Systematic Literature Review of Residual Symptoms and an Unmet Need in Patients With Rheumatoid Arthritis

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Objective. To evaluate the nature and burden of residual disease in rheumatoid arthritis (RA) in patients who meet treatment targets. Second, for those who did not meet targets, to evaluate how much is due to patient symptoms.

Methods. Prospective and retrospective studies were searched in Medline, Embase, and Cochrane Library in the English language from January 1, 2008 to April 18, 2018; conference abstracts (from January 2016 to April 2018) and reference lists of relevant studies were also screened.

Results. Of 8,339 records identified, 55 were included in the review; 53 were unique studies, including 10 randomized controlled trials. Of these, 48 reported on patients who achieved low disease activity (LDA) or remission. Studies varied in population, treatment goals, and outcome reporting. The proportions of patients with residual symptoms in these studies varied by the definitions used for LDA or remission and were more often reported in patients with LDA than those in remission. The most commonly reported outcome measures were functional disability (n = 34 studies), tender or swollen joints (n = 18), pain (n = 17), patient global assessment (n = 15), and fatigue (n = 14). However, few studies reported the percentage of patients achieving a specific threshold, which could then be used to easily define the presence of residual symptoms.

Conclusion. Residual symptoms are present in some patients despite their achieving LDA or remission, highlighting an unmet need, especially with respect to improving pain, fatigue, and function. Standardized reporting in future observational studies would facilitate better understanding of this issue in defined RA populations.

INTRODUCTION

Rheumatoid arthritis (RA), the world’s most prevalent chronic autoimmune inflammatory arthritis, is characterized by joint inflammation directly leading to pain, functional decline, and fatigue, all of which negatively impact health-related quality of life (HRQoL) and reduce a patient’s ability to work (1). A combination of new therapeutic strategies and new treat-to-target strategies has made sustained clinical remission a primary goal in treating patients with RA. Reaching this goal requires regular assessment of RA activity composite measures and the ability to escalate treatment regimens (2–4).

Despite successes from this approach, remission may not be achievable, and a low disease activity (LDA) state may be an acceptable alternative goal (2). However, for some patients, particularly in the more established phase of disease, it may not be possible to achieve and sustain even LDA, as evidenced in real-world data (5,6). In addition, patients with RA and moderate...
SIGNIFICANCE & INNOVATIONS

- Based on a very large literature search, we have identified that there is a paucity of data and lack of standardization for reporting residual patient-centered symptoms and outcomes in those patients attaining a treatment target.
- Reliance solely on traditional rheumatoid arthritis disease activity targets as acceptable treatment goals risks underestimating the impact on patients’ pain, function, health, and true burden of illness for a meaningful proportion of patients.
- The use of patient-reported outcomes in addition to a treat-to-target approach may provide information that will inform a management decision necessary to address residual symptoms.

A systematic literature review (SLR) was performed, following Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines, in Medline, Embase, and the Cochrane Library. Searches were conducted on April 18, 2018 and were limited to articles published in the English language after January 1, 2008. Conference abstracts (European Alliance of Associations for Rheumatology [EULAR], American College of Rheumatology [ACR], International Society of Pharmacoeconomics and Outcomes Research, and International Society for Pharmacoepidemiology) were also searched from January 2016 to April 2018, as well as reference lists of any relevant systematic reviews and meta-analyses published in the previous 5 years. Search terms included combinations of free text and Emtree/MeSH subject headings for terms relating to RA, LDA, treatment targets, treatment outcomes, and disease activity. Full details of the search strategy are in Supplementary Table 1, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract.

Study eligibility criteria were created using the population, interventions, and outcomes process (see Supplementary Table 2, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). Studies of interest were prospective (including randomized controlled trials [RCTs]) and retrospective studies in patients with RA age ≥18 years who were treated with conventional synthetic disease-modifying antirheumatic drugs (csDMARDs), biologic DMARDs (bDMARDs), or JAK inhibitors according to a treat-to-target strategy. We included studies that reported outcomes for patients who did or did not meet treat-to-target goals (LDA or remission). Outcomes of interest included any patient symptoms (e.g., pain, fatigue, functioning) or disease impact (e.g., HRQoL, absenteeism, or presenteeism) for studies where patients met treat-to-target goals. For patients who did not achieve treat-to-target goals, outcomes of interest were disease impact or reasons for not achieving goals. Due to the wealth of literature, studies were limited to the following countries where treat-to-target is known to be implemented: Canada, France, Germany, Italy, Japan, the Netherlands, Spain, Sweden, the UK, and the US. Multinational studies that included these countries were included.

