REPORT

Psoriatic arthritis assessment tools in clinical trials

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In order to measure disease activity, progression, and change with therapy in psoriatic arthritis (PsA), it is important to use accurate, reliable, and feasible outcome measures that can ideally be employed in longitudinal cohorts, clinical trials, and clinical practice. Until recently, there has been little focus on this methodology in PsA. Clinical trials and long term clinical registries have used disparate outcome measures. With emerging therapies, the focus on the methodology of outcome assessment has increased to ensure that discriminant and responsive instruments are used. The Group for Research and Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA), in conjunction with the society, Outcome Measures in Rheumatology (OMERACT), is focused on refining and developing outcome measures for a variety of disease domains reviewed in this report. Key domains to assess include joints, skin, enthesitis, dactylitis, spine, joint damage as assessed radiologically, quality of life, and function. These domains can be assessed by individual and composite measures. A number of measures have been "borrowed" from the fields of rheumatoid arthritis, ankylosing spondylitis, and psoriasis and adapted to PsA. Others are being developed specifically for PsA. Few are validated but most have been shown to perform well in distinguishing placebo from treatment response. This report reviews the current state of the art of assessment in PsA and points toward future directions of development of this field.

Accurate, reliable, and reproducible assessment of disease activity and change with time and/or therapy in psoriatic arthritis (PsA) are important for understanding its natural history and relative effectiveness of therapies. In this report we review assessments that have been used in PsA trials as well as those in development, particularly through the work of the Group for Research and Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA) and Outcome Measures in Rheumatology (OMERACT). (The group was founded as Outcome Measures in Rheumatoid Arthritis Clinical Trials, hence the acronym, but has since expanded its mission across the spectrum of rheumatology intervention studies.) Most of the assessment methodologies discussed have been adapted from clinical trials in rheumatoid arthritis (RA), ankylosing spondylitis, or psoriasis; however, these methodologies have not yet been validated for use in PsA. Skin and radiological assessments are covered in separate articles in this supplement.^{1–3}

Assessment begins with inclusion of patients who have an accurate diagnosis. The problem of classification of PsA has been addressed elsewhere in this supplement.⁴ Until the work of the international CLASSification of Psoriatic Arthritis (CASPAR) study group has developed an updated schema, the traditional Moll and Wright criteria⁵ are being used. A number of challenges regarding the application of outcome measures used in other diseases, such as RA, become readily apparent when considering the subsets of PsA identified by

Moll and Wright: oligoarticular, polyarticular, distal interphalangeal joint arthritis, spondylitis, and arthritis mutilans. For example:

- Will the American College of Rheumatology (ACR) responder criteria or the disease activity score (DAS) based responder criteria, developed for a primarily polyarticular disease such as RA, function in a valid way for oligoarticular variants of PsA?
- Will measurement of spinal disease developed in ankylosing spondylitis, wherein all patients have, by definition, spinal involvement, function in a reliable way in PsA, where spine involvement is generally less severe and variable in expression?
- How does one approach measurement of unique clinical features such as dactylitis and enthesitis?
- Without a well documented understanding of the natural disease progression, in terms of joint damage, as we know in RA, how are we to know what to expect in the placebo arm of a study, which may beneficially impact on disease progression?
- What key core domains of PsA should be measured in every trial (such as joint tenderness and swelling, skin involvement, enthesitis, functional impairment, fatigue, etc.) and how should these be weighted in a composite responder index?

These and other questions are being addressed as outcome measures for PsA are being refined and validated. Two recent reviews also summarise outcome measures for PsA.^{6,7}

ARTICULAR AND COMPOSITE DISEASE ASSESSMENT

A key assessment in rheumatology clinical trials is the counting of the number of tender and swollen joints in a patient. Variable numbers of joints are counted in various systems, such as the ACR joint count of 68 tender and 66 swollen (excluding the hips from assessment of swelling), developed in 1949 for the evaluation of RA.⁸ These joint counts are then combined with other core measures of disease activity, such as patient and physician assessed global health, pain, function, and inflammation markers in different formulas. Such composite outcome measures of clinical response, developed in clinical trials of RA, include the ACR response criteria⁹ and the DAS,¹⁰ which form part of the European League Against Rheumatism (EULAR) response criteria,¹¹ discussed in detail below. The DAS has been validated with several different numbers of assessed joints,

Abbreviations: ACR, American College of Rheumatology; BASDAI, Bath Ankylosing Spondylitis Disease Activity Index; CRP, C-reactive protein; DAS, disease activity score; ESR, erythrocyte sedimentation rate; EULAR, European League Against Rheumatism; GRAPPA, Group for Research and Assessment of Psoriasis and Psoriatic Arthritis; HAQ, Health Assessment Questionnaire; OMERACT, Outcome Measures in Rheumatology; PsA, psoriatic arthritis; PsARC, psoriatic arthritic response criteria; RA, rheumatoid arthritis; VAS, visual analogue scale

now commonly 28. The Ritchie Articular Index is an older articular assessment methodology not currently used.¹²

An early attempt to develop a composite index of PsA disease activity included the Ritchie Index, a visual analogue scale (VAS) pain score, duration of morning stiffness, grip strength, and haemoglobin.¹³ Tested in varied subsets of PsA patients, it was only reliable in those with symmetrical polyarthritis, thus it has not been adopted as a responder index. The Ritchie Index was not validated.

PsARC and ACR response criteria

In the Veterans Administration study of sulfasalazine in PsA, a PsA specific response index was developed. To achieve response, a patient had to achieve two of the following, one of which had to be a joint count, and no worsening of any measure: tender or swollen joint count improvement of at least 30%, patient global improvement by one point on a five point Likert scale, or physician global improvement on the same scale.¹⁴ In this study, none of the individual measures showed statistically significant differences between the treated and placebo groups but the composite measure was able to do so marginally.

