

UCLA

UCLA Previously Published Works

Title

The OMERACT Core Domain Set for Outcome Measures for Clinical Trials in Polymyalgia Rheumatica.

Permalink

<https://escholarship.org/uc/item/6br5s0wz>

Journal

The Journal of Rheumatology, 44(10)

Authors

Mackie, Sarah

Twohig, Helen

Neill, Lorna

et al.

Publication Date

2017-10-01

DOI

10.3899/jrheum.161109

Peer reviewed



Published in final edited form as:

J Rheumatol. 2017 October ; 44(10): 1515–1521. doi:10.3899/jrheum.161109.

The OMERACT Core Domain Set for Outcome Measures for Clinical Trials in Polymyalgia Rheumatica

Sarah L. Mackie, Helen Twohig, Lorna M. Neill, Eileen Harrison, Beverley Shea, Rachel J. Black, Tanaz A. Kermani, Peter A. Merkel, Christian D. Mallen, Frank Buttgerreit, Chetan Mukhtyar, Lee S. Simon, Catherine L. Hill on behalf of OMERACT PMR Working Group
Leeds Institute of Rheumatic and Musculoskeletal Medicine (LIRMM), University of Leeds, and UK National Institute for Health Research (NIHR), Leeds; Academic Unit of Primary Medical Care, University of Sheffield, Sheffield; PMR-GCA Scotland, Dundee; PMR-GCA North East, Gateshead; Primary Care and Health Sciences, Keele University, Keele; Norfolk and Norwich University Hospital, Norwich, UK; Ottawa Hospital Research Institute and School of Epidemiology, Public Health and Preventative Medicine, University of Ottawa, Ottawa, Ontario, Canada; Discipline of Medicine, The University of Adelaide; Rheumatology Unit, The Queen Elizabeth Hospital, Adelaide, Australia; Division of Rheumatology, University of California Los Angeles (UCLA), Los Angeles, California; Division of Rheumatology and Department of Biostatistics and Epidemiology, University of Pennsylvania, Philadelphia, Pennsylvania; SDG LLC, Cambridge, Massachusetts, USA; Department of Rheumatology and Clinical Immunology, Charite University Hospital Berlin, Berlin, Germany.

Abstract

Objective.—To inform development of a core domain set for outcome measures for clinical trials in polymyalgia rheumatica (PMR), we conducted patient consultations, a systematic review, a Delphi study, and 2 qualitative studies.

Methods.—Domains identified by 70% or more of physicians and/or patients in the Delphi study were selected. The conceptual framework derived from the 2 qualitative research studies helped inform the meaning of each domain and its relationship to the others. The draft core domain set was refined by further discussion with patients and physicians who had participated in the Delphi study. At the Outcome Measures in Rheumatology (OMERACT) 2016, the domains were

Address correspondence to Dr. S.L. Mackie, Leeds Institute of Rheumatic and Musculoskeletal Medicine, Chapel Allerton Hospital, Leeds, West Yorkshire LS7 4SA, UK. s.l.mackie@leeds.ac.uk.
S.L. Mackie BM, BCh, PhD Associate Clinical Professor and Honorary Consultant Rheumatologist, LIRM, University of Leeds; H. Twohig, MBChB, General Practitioner, Academic Unit of Primary Medical Care, University of Sheffield; L.M. Neill, BSc, PMR-GCA Scotland and OMERACT Patient Research Partner; E. Harrison, BSc, PMR-GCA North East and OMERACT Patient Research Partner; B. Shea, PhD, Senior Methodologist and Adjunct Professor, Ottawa Hospital Research Institute and School of Epidemiology, Public Health and Preventative Medicine, University of Ottawa; R.J. Black, MBBS, Consultant Rheumatologist and Clinical Lecturer, Discipline of Medicine, The University of Adelaide; T.A. Kermani, MD, MS, Assistant Clinical Professor, Division of Rheumatology, UCLA; P.A. Merkel, MD, MPH, Professor of Medicine and Epidemiology, Division of Rheumatology and Department of Biostatistics and Epidemiology, University of Pennsylvania; C.D. Mallen, PhD, NIHR Research Professor in General Practice, Arthritis Research UK Primary Care Centre, Research Institute for Primary Care, Keele University; F. Buttgerreit, MD, Professor of Rheumatology, Department of Rheumatology and Clinical Immunology, Charite University Hospital Berlin; C. Mukhtyar, MD, Consultant Rheumatologist, Norfolk and Norwich University Hospital; L.S. Simon, MD, Principal, SDG LLC; C.L. Hill, MD, Clinical Professor and Consultant Rheumatologist, Discipline of Medicine, The University of Adelaide, and Rheumatology Unit, The Queen Elizabeth Hospital.

discussed and prioritized by 8 breakout groups. Formal voting took place at the end of the workshop and in the final plenary.