Titles and abstracts and full texts were screened by one researcher to determine eligibility according to the predefined inclusion and exclusion criteria. A separate researcher performed a random 10% quality check of the included and excluded studies; any disagreements or uncertainties about relevance were determined by consensus. Data from eligible studies were extracted from full-text publications where possible. Data extraction was verified against the source by a second researcher.

RESULTS

Of the 8,339 records identified after removal of duplicates, 55 met the inclusion criteria for this review (Figure 1). Three articles reported on the same RCT; hence, 53 unique studies were included.

Of the 53 included studies, 10 were RCTs and 43 were nonrandomized studies; 42 reported outcomes of interest for patients who achieved treat-to-target goals (8 RCTs and 34 nonrandomized studies), and 5 reported on patients who did not achieve goals (5 nonrandomized studies); 6 reported data of interest for both groups of patients (see Supplementary Tables 3 and 4, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). Studies were identified from all 10 specified countries, the greatest number of which were from Japan, the Netherlands, the UK, and the US.
Substantial heterogeneity among the studies made direct comparisons difficult. For example, study designs included RCTs, prospective observational studies, retrospective studies, and patient interviews. Patient characteristics varied in terms of duration and severity of disease and prior treatments, and studies were diverse in their use of tools for assessing outcomes and the manner in which they were reported. Minimum clinically important differences were often not reported, even for those instruments that have them defined. Furthermore, the studies used a variety of treatment goals for remission or LDA; Disease Activity Score in 28 joints (DAS28) was the most commonly used measure, with a DAS28 score of <3.2 used to define LDA, and a DAS28 score of <2.6 to define remission (Figure 2).

Residual symptoms. Studies reported on a range of symptoms among patients who achieved treat-to-target goals; various toolsmeasurements were used (see Supplementary Table 3, available at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). Most studies reported mean or median values; however, a handful reported the proportion of patients achieving a certain threshold (Tables 1 and 2).

The most commonly reported outcomes were functional disability, pain, fatigue, tender or swollen joints, and physician global assessment of disease activity (PhGA) or patient global assessment of disease activity (PtGA) (Figure 3). Other symptoms such as anxiety, depression, sleep disturbances, and morning stiffness were reported less often.

Functional disability. Functional disability was the most commonly reported symptom, reported in 34 studies (including 6 RCTs) (results in Supplementary Table 5, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). The tool most often used was the Health Assessment Questionnaire (HAQ) or its derivatives: the HAQ disability index (HAQ DI; 27 studies), modified HAQ (M-HAQ; 3 studies), and multidimensional HAQ (MD-HAQ; 1 study). Other tools included the Patient-Reported Outcomes Measurement Information System (PROMIS-29) physical function domain, Michigan Hand Outcome Questionnaire, Funktionsfragebogen Hannover patient questionnaire, McMaster Toronto Arthritis Patient Preference Questionnaire, Signals of Functional Impairment, and Steinbrocker Functional Classification.

Of the 6 RCTs that used the HAQ DI, all defined LDA using the DAS28 (C-reactive protein [CRP] or erythrocyte sedimentation rate [ESR]) (9–14). A HAQ DI score of <0.5 was often used by studies to describe normative physical function. Mean ± SD scores for the HAQ DI ranged from 0.23 ± 0.33 for patients who experienced early remission (DAS score <1.6) after 4 months of methotrexate in the IMPROVED study (9) to...
The C-EARLY study reported that, although the mean HAQ DI score was 0.3, ~20% of patients had residual effects on physical function (15). Additionally, 21 nonrandomized studies reported results from the HAQ (5,16–35), 8 of which included patients in LDA or remission with a mean HAQ score of >0.5 (16,17,21,24,25,27,28,32). Mean ± SD scores ranged from 0.10 ± 0.02 in patients with a swollen joint count (SJC) of ≤1, tender joint count (TJC) of ≤1, and PtGA score of ≤1 (20) to 1.4 ± 0.6 in patients with a DAS28 score of <3.2 (28).

