



Sociedad Española  
de Reumatología -  
Colegio Mexicano  
de Reumatología

# Reumatología Clínica

www.reumatologiaclinica.org



Original Article

## Experience and satisfaction with a multidisciplinary care unit for patients with psoriasis and psoriatic arthritis



Ana Urruticoechea-Arana<sup>a</sup>, Marta Serra Torres<sup>b</sup>, Mercedes Hergueta Diaz<sup>b</sup>,  
María Eugenia González Guerrero<sup>c</sup>, Leslie Fariñas Padron<sup>c</sup>, Sara Navarro Martín<sup>c</sup>,  
Kelly Vargas Osorio<sup>c</sup>, Andrés Palacios Abufón<sup>b</sup>, María Jesús García de Yébenes<sup>d</sup>, Estíbaliz Loza<sup>d,\*</sup>

<sup>a</sup> Servicio de Reumatología, Hospital Can Misses, Ibiza, Spain

<sup>b</sup> Servicio de Dermatología, Hospital Can Misses, Ibiza, Spain

<sup>c</sup> Medicina Familiar y Comunitaria, Hospital Can Misses, Ibiza, Spain

<sup>d</sup> Instituto de Salud Musculoesquelética, InMusc, Madrid, Spain

### ARTICLE INFO

#### Article history:

Received 2 June 2017

Accepted 6 July 2017

Available online 24 August 2017

#### Keywords:

Psoriasis

Psoriatic arthritis

Multidisciplinary care

Satisfaction

### ABSTRACT

**Objective:** To describe patient's characteristics, the activity and patient's satisfaction with a multidisciplinary care unit in patients with psoriasis and psoriatic arthritis (PsA).

**Methods:** A retrospective medical records review of patients with psoriasis or PsA attended in a multidisciplinary care unit was performed. Included patients were contacted to fulfill a satisfaction questionnaire. A specific electronic database was set up. Data regarding to patients and their baseline characteristics and the activity of the unit were collected. Descriptive analysis were performed.

**Results:** A total of 112 patients with 154 visits were included in almost 3 years, 54% women, with a mean age of 51 years, 43.7% presented hyperlipidemia and 30.4% arterial hypertension. Half of patients were referred due to diagnostic doubts and the other half for therapeutic problems. After the evaluation of the patients, 66 patients (58.9%) met diagnostic criteria for PsA, and 13 (11.6%) of an inflammatory disease other than PsA, and 95% came back to their usual physician. The most ordered test were laboratory tests (75.6% of patients), followed by X-rays in 57 patients (51.3%). In general the number of patients with different treatments increased, and 55.4% and 42% of patients changed their topic and systemic treatments respectively. The level of satisfaction was very high and all of patients considered that their disease was better controlled in this multidisciplinary care unit.

**Conclusions:** This multidisciplinary care unit has improved the care and satisfaction of patients with psoriasis or PsA, and increased collaboration between rheumatology and dermatology departments.

© 2017 Elsevier España, S.L.U. and Sociedad Española de Reumatología y Colegio Mexicano de Reumatología. All rights reserved.

## Experiencia y satisfacción en una unidad de atención multidisciplinaria para pacientes con psoriasis y artritis psoriásica

### RESUMEN

**Objetivo:** Describir las características de los pacientes, la actividad registrada, así como la satisfacción percibida, de una consulta de atención multidisciplinaria para pacientes con psoriasis o artritis psoriásica (APs).

**Métodos:** Estudio observacional retrospectivo con revisión de historias clínicas de todos los pacientes atendidos en la consulta de atención multidisciplinaria. Se contactó con todos ellos para que contestasen una encuesta de satisfacción. Varios investigadores recogieron datos sociodemográficos y clínicos, así como administrativos incluyendo el número de visitas en una base de datos especialmente generada para este proyecto. Se realizó un análisis descriptivo.

#### Palabras clave:

Psoriasis

Artritis psoriásica

Cuidado multidisciplinaria

Satisfacción

\* Corresponding author.

E-mail address: [estibaliz.loza@inmusc.eu](mailto:estibaliz.loza@inmusc.eu) (E. Loza).

<https://doi.org/10.1016/j.reuma.2017.07.007>

1699-258X/© 2017 Elsevier España, S.L.U. and Sociedad Española de Reumatología y Colegio Mexicano de Reumatología. All rights reserved.