The first published study with a biological agent (etanercept)¹⁵ honoured historical precedent by using the same measure used in the Veterans Administration study as its primary outcome measure, and named it for the first time as the PsA response criteria (PsARC). As a secondary outcome measure, a modified version of the ACR joint count was used. The distal interphalangeal joints of the feet and carpometacarpal joints of the hands were added to the usual ACR joint count of 68 tender and 66 swollen, to yield a 78 and 76 joint count, respectively. Thus, the joints assessed for tenderness included the distal interphalangeal, proximal interphalangeal and metacarpophalangeal joints of the hands, and metatarsophalangeal joints of the feet, the carpometacarpal and wrist joints (counted separately), the elbows, shoulders, acromioclavicular, sternoclavicular, hip, knee, talo-tibial, and mid-tarsal joints. All of these except for the hips were assessed for swelling. Also, in this trial, joint tenderness, and swelling were graded 1–3. The other individual elements in the ACR scoring system, VAS scores of patient pain, patient global, physician global, the Health Assessment Questionnaire (HAQ), and acute phase reactant, C-reactive protein (CRP) or erythrocyte sedimentation rate (ESR) were unchanged from the way they are used in traditional RA trials. To achieve an ACR20, 50, or 70 response, at least 20%, 50%, or 70%, respectively, improvement in tender and swollen joint counts and three of five scores of individual elements (VAS scores of patient pain, physician and patient global assessment, a disability measure (HAQ) and an acute phase reactant (ESR or CRP)). Both the PsARC and ACR scoring systems were clearly able to distinguish patients treated with and without etanercept.

In one other trial, that of leflunomide in PsA,¹⁶ the PsARC was used as the primary outcome measure. This measure was able to discriminate treatment from placebo response, as was the ACR20. As will be seen in data from biological trials,¹⁷ frequency of ACR20 response is typically lower than PsARC response, even though PsARC requires $\geq 30\%$ improvement in joint score, possibly because tender or swollen joint change are required, not both, and perhaps because of the absence of the elements of acute phase reactant and the HAQ score.

Subsequently, in the phase III etanercept trial,¹⁸ and phase II¹⁹ and III infliximab²⁰ and adalimumab trials,²¹ the primary outcome measure was the ACR20 response. This was chosen as a “more stringent” outcome measure than the PsARC (based on lower percentage response in placebo treated patients in previous trials). In all of the trials with antitumour necrosis factor (anti-TNF) agents, there was a

highly statistically significant treatment response with p values <0.001 . ACR 50 and 70 responses were also significantly improved in the treatment groups.

DAS

There has been a great deal of interest in the DAS, developed in Europe for both assessment of disease activity state, as well as change of disease activity with therapy in RA.¹⁰ The original DAS used the Ritchie Articular Index (RAI),¹² swollen joint count (SJC), ESR, and general health status (GH) (VAS). The DAS is calculated as follows:

$$\text{DAS} = 0.54(\sqrt{\text{RAI}}) + 0.065(\text{SJC}) + 0.33(\ln \text{ESR}) + 0.0072(\text{GH})$$

The scale range is approximately 0–6 (since ESR is not bounded). The DAS was derived from a calculated weight of key elements in RA activity as observed in clinical trials. It has been modified to a 28 joint count for simplification purposes, and also has been employed with and without the general health status question, all of which correlate with each other and with the ACR score.^{10 22–28}

$$\text{DAS 28} = 0.56(\sqrt{\text{T28}}) + 0.28(\sqrt{\text{SW28}}) + 0.70(\ln \text{ESR}) + 0.014\text{GH}^{24}$$

As mentioned, the DAS has proved to be a useful instrument because unlike the ACR criteria, which only measures change in disease activity, the DAS can characterise the current amount of disease activity. For example, changing from a “very severe” to a “severe” level of disease may be perceived as having different significance than changing from a “modest” to “low” disease state.

The DAS is being used as a basis for the EULAR response criteria, which are derived from a discriminant function analysis of RA patients with active and non-active disease¹¹ in whom experienced rheumatologists judged whether important treatment changes had occurred. The patients are divided into three groups: non, moderate, and good responders. A non-response is a reduction in the DAS of ≤ 0.6 , with an endpoint DAS of >3.7 . A moderate response is a reduction in DAS between 0.6 and 1.2, with an endpoint DAS of >2.4 and ≤ 3.7 . A good response is an improvement of >1.2 and endpoint DAS of ≤ 2.4 . For DAS 28, the reduction amounts are the same, and the endpoint scores are >5.1 for a non-response, >3.2 and ≤ 5.1 for a moderate response, and ≤ 3.2 for a good response.²⁶ These measures have been highly discriminant and responsive in RA trials. A state of remission in RA is considered to be a DAS score of <1.6 .²⁷

Whereas the ACR criteria and PsARC have been used in recent PsA trial—for example, with leflunomide and the biological agents, the DAS has only been reported in trials with infliximab, where it was shown to be discriminant and responsive.¹⁹ Indeed, in a post hoc analysis of the data from the phase II trials of infliximab and etanercept (where the DAS was calculated from the elements of the ACR criteria), presented at OMERACT 7,²⁹ all of these instruments were found to be discriminant and responsive, including all the variants of the DAS (CE Antoni, personal communication). It is important to note that if a 28 joint count is used as part of entry criteria to a trial, a number of PsA patients would be excluded, especially if they have predominantly DIP or lower extremity disease. Thus, we would suggest that if a DAS 28 is used, that the joint count to determine inclusion in the trial be at least the 68/66 count.

