Clinical Outcomes of Patients with Rheumatoid Arthritis Treated in a Disease Management Program: Real-World Results

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Background: Care models can affect the clinical outcome of patients with rheumatic and musculoskeletal diseases.

Objective: We aimed to compare how an innovative model of a rheumatoid arthritis disease-management program can improve the clinical outcomes of patients compared to a conventional assessment approach.

Methods: We performed a retrospective analysis of real-world data from clinical records of a cohort of 5078 patients diagnosed with rheumatoid arthritis who were followed up at the Center of Excellence in Rheumatoid Arthritis vs the clinical outcomes reported in the Colombian National Registry of Rheumatoid Arthritis.

Results: We found significant differences in the diagnosis and follow-up between the specialized Center program and the usual care reported by the Colombian National Registry ($p < 0.005$), including the evaluation of rheumatoid factor, Anti-citrullinated antibodies Disease Activity Score, Health Assessment Questionnaire, number of visits to the rheumatologist, and clinical outcomes measured by the level of disease activity. In addition, when comparing the Center's clinical outcomes – from baseline to the last follow-up, we found an improvement in the level of disease activity, with patients classified in remission increasing from 20.8% to 58.5% ($p < 0.005$), and a reduction in those with high disease activity from 18% to 4.7% ($p < 0.005$).

Conclusion: Real-world evidence showed that patients with rheumatoid arthritis who underwent follow-up under an innovative disease-management model improved their clinical outcomes compared with those patients in a conventional assessment program. These results could suggest a way of improving health policies for patients with rheumatoid arthritis.

Keywords: disease management, rheumatoid arthritis, quality of life, evidence-based practice, patients

Introduction

Because of the increasing costs of chronic disease care, several organizations have implemented disease management programs (DMP) attempting to improve the quality of care and reduce overall costs.^{1,2} This situation has stimulated innovative healthcare strategies, including the creation of specialized centers of excellence in rheumatoid arthritis (RA) aimed to optimize results with limited resources.³

Conventional care programs for RA patients in most Latin-American countries exhibit various degrees of fragmentation and segmentation, providing low-performing and low-quality services. These programs have suffered from a shortage of qualified

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rheumatology professionals, with restricted access to high-cost medications, rehabilitation programs, and orthopedic interventions.^{4–7}

To reduce the burden of RA, a model of care based on early diagnosis and effective DMP should incorporate quality improvement methodologies and contextual factors dependent on the individual, environment, and health-disease process, as well as pharmacological and non-pharmacologic treatment strategies.^{8,9}

Also, a team of healthcare-related professionals should deliver such an integrated care model with a multidisciplinary approach, within an environment of quality, opportunity, accessibility, and safety.^{1,2,10}

In Colombia, the healthcare system operates through a mixed public-private model regulated by the Ministry of Health. This system covers 94.6% of the population and includes benefits for high-cost diseases including rheumatoid arthritis for diagnosis, treatment and the access to biological drugs if it is required.^{11,12} Over time it has become clear that a large variety of contracts are available between payers and providers, with gaps in the costs in the health market.⁶ As a result, there is fragmentation and a reduction in the quality of care.¹³

To improve healthcare and regulate cost, the Colombian Ministry of Health developed a national policy for integrating the healthcare system, including the creation of a National Administrative Registry of Rheumatoid Arthritis (NARRA) to evaluate the status of patients living with RA as reported by payers and healthcare providers.^{6,14}

Through this process, real-world data offers an opportunity to stakeholders to improve policy and programs with information on healthcare from multiple sources outside typical clinical research settings, including electronic health records (EHR), claims and billing data, and disease registries.¹⁵

Implementing this policy requires high-quality healthcare providers with available DMPs to reduce barriers of access and minimize the fragmentation of medical services through the delivery of integrated healthcare services.⁶ This article aimed to offer some real-world evidence regarding how an integrated DMP may be able to improve the outcomes of patients with RA compared to conventional approaches such as the NARRA.