**Resultados:** Se incluyó a 112 pacientes con 154 visitas en casi 3 años, 54% mujeres, y una edad media de 51 años; el 43,7% presentó hiperlipidemia y el 30,4% hipertensión arterial. La mitad fueron referidos por dudas diagnósticas y la otra por problemas terapéuticos. Tras su evaluación, 66 pacientes (58,9%) cumplieron los criterios diagnósticos de APs y 13 (11,6%) de una enfermedad inflamatoria distinta. El 95% regresó a su médico habitual. La pruebas complementarias más solicitadas fueron analíticas (75,6%) y radiografías simples (51,3%). En general, el número de pacientes con nuevos tratamientos aumentó y el 55,4 y el 42% de los pacientes cambiaron sus tratamientos tópico y sistémico, respectivamente. El nivel de satisfacción fue muy alto y todos los pacientes consideraron que su enfermedad estaba mejor controlada en esta unidad.

**Conclusiones:** Esta consulta de atención multidisciplinar ha mejorado el manejo y satisfacción de pacientes con psoriasis o APs y ha incrementado la colaboración entre los servicios de Reumatología y Dermatología.

© 2017 Elsevier España, S.L.U.

y Sociedad Española de Reumatología y Colegio Mexicano de Reumatología. Todos los derechos reservados.

## Introduction

Psoriasis is a chronic inflammatory skin disease characterized by raised, red scaly plaques. Psoriatic arthritis (PsA) is a heterogeneous chronic inflammatory disease that, apart from the skin may affect the peripheral and axial joints, entheses, skin and nails, and other organs.<sup>1–3</sup>

Epidemiologic studies have also shown that, in patients with psoriasis, but specially in those with PsA, associated comorbidities may occur more frequently than expected.<sup>4,5</sup> Cardiovascular disease, metabolic syndrome, obesity, diabetes, fatty liver disease Crohn's disease, ophthalmic disease, depression and anxiety are common comorbidities in these patients.<sup>6</sup>

Taking into account all of this, psoriasis and PsA are heterogeneous and potentially severe diseases, that negatively impact patients quality of life.<sup>7,8</sup>

On the other hand, many of these patients are evaluated and managed by dermatologists and rheumatologist independently. However, in the dermatology clinics, PsA is not always suspected and therefore the diagnosis of the disease may be delayed.<sup>9,10</sup> Furthermore, skin involvement sometimes has little or no attention for rheumatologists, even though it may have a considerable impact on the patient's physical and psychological health.<sup>11,12</sup>

In order to improve patient's outcomes and prognosis, national and international organizations as well as experts and patients recommend multidisciplinary treatment.<sup>13</sup> In this context, in recent years, experiences from a few multidisciplinary care units or models for patients with psoriasis and PsA have been published in the literature.<sup>14,15</sup> Although data from these units are still poor they suggest that skin and joint improved after the multidisciplinary consultation, were better considered compared with usual care and reported very high satisfaction levels. One of this units also described that after evaluation diagnosis and treatment were modified in 32% and 47% of cases, respectively.<sup>15</sup>

The aim of this study was to describe the experience of a multidisciplinary care unit where dermatologists and rheumatologists collaborate closely following pre-defined referral criteria and management protocols (see methods section for more information).

## Methods

This project was approved by the Ethics Committee of the Hospital Can Misses and patients gave informed consent.

### *Multidisciplinary Care Unit*

A multidisciplinary care unit with specialists with experience and interest in psoriatic disease (2 dermatologists, one rheumatologist) was established to attend routine patients with

psoriasis or PsA. Specific referral criteria were generated including: (1) Diagnostic problems (e.g. patients with psoriasis and arthralgia/arthritis); (2) Therapy related issues (e.g. the prescription of a biologic therapy in refractory patients); (3) Comorbidity management; (4) Safety concerns. Before the unit was implemented, a member of this unit followed a trainee program in another multidisciplinary care unit. Patients are referred from the dermatology and rheumatology departments. Visits (once a month) are made in the rheumatology department in order to take advantage of infrastructure (units and nursing staff). When the patient's problems are solved, they return to their reference specialist for standard follow-up. Patients are evaluated and treated using national and international guidelines. Additional tests as laboratory tests, imaging techniques or biopsies are ordered if necessary.

### *Design*

A retrospective observational study was performed. Medical records of patients attended in the multidisciplinary unit from August 2012 to December 2014 were reviewed. Included patients were contacted by telephone to come to the unit and complete the satisfaction questionnaire. However, if the patients was not able to come, the satisfaction questionnaire was completed by telephone. This was anonymous and patients gave informed consent.

