, Northern Europe, Western Europe, Southern Europe, Eastern Europe and Russia, Australia and New Zealand, Middle East and North Africa.
 - In total, 41 MS neurologists from 22 countries were invited to participate in the **Delphi Consensus Panel** (Figure 1); 29 agreed to participate
 - Thirty-nine MS nurses, people with MS and allied healthcare professionals were invited to participate in the **Reviewing Group** to advise the Chairs; 31 agreed to participate (**Figure 1**).

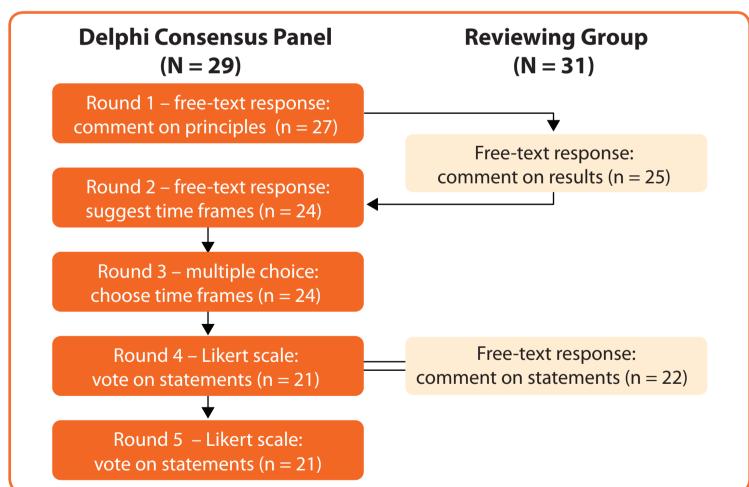


Figure 1. Modified Delphi process flow chart.

Round 1 – principles

- We derived 21 time-related principles from the recommendations in the report Brain health: time matters in multiple sclerosis.³
- The Panel were asked if each principle was 'an appropriate and accurate description of a good standard when considering brain health in people with MS' and were invited to suggest additional principles for inclusion.
- The Reviewing Group reviewed the results and provided feedback.
- Variables describing the principles in clinical practice were developed for round 2. Some principles identified were not considered time dependent; these were taken straight to round 4.

Rounds 2 and 3 – timings

- In round 2, the Panel suggested timings for 'core', 'achievable' and 'aspirational' standards (Table 1) for each variable, by free text.
- In round 3, the Panel were shown box plots of the round 2 data and asked to choose timings from given options.
- Consensus statements related to symptom onset, referral, diagnosis, treatment decisions, lifestyle, monitoring and managing new symptoms were developed based on these results.

Standard	Definition
Core	This should currently be achie of the local healthcare system
Achievable	This is a realistic target for mo of care
Aspirational	This might be achieved by on system allows, but should set

Table 1. Definitions used for consensus standards.

Rounds 4 and 5 – consensus statements

In round 4, the Panel voted on the consensus statements, indicating agreement (or otherwise) on a five-point scale.



DMT, disease-modifying therapy; MRI, magnetic resonance imaging

Disclosures

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- eved by most MS teams worldwide, regardless n, and will provide a **minimum standard**
- ost MS teams and reflects a **good standard**
- ly a few MS teams, where the local healthcare the standard for high-quality care

- The predefined threshold for consensus was at least 75% agreement, with a minimum of 66% of participants from round 1 completing the process.
- In round 5, the Panel were asked to vote again on statements from round 4 for which consensus was not reached, taking the results into consideration. Those who did not agree with the statements were asked to give reasons.
- The Reviewing Group were asked to review the statements and vote regarding the ambition of each using a three-point scale (not ambitious enough, about right or too ambitious).

Results

Defining a good standard of care

- For all 21 principles, over 75% of the 27 Panel members agreed in round 1 that the principle was an appropriate and accurate description of a good standard.
- Three statements gained 100% (27/27) agreement:
- 'Early discussion with patient about the aims of treatment'
- 'Evaluation of suitability/eligibility for treatment shortly after MS diagnosis'
- 'Regular review of the aims of treatment'.