Results.—Ninety-three percent of voters in the final plenary agreed that the inner core of domains considered mandatory for clinical trials of PMR should consist the following: laboratory markers of systemic inflammation, pain, stiffness, and physical function. Patient’s global and fatigue were considered important but not mandatory (outer core). The research agenda included psychological impact, weakness, physical activity, participation, sleep, imaging, and health-related quality of life.

Conclusion.—This core domain set was considered sufficiently well-defined that the next step will be to apply the OMERACT Filter 2.0 Instrument Selection Algorithm to select candidate instruments for a subsequent “deeper dive” into the data. This will allow instruments to be mapped onto each of our core domains to derive a core outcome set for PMR.

Key Indexing Terms:

POLYMYALGIA RHEUMATICA; OUTCOMES; OMERACT

Polymyalgia rheumatica (PMR) is an inflammatory disease of older people, causing pain and stiffness of the shoulders and hip girdles¹. The prevalence of PMR is about 1% in people over 50 years in the United States² and the United Kingdom³. Many patients with PMR are managed by general practitioners/family physicians rather than rheumatologists^{4,5}. The mainstay of treatment is longterm therapy with glucocorticoids. This treatment approach has the potential for toxicity, depending on glucocorticoid dose and patient-specific factors such as age^{6,7}. The most recent PMR treatment guidelines conditionally recommend early addition of methotrexate to glucocorticoids, especially if there are risk factors for relapse, for prolonged therapy, or for glucocorticoid-related adverse effects⁸. A stronger recommendation could not be made because the published randomized trials were small, with partly contradictory results. No high-quality evidence was identified evaluating any other potential glucocorticoid-sparing agent⁸. A systematic review of domains and instruments in 35 PMR trials and longitudinal observational studies, conducted by the Outcome Measures in Rheumatology (OMERACT) PMR Working Group, found inconsistency and poor clarity of outcome measures recorded for PMR⁹. The poor evidence base for management of PMR urgently requires improvement. Our objective is to produce guidance to researchers on a core outcome set for PMR: the minimal common set of outcome measurement instruments that should always be included in clinical trials of PMR, whether conducted in the community or specialist setting. Prior to recommending measurement instruments, it is necessary to define a core domain set of what it is that must be measured.

Here we report on the process that was used to generate a core domain set for clinical trials of PMR based on a combination of personal engagement, evidence synthesis, qualitative research, and a Delphi study. To our knowledge, this is the first core domain set developed for clinical trials of PMR; it has had strong patient involvement throughout. This core domain set will inform selection and validation of instruments to be used in clinical trials of PMR. It will also be relevant to design of observational studies and studies to develop a

PMR-specific patient-reported outcome measure. Our report represents the culmination of a process reported in 2 prior OMERACT Special Interest Group reports^{10,11}, work leading up to and during the 2016 OMERACT Workshop on PMR, and original primary research already published in full elsewhere^{9,12,13}. The new matter in this report includes a description of the methods and results of the Delphi survey and the process that was used to bring together multiple different sources of information (patient consultations, systematic literature review, 1 Delphi survey, 2 qualitative studies, further patient and clinician consultation to refine the draft core domain set, and a workshop at OMERACT 2016) to arrive at a core domain set for PMR that was endorsed by 93% of voters in the final conference plenary, as well as highlighting areas that required further definition, such as psychological impact.

Scoping the Problem

We intend our core outcome set to apply to interventional research studies conducted in any setting, with a study duration of at least 3 months and typically 1 year¹⁴. The selected domains would also be relevant to the design of observational studies, which could be much larger or of longer duration¹⁵. We began by consulting those involved on all outcomes they considered important for patients diagnosed with PMR; in later phases, we asked them to focus on clinical trials to give the context necessary for the prioritization of domains for a parsimonious core domain set.

