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Treatment Satisfaction, Patient Preferences, and the Impact of Suboptimal Disease Control in a Large International Rheumatoid Arthritis Cohort: SENSE Study

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Background: Patients' needs and perspectives are important determinants of treatment success in rheumatoid arthritis (RA). Assessing patients' perspectives can help identify unmet needs and enhance the understanding of treatment benefits.

Objective: The SENSE study assessed the impact of inadequate response to disease-modifying antirheumatic drugs (DMARDs) on treatment satisfaction, disease outcomes, and patient perspectives related to RA disease management.

Methods: SENSE was a noninterventional, cross-sectional study conducted in 18 countries across Europe, Asia, and South America. Adult patients with poorly controlled RA of moderate/high disease activity were eligible. Patient satisfaction was assessed by the Treatment Satisfaction Questionnaire for Medication (TSQM v1.4). Treatment adherence, healthcare resource utilization (HRU), quality of life (QoL), work ability, digital health literacy (DHL), patient preference information, and treatment strategy were also assessed.

Results: A total of 1624 patients were included in the study: most were female (84.2%) and middle-aged, and mean disease duration was 10.5 years. Mean TSQM global satisfaction subscore was 60.9, with only 13.5% of patients reporting good treatment satisfaction (TSQM global ≥ 80). The strongest predictor of good treatment satisfaction was treatment with advanced therapies. Most patients (87.4%) reported good treatment adherence. In general, patients had impaired QoL and work ability, high HRU, and 67.4% had poor DHL. Leading treatment expectations were "general improvement of arthritis" and "less joint pain". Most patients preferred oral RA medications (60.7%) and rapid (≤ 1 week) onset of action (71.1%). "Increased risk for malignancies" and "increased risk for cardiovascular disease" were the least acceptable side effects. Despite suboptimal control, advanced therapies were only used in a minority of patients, and DMARD switches were planned for only half of the patients.

Conclusion: Suboptimal disease control negatively impacts treatment satisfaction, work ability, QoL, and HRU. Data collected on patient perspectives may inform shared decision-making and optimize treat-to-target strategies for improving patient outcomes in RA.

Keywords: adherence, digital health literacy, patient preference, rheumatoid arthritis, treatment satisfaction

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Introduction

Rheumatoid arthritis (RA) is a treatable condition, largely due to the expansion of the number of available, effective disease-modifying antirheumatic drugs (DMARDs). However, in the real world, rates of remission or low disease activity

range from 32% to 55% at 12 months, suggesting that approximately half of patients with RA may have high unmet needs for more effective disease management,¹ at least in part because treat-to-target (T2T) principles are not universally applied. Barriers to implementing T2T principles are numerous, including treating physicians' clinical inertia, patients' concerns related to treatment intensification, and reimbursement of effective medications.²

As well as effective therapies, patients' characteristics and perspectives, including treatment preferences and expectations, are important determinants of treatment success in RA.³ Patient-reported outcomes (PROs) and patient preference information (PPI) are of growing importance and can provide meaningful insights into patients' perspectives, reflecting an improved patient–physician partnership in RA care. PROs are measurements of patients' health conditions as reported directly by patients, without amendment or interpretation by clinicians. PPI refers to qualitative or quantitative assessments of patients' relative preference or acceptability of specified choices among treatment options (eg, risks and benefits) or other attributes (eg, route of drug administration and need for combination with other drugs).^{4–6}

Adherence to RA therapy is a key component in the effective management of RA.⁷ Patients' satisfaction with therapy has an impact on medication adherence, treatment continuation, and future treatment choices, but these aspects of satisfaction have, as yet, been studied only in small populations of patients with RA.^{8,9} Satisfaction is also closely linked to patients' treatment expectations,¹⁰ which relate to patients' everyday lives, personal concerns, and levels of pain. However, patients' expectations can be very different from rheumatologists' treatment goals.¹¹

Adherence may be improved via patient support programs (PSPs)¹² and health education, including easily accessible digital information, which may contribute to patient empowerment and more effective self-care.⁷ The effectiveness of health education via digital information relies on digital health literacy (DHL), ie, on patients' abilities to access and use credible online health information, so they may become better informed to make decisions about their treatment, which, in turn, may positively influence their health behavior.¹³

Evaluating PPI and PROs and understanding what drives patient satisfaction with therapy can provide valuable insight into the impact of RA on patients' lives and a more comprehensive assessment of unmet needs and treatment benefits.^{3,14–16} The SENSE study was designed to better understand the impact of inadequate response to

DMARDs on treatment satisfaction and various disease outcomes and to analyze patient attitudes and perspectives toward treatment and their disease. SENSE also provided an opportunity to assess DHL in a large multinational cohort of patients with RA using a validated tool, the eHealth Literacy Scale (eHEALS).

Methods

Design

SENSE was a noninterventional, cross-sectional study conducted in 18 countries across Europe, Asia, and South America, including Argentina, Brazil, Bulgaria, Chile, Croatia, Czech Republic, Estonia, Greece, Hungary, Japan, Latvia, Lithuania, Poland, Romania, Russian Federation, Slovakia, Turkey, and Uruguay.

Patients

Eligible patients were adults (≥ 18 years old) diagnosed with RA and fulfilling the 1987-revised American College of Rheumatology (ACR) or the 2010 ACR/European League Against Rheumatism (EULAR) classification criteria for RA, who attended a routine visit (previously scheduled as a follow-up of their RA disease), and were receiving treatment with an approved conventional synthetic DMARD (csDMARD), targeted synthetic DMARD (tsDMARD), or biologic DMARD (bDMARD). At the time of enrollment, patients had been exposed to ≤ 2 bDMARDs. Eligible patients had suboptimal disease control, defined as moderate or high disease activity (Disease Activity Score in 28 joints [DAS28] > 3.2 for ≥ 1 month but for ≤ 4 months prior to enrollment) despite receiving a full tolerable dose of DMARD therapy that had been administered for ≥ 3 months.

Patients had to understand the language of their country of residence, be willing and able to complete PRO questionnaires, and provide written authorization to the investigator to use/disclose personal and/or health data, or to provide informed consent if required by local regulations. Community- or hospital-based medical centers experienced in the treatment of RA were selected as research sites. Convenience sampling was employed; all consecutive patients attending a routine visit and fulfilling enrollment criteria could be included.