Methods

We developed a retrospective real-world data analysis from a detailed review of electronic health records of 5078 patients with RA who regularly visited a specialized

Center under a model of DMP dedicated to RA care in Bogotá, Colombia, between July 1, 2015 and June 30, 2016, patients were diagnosed in the last year or before. A rheumatologist confirmed the diagnosis of RA according to the American College of Rheumatology and European League Against Rheumatism 2010 Classification criteria. All patients were followed up on and received treatment in a DMP protocol under a treat-to-target (T2T) strategy.¹⁶ The data collected were anonymized to protect the confidentiality and privacy of patients.

We then compared the gathered information with the results obtained from the “Situación de la artritis reumatoide en Colombia 2016” (“Situation of rheumatoid arthritis in Colombia 2016”) a document developed and edited by Colombian High-Cost Disease Office (Spanish acronym – Cuenta de Alto Costo) nominated in this work as the National Registry of Rheumatoid Arthritis (NARRA), this registry report data for 68,247 patients with RA.

There were 89 variables collected on demographics, concomitant diagnoses, treatment, and clinical outcomes; including age, gender, disease duration, comorbidities, type of pharmacologic treatment, measurements of disease activity, the disease activity score (DAS28),¹⁷ and functionality by the Health assessment Questionnaire (HAQ).¹⁸ The reporting of this information by healthcare providers to the Colombian High-Cost Disease Office is mandatory by Colombian law.

The Disease Management Program

We treated patients under DMP to provide a multidisciplinary healthcare approach, comprising of the following healthcare areas: rheumatology, nutrition, psychology, physical and occupational therapy, physical medicine, and rehabilitation according to their RA activity status (Table 1).

We followed patients under standard Treat-to-Target (T2T) strategies. We considered a DAS28 score <2.6, ≥2.6 and <3.2, ≥3.2 and <5.1, and ≥5.1 as an indicator of remission, low disease activity (LDA), moderate (MDA), and high disease activity (HDA), respectively.

The clinical follow-up program, according to the DAS28 cut-off values, was designed and followed in the specialized Center, with rheumatologist evaluations every 3–5 weeks (DAS28 ≥5.1); 6–8 weeks (DAS28 between ≥3.2 and <5.1), and 9–12 weeks (DAS28 <3.2). Tender and swollen joint counts and DAS28 using the erythrocyte sedimentation rate were calculated on each visit.

The pharmacologic and non-pharmacologic treatment program required adjustments with values based on the

Table 1 Specialized Center Disease-Management Program Number of Visits per Year

Speciality	Disease Activity		
	High	Moderate	Low
Rheumatology	12	6	4
Physical Medicine and Rehabilitation	4	3	2
Nutrition	2	1	1
Psychology	4	3	2
Physical therapy	6	4	2
Occupational therapy	6	4	2
Educational sessions	6	4	2

standards developed for the specialized Center, according to REAL-PANLAR (Project of implementation and accreditation of centers of excellence in rheumatoid arthritis throughout Latin America)².

The frequency of visits to the multidisciplinary health-care team in the previous year was established according to the results of disease activity and adjusted to the needs and availability of the individual patient. We also included educational programs for patients and blood tests in the DMP (Table 1).

Statistical Analysis

We performed statistical analysis in 3 phases. First, quantitative data of normal distributions are presented as the means and standard deviations (SDs). Non-normally distributed data are presented as the medians and ranges. Qualitative data are presented as frequencies and percentages. Second, we calculated the difference between the principal characteristics of the population and DAS28 and HAQ values between the specialized Center and NARRA

using Chi2 test. Third, we compared the median DAS28 values at baseline and the last visit using Fisher's exact test, assuming a non-normal distribution for DAS28 values. Differences were considered significant at p-values <0.05. We used the STATA v.13 software for statistical analysis.

Results

Table 1 shows the frequency of visits per year for any specialization according to the level of disease activity (at least 4 visits per year to the rheumatologist for patients with LDA and 1 or 2 visits to the interdisciplinary team). Table 2 summarizes the differences of the model based on DMP and conventional methods in the country reported by NARRA.