### *Patient's Selection and Data Acquisition*

Patients aged 18 or above attended in the unit were included. A specific electronic database was set up for the purpose of this project. The following data were collected: (a) sociodemographics (age, sex), smoking status; (b) clinical variables including comorbidities (cardiovascular disease, hyperlipidemia, arterial hypertension, Diabetes Mellitus, obesity, hyperuricemia, gout, depression, anxiety, uveitis, hepatic disease, others), baseline disease duration; (c) multidisciplinary care unit related variables like number of visits, follow-up, speciality of referral, main referral reason, referral reason by symptoms, final diagnosis (following international diagnostic criteria), final destination, laboratory tests, imaging techniques and others required during the study period, treatment and treatment changes (topical therapy, ultraviolet phototherapy, NSAIDs, systemic steroids, DMARDs, biologics); (d) satisfaction questionnaire. Patients were asked to (1) qualify from 10 (excellent) to 0 (very bad) the general attention in the multidisciplinary care unit, (2) describe the care of the multidisciplinary unit compared with your usual care (much better, better, the same, worse, much worse), (3) describe the quality of the information given in the multidisciplinary unit (very good, good, normal, bad, very bad), and (4) answer the following question: Was your disease better controlled in this multidisciplinary care unit?

## Statistical Analyses

To describe the sample, we used the distribution of frequencies, the mean and standard deviation, or the median and interquartile range, depending on the distribution of the variable.

## Results

A total of 112 patients were analyzed that produced 154 visits (range 1–7, 77% of patients only 1 visit, 16% 2, 7%  $\geq 3$  visits) in almost 3 years. There was no waiting list during this period. Mostly women (54%), with a mean age of 51 years  $\pm$  12 years, and 20.5% were smokers (Table 1). Mean psoriasis and PsA duration were 16  $\pm$  15 years and 6  $\pm$  7 years respectively. Interestingly, in our study population, 43.7% of patients presented hyperlipidemia, 30.4% arterial hypertension, and 27.7% anxiety. A total of 88 patients (78.6%) were referred from the rheumatology unit. We found that 95% of patients eventually returned to their responsible physician.

The main reasons for being referral to the multidisciplinary unit (Table 2) were equally distributed between diagnostic and therapeutic doubts. When the reasons of the referral were analyzed according to the patient's symptoms, most of them were evaluated because of cutaneous symptoms (almost 77%), followed by articular symptoms (17.8%). After the evaluation of the patients, we found that 66 patients (58.9%) met diagnostic criteria for PsA, and 13 (11.6%) presented an inflammatory disease other than PsA.

During the study period (see Table 3), the following tests were ordered: laboratory tests in a total of 85 patients (75.6%), X-rays in 57 patients (51.3%), musculoskeletal US in 4 (3.6%), musculoskeletal MRI in another 4 patients (3.6%), biopsies in 10 (8.9%) and cultures (blood, tissues, etc.) in 4 (3.6%) patients.

Following, after the evaluation of patients, the number of patients with different treatments increased (see Table 4). In the case of topical therapy from 44.6% to 88.4%, the DMARDs from 44% to 54% and biologic treatments from 17% to 29%. The number of patients with systemic steroids slightly increased, from 28.6% to 29.5%. We found that 42% of patients changed their systemic treatment, 33 patients (29.5%) a molecule change and 12 patients (10.7%) a dose change. Only 2 patients stopped their therapy.

**Table 1**  
Baseline Characteristics of Study Sample.<sup>a</sup>

|   |                 |
|---|-----------------|
| Sex (woman)   | 61 (54%)        |
| Age (years) <sup>b</sup>  | 51.4 $\pm$ 12.8 |
| Psoriasis duration (years) <sup>b</sup>                             | 16 $\pm$ 15     |
| PsA duration (years) <sup>b</sup>                                   | 6 $\pm$ 7       |
| Smoking status  |                 |
| Yes   | 23 (20.5%)      |
| No  | 82 (73.2%)      |
| Former smoker   | 7 (6.2%)        |
| Comorbidity   |                 |
| Cardiovascular  | 4 (3.6%)        |
| Hyperlipidemia  | 49 (43.7%)      |
| Arterial Hypertension   | 34 (30.4%)      |
| Diabetes Mellitus   | 10 (8.9%)       |
| Obesity   | 24 (21.4%)      |
| Hyperuricemia   | 12 (10.7%)      |
| Gout  | 3 (2.7%)        |
| Depression  | 19 (17.0%)      |
| Anxiety   | 31 (27.7%)      |
| Uveitis   | 1 (0.9%)        |
| Hepatic disease   | 18 (16.1%)      |
| Others  | 17 (15.2%)      |
| Visits to the multidisciplinary care clinic <sup>b</sup>            | 1.4 $\pm$ 0.9   |
| Speciality of referral (rheumatology)                               | 88 (78.6%)      |
| Follow-up in the multidisciplinary care clinic (years) <sup>b</sup> | 0.2 $\pm$ 0.5   |

Abbreviation: PsA, psoriatic arthritis.