Patients were enrolled from September 14, 2018 to May 31, 2019. Data available in the medical records of each patient were collated and documented in an electronic case-report form. For each patient, the study consisted of

one visit in which the patient was asked to complete PRO and PPI questionnaires. According to the requirements for noninterventional or observational studies, no additional diagnostic or monitoring procedures were applied to the patients included in the study other than those that would ordinarily be applied in the course of the therapeutic strategy.

The study was conducted in accordance with the Declaration of Helsinki and has been approved by local ethics committees according to local laws and regulations for participating countries ([Supplementary Table 1](#)). All patients provided written authorization to the investigator to use/disclose personal and/or health data or to provide informed consent if required by local regulations.

Assessments

Clinical parameters, including comorbidities at the Medical Dictionary for Regulatory Activities system organ class level, and sociodemographic characteristics were collected for all patients, along with details of their current treatment and planned treatment strategy. Physicians were asked to report if switching to a different DMARD was planned for their patient, and the mode of action of planned treatment switches was captured.

The primary outcome was patients' RA treatment satisfaction, which was assessed using the Treatment Satisfaction Questionnaire for Medication (TSQM; v1.4).¹⁵ The TSQM provides a validated score for four subscales: effectiveness, side effects, convenience, and global satisfaction. Good treatment satisfaction was defined as a global satisfaction score ≥ 80 out of a possible 100.¹⁵

Health-related QoL (HRQoL) was assessed using validated PRO measures: the Functional Assessment of Chronic Illness Therapy-Fatigue (FACIT-F) scale;¹⁷ Health Assessment Questionnaire-Disability Index (HAQ-DI);¹⁸ 36-item Short Form Health Survey (SF-36; Version 2) Physical and Mental Component Summaries (PCS, MCS);¹⁹ and the Work Productivity and Activity Impairment (WPAI)-RA (Version 2.0) questionnaire.²⁰ Visual analog scales (VAS) using numeric rating scales (NRS) were used to assess severity of morning stiffness and pain in the past 7 days (range 0 = "no stiffness/pain" to 10 = "worst possible stiffness/pain"). Duration of morning stiffness in the past 7 days (in minutes) was also recorded. Patients also reported their adherence to medication, which was assessed using a VAS, a validated measure

that highly correlated with electronic monitoring results in patients with chronic conditions, including RA.^{21,22} Good adherence was defined as $\geq 80\%$ on the VAS.²³

DHL was determined using the eHEALS,¹³ a self-report tool of 10 questions based on individuals' perceptions of their own skills and knowledge within each measured domain, providing scores ranging from 8 to 40. The instrument is designed to provide a general estimate of consumer eHealth-related skills and offers insight into the ability of patients to seek, find, understand, use, and appraise health information from electronic sources and to apply the knowledge gained to addressing or solving health problems.¹³ A higher total eHEALS indicates greater perceived skills at using online health information to help solve health problems; a score of < 26 was considered to represent poor DHL.²⁴

Patient perspectives were assessed using PPI questionnaires developed by AbbVie ([Supplementary Text 1](#)), based on the available literature^{3,16,25} and patient research due to limited availability of appropriate validated tools. Preferences for pharmacologic treatment were assessed via a 6-item direct preference assessment questionnaire related to different treatment attributes. Respondents were asked to select only one alternative at a time, without providing information on the order of the unselected alternatives. The frequency (%) of answer options given by study patients for each question – which included preferences for route of administration, combination therapy, time to effect, and acceptable side effects – were recorded. These attributes were not combined in a single scenario as a discrete choice experiment.²⁶ Expectations for pharmacologic treatment were assessed using an 11-item questionnaire with a 7-point NRS (1 = "no improvement needed" to 7 = "the most improvement needed"). Need for a PSP was assessed using a 17-item questionnaire with a 7-point NRS (1 = "not needed at all" to 7 = "very much needed").

Statistical Analysis

Approximately 1500 patients with RA were planned to be included in the study, with the sample size calculation based on information of global satisfaction measured by TSQM. Based on a study conducted in patients treated with tumor necrosis factor inhibitor (TNFi) for 3 months, it seemed reasonable to assume that the standard deviation (SD) for the global satisfaction was about 20, with mean values ranging from 79 to 83.²⁷

Descriptive statistics were used to describe treatment satisfaction, expectations, and patient preferences using the full analysis set (FAS), which included all patients who fulfilled all inclusion criteria. Patient data and all PROs were assessed by subgroups, which were stratified by treatment, disease activity, and presence of comorbidities. For subgroup analyses of continuous variables, Wilcoxon–Mann–Whitney tests were used; and for categorical variables, Chi-square tests or Fisher exact tests were used. Analysis of variance tests were conducted for each individual score of a questionnaire to assess treatment expectations (dependent variable) and with the grouping variable csDMARD vs bDMARD vs tsDMARD. Spearman's rank correlation coefficient (ρ) was calculated for selected associations (self-reported adherence, DAS28, need for PSP in general, and individual scores of the questionnaire to assess treatment expectations) with various parameters. Multiple logistic regression analyses were conducted to determine variables that might predict good treatment satisfaction, patients' preference for oral therapy, preference for monotherapy, good adherence, and willingness of the treating physician to switch medication. Prespecified prognostic factors were used in the regression models with dichotomized or grouped categorical independent variables ([Supplementary Text 2](#); [Supplementary Table 2](#)). Backward elimination with a significance level of 0.05 was applied to stepwise automatically select relevant predictors. Effect estimates and *P*-values using the Chi-square test are provided for each prespecified predictor.

Results

A total of 1629 patients were enrolled in this study, of which 1624 were included in the FAS.

Clinical Parameters and Sociodemographic Characteristics

Most patients were female (84.2%) with a mean age of 58.4 years (range 18–90) and a mean (SD) disease duration of 10.5 (9.3) years. The education level for most patients (64.6%) was secondary school education or less ([Table 1](#)). About one-quarter (26.2%) of patients were in full-time work, 16.0% were unemployed (and seeking work), and 51.1% were retired. RA had had a considerable impact on patients' work-life, as 11.9% had retired early due to RA, while 6.0% were unemployed due to RA.