Patient Involvement

Clinical management decisions relating to patients diagnosed with PMR are highly dependent on the patient's symptoms; acute-phase laboratory markers are used as supportive evidence¹. Defining what these symptoms are is therefore essential. Some of the patient research partners, including both co-authors of our current report, were involved over the life of this project and were deeply involved in patient support groups (telephone and/or Internet forums). Patient support groups were also helpful in identifying participants for our Delphi study.

Patient Consultations

To inform the scope of the problem, we started with a patient-driven consultation exercise¹¹. A convenience sample of 104 English-speaking patients with PMR under the care of rheumatologists from the United Kingdom and elsewhere in Europe were included and a modified nominal group technique was used, involving group discussions about 3 prespecified topics (symptoms, diagnosis, and treatment), followed by sorting of cards to identify each patient's "top ten" items for each topic. We reported these within the International Classification of Functioning, Disability and Health framework of impairments, disability, and participation¹¹.

Comparing Outcome of Patient Consultations with Systematic Review

Findings

Using the OMERACT Filter 2.0 framework¹⁶ we identified that outcomes reported in trials and observational studies of patients with PMR⁹ did not always map well onto the messages emerging from our patient consultations (Table 1). For example, patients preferred “stiffness” to “morning stiffness” and also considered fatigue to be important. Patients preferred to describe their experience of PMR in terms of its effect on activities such as getting out of bed, turning over in bed, getting up from the sofa or toilet, driving, picking items up from the floor, opening doors, walking, and dressing. They found the symptoms themselves hard to describe. The psychological impact of their condition was also mentioned. We noted that research studies had no standard definitions of key PMR symptoms; for example, in the literature it was frequently unclear exactly how patients had been asked about their pain severity, where that pain was, and what period of time was being asked about⁹. Similarly, the precise definition and meaning of morning stiffness in PMR appeared unclear in many published studies⁹. There was also no standard method used for reporting outcomes related to the burden of glucocorticoid therapy. Even the main daily dose and cumulative dose of glucocorticoids were not always well reported.

Analysis of composite outcomes used in studies of PMR¹⁷ showed that many included domains from both Patho-physiological Manifestations (acute-phase markers and/or ability to elevate upper limbs) and Life Impact (symptom or patient-reported component). Although none of these composite outcomes has yet been completely validated according to the OMERACT Filter, they are informative regarding what aspects of PMR are considered important by experts in PMR.

Delphi Study

To understand the differing perspectives of patients and physicians in prioritizing outcomes, we carried out a 3-round Delphi study¹⁰. We were advised by the National Research Ethics Service that ethical approval was not required. Although the disease (PMR) and its Life Impact may well be similar across countries, there are differences in the language used to describe this by patients. Whereas international English-speaking physicians are accustomed to using a common dialect (medical English) for accessing research studies and educational material, this is not necessarily the case for patients. To avoid potential misunderstanding arising from international differences in English vocabulary and usage, for our Delphi study we chose to recruit English-speaking patients from the United Kingdom.

The Delphi study started with 2 groups: patients (from UK patient organizations, self-identifying as diagnosed with PMR) and clinicians. Fifty-five patients with PMR took part. Of these, 46 completed round 2 and 34 completed round 3. Eighty-five clinicians with an interest in PMR were identified from PubMed searches and attendance at relevant sessions at international meetings (American College of Rheumatology, European League Against Rheumatism). Sixty clinicians replied to round 1, 55 to round 2, and 53 to round 3. Among the 60 clinicians in round 1, 21 were from the United Kingdom, 28 from elsewhere in Europe, 6 from North America, and 5 from Australasia. Self-reported expertise, other than

clinical rheumatology and an interest in PMR, was clinical trials research (26), outcomes research (19), epidemiology (11), qualitative research (5), general practice (5), and the allied health professions (2). Potential domains were grouped using the framework of Filter 2.0 (including “Resource Use,” but omitting “Death” from the list, because the latter is always mandatory in Filter 2.0) and informed by the prior patient consultations and systematic review findings.