The M-HAQ was used in 3 studies; mean scores ranged from 0.04 for patients in Clinical Disease Activity Index (CDAI) remission (CDAI score ≤2.8) at year 5 (36) to 0.45 for patients with DAS28-ESR scores of >2.6 and ≤3.2 (37). M-HAQ scores were lower for patients in remission than for those with LDA and varied by target used (Table 1). In the one study that used the MD-HAQ, patients in remission (DAS28-CRP score <2.6) had a mean ± SD score of 0.32 ± 0.32 (38).

The level of functional disability depends on the target criteria used. Two studies reported a higher proportion of patients with residual functional impairment (HAQ score >0.5) when a target DAS28 score of <2.6 was used compared with more stringent targets such as ACR/EULAR-defined remission (18,19) (Table 1). Additionally, an analysis of 753 patients from a post-marketing registry (16) also found that residual HAQ scores in patients treated with infliximab or golimumab were higher when the definition of remission used was a DAS28 score of <2.6 than when the definition was a CDAI score of ≤2.8 or an SDAI score of ≤3.3, with mean ± SD scores of 0.76 ± 0.67, 0.57 ± 0.56, and 0.57 ± 0.57, respectively. Additionally, patients with LDA had higher residual HAQ scores than those in remission: 0.86 ± 0.67 for DAS28 score of ≤3.2, 0.94 ± 0.70 for CDAI score of ≤10.0, and 0.89 ± 0.69 for SDAI score of ≤11.0 (16).

Whether remission is sustained also impacts the presence of functional disability. An observational study of patients receiving anti–tumor necrosis factor (anti-TNF) therapy over 6 years (25) observed that patients in sustained remission (defined as DAS28 score <2.6 on at least 2 consecutive occasions and for ≥6 months) had lower HAQ scores (better physical functioning) than those who had only occasional remission. Full physical function (HAQ score = 0) was achieved by 43% of patients with DAS28-defined sustained remission, 60% with SDAI-defined sustained remission, and only 12% with no DAS28-defined sustained remission. Furthermore, an analysis of the Better Anti-Rheumatic PharmacOTherapy (BARFOT) trial (26) reported that 17.5% of patients in remission (DAS score <2.6) at years 1, 2, 5, and 8 had a HAQ score of ≥1 at 8 years, indicating that some patients still experienced significant disability despite sustained remission.

Pain. Pain was reported in 17 studies, including 3 RCTs. Thirteen reported pain using a visual analog scale (VAS), 3 used the 36-item Short Form 36 (SF-36) health survey bodily pain domain, and 1 each used the PROMIS-29 pain interference domain and an 11-point scale (results in Supplementary Table 6, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract).
Table 1. Studies reporting percentage of patients achieving a threshold value for functional disability, pain, and fatigue*