In several recent RA trials, instruments which demonstrate more continuous data than the ACR 20, 50, 70, such as the

ACR-N, or summary measures, such as the area under the curve analysis of the ACR-N, have been used.^{30–31} Schiff has provided a review of this methodological approach,³² however, this methodology has not yet been applied in PsA. A committee authorised by the American College of Rheumatology is working on updating the ACR scoring system and incorporating informative elements from alternative criteria sets. This work will be reviewed and considered for incorporation into assessments of PsA.

An interesting observation is that PsA patients may experience less tenderness than RA patients when their joints are palpated. This was noted in a dolorimeter based study in Toronto.³³ This should be kept in mind as we observe the rating of both the patient and physician regarding the relative severity of PsA.

SPINE ASSESSMENTS

Spine involvement has been reported in 51% of PsA patients.³⁴ Sacroiliitis has been reported in up to a quarter of patients in several series.^{35–37} However, in one study of 221 patients, all with pelvis films, sacroiliac involvement was noted in up to 78%.¹⁴ Unlike ankylosing spondylitis, wherein axial involvement is present in all patients and tends to be more consistent in severity and in imaging findings, axial PsA is more inconsistent and heterogeneous. Because of this, current studies with biologicals have enrolled patients with active peripheral disease and have not attempted to measure axial involvement, even though the previous study by Clegg *et al*¹⁴ did attempt to measure spinal pain via the Douglas Spondylitis Articular Index.³⁸

The ASAS in Ankylosing Spondylitis (ASAS) working group has recommended the use of a number of outcome measures for spinal involvement in ankylosing spondylitis, which includes the Bath Ankylosing Spondylitis Disease Activity Index (BASDAI),³⁹ the Bath Ankylosing Spondylitis Function Index (BASFI)⁴⁰ and the Bath Ankylosing Spondylitis Metrology Index (BASMI).⁴¹ A more detailed description of spinal physical examination has been recently reviewed.⁶ The ASAS group has developed a responder index—the ASAS 20, 50, and 70, based on elements of the above indices and other patient assessment scales for use in ankylosing spondylitis trials. It is not clear if these instruments will perform well in PsA.

There are a number of difficulties in measuring spine disease in PsA clinical trials. As mentioned before PsA occurs less frequently and with greater variability of expression than is seen in ankylosing spondylitis. In an older patient, it is often difficult to distinguish pain from inflammatory spine disease versus pain from the very common presence of degenerative spine disease, even with imaging studies. Thus, these measures have not been attempted in most recently reported studies. In Clegg's study of sulfasalazine,¹⁴ duration of morning stiffness, spinal and nocturnal pain, and the Dougados Spondylitis Articular Index³⁸ were measured. The BASDAI items, as mentioned, have not been used in intervention trials in PsA. When PsA patients with and without spinal disease were assessed in a clinic cohort, the BASDAI scores did not differ.⁴² A principal components factor analysis in a different cohort of PsA patients showed that the BASDAI correlated more with measures of self reported wellbeing than with measures of disease activity, whereas the Dougados Articular Index showed the opposite, leading to a suggestion that this index should be further evaluated as a measure of spine inflammation in PsA.⁴³ At the present time, the GRAPPA initiative is making an effort to determine a feasible and appropriate measure to distinguish the presence of axial disease due to PsA in these patients and to measure its change in clinical trials.

ENTHESITIS

Enthesitis, a common feature of PsA, is characterised by inflammation at sites of tendon, ligament, and joint capsule insertion into bone. Common symptomatic areas include insertions of the Achilles' tendon and plantar fascia to the calcaneus, ligamentous insertions, and insertions into the pelvis and to the bones of the thorax and spine. The enumeration of patients with or without enthesitis, before and after treatment, has been studied in three PsA trials.^{14 19 44} Neither sulfasalazine nor azathioprine showed response in this domain, whereas in both studies with the anti-TNF medicine, infliximab, there was. This particular domain has not been studied with etanercept, and results from trials of adalimumab are pending.

For spondyloarthropathies in general, two enthesitis measures have been developed, the Mander index⁴⁵ and the Maastricht Ankylosing Spondylitis Enthesis Score (MASES).⁴⁶ The Mander index assesses 66 sites; it is generally accepted that this is too many to feasibly and reliably assess in clinical trials. The MASES index, which represents a culling of the 13 most specific and sensitive sites from the Mander index in an assessment of patients with ankylosing spondylitis over two years, correlated well with the Mander. The MASES is a more feasible instrument, but it has not yet been assessed in PsA trials. A theoretical issue with these instruments is whether they will adequately discriminate from fibromyalgia tender points in patients with this as a comorbid condition with PsA. Gladman *et al* have reported on the performance of investigators from the Canadian Spondyloarthropathy Group (SPARCC) in their ability to reliably assess enthesitis areas: the plantar fascia, Achilles' tendon, tibial tuberosity, and rotator cuff insertions. In the first three locations, observer agreement was "moderate" and in the fourth, "poor".^{6 47} Both the degree to which this element of PsA contributes to disease burden and the prospect of its meaningful improvement with newer therapies argue for focus on reliable assessment and measurement of change to further understand the impact of therapy.

DACTYLITIS

Dactylitis is characterised by swelling of a whole digit and represents a combination of synovitis and inflammation of tendon and ligament insertions. It occurs in approximately half of all patients with PsA.⁴⁸ Presence or absence of dactylitis before and after drug treatment has been reported in three trials,^{9 18 19} with statistically significant change demonstrated in the two anti-TNF trials. A simple quantitative assessment, on a 0–3 scale is being employed in a current trial in PsA (PJ Mease, personal communication). A more quantitative scoring system is being developed, but has not yet been used in a clinical trial (P Helliwell, personal communication).