To avoid influence of the patients on the clinicians or vice versa, rounds 1 and 2 were conducted separately. However, to identify areas of consensus and disagreement, we started with the same list of domains for everyone, using plain language rather than rheumatology jargon wherever possible. In round 1, respondents selected their “top ten” domains and had the option of adding any further domains to generate an expanded list. In round 2, each group was presented with the domains selected by > 70% of respondents and were asked which other domains from the expanded list they considered essential for a core domain set for clinical trials of PMR. Those new domains selected by > 70% of respondents in round 2 were added to that group’s list. The 70% cutoff, while arbitrary, is conventional for Delphi studies as well as being the usual level of consensus for OMERACT voting. Because of the variety of potential domains that seemed more relevant to glucocorticoid exposure, a separate item for glucocorticoid-related adverse effect was added in round 2. Results of rounds 1 and 2 are given in Table 2. In round 3, the domains finally selected by both groups were presented and opinions sought on the combined domain set. Free-text feedback at each stage allowed participants to give their reasoning for including or not including particular domains. A total of 91% of respondents (85% clinicians, 97% patients) agreed with the draft core domain set, with the major divergence of opinion appearing to be in relation to different perceptions of the meaning of the words *muscle weakness* in medical English versus everyday English. It also became clear that *morning stiffness [duration]*, a technical diagnostic term in rheumatology, is a different domain from “stiffness” as conceptualized by patients, who said that stiffness severity (rather than duration) was of key importance.

Qualitative Research on Core PMR Symptoms of Pain and Stiffness

A qualitative study¹³ analyzed in more depth what stiffness means to patients and how it relates to pain. Fifty patients with a clear, rheumatologist-confirmed diagnosis of PMR took part in 8 focus groups; this convenience sample was recruited from 3 UK rheumatology clinics. Pain and stiffness usually represented related but different symptoms. Pain (“ache, hurt”) was an unpleasant experience, not necessarily related to movement. Stiffness (the experience of being prevented from movement) had profound consequences for daily functioning. Many patients suggested that measuring physical function would be the best way to measure stiffness itself. Fatigue was seen as separate from either pain or stiffness, but affecting the broader experience of PMR.

Qualitative Research on the Broader Patient Experience in PMR

A second qualitative study analyzed the broader experience of PMR for patients treated in the community¹². The analysis of the study proceeded in parallel with the activities of the PMR Working Group and discussions before its publication informed the group’s thinking.

At OMERACT 2016, the methodology and findings were presented. Based on the conceptual framework derived from the qualitative data, we added the domain “Psychological Impact,” which had emerged as a surprisingly strong theme from the interviews.

Domain Prioritization

OMERACT presents domains using an “onion” diagram of 3 nested circles, with the domains in the innermost circle (“Inner Core”) being mandatory for every clinical trial; the middle circle is labeled “Important” and the outer circle “Research Agenda”¹⁸. The Inner Core should contain at least 1 domain chosen from each of the core areas including Pathophysiological Manifestations and Life Impact. It was recognized that the list of candidate domains derived from the Delphi was likely too long to be suitable for an Inner Core. Therefore, in the run-up to OMERACT 2016, informal e-mail engagement was carried out with patients and physicians who had participated in the Delphi study. A longlist of domains that might be eligible for the Inner Core was proposed, based on all of the evidence presented above, and feedback was invited. This resulted in removal of the domain of Physician Global because several physicians said it is a composite construct, principally consisting of information from laboratory markers of inflammation and the patient’s global (both of which were already on the longlist of domains). There were also questions about whether the underlying construct of Physician Global would genuinely be a scalar quantity or if it was better conceptualized as a binary decision to escalate or reduce glucocorticoid dose, closer to the concept of relapse/remission. Because the only remaining “Pathophysiological Manifestations” domain was Systemic Inflammation (Laboratory Blood Tests), the breakout discussions at the OMERACT Workshop focused on the Life Impact aspect of PMR.

Breakout Group Discussions

To encourage the discussion at breakout groups to draw on authentic patient experience, quotes from the qualitative interview were printed onto cards; each individual participant in the breakout group received a randomly chosen card. Breakout group facilitators then asked their groups to arrange the domains by priority, based on the results of the research described and cited in the preconference reading, the work presented in the plenary, and the quotes they had on their individual cards.