| Author, year (ref.) | Country | Outcome definition and assessment tool | Definition of remission or LDA | Results, no./total no. (%) of patients |
|---------------------|---------|---------------------------------------|--------------------------------|---------------------------------------|
| **Functional disability** |         |                                       |                                |                                       |
| C- EARLY study (13,15,55) |        | HAQ DI score >0.5 at week 52          | Sustained LDA (DAS28-ESR score ≤3.2 both at weeks 40 and 52) | CZP standard dose 17/84 (19.3) CZP reduced frequency 25/126 (19.8) CZP stopped 19/79 (24.1) |
| Perrotta et al, 2018 (18) | Italy  | Residual functional impairment (HAQ DI score >0.5) | DAS28-CRP score <2.6 ACR/EULAR remission† CDAI score <2.8 SDAI score <3.3 DAS28-CRP score <3.2 CDAI score <10 SDAI score <11 DAS28-ESR score <1.6 DAS28-ESR score <2.6 DAS28-ESR score <2.0 ACR/EULAR remission SDAI score ≤3.3 | 9/47 (19.1) 0/12 (0) 1/16 (6.25) 2/19 (10.5) 15/59 (25.4) 14/51 (27.4) 14/56 (25) 10/72 (13.9) 7/56 (12.5) 1/22 (4.5) 1/23 (4.3) 1/28 (3.6) |
| Sakellariou et al, 2013 (19) | Italy  | HAQ DI score >0.5                     | DAS28-ESR score <1.6 DAS28-ESR score <2.6 DAS28-ESR score <2.0 ACR/EULAR remission SDAI score ≤3.3 | 10/72 (13.9) 7/56 (12.5) 1/22 (4.5) |
| Einarsson et al, 2016 (25) | Sweden | Full physical function (HAQ DI score = 0) after 6 years | DAS28 sustained remission No DAS28 sustained remission§ SDAI sustained remission¶ | 43‡ 12‡ |
| Svensson et al, 2016 (26) | Sweden | HAQ DI score ≥1 at 8 years             | DAS28 sustained remission No DAS28 sustained remission§ SDAI sustained remission¶ | 60‡ 17.5‡ |
| Navarro-Millán et al, 2013 (37) | US    | M-HAQ score = 0                      | ACR/EULAR remission† M-HAQ score >0 and ≤0.5 M-HAQ score >0.5 | 423/2,351 (18) 71/2,351 (3) |
| **Pain** |         |                                       |                                |                                       |
| Perrotta et al, 2018 (18) | Italy  | Residual pain defined as VAS score >10 mm (range 0–100 mm) | DAS28-CRP score <2.6 CDAI score <2.8 SDAI score <3.3 ACR/EULAR remission SDAI score <11 | 24/47 (51.1) 2/19 (10.5) 0/12 (0) 34/59 (57.6) 3/56 (5.4) |
| Navarro-Millán et al, 2013 (37) | US    | VAS score = 0 (range 0–10 cm) VS score = 1 (range 0–10 cm) VAS score = 2 (range 0–10 cm) VAS score ≥3 (range 0–10 cm) | ACR/EULAR remission† | 1,364/2,351 (58) 212/2,351 (9) 118/2,351 (5) |
| Altawil et al, 2016 (42) | Sweden | Significant remaining pain, VAS score >20 (range 0–100 mm) | Good EULAR response Moderate EULAR response | 123/421 (29) 280/402 (70) |
| **Fatigue** |         |                                       |                                |                                       |
| Druce et al, 2016 (47) | UK    | Partial remission of fatigue: SF-36 VT domain >5th percentile from a matched general population (>12.5 on the SF-36 VT [scale 0–100]) Complete remission of fatigue: SF-36 VT domain >25th percentile from a matched general population (>50 on the SF-36 VT [scale 0–100]) Nonremission of fatigue >62.5 on the SF-36 VT (score ≥ general population median) | DAS28 score <2.6 | 255/271 (83) |
| Navarro-Millán et al, 2013 (37) | US    | 0 on VAS (range 0–10 cm) 1 on VAS (range 0–10 cm) 2 on VAS (range 0–10 cm) ≥3 on VAS (range 0–10 cm) | ACR/EULAR remission† | 682/2,351 (29) 1,011/2,351 (43) 282/2,351 (12) 353/2,351 (15) |

* ACR = American College of Rheumatology; CDAI = Clinical Disease Activity Index; CZP = certolizumab pegol; DAS28-CRP = Disease Activity Score in 28 joints using the C-reactive protein level; DAS28-ESR = DAS28 using the erythrocyte sedimentation rate; EULAR = European Alliance of Associations for Rheumatology; HAQ = Health Assessment Questionnaire; HAQ DI = Health Assessment Questionnaire disability index; LDA = low disease activity; M-HAQ = modified Health Assessment Questionnaire disability index; SDAI = Simplified Disease Activity Index; SF-36 = Short Form 36 (health survey); SF-36 VT = SF-36 vitality scale; VAS = visual analog scale.
† Swollen joint count of 28 joints, tender joint count of 28 joints, CRP level, and patient global assessment of disease activity VAS ≤1.
§ DAS28 score of <2.6 on at least 2 consecutive occasions and for at least 6 months.
¶ SDAI score of <3.3 on at least 2 consecutive occasions and for at least 6 months.
Ten studies (9,14,22–24,26,35,39–41) reported mean or median VAS scores. For patients in remission, the average VAS score ranged from 3 mm (range 0–12; scale 0–100 mm) for patients in Boolean remission (40) to 3 cm (interquartile range 2–4, scale 0–10 cm) for patients with a DAS in 44 joints score of ≤2.4 and SJC score of ≤1 plus ultrasound remission (22). For patients
with LDA, the mean ± SD VAS score ranged from 12.8 ± 15.5 for patients with sustained LDA (14) to 28.3 (scale 0–100 mm) for patients with moderate or good EULAR response who perceived their overall health not to have improved (41). Only 4 studies reported an average pain VAS score of <1 cm (10 mm), and these were all for patients in remission (23,35,40,41).