Table 1 Sociodemographic Characteristics

Characteristics	Full Analysis Set (n=1624)
Female, n (%)	1368 (84.2)
Age, years, mean (SD)	58.4 (13.1)
Race, n (%)	
White	1431 (88.1)
Black	7 (0.4)
Asian	139 (8.6)
Mixed	46 (2.8)
Other	1 (0.1)
Occupation, n (%)	
Employed full-time	426 (26.2)
Employed part-time (not due to RA)	28 (1.7)
Employed part-time due to RA	53 (3.3)
Attending school or university	12 (0.7)
Unemployed (not due to RA)	163 (10.0)
Unemployed due to RA	97 (6.0)
Early retired (not due to RA)	87 (5.4)
Early retired due to RA	194 (11.9)
Regularly retired	549 (33.8)
Education, n (%)	
No formal education	27 (1.7)
Primary school	388 (23.9)
Secondary school (eg high school)	633 (39.0)
Non-university, professional education	311 (19.2)
University	263 (16.2)
Residence, n (%)	
Urban center (>80,000 inhabitants)	814 (50.1)
Town (10,000–80,000 inhabitants)	432 (26.6)
Rural area (<10,000 inhabitants)	378 (23.3)

Abbreviations: RA, rheumatoid arthritis; SD, standard deviation.

Patients had moderate-to-high disease activity, with mean tender and swollen joint counts of 28 joints (TJC28 and SJC28) of 7.8 and 4.6, respectively, and a mean DAS28 with C-reactive protein (CRP) of 4.7 ([Table 2](#)). Radiographic structural damage (based on X-ray data ≤ 6 months from enrollment) was apparent in 76.9% of patients with valid data (n=934). Rheumatoid factor (RF) status was positive for 78.4% of patients with valid data (n=1044), as was anti-citrullinated protein antibody (ACPA) status for 74.1% (valid data n=717), with a total of 82.2% (n=1060 of patients with valid data < 3 months from enrollment) of seropositive patients, defined as RF and/or ACPA positive. Most patients had

Table 2 Rheumatoid Arthritis Disease Activity and Joint Assessment

Measure, Mean (SD)	Full Analysis Set (n=1624) ^a
TJC28 (0–28)	7.8 (5.4)
SJC28 (0–28)	4.6 (4.2)
PtGA (VAS, 0–10 cm)	5.7 (2.1)
PhGA (VAS, 0–10 cm)	5.2 (1.9)
DAS28-CRP ^b	4.7 (1.1)
DAS28-ESR ^c	5.0 (1.1)
CDAI (0–76)	23.3 (10.6)
SDAI (0–86) ^b	26.9 (12.9)

Notes: ^aUnless otherwise stated. ^bn=1500 patients with valid data. ^cn=1515 patients with valid data.

Abbreviations: CDAI, Clinical Disease Activity Index; DAS28-CRP, Disease Activity Score in 28 joints with C-reactive protein; DAS28-ESR, Disease Activity Score in 28 joints with erythrocyte sedimentation rate; PhGA, Physician Global Assessment; PtGA, Patient Global Assessment; SD, standard deviation; SDAI, Simplified Disease Activity Index; SJC28, swollen joint count in 28 joints; TJC28, tender joint count in 28 joints; VAS, visual analog scale.

comorbidities (73.3%; n=1191), the most common of which were cardiac disorders (36.6%; n=595), musculoskeletal connective tissue disorders (25.6%; n=415), endocrine disorders (19.8%; n=322), and metabolic/nutrition disorders (18.5%; n=301). Vascular disorders were present in 9.9% (n=160), and psychiatric disorders in 6.8% (n=111) of patients.

Healthcare resource utilization (HRU) was relatively high in this population, with frequent visits to healthcare professionals. Most (98%; n=1153) of these visits were outpatient visits, although 7.4% (n=87) of patients had an emergency room visit, and 13.5% (n=159) had been hospitalized for a mean (SD) length of stay of 7.5 (6.1) days in the 3 months prior to the study. For patients who had had an outpatient or emergency room visit in the 3 months prior to study visit, the mean (SD) number of visits was 2.2 (2.5) and 1.6 (1.3), respectively.

Medication and Treatment Strategy

csDMARDs were the most frequently prescribed medications, used by 1447 (89.1%) patients, followed by bDMARDs (32.6%; n=530) and tsDMARDs (2.4%; n=39; [Table 3](#)). Across all classes of DMARDs, more than half of the patients (53.9%; n=875) received their DMARD as monotherapy. Within each of the DMARD classes, 709/1447 (49.0%) patients received csDMARDs as monotherapy, 150/530 (28.3%) received bDMARDs as

Table 3 Current Medications Administered for Rheumatoid Arthritis

Medication, n (%)	Full Analysis Set (n=1624)
csDMARD	1447 (89.1)
Methotrexate	991 (61.0)
Leflunomide	363 (22.4)
Hydroxychloroquine	244 (15.0)
Sulfasalazine	212 (13.1)
Chloroquine	22 (1.4)
bDMARD	530 (32.6)
Adalimumab	116 (7.1)
Etanercept	115 (7.1)
Rituximab	73 (4.5)
Tocilizumab	72 (4.4)
Abatacept	52 (3.2)
Infliximab	39 (2.4)
Certolizumab pegol	32 (2.0)
Golimumab	30 (1.8)
tsDMARD	39 (2.4)
Tofacitinib	34 (2.1)
Baricitinib	5 (0.3)
Systemic glucocorticoids	163 (10.0)

Abbreviations: bDMARD, biologic DMARD; csDMARD, conventional synthetic DMARD; DMARD, disease-modifying antirheumatic drug; tsDMARD, targeted synthetic DMARD.

monotherapy, and 16/39 (41.0%) received tsDMARDs as monotherapy. Out of all patients, 21.7% (n=352) and 2.7% (n=44) of patients received double or triple csDMARDs, respectively. A total of 61.0% (n=991) received the anchor drug methotrexate as part of their treatment regime: of those with valid data (n=929), 54.4% received 10 to 15 mg/week, while 39.3% received >15 mg/week. Only 10.0% (n=163) of patients used systemic glucocorticoids for RA (FAS).

Only half (51.8%; n=840) of all patients had a planned switch of medication, despite long-standing disease and suboptimal symptom control with current DMARD therapy. Among these patients, a planned switch to a TNFi was most frequently reported (42.4%; [Figure 1](#)).