Regarding DMP, we found statistical differences in RF and ACPA evaluation in the diagnosis and performance of DAS28 and HAQ in a specialized center compared to those in NARRA. The percentage of patients in whom we calculated the DAS28 in the Center was 99%. This figure was much higher than the Colombian national average (39.8%) (Table 2).

In addition, the application rate of HAQ was higher for the specialized center group (71.7%) than that reported by NARRA (21.2%). Finally, we found differences in the number of visits to the rheumatologist per year: 4 for the specialized center compared with 1 visit per year in the NARRA report.

Table 3 summarizes the demographic and clinical characteristics of the 5078 patients with RA evaluated and prospectively followed up at the specialized center between 2015 and 2016, with 1380 patients newly diagnosed during this period. The age group of 50–65 years made up 45% of the total population. Disease duration was 3 years on

Table 2 Comparision of the DMP in the Specialized Center and the Usual Care for RA Reported by NARRA

	Specialized Center		NARRA		P value
	Freq.	Percent	Freq.	Percent	
Diagnosis*					
Performance of RF at diagnosis time	1251	92.5	2819	76	0.0001
Performance of ACPA at diagnosis time	1315	96.1	890	24	0.0001
Follow-up**					
Percentage in which DAS28 was obtained	5027	99.0	27,191	39.8	0.0057
HAQ measurement	3641	71.7	14,519	21.3	0.0051
Number of rheumatology visits per year	4	***	1	***	***
Number of visits with Interdisciplinary team	2	***	Not reported	***	***

Notes: *Specialized center n=1385 NARRA n=3709. **Specialized center n=5078 NARRA n=68,247. ***Not applicable, Number of visits to interdisciplinary team are not available in the NARRA.

Table 3 Demographic and Clinical Characteristics of the Population in the Specialized Center and NARRA

Variable	Specialized Center		NARRA		P-value
	n=5078		n=68,247		
	n/Mean	Percent/SD	n/Mean	Percent/SD	
Age*	59	1.28	57	13.48	0.1320
Sex					
Female	4155	81.8	57,372	84.1	0.1392
Male	923	18.2	10,875	15.9	0.1394
Disease duration (years)*	3	3.4	8	8.44	0.0020
Comorbidities at cohort entry+					
Overweight	1651	32.5	1032	27.8	0.0031
Obesity	812	16	452	12.2	0.0410
Hypertension	1263	24.9	839	22.6	0.0370
Diabetes	231	4.6	238	6.4	0.0065
Chronic Kidney disease	73	1.4	155	4.2	0.0001
Cerebrovascular disease	49	1	90	2.4	0.0153
Type of Treatment					
Only corticoids	534	10.5	13,226	19	0.0010
cDMARD	3508	69.1	44,450	65.1	0.0330
bDMARD	204	4	10,571	15.5	0.0010
cDMARD combined bDMARD	832	16.4	**	**	**

Notes: *Results in means and Standard Deviation (SD). Age reported in the last visit. +Data of comorbidities of NARRA was calculated just in incident population 3709.

**No data available.

average, and the mean follow-up period was 2 years. Regarding comorbidities, we found that being overweight, obesity, and high blood pressure were the most common conditions present at the time of diagnosis or developed during follow-up.

Up to 2/3rds (n=3508; 69.1%) of patients were treated with conventional disease-modifying antirheumatic drugs (cDMARDs), and 204 (4.0%) were treated with biologic DMARDs (bDMARDs), whereas 16.4% were treated with a cDMARD-bDMARD combination. Thus, 20.4% (n=1036) of the patient population was treated with biologic agents. A significant difference was found in the use of DMARD and bDMARD in the RA specialized center compared to that in NARRA (Table 3).

Likewise, in the specialized center, the proportion of patients classified as remission increased from 20.8% to 50.5% between baseline and the last assessment, while the proportion of persons with HDA showed a significant reduction from 18.0% to 4.7% ($p<0.005$) (Table 4).

At last, a comparison of disease activity status between patients who received their healthcare in the specialized center compared to that in the NARRA systems revealed a significant difference in the percentage of patients classified

as LDA over the Colombian national registry. The Colombian NARRA (the conventional healthcare model) showed a higher percentage of patients with HDA compared with the specialized center with DMP results (Table 5).