<sup>a</sup> Results are expressed as number and percentage (%) otherwise is indicated.

<sup>b</sup> Mean  $\pm$  standard deviation.

**Table 2**  
Referral Reasons and Final Diagnosis of Patients Attended in the Multidisciplinary Care Clinic.

|                                     | n (%)      |
|-------------------------------------|------------|
| <i>Main referral reason</i>         |            |
| Diagnostic                          | 61 (50%)   |
| Treatment                           | 61 (50%)   |
| <i>Referral reason by symptoms</i>  |            |
| Arthralgia                          | 12 (10.7%) |
| Arthritis                           | 8 (7.1%)   |
| Dactylitis                          | 2 (1.8%)   |
| Axial symptoms                      | 1 (0.9%)   |
| Cutaneous psoriasis                 | 41 (36.6%) |
| Nail psoriasis                      | 3 (2.7%)   |
| Cutaneous problem                   | 45 (40.2%) |
| <i>Final diagnosis</i>              |            |
| PsA                                 | 66 (58.9%) |
| Inflammatory disease other than PsA | 13 (11.6%) |
| Degenerative arthritis              | 5 (4.5%)   |
| Cutaneous psoriasis                 | 15 (13.4%) |
| Other articular diseases            | 1 (0.9%)   |
| Other cutaneous diseases            | 12 (10.7%) |

Abbreviation: PsA, psoriatic arthritis.

**Table 3**  
Patients With Laboratory Tests, Imaging Techniques and Others Required in the Multidisciplinary Care Clinic During the study Period.<sup>a</sup>

| Test                            | n (%)      |
|---------------------------------|------------|
| Laboratory tests                | 85 (75.6%) |
| X-rays                          | 57 (51.3%) |
| Musculoskeletal US              | 4 (3.6%)   |
| Musculoskeletal MRI             | 4 (3.6%)   |
| Biopsy                          | 10 (8.9%)  |
| Cultures (blood, tissues, etc.) | 4 (3.6%)   |

<sup>a</sup> Results are expressed as number and percentage (%).

Abbreviations: US, ultrasound; MRI, magnetic resonance imaging.

**Table 4**  
Treatment Changes in the Multidisciplinary Care Clinic.<sup>a</sup>

| Treatment                             | Prior      | After visit |
|---------------------------------------|------------|-------------|
| <i>Topical therapy</i>                | 51 (44.6%) | 99 (88.4%)  |
| <i>Ultraviolet phototherapy</i>       | 1 (0.9%)   | 1 (0.9%)    |
| <i>NSAID</i>                          | 74 (68.7%) | 83 (74.1%)  |
| <i>Systemic steroids</i>              | 31 (28.6%) | 33 (29.5%)  |
| <i>DMARD</i>                          |            |             |
| Methotrexate                          | 34 (30.4%) | 44 (39.3%)  |
| Leflunomide                           | 9 (8%)     | 9 (8%)      |
| Cyclosporine                          | –          | 1 (0.9%)    |
| Sulfasalazine                         | 4 (3.6%)   | 3 (2.7%)    |
| Others                                | 4 (3.6%)   | 3 (2.7%)    |
| <i>Biologic therapy</i>               |            |             |
| Etanercept                            | 7 (6.2%)   | 9 (8%)      |
| Adalimumab                            | 5 (4.5%)   | 13 (11.6%)  |
| Infliximab                            | 4 (3.6%)   | 4 (3.6%)    |
| Golimumab                             | 1 (0.9%)   | 3 (2.7%)    |
| Ustekinumab                           | 1 (0.9%)   | 2 (1.8%)    |
| Others                                | 1 (0.9%)   | 1 (0.9%)    |
| <i>Changes in topic treatments</i>    |            |             |
| Without changes                       | 50 (44.6%) |             |
| Molecule change                       | 59 (52.7%) |             |
| Doses change                          | 2 (1.8%)   |             |
| Treatment cessation                   | 1 (0.9%)   |             |
| <i>Changes in systemic treatments</i> |            |             |
| Without changes                       | 65 (58.0%) |             |
| Molecule change                       | 33 (29.5%) |             |
| Doses change                          | 12 (10.7%) |             |
| Treatment cessation                   | 2 (1.8%)   |             |

Abbreviations: NSAID, non-steroidal anti-inflammatory drugs; DMARD, disease modifying antirheumatic drugs.