Regression analysis revealed that predictors for a decision to switch to a different DMARD by the treating physician included younger patient age ($P=0.0012$), having a high disease activity by DAS28 with erythrocyte sedimentation rate (DAS28-ESR) ($P<0.0001$ for high disease activity >5.1 vs moderate/low disease activity ≤5.1), worse HAQ-DI physical function ($P=0.0062$), lower number of current DMARDs ($P<0.0001$), lower global satisfaction score

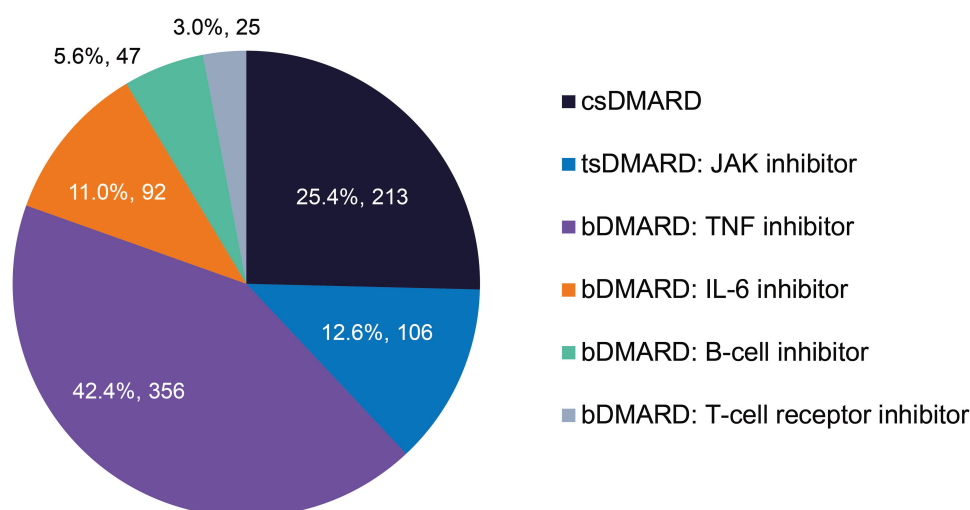


Figure 1 Mode of action for planned DMARD switches.

Note: Treatment switch to a different DMARD was planned for 840 patients (51.8%) out of 1623 patients with valid data.

Abbreviations: bDMARD, biologic DMARD; csDMARD, conventional synthetic DMARD; DMARD, disease-modifying antirheumatic drug; IL-6, interleukin-6; JAK, Janus kinase; TNF, tumor necrosis factor.

($P<0.0001$), and patient preference for oral administration ($P=0.0019$). Predictors for not switching to a different DMARD included a higher number of comorbidities ($P<0.0001$), lower current DAS28-ESR score ($P<0.0001$), lower number of concomitant medications ($P=0.0002$), current treatment with ts/bDMARDs ($P<0.0001$), and higher TSQM effectiveness subscore ($P<0.0001$). In addition, physicians in Asia were significantly less likely to switch to a different DMARD than physicians in South America ($P=0.0069$) or Europe ($P=0.0136$).

A switch to the recently introduced tsDMARD class was reported in 12.6% of patients with planned switching and was significantly associated with RA disease duration ≥ 2 years, DAS28-ESR >5.1 , higher HAQ-DI, and patients' preference for oral administration.

Overall, 61.4% of FAS patients took any medication for their current comorbidities; the mean (SD) number of medications administered for comorbidities was 2.1 (1.9). The most frequent medications for comorbidities were selective beta blocking agents (12.6%), vitamin D and analogs (12.0%), plain angiotensin-converting enzyme inhibitors (10.9%), and proton pump inhibitors (10.3%). Platelet aggregation inhibitors (excluding heparin) were used by 4.2% of patients.

Primary Outcome: Patients' RA Treatment Satisfaction

Mean (SD) TSQM global satisfaction subscore was 60.9 (20.9) (Figure 2). Based on a TSQM global satisfaction

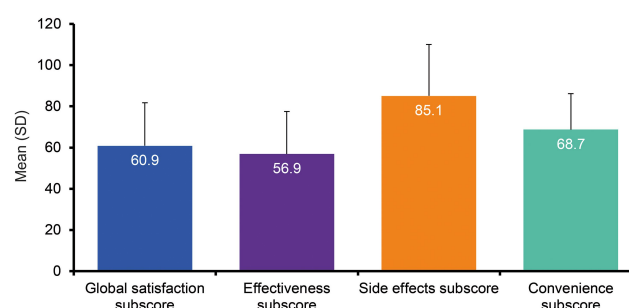


Figure 2 Treatment Satisfaction Questionnaire for Medication subscores.

Notes: Full analysis set ($n=1624$). Each TSQM subscore ranges from 0 to 100, with higher scores denoting greater satisfaction.

Abbreviations: SD, standard deviation; TSQM, Treatment Satisfaction Questionnaire for Medication.

subscore cut-off value of ≥ 80 , as few as 13.5% of patients reported good treatment satisfaction.

In a subgroup analysis of those treated with advanced therapies (b/tsDMARDs), patients treated with an oral DMARD ($n=52$) had significantly lower mean TSQM global satisfaction (60.4 vs 68.9, $P=0.0212$) and effectiveness (54.2 vs 63.1, $P=0.0023$) subscores compared with patients treated with parenteral DMARD ($n=517$). Patients on DMARD monotherapy ($n=875$) had significantly lower mean TSQM global satisfaction (59.7 vs 62.2, $P=0.0262$) and effectiveness (55.9 vs 58.0, $P=0.0268$) subscores compared with patients using DMARD combination therapy ($n=749$).

Multivariate logistic regression analysis indicated that high disease activity (DAS28-ESR >5.1) was a negative predictor of good treatment satisfaction, while current

Table 4 Predictors of Good Treatment Satisfaction (TSQM Global Satisfaction Score ≥ 80)

Variables	Effect Estimate	P-value ^a	OR (95% CI)
High disease activity (DAS28-ESR >5.1)	-0.6045	0.0026	0.546 (0.368, 0.810)
SF-36 PCS	0.0554	<0.0001	1.057 (1.036, 1.078)
SF-36 MCS	0.0340	<0.0001	1.035 (1.019, 1.050)
Psychiatric disorders	0.9475	0.0004	2.579 (1.531, 4.344)
Current treatment with ts/bDMARDs	1.2665	<0.0001	3.548 (2.593, 4.856)

Note: ^aChi-square test, multivariate logistic regression model using the full analysis set (n=1624).

Abbreviations: bDMARD, biologic DMARD; CI, confidence interval; DAS28-ESR, Disease Activity Score in 28 joints with erythrocyte sedimentation rate; DMARD, disease-modifying antirheumatic drug; MCS, Mental Component Summary; OR, odds ratio; PCS, Physical Component Summary; SF-36, 36-item Short Form Health Survey; tsDMARD, targeted synthetic DMARD; TSQM, Treatment Satisfaction Questionnaire for Medication.

treatment with ts/bDMARD, higher SF-36 PCS and SF-36 MCS scores, and the presence of a psychiatric disorder (not further specified) were positive predictors of good treatment satisfaction (Table 4).