Synthesis of Advice from Breakout Groups

Consistent with the conceptual model that emerged from both qualitative studies, breakout groups gave the highest priority to pain/ache, stiffness, and physical function in regard to Life Impact (Table 3).

Feedback from several breakout groups suggested that including Patient’s Global in addition to the “top three” Life Impact domains could introduce redundancy, because the qualitative data suggested such a strong overlap with physical function. Given the strong drive toward parsimony for this patient population, and given the lack of quantitative evidence to confirm

or refute this suggestion, it was decided to provisionally rank this as important rather than core.

Psychological Impact was considered important, but to require further clarification of its meaning before inclusion in the Inner Core. The 2 candidate “psychological” domains that were drawn from the literature and entered into the Delphi (Mood problems — low or “high,” Anxiety) reached the 70% threshold in the patient arm of the Delphi study. However, the qualitative study data suggested that Psychological Impact goes beyond the clinical constructs of simple anxiety or mood disturbance and in fact describes complex, evolving, and pervasive effects on patients’ psychological state (for example, pre-diagnosis fears, relief at diagnosis followed by an ongoing sense of loss¹², and “PMR always on one’s mind”¹³) that are not necessarily well described by the clinical constructs of anxiety or depression or indeed well understood by clinicians. This was identified as a clear priority for further patient-centered research, perhaps with a view to developing a PMR-specific patient-reported outcome measure encompassing the psychological impact relating to this disease.

Breakout groups also advised adding to the research agenda the following domains: Participation, Weakness, Glucocorticoid Exposure, Physical Activity, Sleep, Imaging, and Health-related Quality of Life. Some attendees also pointed out that some caution was required in the interpretation of the qualitative research because of the limited geographical area (United Kingdom) from which the participants were drawn.

The Workshop concluded with a formal vote on whether each of our longlist domains should be included in the inner core for clinical trials (Table 3). Based on these votes, which was also in line with the results of our qualitative studies, we entered the 3 Life Impact domains plus Systemic Inflammation (Laboratory Blood Tests) into the proposed Inner Core.

Summary

Based on all the quantitative and qualitative feedback received during the whole process, a diagram (Figure 1) was presented at the final plenary session of the conference. Ninety-three percent of voters agreed with the final proposed Inner Core Domain Set (laboratory markers of systemic inflammation, pain, stiffness, physical function).

Future Work

Although there was substantial agreement on the inner core domains, the limitations of the voting procedure should be acknowledged; the system of 1 vote per attendee meant that clinicians’ votes outnumbered patients’ votes. The process also identified a substantial list of potential outcomes requiring further research in PMR. It will also be important to conduct further work with patients outside the United Kingdom, including non-English speakers, to assess generalizability of the concepts presented here. *The OMERACT Handbook* describes the next step, which will be to apply the OMERACT Filter 2.0 Instrument Selection Algorithm (the “eyeball test”), a systematic screening process to select candidate instruments for a subsequent “deeper dive” into the data to finally determine whether each selected instrument should be included in the core outcome set.

ACKNOWLEDGMENT

We thank all the patients who so generously contributed in so many ways to all stages of this project. We also thank all the members of the rheumatology research community who have provided important feedback during the latter stages of the development of this core domain set. Those not already listed as co-authors include (but are not limited to): Drs. Toby Helliwell, Elisabeth Brouwer, Joanna Robson, Christina Duftner, Mar Pujades-Rodriguez, Pereira da Silva, Maria Cid, Lyn March, and Maarten Boers. Dr. John Kirwan provided invaluable leadership, expertise, and guidance, including in the design of the Delphi.

Supported by the NIHR Clinician Scientist Fellowship to Sarah Mackie; unrestricted grant from Horizon Pharmaceuticals. Rachel Black is the recipient of an Australian Rheumatology Association Outcome Measures in Rheumatology (OMERACT) Fellowship 2016. Financial support for patient travel was received from the OMERACT charity DINORA, which received an unrestricted grant from Horizon Pharmaceuticals.