QUALITY OF LIFE, FUNCTION, "PARTICIPATION"

Increasingly, assessment of quality of life, physical function, and "participation"—the capacity to engage meaningfully and capably in activities of life—is important when judging the impact of disease and improvement with treatment. New therapies have shown significant effect in this arena. Treatment expense increasingly requires justification: Does it not only ameliorate symptoms and prevent disease progression but does it also enhance quality of life, reduce work absenteeism, improve functional capacity, and allow for return to a "normal life"? The methodology in this field is complex and draws input from such diverse fields as psychology, ergonomics, pharmacoeconomics, psychometrics, and sociology.

Quality of life measures can be generic, crossing multiple disease states, or specific, developed for a single condition. An

example of the former, across multiple disease states and thus comparable between them, is the Medical Outcomes Study Short Form 36 (SF-36).⁴⁹ This patient self-administered questionnaire assesses eight domains of health status: physical functioning, pain, vitality, social functioning, psychological functioning, general health perceptions, and role limitations due to physical and emotional problems. This has been employed in most recent rheumatology trials, including PsA, has shown significant improvements in PsA patients with effective treatment,^{15, 18} and has been validated in PsA.^{50–51} The EuroQol-5D⁵² has shown no difference in the impact of RA and PsA on quality of life despite greater physical damage in the comparator RA group.⁵³ The Arthritis Impact Measurement Scales (AIMS and AIMS2)⁵⁴ instruments assess, in arthritis patients, physical, emotional, and social wellbeing, and have been validated and critiqued in PsA.^{55–56} Gladman *et al* suggest that although broadly used, generalised questionnaires allow comparison of patient groups and are worthwhile to use in both clinical trials and longitudinal cohorts, it is disease specific quality of life instruments that are likely to yield larger effects in clinical trials.⁶

The Psoriatic Arthritis Quality of Life (PsAQoL) measure, the first patient derived instrument specific for PsA, has shown reliability and construct validity,⁵⁷ but has not yet been used in a clinical trial. Several measures have been developed to assess quality of life in patients with psoriasis and/or dermatological disease in general. The Dermatology Life Quality Index (DLQI)⁵⁸ is a 10 item instrument developed as a measure of disability for a wide range of dermatological conditions. It has been the most used and validated instrument in psoriasis, and has been used in some studies of PsA to consistently show discriminant ability.¹⁸ A complementary instrument used to assess psychosocial wellbeing is the Dermatology Quality of Life Scale (DQoLS).⁵⁹ This has not been used in PsA studies. Some psoriasis specific instruments, also not yet used in PsA trials, are the Koo-Menter Psoriasis Instrument (KMPI),⁶⁰ the Psoriasis Disability Index (PDI),⁶¹ the Psoriasis Life Stress Inventory (PLSI),⁶² Psoriasis Quality of Life instrument (PsoriQoL)⁶³ and the Salford Psoriasis Index (SPI).⁶⁴ Quality of life, its measurement, and effects of therapy in both psoriasis and PsA have been recently reviewed.⁶⁵

The Health Assessment Questionnaire (HAQ), originally developed to assess disability in RA,⁶⁶ by focusing on physical disability and pain, has been used widely in inflammatory arthritis clinical trials, including PsA. The HAQ has been modified for spondyloarthropathies, which includes two spinal domains (HAQ-S),⁶⁷ as well as a further skin modification for patients with psoriasis (HAQ-SK).^{68–69} These were shown to perform similarly, and thus the original instrument has been used in clinical trials. Gladman *et al* have emphasised that not only should this instrument be used in clinical trials, but it is also useful as a measure of health status in longitudinal clinical cohorts.⁶ Since the HAQ focuses on physical disability, even with the skin modification, it may not adequately capture disability in patients with predominantly skin disease. The HAQ will presumably show less change in the context of treatment that has a predominant effect on the skin and not the joints. Taylor has further observed that it may not adequately measure the activities affected in some subsets of PsA patients—by applying a measurement modelling technique known as Rasch analysis,⁷⁰ we may better determine if the HAQ or its modifications are performing adequately in patients with different patterns of PsA.⁷

An important question in the field of functional evaluation is how much improvement in functional status is considered important by patients? Determining the “minimal clinically

important difference” (MICD), for a measurement instrument is a key construct, which helps us determine if a change with treatment is not only statistically significant, but is also meaningful to a patient. The methodological science of determining MCID is complex and can be approached in a number of ways.⁷¹ A generally accepted value for MCID of the HAQ in RA is 0.22.⁷² The first attempt at determining such a value for PsA, based on analysis of the HAQ data from the phase III trial of etanercept on PsA, found that the MCID calculated on the basis of a patient rating method, conjectured as being more valid was 0.3, and when based on standard error of measurement method, was 0.4.⁷³ This observation needs to be further confirmed by analysis in other trials and across trials before it can be fully accepted.

As noted earlier, there is a new domain in disease assessment, “participation”. This domain addresses not just a person’s ability to perform a task or action, but it takes into account their actual performance in life situations. For example, if a person is physically capable of working in a certain job capacity or visiting friends, but is embarrassed to do so because of unsightly psoriasis lesions, then their “participation” may be impaired even if not due to change in body function or structure. A World Health Organization initiative, the International Classification of Functioning, Disability and Health,⁷⁴ is stimulating development of assessment tools, not just a self-report questionnaire, to capture this complex interaction between disease state, body function and structure, activities, participation, and environmental and personal factors.⁷⁵ It is anticipated that such measurement will give us a better idea of the true impact of a disease on the individual, their family, and society, as well as inform us of the full impact of therapies in this global domain. The GRAPPA group was charged with developing such an instrument at OMERACT 7 and is proceeding forthwith.