Secondary Outcomes: Other PROs

RA was shown to have an impact on measures of HRQoL and work productivity, as shown in Table 5. Work ability showed significant impairment: among patients who were employed (n=413), the median total work productivity impairment score was 50, representing a 50% work productivity loss because of RA. Similarly, patients reported 60% total activity impairment (Table 5).

Of the 1601 patients with valid data, mean (SD) total eHEALS score was 21.3 (8.4). Poor DHL, defined as eHEALS <26 , was evident in two-thirds of patients (67.4%; n=1079). In general, 35.4% of patients found the internet “not useful” or “not useful at all” in helping to inform decisions about their own health, 33.2% found it “useful” or “very useful”, and 31.4% were unsure of its impact. A total of 38.6% of patients believed it is “important” or “very important” to be able to access health resources on the internet, 36.9% considered it as “not important” or “not important at all”, and 24.5% were unsure.

A total of 1623 patients reported their adherence to medication; most (87.4%) reporting good adherence, defined as

Table 5 Patient-Reported Outcomes

Patient-Reported Outcome	n ^a	Mean (SD) ^b
Worst joint pain (0–10)	1623	6.0 (2.5)
Severity of morning stiffness (0–10)	1623	5.0 (2.8)
Duration of morning stiffness, hours, median (IQR)	1488	1.0 (0.5–2.0)
FACIT-F score (0–52)	1623	28.1 (11.2)
HAQ-DI score (0–3)	1624	1.3 (0.7)
SF-36 PCS (0–100)	1623	36.6 (7.9)
SF-36 MCS (0–100)	1623	41.9 (11.0)
WPAI-RA scores, %, median (IQR)		
Presenteeism ^c	418	40.0 (20.0–70.0)
Absenteeism ^c	461	0.0 (0.0–16.7)
Total work productivity impairment ^c	413	50.0 (20.0–70.0)
Total activity impairment	1619	60.0 (30.0–80.0)

Notes: ^aValid data from the full analysis set (n=1624). ^bUnless otherwise stated. ^cEmployed patients only.

Abbreviations: FACIT-F, Functional Assessment of Chronic Illness Therapy-Fatigue; HAQ-DI, Health Assessment Questionnaire-Disability Index; IQR, interquartile range; MCS, Mental Component Summary; PCS, Physical Component Summary; SD, standard deviation; SF-36, 36-item Short Form Health Survey; WPAI-RA, Work Productivity and Activity Impairment-Rheumatoid Arthritis.

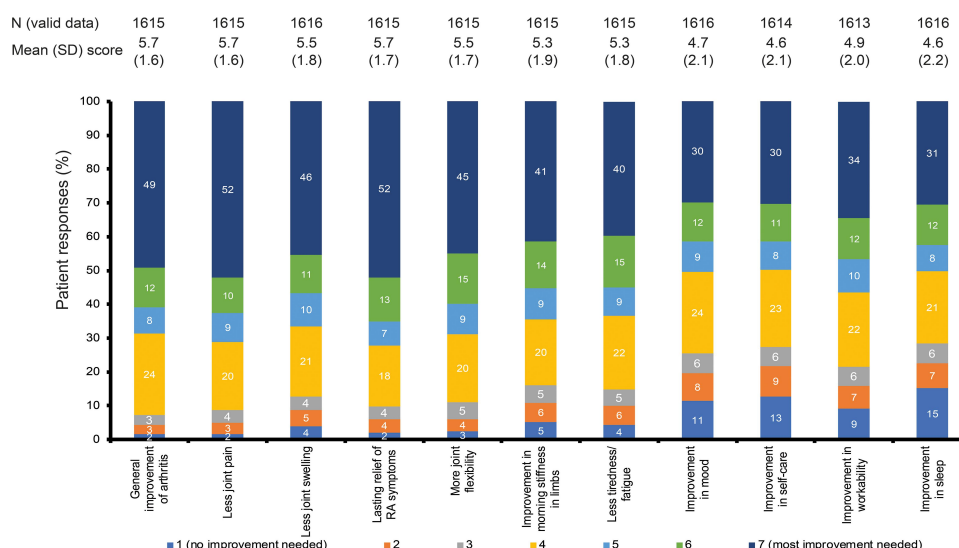


Figure 3 Patients' expectations for RA treatments.

Notes: Patients scored their expectations based on a 7-point numeric rating scale where 1 = no improvement needed to 7 = the most improvement needed. The mean (SD) score and distribution of scores are shown for each treatment expectation. Due to rounding, percentages may not sum to 100%.

Abbreviations: RA, rheumatoid arthritis; SD, standard deviation.

≥80% on the VAS. Mean (SD) self-reported adherence was 91.4% (16.9). After correcting for confounders, the level of self-reported adherence was positively associated with TSQM global satisfaction subscore ($P=0.0059$) and with lower levels of joint pain ($P=0.0473$). There was no significant association with the other variables evaluated in the model ([Supplementary Table 2](#)).

PPI Outcomes: Patient Expectations for Pharmacologic Treatment

When asked about treatment expectations, patients rated “general improvement of arthritis”, “less joint pain”, and “lasting relief of RA symptoms” highest for their treatment expectations, with mean (SD) scores of 5.7 (1.6–1.7) for each ([Figure 3](#)). The strongest association between the top three highest-rated treatment expectations and outcomes was worst joint pain ($\rho=0.31$, $\rho=0.35$, and $\rho=0.31$, respectively; all $P<0.0001$).

PPI Outcomes: Patient Preferences for Pharmacologic Treatment

Most patients preferred oral administration of medication (60.7%) and rapid (≤ 1 week) onset of treatment action (71.1%) ([Figure 4](#)). After correcting for confounders, patient preference for oral administration was significantly associated with older age ($P=0.0005$), duration of RA <2 years ($P<0.0001$), and a greater number of

comorbidities ($P=0.0014$), however, we did not find any significant relationships with the other variables evaluated in the model ([Supplementary Table 2](#)).

While 37.4% of patients felt treatment requiring combination with another drug once a week was acceptable ([Figure 4](#)), 31.3% of patients preferred not to use drug combinations for RA. After correcting for confounders, patient preference for monotherapy was significantly associated with a lower number of current DMARDs ($P=0.0054$) and lower TSQM global satisfaction subscore ($P<0.0001$). We did not find any significant association with the other variables evaluated in the model ([Supplementary Table 2](#)).