REFERENCES

1. Mackie SL, Mallen CD. Polymyalgia rheumatica. *BMJ* 2013;347:f6937. [PubMed: 24301266]
2. Lawrence RC, Felson DT, Helmick CG, Arnold LM, Choi H, Deyo RA, et al.; National Arthritis Data Workgroup. Estimates of the prevalence of arthritis and other rheumatic conditions in the United States. Part II. *Arthritis Rheum* 2008;58:26–35. [PubMed: 18163497]
3. Hayward RA, Rathod T, Muller S, Hider SL, Roddy E, Mallen CD. Association of polymyalgia rheumatica with socioeconomic status in primary care: a cross-sectional observational study. *Arthritis Care Res* 2014;66:956–60.
4. Barraclough K, Liddell WG, du Toit J, Foy C, Dasgupta B, Thomas M, et al. Polymyalgia rheumatica in primary care: a cohort study of the diagnostic criteria and outcome. *Fam Pract* 2008;25:328–33. [PubMed: 18687983]
5. Kremers HM, Reinalda MS, Crowson CS, Zinsmeister AR, Hunder GG, Gabriel SE. Use of physician services in a population-based cohort of patients with polymyalgia rheumatica over the course of their disease. *Arthritis Rheum* 2005;53:395–403. [PubMed: 15934100]
6. Harris E, Tiganescu A, Tubeuf S, Mackie SL. The prediction and monitoring of toxicity associated with long-term systemic glucocorticoid therapy. *Curr Rheumatol Rep* 2015;17:513. [PubMed: 25903665]
7. Strehl C, Bijlsma JW, de Wit M, Boers M, Caeyers N, Cutolo M, et al. Defining conditions where long-term glucocorticoid treatment has an acceptably low level of harm to facilitate implementation of existing recommendations: viewpoints from an EULAR task force. *Ann Rheum Dis* 2016;75:952–7. [PubMed: 26933146]
8. Dejaco C, Singh YP, Perel P, Hutchings A, Camellino D, Mackie S, et al.; European League Against Rheumatism; American College of Rheumatology. 2015 Recommendations for the management of polymyalgia rheumatica: a European League Against Rheumatism/American College of Rheumatology collaborative initiative. *Ann Rheum Dis* 2015;74:1799–807. [PubMed: 26359488]
9. Duarte C, Ferreira RJ, Mackie SL, Kirwan JR, Pereira da Silva JA; OMERACT Polymyalgia Rheumatica Special Interest Group. Outcome measures in polymyalgia rheumatica. A systematic review. *J Rheumatol* 2015;42:2503–11. [PubMed: 26568600]
10. Helliwell T, Brouwer E, Pease CT, Hughes R, Hill CL, Neill LM, et al. Development of a provisional core domain set for polymyalgia rheumatica: report from the OMERACT 12 Polymyalgia Rheumatica Working Group. *J Rheumatol* 2016;43:182–6. [PubMed: 26568595]
11. Mackie SL, Arat S, da Silva J, Duarte C, Halliday S, Hughes R, et al. Polymyalgia rheumatica (PMR) special interest group at OMERACT 11: outcomes of importance for patients with PMR. *J Rheumatol* 2014;41:819–23. [PubMed: 24488422]
12. Twohig H, Mitchell C, Mallen C, Adebajo A, Mathers N. “I suddenly felt I’d aged”: a qualitative study of patient experiences of polymyalgia rheumatica (PMR). *Patient Educ Couns* 2015; 98:645–50. [PubMed: 25638304]
13. Mackie SL, Hughes R, Walsh M, Day J, Newton M, Pease C, et al. “An impediment to living life”: why and how should we measure stiffness in polymyalgia rheumatica? *PloS One* 2015;10:e0126758. [PubMed: 25955770]

14. Hutchings A, Hollywood J, Lamping DL, Pease CT, Chakravarty K, Silverman B, et al. Clinical outcomes, quality of life, and diagnostic uncertainty in the first year of polymyalgia rheumatica. *Arthritis Rheum* 2007;57:803–9. [PubMed: 17530680]
15. Mackie SL, Hensor EM, Haugeberg G, Bhakta B, Pease CT. Can the prognosis of polymyalgia rheumatica be predicted at disease onset? Results from a 5-year prospective study. *Rheumatology* 2010;49:716–22. [PubMed: 20064872]
16. Boers M, Kirwan JR, Gossec L, Conaghan PG, D'Agostino MA, Bingham CO 3rd, et al. How to choose core outcome measurement sets for clinical trials: OMERACT 11 approves Filter 2.0. *J Rheumatol* 2014;41:1025–30. [PubMed: 24584913]
17. Leeb BF, Bird HA. A disease activity score for polymyalgia rheumatica. *Ann Rheum Dis* 2004;63:1279–83. [PubMed: 15361387]
18. Boers M, Kirwan JR, Tugwell P, Beaton D, Bingham CO III, Conaghan PG, et al. The OMERACT Handbook. [Internet. Accessed May 17, 2017.] Available from: www.omeract.org/pdf/OMERACT_Handbook.pdf