Three studies (18,37,42) reported patients achieving a certain VAS value (Table 1). First, a prospective study of patients taking biologics defined residual pain as a VAS score of >10 mm (18). Residual pain was reported by 51.1% of patients with a DAS28-CRP score of <2.6 compared with 6.25% with a CDAI score of <2.8, 10.5% with an SDAI score of <3.3, and 0% meeting ACR/EULAR remission criteria. The proportion of patients experiencing residual pain was higher in patients who achieved only LDA. Second, a cross-sectional analysis of the Corrona registry (37) also found that most patients in ACR/EULAR remission reported low pain scores on the VAS (scale 0–10 cm). However, 9% had a VAS score of 2, and 5% reported pain with a score of ≥3. Finally, a case–control study of patients with early RA treated with methotrexate for 3 months defined remaining pain as a VAS score of >20 (scale 0–100 mm) (42). The study reported that 29% and 70% of patients with a good or moderate EULAR response, respectively, experienced remaining pain. In the good-response group, remaining pain was significantly associated with high baseline HAQ score and low ESR.

Of the 3 studies (12,41,43) that used the SF-36 bodily pain domain, mean scores ranged from 16.3, for patients with moderate or good EULAR response and who considered their health to have improved after treat-to-target strategy aimed at achieving fast remission (41), to 72.1 ± 19.3 for patients with stable LDA (defined as a DAS28 score <3.2 for >6 months prior) (12).

Fatigue. Fatigue was reported in 14 studies (including 5 RCTs); 5 used a VAS, 4 used the Functional Assessment of Chronic Illness–Fatigue (FACIT-F) subscale; 2 used the Bristol RA Fatigue Multidimensional Questionnaire (BRAF-MDQ); and 1 study each used the SF-36 vitality domain, PROMIS-29 fatigue T score, and an 11-point scale (results in Supplementary Table 7, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract).

Of the 5 studies using a VAS (17,35,37,39,41), 3 reported the average score, which was >1 cm (10 mm) in all 3 studies. Notably, a study of patients at the Leiden Early Arthritis Clinic (35) showed that patients who achieved early sustained DMARD-free remission experienced less fatigue than those who achieved remission late or had intermediate remission, with median VAS scores of 3, 13, and 21 (scale 0–100 mm), respectively. Only 1 study reported the proportion of patients achieving certain VAS values (37) (Table 1). In this study, only 29% of patients who met the ACR/EULAR remission criteria reported a fatigue score of 0 (scale 0–10 cm), while 15% reported a score of ≥3.

Four RCTs used the FACIT-F (scale 0–52) to assess fatigue, with a higher score indicating less fatigue. Two studies reported mean ± SD FACIT-F scores, which ranged from 42.0 ± 8.7 for patients with sustained LDA (DAS28 score ≤3.2 at week 36 and average DAS28 score ≤3.2 in weeks 12–36) in the PRESERVE study (14) to 44.4 ± 7.3 for responders (DAS28-ESR score ≤3.2 at week 39 and DAS28-ESR score <2.6 at week 52) in the PRIZE study (44). The REFLEX study (45) reported that 20.6% of responders (ACR 20% improvement criteria) did not achieve a minimum clinically important difference of ≥3.56. The final study (46) reported only the change from baseline in ACR and EULAR responders.

The one study using the SF-36 vitality domain (scale 0–100) found that few patients with a DAS28 score of <2.6 achieved complete fatigue remission (47). Partial fatigue remission was achieved by 83% of patients, while complete fatigue remission was achieved by only 37.3% of patients (Table 1). Those with non-remission of fatigue had a higher proportion of steroid use, stroke, and depression and lower scores for pain and function than the remission group.