In terms of potential side effects of RA therapies, patients indicated that the “most acceptable” side effect was “weight gain”, while “increased risk for malignancies” and “increased risk for cardiovascular disease” were the “least acceptable” ([Figure 4](#)).

PPI Outcomes: Need for PSP

When patients were asked about the need for PSPs, patients assigned the greatest importance for educational materials about RA disease and therapy, followed by the need for a PSP in general ([Figure 5](#)). Conversely, the need for appointment reminders, digital/lifestyle interventions, and medication administration reminders were rated as least important. No strong correlations were found for the need for PSP and other variables.

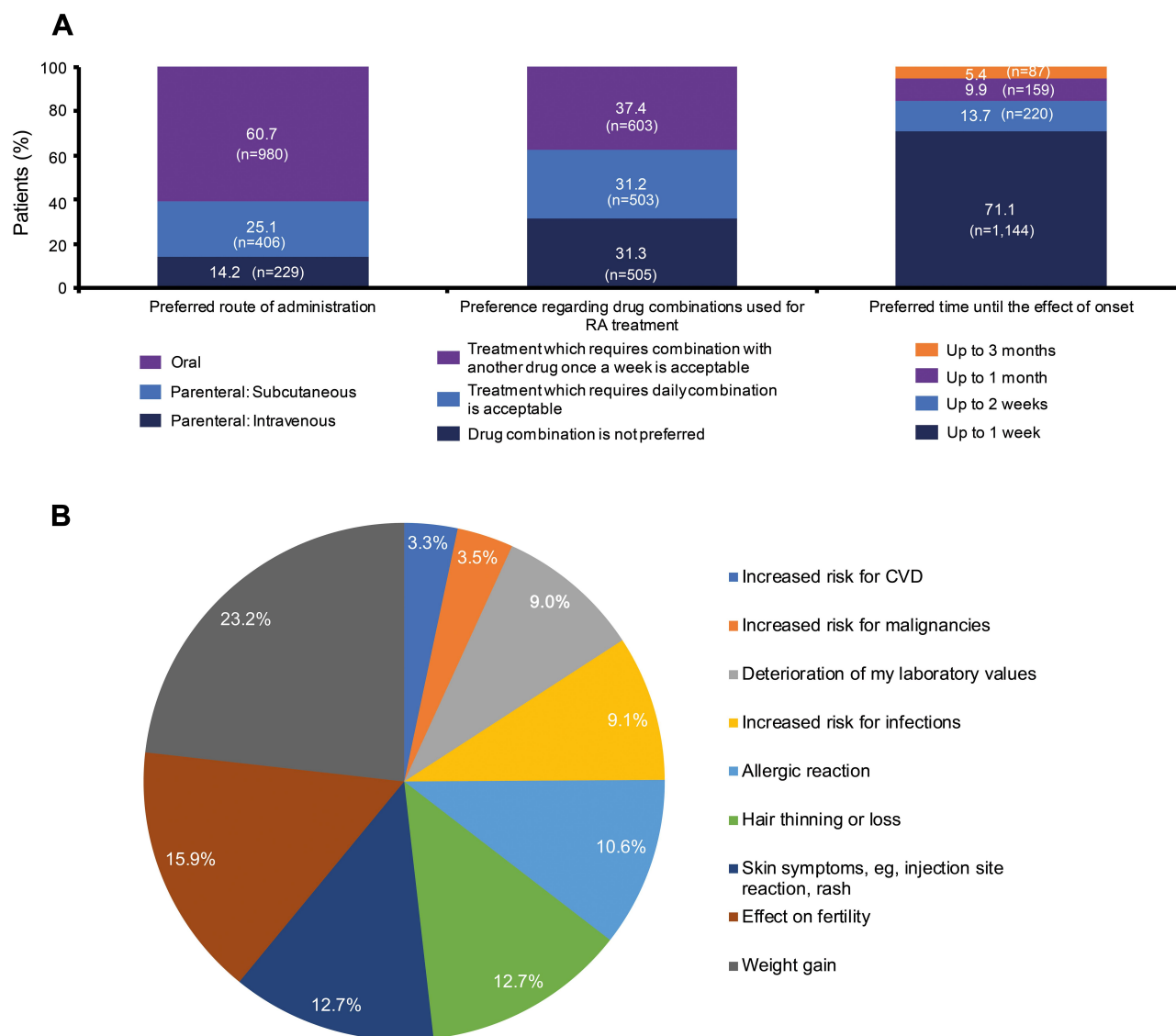


Figure 4 Patients' preferences for RA treatments: **(A)** route of administration, drug combinations, and time to effect; **(B)** acceptability of potential side effects of RA treatments.

Note: **(B)** n=1608.

Abbreviations: CVD, cardiovascular disease; RA, rheumatoid arthritis.

Discussion

In this large international study of patients with poorly controlled RA despite DMARD treatment, only 13.5% of patients reported good treatment satisfaction as measured by a validated tool, despite 87.4% of patients reporting good treatment adherence. Only half of these patients had a planned switch of medication despite suboptimal disease control, impairments in HRQoL and work ability, and high levels of HRU. We found that patients preferred treatments that were administered orally and/or had a rapid onset of action; sustained and general improvement of RA and improvement in joint pain were the leading treatment expectations.

The low treatment satisfaction scores reported in this international population of patients with suboptimal disease control was consistent with previous findings specific to a single country (USA and Germany).^{9,15} In a multivariate model, we identified high vs low disease activity as a negative predictor of good treatment satisfaction, whereas better HRQoL scores and administration of advanced treatment (b/tsDMARD) were positive predictors of good treatment satisfaction. The positive impact of treatment with b/tsDMARDs on patient satisfaction was consistent with previous studies in Germany and Japan, which included patients with RA irrespective of disease control.^{9,28} Consistent with our findings, the German study also reported a positive correlation between treatment