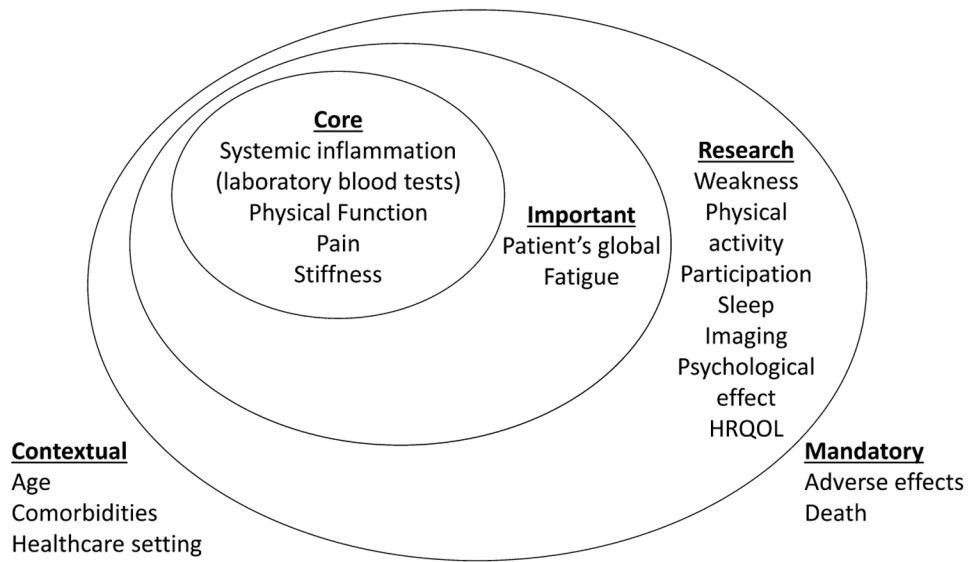


Figure 1.

Proposed core domain set for PMR clinical trials. This “onion” diagram uses nested circles with the innermost circle denoting the Inner Core (mandatory to measure in all clinical trials of PMR), the middle circle denoting Important Outcomes (strongly recommended to measure in PMR), and the outer circle denoting the Research Agenda (domains that require further investigation in PMR). Mandatory domains (bottom right) are those that should be reported by default in all clinical trials of any condition. The proposed contextual factors (bottom left) are suggestions we received regarding possible contextual factors and represent hypothesized factors only. PMR: polymyalgia rheumatica; HRQOL: health-related quality of life.

Table 1.

Comparison of domains reported as being important to patients with PMR. Outcomes measured in 35 clinical trials and longitudinal observational studies⁹, grouped according to the OMERACT Filter 2.0 Framework. Adapted from Duarte, *et al.* J Rheumatol 2015;42:2503–11; with permission⁹.

Core Area	Domain	Instrument Used in Published Studies	No. Studies of PMR Reporting Data on This Domain
Pathophysiological manifestations	Laboratory markers of systemic inflammation	ESR, CRP, IL-6, fibrinogen	30
	Elevation of upper limbs	Part of composite disease activity score	5
Life impact	Ability to carry out physical function tests	Grip strength, chair stand, 10-m walk	1
	Ultrasonography	Synovitis, bursitis in shoulder and hip regions	4
Physician's global	Physician's global	VAS, NRS	14
	Pain	VAS, NRS	17
Morning stiffness	Morning stiffness	Duration (min), grade, severity (VAS)	26
	Fatigue	VAS	1
Sleep disturbance	Sleep disturbance	Not reported	0
Patient's global	Patient's global	VAS, NRS	9
	Physical functioning (in daily living)	HAQ	9
Anxiety, depression	Anxiety, depression	Part of SF-36	2
	Health-related quality of life	SF-36, VAS	3
Death	Mortality	Death registries, patient files	1
Resource use	Not reported	Not reported	0
Adverse events	Adverse effects of medication	Standard methods for clinical trials	14

Bold face indicates highlighted in initial patient consultations¹¹. PMR: polymyalgia rheumatic; OMERACT: Outcome Measures in Rheumatology; ESR: erythrocyte sedimentation rate; CRP: C-reactive protein; IL-6: interleukin 6; VAS, visual analog scale; NRS: numerical rating scale; HAQ: Health Assessment Questionnaire; SF-36: Medical Outcomes Study Short Form-36.