Other symptoms. Very few studies reported on symptoms of anxiety, depression, sleep disturbance, or morning stiffness. PtGA and PhGA scores, TJCs, and SJCcs were generally low when reported (see Supplementary Tables 8–10, available on the Arthritis Care & Research website at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). These outcomes are often required to be low according to criteria that define LDA and remission. However, many remission criteria still allow for a TJC, SJC, PhGA, or PtGA score of 1.

For example, an observational study in patients with early RA who were treated with DMARDs (23) defined residual disease activity as ≥1 swollen or tender metatarsophalangeal joints. Residual disease activity was present in 10 of 38 patients (26.3%) in Boolean-based remission (TJC, SJC, CRP, and PtGA ≤1), 12 of 42 patients (28.6%) in Boolean clinical practice remission (TJC, SJC, and PtGA ≤1), 22 of 61 patients (36.1%) with an SDAI score of ≤3.3, and 25 of 63 patients (39.7%) with a CDAI score of ≤2.8. Similarly, an analysis of patients with early RA who were treated with anti-TNF agents (21) reported on those with an SJC and TJC of ≥1. Fifty-one percent of patients who met a DAS28-ESR score of <2.6, and 34.4% who were in ACR/EULAR remission had an SJC in 28 joints of >0. Additionally, 25.2% of patients in remission defined as a DAS28-ESR score of <2.6, and 21.5% who were in ACR/EULAR remission had a TJC in 28 joints of >0.

Disease impact. The disease impact of illness may include a number of factors such as quality of life, activity impairment, and health care resource utilization. In this SLR, we found that HRQoL and work or activity impairment were 2 aspects of disease impact that were reported in patients who achieved treat-to-target goals in 19 studies (8 RCTs and 11 nonrandomized studies) (see Supplementary Table 3, available at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). However, the nature of how
residual symptoms affected disease impact was generally not reported, and the considerable heterogeneity between the studies makes comparisons difficult.

Patients not achieving treatment goals. Only 5 studies (all nonrandomized, 3 of which were abstracts) reported reasons why patients did not achieve treatment goals (see Supplementary Table 4, available at http://onlinelibrary.wiley.com/doi/10.1002/acr.24369/abstract). No studies were identified that reported how patients’ symptoms contributed to disease burden.

One abstract reported (48) that patients who did not achieve their goals were significantly older, had higher rates of disease activity, had a greater number of TJC, and had higher PtGA and HAQ DI scores than those who did achieve their goals. Similarly, an analysis of the BARFOT cohort (26) reported that female sex, current smoker, disease activity at baseline, and nonremission status at 6 months were predictive of persistent disease.

Three of the studies linked not achieving treatment goals to the treatment regimens that were used. One abstract (49) reported that, of patients with high disease activity (DAS28-ESR score >5.1), only 64 patients (9%) were receiving bDMARDs, a much lower proportion than the 18–20% of patients with less active RA. This was reported to be due to failed response to bDMARDs, unwillingness, or contraindications. Yamazaki and Takanashi (48) found that 21 of 60 patients (35%) not achieving LDA or remission were thought to have been insufficiently treated, and 18 of 60 (30%) were insufficiently treated for their complications. Furthermore, 6 of 9 patients from focus groups and interviews (50) stated that they “did not think they had found a treatment regime that controlled their RA properly and expressed disappointment about past drug combination which did not lead to symptom relief.” Three participants reported that they were wary of trying more intensive treatments due to potential side effects.