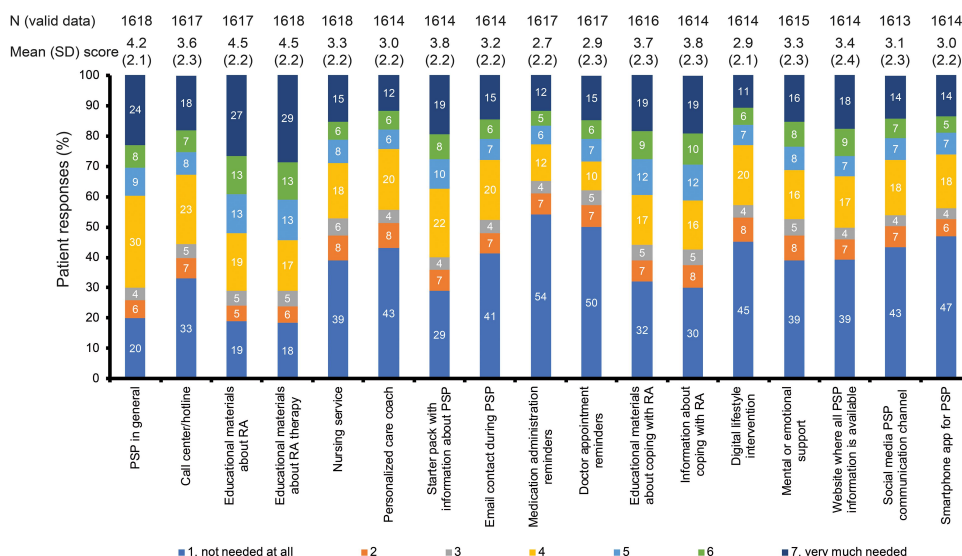


Figure 5 Patients' need for patient support programs.

Notes: Patients scored their perceived need for PSP items based on a 7-point numeric rating scale where 1 = not needed at all to 7 = very much needed. Patients received a short description of PSPs as part of the questionnaire. The mean (SD) score and distribution of scores are shown for each PSP item. Due to rounding, percentages may not sum to 100%.

Abbreviations: PSP, patient support program; RA, rheumatoid arthritis; SD, standard deviation.

satisfaction and improved disease activity and/or physical function.⁹ Of note, we also found a positive association between good treatment satisfaction and the presence of psychiatric disorders. This could imply that a perceived good articular response to therapy may be associated with improving mental state, or that those with mental health disorders are less discerning with respect to expectations of therapeutic response; further research is needed to confirm these hypotheses. Previous reports have suggested that treatment satisfaction may be influenced by the mental state of patients although, contrary to our findings, patients with personality disorders and depression, in general^{29,30} or in a rheumatology setting,^{9,28} experienced lower levels of treatment satisfaction. However, these results must be interpreted with caution, because the specific psychiatric diagnoses were not captured in this study. Moreover, these results do not rule out the influence of other unknown confounding factors, as satisfaction can include other aspects that were not evaluated in our study such as level and access to service, changes in disease activity during the particular therapy, unmet expectations, patient–physician relationship, and patients' beliefs and concerns.¹⁰ Assessing and understanding these patient characteristics can help toward identifying patients who may be unwilling to change medication despite suboptimal control.³¹

Study patients had poorly controlled RA with moderate-to-high disease activity for 1 to 4 months despite full

tolerable dosages of DMARD therapy, which should trigger treatment adjustment under a T2T strategy. For such patients, RA guidelines recommend that treatment should be adjusted,^{32,33} yet >40% of patients were still receiving a csDMARD as monotherapy rather than adding or switching to a b/tsDMARD or adding a second csDMARD to therapy. A switch to a different DMARD was planned for only half of the patients and, of those patients who planned a switch, only three-quarters considered stepping up to advanced therapies with a bDMARD or tsDMARD. Although some of these treatment decisions may have been driven by reimbursement restrictions in participating countries with limited healthcare resources, this failure to plan treatment switches despite inadequate disease control suggests a degree of “treatment inertia”. Other real-world data indicated that many patients with poorly controlled RA remain on their current regimens rather than escalating therapy.³⁴ T2T approaches can be demanding for patients and physicians: some patients may not be willing to commit to more frequent and necessary clinic visits, while others may not have easy access to rheumatologists.³⁵ Furthermore, as non-rheumatologists have limited experience with and confidence in administering DMARD therapy, treatment escalation is rarely performed outside of the rheumatology setting. Therefore, more support should be provided to prevent treatment inertia, eg, through patient

education and by raising awareness of the importance of T2T approaches with physicians.

In this study, high disease activity defined by DAS28-ESR >5.1 predicted the decision to switch treatments, while a lower DAS28-ESR score predicted the decision not to switch. A recent study has found that DAS28-ESR is associated with a numerically higher value than DAS28-CRP; such that using 5.1 as an absolute cut-off for DAS28-CRP may underestimate the number of patients with high disease activity.³⁶ Therefore, the distinction between DAS28-ESR and DAS28-CRP as measures of disease activity should be considered, especially when treatment access is dependent on DAS28 disease activity.

Adherence to therapy is essential if patients are to gain the most benefit from their treatments. In the current study, almost 90% of patients reported good adherence to their medication. This is consistent with the upper limits of adherence observed in the international ALIGN study (57–76%)³⁷ and in a systematic review of US studies (16–81% with TNFis including etanercept, adalimumab, and infliximab).³⁸ The high adherence rate in the current study suggests that suboptimal disease control in this population is unlikely to be due to poor adherence, although there may be a possibility of ascertainment or response bias. We also found that treatment adherence was significantly associated with treatment satisfaction and lower levels of pain, one of the leading patient expectations identified in this study. This finding underlines the importance of effective pain control in RA.

Although the SENSE population largely comprised older persons, with only 26.2% in full-time employment, suboptimally controlled RA had a substantial impact on working life: Of those employed, based on responses to the WPAI-RA questionnaire, 50% of total work productivity loss and 40% of presenteeism were attributable to RA. The levels of impairment in SENSE appear greater than the findings from a previous cross-sectional study in a similar setting using the same validated tool (WPAI-RA).³⁹ Reduced work productivity is associated with poor HRQoL driven by greater disease activity, worse mental health, and greater physical disability and pain.^{40,41} The impact of worsening disease can be profound, so it is important that a strategy to achieve the tightest disease control possible is adopted to preserve work ability and decrease the indirect societal cost of the disease.

Consistent with previous research, most patients in SENSE preferred oral monotherapy,^{3,16,25} largely because

of convenience and ease of use. Significant predictors for oral preference included older age, shorter disease duration, and a greater number of comorbidities. Although many patients were open to combination therapy, 31% preferred not to use drug combinations, and the latter preference was associated with a lower current number of DMARDs and a lower treatment satisfaction. It has been suggested that psychosocial factors, such as education, may be associated with a belief in a stronger treatment effect being associated with parenteral and/or combination therapies, but our data do not support this theory.