Table 2.

Results from rounds 1 and 2 of the Delphi study of core domains for PMR. Numbers given are percentages of respondents selecting each outcome. Domains with > 70% agreement are highlighted in bold face; for each group (clinicians or patients), only domains with 20%–69% agreement in round 1 went forward to round 2. Bold face denotes that with > 70% agreement from either group in round 1 or round 2, the domain went forward into the combined round 3.

Domain	Round 1 Clinicians, n = 60	Round 1 Patients, n = 55	Round 2 Clinicians, n = 55	Round 2 Patients, n = 46
Pain/ache	90	78	—	—
Stiffness severity	56	56	53	74
Morning stiffness duration	68	36	85	43
Muscle weakness	12	53	—	80
Fatigue/tiredness	49	73	65	—
Sleep disturbance	10	35	—	57
Mood problems, low or “high”	3	20	—	37
Anxiety	3	33	—	30
Weight loss/gain	25	33	27	48
Appetite loss/gain	3	11	—	—
Balance problems	2	7	—	—
Fevers/shivers/sweats/flu-like symptoms	12	15	—	—
Ability to do everyday activities	51	55	65	74
Lack of mobility	25	35	7	59
Dependence on stick, wheelchair, etc.	0	2	—	—
Dependence on other people	5	9	—	—
Ability to carry out usual roles (work, caring for others, etc.)	20	35	16	37
Health-related quality of life	50	24	76	39
Overall quality of life	27	51	13	72
Change in appearance of face or body	8	22	—	41
Fluid retention/ankle swelling	2	9	—	—
Bruising, poor healing, or other skin change	5	24	—	50
Doctor’s assessment of activity/severity of PMR	59	60	71	83
Patient’s assessment of activity/severity of PMR	76	42	—	76
Blood tests	86	40	—	70

Domain	Round 1 Clinicians, n = 60	Round 1 Patients, n = 55	Round 2 Clinicians, n = 55	Round 2 Patients, n = 46
Abnormalities identified by physical examination by a doctor	22	7	20	—
Abnormalities identified by imaging tests	36	20	29	33
Bone fragility	27	13	29	—
New or worsening diabetes mellitus	14	5	—	—
High blood pressure	12	11	—	—
Cost of treatments used in the study	31	18	13	—
Overall costs to the healthcare provider	36	24	22	11
Overall costs to society	56	40	55	20
Any glucocorticoid-related adverse effect in judgment of doctor	—	—	46	63
Any glucocorticoid-related adverse effect in judgment of patient	—	—	23	80

“—” means the question was not asked in that round. PMR: polymyalgia rheumatica.

Table 3.

Votes for inner core at OMERACT PMR workshop. Percentage votes are presented to the nearest whole number.

Domain	No. Breakout Groups Selecting Domain among "Top 3" Life Impact Domains	Voted Yes	Voted No	Voted Insufficient Evidence or Information
Systemic inflammation, laboratory blood tests	Not asked	123/142 (87%)	4/142 (3%)	15/142 (11%)
Physical function	8/8	137/145 (94%)	3/145 (2%)	5/145 (3%)
Patient global	1/8	66/144 (46%)	41/144 (28%)	37/144 (26%)
Pain/ache	8/8	133/145 (92%)	4/145 (3%)	8/145 (6%)
Stiffness	7/8	131/145 (90%)	3/145 (2%)	11/145 (8%)
Fatigue	0/8	74/145 (51%)	35/145 (24%)	36/145 (25%)
Psychological impact	0/8	50/146 (34%)	57/146 (39%)	39/146 (27%)

OMERACT: Outcome Measures in Rheumatology; PMR: polymyalgia rheumatica.