FATIGUE

Although some would argue that fatigue is a domain subsumed under global patient assessment, others, including patients, would suggest that it is an important domain in and of itself. It is notable that when asked about improvement with the newer biological agents, fatigue is a dimension, along with pain and function, often mentioned by patients as demonstrating significant improvement. A number of instruments to measure fatigue have been developed. One is to have the patients rate fatigue on a VAS scale. Multidimensional measures have been developed to try to capture various aspects of fatigue, such as physical and emotional fatigue. Examples of these include the Multidimensional Fatigue Inventory (MFI),⁷⁶ the Fatigue Severity Scale (FSS),⁷⁷ the Functional Assessment of Chronic Illness Therapy (FACIT) scale,⁷⁸ and the Multidimensional Assessment of Fatigue (MAF) index, referred to as MFI.⁷⁹ The MFI has been used in patients with ankylosing spondylitis and was found to be highly correlated with the single question on fatigue in the BASDAI. It also provided more complex information about fatigue in this patient group. The FSS has been studied in PsA patients, and it was able to distinguish patients from controls and showed correlation with disease activity.⁸⁰ The FACIT is currently being employed in clinical trials of PsA patients. An advantage of the FACIT is that it can be adapted to a variety of chronic disease states, so can be compared across diseases.

BIOMARKERS

Simple measures of inflammation, such as the acute phase reactants, ESR and CRP, are not reliably elevated in patients with PsA, even with active inflammation. In a recent analysis of the phase II etanercept and infliximab studies in PsA, a

receiver operator curve analysis showed that these measures were not highly specific in discriminating placebo from treatment response (CE Antoni, personal communication). Thus, these may not be reliable markers to assess baseline disease activity or response to therapy. However, it has been noted that high ESR at presentation does correlate with progression of joint disease and early mortality.⁸¹

Translational studies of histological and immunohistochemical changes in PsA synovium and skin due to therapeutic intervention, recently reviewed,⁸² are a mode of outcome assessment, albeit impractical for general clinical use. It is through this work that we are learning more about the basic pathophysiology of joint and skin inflammation in PsA and the specific change that accrues from targeted therapy. A subgroup of GRAPPA is chartered to develop this area and standardise specific biomarkers assessed in different research centres.

FUTURE DEVELOPMENT OF PsA SPECIFIC COMPOSITE RESPONDER CRITERIA

It will be noted that the ACR criteria, the PsARC, and the EULAR response criteria are all considered composite responder indices and have proved effective in discriminating between placebo and treatment response. However, they do not incorporate what some might consider a full set of core domains to be assessed, such as skin, spine, and enthesal involvement. A future project in the development of a more comprehensive responder index will be to measure individual core domains and then to statistically derive a more full combination of measures that is both responsive and discriminant.

CONCLUSION

Outcome measures which measure clinical response, disease progression, and quality of life in PsA have been shown to be discriminant and responsive in recent clinical trials of emerging therapeutics. These measures will be undergoing further refinement through the efforts of GRAPPA and OMERACT. Methods to assess less fully explored domains, such as enthesitis, dactylitis, and "participation", will be developed and applied in future clinical trials. The availability of effective outcome measures allows us to define more fully the impact of therapy, compare therapeutic approaches, and understand the course of PsA.