When considering patient expectations, the highest-rated treatment expectations related to treatment benefit: improvement of arthritis, less joint pain, and lasting relief of symptoms. This is consistent with the findings of a systematic review of treatment preferences in which patients with RA placed most importance on treatment benefit over other treatment attributes, including serious or minor side effects and cost or route of administration.¹⁴ However, patient preferences are variable and mutable, so understanding patient preferences on an individual level is likely to aid shared decision-making between patients and their physicians.

Patient concerns relating to side effects are of growing importance, especially as patients' beliefs about harms associated with medications are significant predictors of adherence in patients with RA.³⁷ Previous reports have suggested that patients were more willing to accept side effects associated with temporary cosmetic changes, such as weight gain or allergic reactions, and least willing to take medications associated with deterioration of laboratory values, infections, or a risk of major toxicity such as cancer.⁴² Similar findings were evident in SENSE, with weight gain being the most acceptable side effect reported by patients, whereas increased risk of malignancies or cardiovascular disease were the least acceptable. A clear understanding of the most acceptable side effects can support shared decision-making at the individual level, as well as the development of new medications at the population level.

Patients recognized a need for a PSP in general, although reminders and digital/lifestyle interventions were considered less important. Adoption of eHealth technologies may be a key strategy in supporting and providing care for patients with chronic diseases, such as RA, and good DHL is a prerequisite for the optimum use of eHealth tools. Among the SENSE patient population, two-thirds had poor DHL, and only one-third found the internet

important or useful in helping them access health resources or make decisions about their own health, which may be a reflection of the relatively advanced age of the population. Other studies have reported poor DHL, highlighting a need to assist patients with poor DHL, so they may realize the potential benefits of eHealth tools as part of PSPs.^{43,44}

There are several limitations to this study. Noninterventional studies have inherent limitations on data quality compared with randomized controlled trials. For instance, patients were selected based on convenience (resulting in potential selection bias) and willingness to complete time-consuming PRO questionnaires, with all PROs having inherent response and assessment bias. Likewise, patient-reported HRU is prone to recall bias. PPI questionnaires used in the study have not been validated, which may limit their reliability and generalizability. Furthermore, while the direct preference assessment method used for the PPI questionnaires is considered a valid tool to determine treatment characteristics that are important to patients, it does not provide information about the degree of importance of these characteristics and the trade-offs patients are willing to make.¹⁶ Data from this observational study portray a diverse, multinational study cohort under real-world conditions. Country-specific analyses are warranted to determine the impact of local culture, and healthcare-related and socioeconomic factors on patient outcomes. Patient selection criteria limit the generalizability of results across all RA patient populations, as often patients with better disease control and shorter duration of disease are recruited for studies. However, this study provides important insight into the proportion of patients with RA who are in need of treatment adjustment, both in line with recent treatment standards and taking into account T2T principles.

Conclusion

The findings from SENSE demonstrate that suboptimal disease control has a significant impact on patients' treatment satisfaction, working life, and HRU. Patient preferences and attitudes may influence treatment satisfaction and adherence to DMARDs, which ultimately will impact real-world efficacy. Poor DHL may represent an obstacle to patients' understanding of information critical to decision-making. Taken together, these results support the commitment to current T2T strategies, and they also highlight the critical need to individualize treatment decision-making.

Abbreviations

ACPA, anti-citrullinated protein antibody; ACR, American College of Rheumatology; bDMARD, biologic DMARD; CDAI, Clinical Disease Activity Index; CRP, C-reactive protein; CVD, cardiovascular disease; csDMARD, conventional synthetic DMARD; DAS28, Disease Activity Score in 28 joints; DHL, digital health literacy; DMARD, disease-modifying antirheumatic drug; eHEALS, eHealth Literacy Scale; ESR, erythrocyte sedimentation rate; EULAR, European League Against Rheumatism; FACIT-F, Functional Assessment of Chronic Illness Therapy-Fatigue; FAS, full analysis set; HAQ-DI, Health Assessment Questionnaire-Disability Index; HRQoL, health-related quality of life; HRU, healthcare resource utilization; IQR, interquartile range; MCS, Mental Component Summary; NRS, numeric rating scale; PCS, Physical Component Summary; PhGA, Physician Global Assessment; PPI, patient preference information; PRO, patient-reported outcome; PSP, patient support program; PtGA, Patient Global Assessment; QoL, Quality of Life; RA, rheumatoid arthritis; RF, rheumatoid factor; SD, standard deviation; SDAI, Simplified Disease Activity Index; SF-36, 36-item Short Form Health Survey; SJC28, swollen joint count of 28 joints; T2T, treat-to-target; TJC28, tender joint count of 28 joints; TNFi, tumor necrosis factor inhibitor; tsDMARD, targeted synthetic DMARD; TSQM, Treatment Satisfaction Questionnaire for Medication; VAS, visual analog scale; WPAI, Work Productivity and Activity Impairment.

Data Sharing Statement

AbbVie is committed to responsible data sharing regarding the clinical trials we sponsor. This includes access to anonymized, individual, and trial level data (analysis datasets), as well as other information (eg, protocols and Clinical Study Reports), provided the trials are not part of an ongoing or planned regulatory submission. This includes requests for clinical trial data for unlabeled products and indications. These clinical trial data can be requested by any qualified researchers who engage in rigorous, independent scientific research and will be provided following review and approval of a research proposal and statistical analysis plan and execution of a Data Sharing Agreement. Data requests can be submitted at any time, and the data will be accessible for 12 months, with possible extensions considered. For more information on the process, or to submit a request,

visit <https://www.abbvie.com/our-science/clinical-trials/clinical-trials-data-and-information-sharing/data-and-information-sharing-with-qualified-researchers.html>.

Ethics Approval

The study was conducted in accordance with the Declaration of Helsinki and has been approved by local ethics committees according to local laws and regulations for participating countries ([Supplementary Table 1](#)). All patients provided written authorization to the investigator to use/disclose personal and/or health data or to provide informed consent if required by local regulations.

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Author Contributions

All authors made substantial contributions to the conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; agreed to submit to the current journal; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

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AbbVie funded this study and participated in the study design, research, analysis, data collection, interpretation of data, review, and approval of the publication. All authors had access to relevant data and participated in the drafting, review, and approval of this publication. No honoraria or payments were made for authorship.

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