Timings for key steps in the patient pathway

- Rounds 4 and 5 were completed by 21/27 (78%) of the Delphi Consensus Panel.
- Here, we present standards related to treatment decisions, monitoring and managing new symptoms, which the Panel agreed should be achievable (Figure 2).

2 weeks		The MS team should discuss the aims of treatment with each patient within 2 weeks of MS diagnosis
3 weeks		The MS team should discuss the pros and cons of early treatment with a DMT with patients within 3 weeks of diagnosis
	•	The MS team should assess within 3 weeks of an MS diagnosis whether the patient is eligible for treatment with a suitable DMT
nes eligible	fo	DMT
3 weeks		A DMT should be offered to a patient with MS within 3 weeks of their becoming eligible for one
es to start D	DM-	
2 weeks	•	Treatment with a DMT should commence within 2 weeks of a patient with MS agreeing this approach with their neurologist
Every 5 months	1	The MS team should review at least once every 6 months whether each patient with MS who is not receiving a DMT is eligible for one, based on applicable guidelines
ultations		
Every		The MS team should perform a follow-up clinical evaluation of each patient at least once every 6 months
5 months	•	The MS team should review with each patient at least once every 6 months the aims of their treatment for MS
	•	The MS team should review with each patient at least once every 6 months their currently prescribed DMT and consider alternatives if appropriate
Every 1 year		All patients with MS should be offered an MRI scan at least once every year
Regularly		The MS team should regularly enter patient data into an MS database
esponse to	DN	Т
l weeks	•	If a patient's response to their current DMT is judged to be suboptimal, an appropriate, alternative DMT should be offered within 4 weeks
ned sympto	om	s s
' days	1	Patients with MS should report new or worsened symptoms to their MS team within 7 days of experiencing these symptoms
w or worser	ned	l symptoms
2 days		The MS team should respond within 2 days to a patient with MS reporting an acute deterioration of symptoms
8 days	•	Patients with MS who experience an acute deterioration of symptoms should be seen by the relevant member of their MS team within 3 days of reporting these symptoms
c that gain	ad -	at least 75% agreement from the Delphi Consensus Panel

Figure 2. Subset of 'achievable' standards that gained at least 75% agreement from the Delphi Consensus Panel.





Using standards to improve care

Our vision

- The quality standards from the Delphi process will be used as the basis of practical tools to support the implementation of recommendations from *Brain health: time* matters in multiple sclerosis.³
 - Established and developing MS clinics in different countries will be encouraged to compare their services to the core, achievable or aspirational standards, as appropriate.

Quality improvement tool for clinics

- An MS Brain Health quality improvement tool is proposed that will help MS clinics strive for the best possible standard of patient care (Figure 3).
- Leading MS specialist neurologists will be part of a collaborative effort to design the tool and pilot it in their clinics.

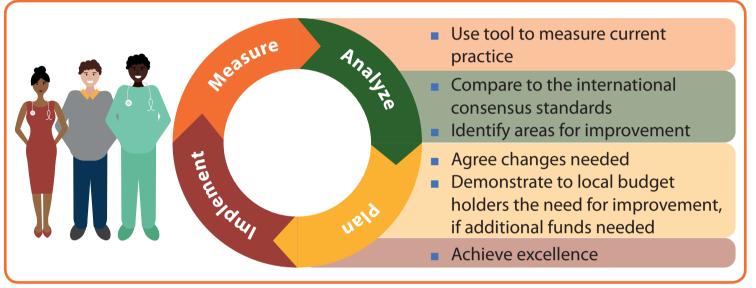


Figure 3. Proposed MS Brain Health quality improvement cycle.

Tool to empower people with MS

 Resources for people with MS will be developed in close consultation with representatives from patient organizations (Figure 4) • We will explore whether it would be possible to collect information on what patients actually experience and provide feedback to participating clinics.



Figure 4. Our vision for planned MS Brain Health tools for people with MS.

Conclusions

- An international group of MS neurologists, MS nurses, allied healthcare professionals and people with MS have been involved in a modified Delphi process to develop quality standards for MS care.
- These quality standards describe the timings of key steps in the MS care pathway and will provide a new benchmark for MS clinics globally. Tools will be developed to help multiple stakeholders improve care and deliver these standards in practice.

References

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