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REFERENCES

- Feldman SR, Krueger GG. Psoriasis assessment tools in clinical trials. *Ann Rheum Dis* 2005;**64**(suppl II):ii65-8.
- Ory PA, Gladman DD, Mease PJ. Psoriatic arthritis and imaging. *Ann Rheum Dis* 2005;**64**(suppl II):ii55-7.
- van der Heijde D, Sharp J, Wassenberg S, Gladman DD. Psoriatic arthritis imaging: a review of scoring methods. *Ann Rheum Dis* 2005;**64**(suppl II):ii61-4.
- Helliwell PS, Taylor WJ. Classification and diagnostic criteria for psoriatic arthritis. *Ann Rheum Dis* 2005;**64**(suppl II):ii3-8.
- Moll J, Wright V. Psoriatic arthritis. *Semin Arthritis Rheum* 1973;**3**:55-78.
- Gladman DD, Helliwell P, Mease PJ, Nash P, Ritchlin C, Taylor W. Assessment of patients with psoriatic arthritis: a review of currently available measures. *Arthritis Rheum* 2004;**50**:24-35.
- Taylor WJ. Assessment of outcome in psoriatic arthritis. *Curr Opin Rheumatol* 2004;**16**:350-6.
- Deandrade JR, Casagrande PA. A seven-day variability study of 499 patients with peripheral rheumatoid arthritis. *Arthritis Rheum* 1965;**8**:302-34.
- Felson DT, Anderson JJ, Boers M, Bombardier C, Furst D, Goldsmith C, et al. American College of Rheumatology preliminary definition of improvement in rheumatoid arthritis. *Arthritis Rheum* 1995;**38**:727-35.
- van der Heijde DM, van't Hof MA, van Riel PL, Theunisse LA, Lubberts EW, van Leeuwen MA, et al. Judging disease activity in clinical practice in rheumatoid arthritis: first step in the development of a disease activity score. *Ann Rheum Dis* 1990;**49**:916-20.
- van Gestel AM, Prevoo ML, van't Hof MA, van Rijswijk MH, van de Putte LB, van Riel PL. Development and validation of the European League Against Rheumatism response criteria for rheumatoid arthritis. Comparison with the preliminary American College of Rheumatology and the World Health Organization/International League Against Rheumatism Criteria. *Arthritis Rheum* 1996;**39**:34-40.
- Ritchie DM, Boyle JA, McInnes JM, Jasani MK, Dalakas TG, Grieve P, et al. Clinical studies with an articular index for the assessment of joint tenderness in patients with rheumatoid arthritis. *Q J Med* 1968;**37**:393-406.
- Daunt AO, Cox NL, Robertson JC, Cawley ML. Indices of disease activity in psoriatic arthritis. *J R Soc Med* 1987;**80**:556-8.
- Clegg DO, Reda DJ, Mejias E, Cannon GW, Weisman MH, Taylor T, et al. Comparison of sulfasalazine and placebo in the treatment of psoriatic arthritis. A Department of Veterans Affairs Cooperative Study. *Arthritis Rheum* 1996;**39**:2013-20.
- Mease PJ, Goffe BS, Metz J, VanderStoep A, Finck BK, Burge DJ. Etanercept in the treatment of psoriatic arthritis and psoriasis: a randomised trial. *Lancet* 2000;**356**:385-90.
- Kaltwasser JP, Nash P, Gladman D, Rosen CF, Behrens F, Jones P, et al. Efficacy and safety of leflunomide in the treatment of psoriatic arthritis and psoriasis. *Arthritis Rheum* 2004;**50**:1939-50.
- Mease PJ, Antoni CE. Psoriatic arthritis treatment: biological response modifiers. *Ann Rheum Dis* 2005;**64**(suppl II):ii78-82.
- Mease P, Kivitz A, Burch F, Siegel E, Cohen S, Ory P, et al. Etanercept treatment of psoriatic arthritis: safety, efficacy, and effect on disease progression. *Arthritis Rheum* 2004;**50**:2264-72.
- Antoni C, Kavanaugh A, Kirkham B, Burmester G, Weisman MH, Keystone E, et al. The infliximab multinational psoriatic arthritis controlled trial: substantial efficacy on synovitis and psoriatic lesions with or without concomitant DMARD therapy. *Arthritis Rheum* 2005 (in press).
- Kavanaugh A, Krueger G, de Vlam K, Barbra C, Beutler A, Guzzo C, et al. Infliximab improves arthritis and psoriasis in patients with active polyarticular psoriatic arthritis: results of the IMPACT II trial. *Ann Rheum Dis* 2004;**63**(suppl 1):402.
- Mease P, Gladman DD, Ritchlin C, Ruderman EM, Steinfeld D, Choy E, et al. Adalimumab therapy in patients with psoriatic arthritis: 24-week results of a phase II study. *Arthritis Rheum* 2004;**50**:A097.
- van der Heijde DM, van't Hof M, van Riel PL, van Leeuwen MA, van de Putte LB. A comparison of validity of single variables and composed indices for measuring disease activity in rheumatoid arthritis. *Ann Rheum Dis* 1992;**51**:177-81.
- Prevoo ML, van Riel PL, van't Hof MA, van Rijswijk MH, van Leeuwen MA, Kuper HH, et al. Validity and reliability of joint indices. A longitudinal study in patients with recent onset rheumatoid arthritis. *Br J Rheumatol* 1993;**32**:589-94.
- Prevoo ML, van't Hof MA, Kuper HH, van Leeuwen MA, van de Putte LB, van Riel PL. Modified disease activity scores that include twenty-eight-joint counts. Development and validation in a prospective longitudinal study of patients with rheumatoid arthritis. *Arthritis Rheum* 1995;**38**:44-8.
- Fuchs HA. The use of the disease activity score in the analysis of clinical trials in rheumatoid arthritis. *J Rheumatol* 1993;**20**:1863-6.
- van Gestel AM, Haagsma CJ, van Riel PL. Validation of rheumatoid arthritis criteria that include simplified joint counts. *Arthritis Rheum* 1998;**41**:1845-50.
- Prevoo ML, van Gestel AM, van't Hof MA, van Rijswijk MH, van de Putte LB, van Riel PL. Remission in a prospective study of patients with rheumatoid arthritis. American Rheumatism Association preliminary remission criteria in relation to the disease activity score. *Br J Rheumatol* 1996;**35**:1101-5.
- van Gestel AM, Anderson JJ, van Riel PL, Boers M, Haagsma CJ, Rich B, et al. ACR and EULAR improvement criteria have comparable validity in rheumatoid arthritis trials. American College of Rheumatology European League of Associations for Rheumatology. *J Rheumatol* 1999;**26**:705-11.
- Gladman DD, Strand V, Mease PJ, et al. OMERACT 7 psoriatic arthritis workshop: synopsis. *Ann Rheum Dis* 2005;**64**(suppl II):ii115-6.
- Bathon JM, Martin RW, Fleischmann RM, Tesser JR, Schiff MH, Keystone EC, et al. A comparison of etanercept and methotrexate in patients with early rheumatoid arthritis. *N Engl J Med* 2000;**343**:1586-93.
- Smolen J, Emery P, Bathon JM, Keystone EC, Maini R, Kalden J, et al. Treatment of early rheumatoid arthritis with infliximab plus methotrexate or methotrexate alone: preliminary results of the ASPIRE trial. *Ann Rheum Dis* 2003;**62**(suppl 1):64.
- Schiff M. A rationale for the use of summary measurements for the assessments of the effects of rheumatoid arthritis therapies. *Clin Ther* 2002;**25**:993-1001.