DISCUSSION

The management of RA has changed greatly over the last 2 decades. This evolution reflects a number of factors. First, earlier intervention in the course of disease was facilitated by the formulation of the ACR/EULAR 2010 classification criteria for RA (51). Second, researchers came to understand that best outcomes are achievable when systemic and local inflammatory disease activity is optimally suppressed over time. This was facilitated by the treat-to-target approach (2), which is recommended for use and widely adopted in clinical practice in the geographic regions from which the studies included in this SLR were undertaken. Third, the choices for effective pharmacotherapy have expanded, with introduction of a variety of classes of targeted biologic therapies with differing mechanisms of action (i.e., TNF inhibitor and other bDMARDs) and, more recently, introduction of small molecule, orally available, JAK inhibitors. Because of these advances, the outlook for a contemporary patient presenting with RA is genuinely better than was the case a generation ago. The presence and magnitude of residual symptoms may depend on the stringency of the remission or LDA measure attained as well as on the durability of this level of response. But even when remission criteria are met, it is not straightforward to interpret the cause underlying any symptomatic deficit. In the case of functional disability, it may be the consequence of aging given that HAQ scores increase with age in long-standing disease; and furthermore, there may be an irreversible component of the HAQ due to past structural damage (52). Everyday clinical experience illustrates that unmet need remains (1). It also raises the question of the underlying pathophysiologic or other causes of residual symptomatology and how this knowledge can be best harnessed to inform a management decision. In this SLR, we sought to better understand the evidence regarding ongoing symptomatology and its relationship to unresolved disease activity and to gain insight into the nature of remaining symptoms experienced by patients despite their attaining the recommended therapeutic target of remission or LDA. These questions are of importance because we have long known that many of the symptoms associated with active RA represent generic features of inflammation, such as pain, fatigue, and functional deficit. But it is also clear that such symptoms have a multifactorial etiology and may be the result of both inflammatory and noninflammatory processes. Furthermore, with respect to inflammatory causes, it might be the case that symptoms in any given individual respond differentially to distinct classes of targeted therapies.

This SLR is notable for revealing a relative paucity of data relevant to these questions. This in part reflects the challenges of extrapolating cohort-level data to the needs of an individual. It is also a reflection of the tendency to limit data capture to a small set of end points in clinical trials. But perhaps most revealing, in the case of observational studies, is the lack of standardization with respect to the outcome measures and scales employed, therefore complicating the ability to make comparisons. Nevertheless, with respect to the range of symptoms and/or outcomes reported in this SLR, the best symptomatic outcomes show relationships to the attainment and maintenance of the more robust measures of remission. These observations support the treat-to-target principle as being an effective approach to the overall goal of patient well-being in RA. However, it also has to be acknowledged that only a minority of patients both attain and maintain a robust remission over the longer term (53,54), and the question arises as to whether we can employ patient-reported outcome measures to identify aspects of life that matter most to the individual and then make subsequent use of this information to inform a management choice that will achieve an overall goal of well-being. The current treat-to-target recommendations have limited utility in a subgroup of patients for whom the desirable target cannot be attained despite a change in pharmacotherapy up to every 3 months. For such patients, it may be more appropriate to focus on personalized treatment
goals such as a meaningful reduction in pain or fatigue or patient-defined functional goals for activities that matter to them. This may require a careful choice of pharmacologic intervention as well as nonpharmacologic approaches that incorporate patient-centered approaches. The practice of protocol-driven medicine in a treat-to-target approach has the advantage of optimizing outcomes at a cohort level and for the minority who attain and maintain the target, particularly remission. This has the potential disadvantage of treating the disease activity while ignoring the patient who has the disease, thus making care less personalized.

The main limitation of this review arises from the multiple sources of heterogeneity among the studies. The range of treatment goals makes comparisons difficult, as the differing requirements lead to varying levels of residual symptoms that can be present. Additionally, outcomes from different countries may not be comparable due to geographic differences in responses to symptoms such as pain.

In conclusion, despite evidence to support adoption of a treat-to-target paradigm in the routine management of RA, residual symptoms still occur in patients achieving LDA or remission. This SLR confirms that there is an unmet need, especially with respect to improving pain, fatigue, and function where possible, even when a target of LDA or remission has been met. Standardized reporting in future observational studies and use of measures that inform on the interference of these symptoms in the daily lives of patients would facilitate better understanding of this issue in defined RA populations. From a pragmatic perspective, these findings suggest that setting personalized goals for the individual in addition to the practice of treat-to-target may inform individualized management as part of holistic care.

AUTHOR CONTRIBUTIONS

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be submitted for publication. Dr. Taylor had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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Analysis and interpretation of data. Michaud, Pope, van de Laar, Curtis, Kannowski, Mitchell, Bell, Workman, Paik, Cardoso, Taylor.

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