Discussion

The deficiencies of most Latin American health systems relegate the quality of patients' care to the background, as

Table 4 Outcomes of Disease Management at Baseline and the Last Measurement of Patients with More Than Six Months in the Specialized Center

	Diagnosis		Follow-Up		p-value
	Freq.	Percent	Freq.	Percent	
Disease activity according to DAS28					
Remission	835	20.8	2346	58.5	0.004
Low	1045	26.1	692	17.3	0.013
Moderate	1406	35.1	781	19.5	0.002
High	722	18.0	189	4.7	0.001
Total	4008	100.00	4008	100	

Notes: DAS28 <2.6 Remission; DAS28 ≥2.6 and <3.2 low disease activity; moderate between DAS28 ≥3.2 and <5.1; a DAS28 ≥5.1 high disease activity. Follow-up: Last measurement in the study year (July 2015–June 2016).

Table 5 Comparison of Disease Activity Status Between Different Rheumatology Practices

Disease Activity Classification	Specialized Center	NARRA	P-value
	n=5078	n=27,191	
	%	%	
LDA (Low Disease Activity)	75.8	70.1	0.001
MDA (Moderate Disease Activity)	19.5	22.5	0.012
HDA (High Disease Activity)	4.2	7.3	0.001

Notes: DAS28 score <3.2 low disease activity; a DAS28 score ≥3.2 and <5.1 moderate disease activity; a DAS28 score ≥5.1 high disease activity.

some of their economic-administrative features received priority over chronic and potentially disabling conditions.¹ This situation has stimulated the emergence of innovative healthcare strategies, like the creation of specialized centers with high standards, aimed to seek quality results.¹⁹

This study supports the rising importance of evidence-based, real-world data in Latin American countries like Colombia, through the follow-up of a significant cohort of patients under a DMP. This first exercise allows us to compare the clinical outcomes between innovative and more conventional practices.

We found that the specialized center results in terms of DAS28 (99%) performance, was higher than the Colombian NARRA average (39.8%) and even above the American College of Rheumatology (ACR) Rheumatology Informatics System for Effectiveness (RISE) in the United States results (52.1%).¹⁵ Consequently, our data clearly showed a higher number of patients with LDA and compared to the Colombian registry.

These results also suggest that the implementation of a DMP, incorporating a team of allied health professionals and flexible healthcare model adaptable to the disease activity status, resulted in a significant gap regarding the mean number of follow-up visits to the rheumatologist (4 per year in the specialized center compared to 1 in the NARRA). We assumed this gap may be related to access barriers to rheumatology services, such as a lack of timely and appropriate referral programs, workforce shortages, the concentration of specialists in the metropolitan area, and underserved country areas with a lower likelihood of applying T2T strategies. In addition, the study about integrated care in musculoskeletal disorders suggest that patients attended under an integrative approach with more focus on patients needs, improves the clinical outcomes as the quality of life.²⁰

We are aware that most costs of RA medical attention are due to pharmacologic treatment. In this field, the advent of new therapies (biologic medicines and small-molecule therapeutics) has pushed healthcare expenses.^{21,22} Therefore, improvements in disease activity, functional status, and reducing disability through a comprehensive DMP may contribute to savings in direct and indirect costs associated with early diagnosis, systematic medical management, and RA treatment.^{23,24} Health insurers will need to become more cautious about specialty drug use, focusing on cost-saving options such as DMP, including drug distribution channels, contracting activities, and care coordination.²⁵

Also, most patients with RA are over 50 years of age. This aging population will develop greater risks and have a higher number of hospitalizations in contrast with a younger population. Evidence shows that patients with multiple comorbidities can cost up to 7 times as much as required for patients with a single chronic condition.^{23,26}

In this context, providers and stakeholders need to develop healthcare services to anticipate and prevent complications and maintain stable underlying chronic conditions to preserve functional capacity, minimizing the disease impact on patients and lowering the burden on the national healthcare system.²⁷