- 33 **Buskila D**, Langevitz P, Gladman DD, Urowitz S, Smythe HA. Patients with rheumatoid arthritis are more tender than those with psoriatic arthritis. *J Rheumatol* 1992;**19**:1115-19.
- 34 **Gladman DD**, Brubacher B, Buskila D, Langevitz P, Farewell VT. Psoriatic spondyloarthropathy in men and women: a clinical, radiographic, and HLA study. *Clin Invest Med* 1992;**15**:371-5.
- 35 **Gladman DD**, Shuckett R, Russell ML, Thorne JC, Schachter RK. Psoriatic arthritis (PSA)—an analysis of 220 patients. *Q J Med* 1987;**62**:127-41.
- 36 **Torre-Alonso JC**, Rodriguez Perez A, Arribas Castrillo JM, Ballina Garcia J, Riestra Noriega JL, Lopez Larrea C. Psoriatic arthritis (PA): a clinical, immunological and radiological study of 180 patients. *Br J Rheumatol* 1991;**30**:245-50.
- 37 **Veale D**, Rogers S, Fitzgerald O. Classification of clinical subsets in psoriatic arthritis. *Br J Rheumatol* 1994;**33**:133-8.
- 38 **Dougados M**, Gueguen A, Nakache JP, Nguyen M, Mery C, Amor B. Evaluation of a functional index and an articular index in ankylosing spondylitis. *J Rheumatol* 1988;**15**:302-7.
- 39 **Garrett S**, Jenkinson T, Kennedy LG, Whitelock H, Gaisford P, Calin A. A new approach to defining disease status in ankylosing spondylitis: the Bath Ankylosing Spondylitis Disease Activity Index. *J Rheumatol* 1994;**21**:2286-91.
- 40 **Calin A**, Garrett S, Whitelock H, Kennedy LG, O'Hea J, Mallorie P, et al. A new approach to defining functional ability in ankylosing spondylitis: the development of the Bath Ankylosing Spondylitis Functional Index. *J Rheumatol* 1994;**21**:2281-5.
- 41 **Jenkinson TR**, Mallorie PA, Whitelock HC, Kennedy LG, Garrett SL, Calin A. Defining spinal mobility in ankylosing spondylitis (AS). The Bath AS Metrology Index. *J Rheumatol* 1994;**21**:1694-8.
- 42 **Brockbank J**, Schimmer J, Schentag C, Hyrich K, Gladman DD. Spinal disease assessment in psoriatic arthritis (PsA). *J Rheumatol* 2001;**28**(suppl 63):62.
- 43 **Taylor WJ**, Harrison AA. Could the Bath Ankylosing Spondylitis Disease Activity Index (BASDAI) be a valid measure of disease activity in patients with psoriatic arthritis? *Arthritis Rheum* 2004;**51**:311-15.
- 44 **Antoni C**, Kavanaugh A, Kirkham B, Bermester G, Manger B, Schneider U, et al. The one year results of the infliximab multinational psoriatic arthritis controlled trial (IMPACT): substantial efficacy on synovitis and psoriatic lesions with or without concomitant DMARD therapy. *Arthritis Rheum* 2003;**48**(suppl 9):S265.
- 45 **Mander M**, Simpson JM, McLellan A, Walker D, Goodacre JA, Dick WC. Studies with an enthesitis index as a method of clinical assessment in ankylosing spondylitis. *Ann Rheum Dis* 1987;**46**:197-202.
- 46 **Heuft-Dorenbosch L**, Spoorrenberg A, van Tubergen A, Landewe R, van der Tempel H, Mielants H, et al. Assessment of enthesitis in ankylosing spondylitis. *Ann Rheum Dis* 2003;**62**:127-32.
- 47 **Gladman DD**, Cook RJ, Schentag C, Feletar M, Inman RI, Hitchon C, et al. The clinical assessment of patients with psoriatic arthritis: results of a reliability study of the spondyloarthritis research consortium of Canada. *J Rheumatol* 2004;**31**:1126-31.
- 48 **Brockbank J**, Stein M, Schentag C, Gladman DD. Characteristics of dactylitis in psoriatic arthritis (PsA). *J Rheumatol* 2001;**28**(suppl 6):1416.
- 49 **Ware JE Jr**, Sherbourne CD. The MOS 36-item short-form health survey (SF-36). I. Conceptual framework and item selection. *Med Care* 1992;**30**:473-83.
- 50 **Husted JA**, Gladman DD, Farewell VT, Long JA, Cook RJ. Validating the SF-36 health survey questionnaire in patients with psoriatic arthritis. *J Rheumatol* 1997;**24**:511-17.
- 51 **Husted JA**, Gladman DD, Cook RJ, Farewell VT. Responsiveness of health status instruments to changes in articular status and perceived health in patients with psoriatic arthritis. *J Rheumatol* 1998;**25**:2146-55.
- 52 **Williams A**. The EuroQol Group: EuroQol—a new facility for the measurement of health-related quality of life. *Health Policy* 1990;**19**:199-208.
- 53 **Sokoll KB**, Helliwell PS. Comparison of disability and quality of life in rheumatoid and psoriatic arthritis. *J Rheumatol* 2001;**28**:1842-6.
- 54 **Mennan R**. Measuring health status in arthritis: the Arthritis Impact Measurement Scales. *Arthritis Rheum* 1980;**23**:146-52.
- 55 **Husted J**, Gladman DD, Farewell VT, Long JA. Validation of the revised and expanded version of the Arthritis Impact Measurement Scales for patients with psoriatic arthritis. *J Rheumatol* 1996;**23**:1015-19.
- 56 **Husted J**, Gladman DD, Long JA, Farewell VT. Relationship of the Arthritis Impact Measurement Scales to changes in articular status and functional performance in patients with psoriatic arthritis. *J Rheumatol* 1996;**23**:1932-7.
- 57 **McKenna SP**, Doward LC, Whalley D, Tennant A, Emery P, Veale DJ. Development of the PsAQoL: a quality of life instrument specific to psoriatic arthritis. *Ann Rheum Dis* 2004;**63**:162-9.
- 58 **Finlay AY**, Khan GK. Dermatology Life Quality Index (DLQI)—a simple practical measure for routine clinical use. *Clin Exp Dermatol* 1994;**19**:210-16.
- 59 **Morgan M**, McCreedy R, Simpson J, Hay RJ. Dermatology quality of life scales—a measure of the impact of skin diseases. *Br J Dermatol* 1997;**136**:202-6.
- 60 **Koo J**, Menter A. The Koo-Menter instrument for identification of psoriasis patients requiring systemic therapy. *National Psoriasis Foundation Psoriasis Forum* 2003;**9**:6-9.
- 61 **Finlay AY**, Kelly SE. Psoriasis—an index of disability. *Clin Exp Dermatol* 1987;**12**:8-11.
- 62 **Gupta MA**, Gupta AK. The Psoriasis Life Stress Inventory: a preliminary index of psoriasis-related stress. *Acta Derm Venereol* 1995;**75**:240-3.
- 63 **McKenna SP**, Cook SA, Whalley D, Doward LC, Richards HL, Griffiths CE, et al. Development of the PSORIQoL, a psoriasis-specific measure of quality of life designed for use in clinical practice and trials. *Br J Dermatol* 2003;**149**:323-31.
- 64 **Kirby B**, Fortune DG, Bhushan M, Chalmers RJ, Griffiths CE. The Salford Psoriasis Index: a holistic measure of psoriasis severity. *Br J Dermatol* 2000;**142**:728-32.
- 65 **Mease P**, Menter A. Quality of life issues in psoriasis and psoriatic arthritis: Outcome measures and therapies from a dermatological perspective. *J Am Acad Dermatol* 2005 (in press).
- 66 **Fries JF**, Spitz P, Kraines RG, Holman HR. Measurement of patient outcome in arthritis. *Arthritis Rheum* 1980;**23**:137-45.
- 67 **Daltroy LH**, Larson MG, Roberts NW, Liang MH. A modification of the Health Assessment Questionnaire for the spondyloarthropathies. *J Rheumatol* 1990;**17**:946-50.
- 68 **Blackmore MG**, Gladman DD, Husted J, Long JA, Farewell VT. Measuring health status in psoriatic arthritis: the Health Assessment Questionnaire and its modification. *J Rheumatol* 1995;**22**:886-93.
- 69 **Husted JA**, Gladman DD, Long JA, Farewell VT. A modified version of the Health Assessment Questionnaire (HAQ) for psoriatic arthritis. *Clin Exp Rheumatol* 1995;**13**:439-43.
- 70 **Tennant A**, Hillman M, Fear J, Pickering A, Chamberlain MA. Are we making the most of the Stanford Health Assessment Questionnaire? *Br J Rheumatol* 1996;**35**:574-8.
- 71 **Beaton DE**, Boers M, Wells GA. Many faces of the minimal clinically important difference (MCID): a literature review and directions for future research. *Curr Opin Rheumatol* 2002;**14**:109-14.
- 72 **Wells GA**, Tugwell P, Kraag GR, Baker PR, Groh J, Redelmeier DA. Minimum important difference between patients with rheumatoid arthritis: the patient's perspective. *J Rheumatol* 1993;**20**:557-60.
- 73 **Mease P**, Ganguly L, Wanke E, Yu E, Singh A. How much improvement in functional status is considered important by patients with active psoriatic arthritis: applying the outcome measures in rheumatoid arthritis clinical trials. *Ann Rheum Dis* 2004;**63**(suppl 1):391.
- 74 **World Health Organization**. International classification of functioning, disability and health: ICF. Geneva: WHO, 2001.
- 75 **Cieza A**, Ewert T, Ustun TB, Chatterji S, Kostanjsek N, Stucki G. Development of ICF Core Sets for patients with chronic conditions. *J Rehabil Med* 2004;(suppl 44):9-11.
- 76 **Smets E**, Garssen B, Bonke B, De Haes J. The multidimensional fatigue inventory (MFI) psychometric qualities of an instrument to assess fatigue. *J Psychosom Res* 1995;**39**:315-25.
- 77 **Krupp LB**, LaRocca NG, Muir-Nash J, Steinberg AD. The fatigue severity scale. Application to patients with multiple sclerosis and systemic lupus erythematosus. *Arch Neurol* 1989;**46**:1121-3.
- 78 **Webster K**, Cella D, Yost K. The Functional Assessment of Chronic Illness Therapy (FACIT) Measurement System: properties, applications, and interpretation. *Health Qual Life Outcomes* 2003;**16**:79.
- 79 **Belza B**, Henke C, Yelin E, Epstein W, Gilliss C. Correlates of fatigue in older adults with rheumatoid arthritis. *Nurs Res* 1993;**42**:93-9.
- 80 **Schentag CT**, Cichon J, MacKinnon A, Gladman DD, Urowitz MB. Validation and normative data for the 0-10 point scale version of the fatigue severity scale (FSS). *Arthritis Rheum* 2000;**43**:S177.
- 81 **Gladman DD**, Farewell VT, Wong K, Husted J. Mortality studies in psoriatic arthritis: results from a single outpatient center. II. Prognostic indicators for death. *Arthritis Rheum* 1998;**41**:1103-10.
- 82 **Mease P**. TNF α therapy in psoriatic arthritis and psoriasis: How and why does it work? *Ann Rheum Dis* 2004;**63**:755-8.