The DMP may represent a potential solution to the fragmentation of RA care at the national level, reducing timely access to the healthcare system, and improving RA management (with attention to comorbidities and risk factors). This study aligned with other analyses revealing that when the disease course is monitored, and medications are adjusted accordingly in conjunction with rehabilitation programs addressed by a multidisciplinary healthcare team, the comorbidities often observed during RA may be prevented or, if already present, may have a lower impact on clinical outcomes.²⁸

An important factor increasing the cost of RA care is the use of newer biologic medications. Our results showed that 20% of patients were treated with one of these agents, a percentage higher than the 16% reported by the NARRA at the national level. This may be explained by the fact that the NARRA is a relatively new survey instrument and may under-report data.

Another possible reason may entail fewer access barriers to high-cost treatments in the Center program, compared to individuals seen in primary-care settings or at non-specialized medical clinics for RA, who are required to wait for extended periods for administrative authorization and experience decreased availability of biologic

medications from their insurance companies because of the channel of distribution and others barriers imposed by insurers, this is characteristic in other countries of the region.^{29,30}

Objectives of DMP include the evaluation of functional disability, with remission goals quantified using clinical instruments.^{5,31} HAQ and DAS28 are the most used instruments in RA clinical trials and are also important in DMP patient-centered care.^{32,33}

In the conventional NARRA, which includes data from primary-care physicians, independent rheumatologists, and hospital-based clinics, DAS28 and HAQ may not be regularly obtained, since the importance of these metrics in daily clinical practice may not be recognized as a measurement of the effectiveness of healthcare programs or as an outcome of quality from the patient's perspective.^{34,35}

This first analysis included 3 remarkable limitations. First, there was no remission data of the patients in the Center and NARRA. Second, the differences in the clinical status as the time of disease evolution between the Specialized center and the NARRA could be a limitation; however, this condition has an effect to achieve remission and remission is not compare, for this comparison all patients with DAS28 <3.2 were classified as LDA. Other clinical significance differences as the percentage of chronic kidney disease has not greater effects in the disease activity. Since this is a clinically relevant finding, its absence represents an additional weakness in this analysis which was derived from their retrospective approach. Third, it is also important to point out that this is one of the first comparisons, so there are risks of bias in results analysis that we have not assessed.

Notwithstanding preceding data, these outcomes derived from real-world data may offer stakeholders and clinicians an opportunity for reflection on how to improve policies involving the integration of healthcare components, including patient access to medical attention and drugs, disease-activity and functional monitoring, and the appropriate adjustment of treatment based on a treat-to-target strategy.

Conclusions

We have shown some benefits of DMP for improving the clinical status in patients with RA. Since systematic measurements of the disease activity and functional status of RA patients provides indicators of good clinical practice and quality of care, this process can be an option to consider for RA healthcare providers.

Results from RA patients followed up on in the specialized Colombian Center evidenced that a healthcare model based on a DMP may improve clinical outcomes, reduce access barriers and time for receiving medical attention by a rheumatologist and other allied health professionals, compared with other conventional assessment practices as the NARRA.

Abbreviations

Center, Center of Excellence in Rheumatoid Arthritis; NARRA, National Registry of Rheumatoid Arthritis; RISE, Rheumatology Informatics System for Effectiveness; EHR, electronic health records; T2T, treat to target; DAS28, Disease Activity Score; HAQ, Health Assessment Questionnaire; LDA, low disease activity; MDA, moderate disease activity; HDA, high disease activity; ACR, American College of Rheumatology; cDMARD, conventional disease-modifying activity drugs; bDMARD, biologic disease-modifying activity drugs; PANLAR, Pan American League of Associations of Rheumatology; EBP, evidence based on practice.

Ethics Disclaimer

This study complied with the ethical requirements of the Declaration of Helsinki. According to Resolution 8430 (1993) of the Ministry of Health, this research presents no risks to patients; the study was approved by the Institutional Review Board of BIOMAB, act 004/2018 (code GC.IN.01.FR.03). All patients had previously signed an informed consent for data use; the database was anonymized to protect the confidentiality and privacy of patients.

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