<sup>a</sup> Results are expressed as number and percentage (%).

**Table 5**  
Satisfaction Questionnaire Results.<sup>a</sup>

| Question  | Result     |
|---|------------|
| Qualify from 10 (excellent) to 0 (very bad) the general attention in the multidisciplinary care unit <sup>b</sup> | 9 ± 1.2    |
| Describe the care of the multidisciplinary unit compared with your usual care                                     |            |
| Much better   | 52 (46.8%) |
| Better  | 51 (45.9%) |
| The same  | 8 (7.2%)   |
| Worse   | 0 (0%)     |
| Much worse  | 0 (0%)     |
| Quality of the information given in the multidisciplinary care unit   |            |
| Very good   | 58 (52.2%) |
| Good  | 48 (43.2%) |
| Normal  | 4 (3.6%)   |
| Bad   | 1 (0.9%)   |
| Very bad  | 0 (0%)     |
| ¿Was your disease better controlled in this multidisciplinary care unit?  | 112 (100%) |
| YES   |            |

<sup>a</sup> Results are expressed as number and percentage (%) otherwise is indicated.

<sup>b</sup> Mean ± standard deviation.

Finally, all but 10 patients answered the satisfaction survey (92% response rate). The level of satisfaction was in general very high (Table 5).

The care of the multidisciplinary unit compared with your their care was considered much better of better for 103 patients (almost 93% of the study sample). The same way, the quality of the information given in the multidisciplinary care unit was good or very good for 106 patients (95.4%). All of patients considered that their disease was better controlled in this multidisciplinary care unit.

## Discussion

We have shown the experience of a multidisciplinary care unit for patients with psoriasis or PsA. The main objectives of this unit are to facilitate an early diagnosis of PsA, improve management of patients with psoriasis and PsA, and increase cooperation between the rheumatology and dermatology departments. Referred patients are attended (at the same by a dermatologist and a rheumatologist) in a few days (maximum 1 month), are carefully evaluated including the assessment of comorbidities and the treatment is agreed by the specialists following national and international guidelines.

One of the positive contributions and target of this unit is an early diagnosis of PsA. This is especially relevant since a diagnostic delay of more than 6 months might contribute to poor radiographic and functional outcome in PsA patients.<sup>10</sup> According to our data, an important rate of patients eventually met PsA criteria. Although we cannot demonstrate this, it is possible that since the unit was set up dermatologist feel more comfortable and referral patients earlier with any kind of diagnostic doubts. In fact, it has been shown that traditional care, where patients are attended by rheumatologists and dermatologist independently, might lead to diagnosis delay.<sup>9,10</sup> In the literature other multidisciplinary care units have shown their ability to make at least an earlier diagnosis in these patients, and have shown that there were many discrepancies with the previous diagnosis.<sup>14–16</sup> Moreover, like in our cohort, this multidisciplinary approach facilitate a differential diagnosis for cutaneous and joint problems as well.<sup>14,15</sup>

The same way, we also expected changes in the patient's treatments. Around half of patients changed their topic or systemic treatment (molecule or dose). We could not demonstrate neither that all of these changes improved patients outcomes and prognosis, but we would like to highlight that we increased the percentage of topic and systemic drugs, probably optimizing the

previous ones. Only the rate of patients on corticosteroids slightly increased after the multidisciplinary evaluation. The same results have been described in other multidisciplinary care units.<sup>14–16</sup> Another positive point of these multidisciplinary care units is the sharing decision making between two specialists with different but complementary visions. This way, discrepancies between specialists that can led to medical errors or compliance problems, for example, could be avoided.<sup>17,18</sup>

On the other hand, we would like to point out that apart from labs and X-ray, just a few of patients required MRI or biopsies probably reflecting that clinical evaluation made by the experts in psoriasis and PsA (who follow national and international guidelines) solved most of the referred problems.

Regarding to comorbidity, we have exposed that more than a half of patients evaluated in the unit presented comorbidities. Patients with psoriasis and PsA show a high prevalence of comorbidities and risk factors, among which the most important are cardiovascular disease and metabolic syndrome.<sup>19–21</sup> Other frequent comorbidities are hyperuricemia and gout.<sup>22</sup> We found the same results in our patients. An adequate management of comorbidities is of great importance for the specialists, as comorbidity influences diagnosis, prognosis, and treatment decisions, and has a great impact on health care resources.<sup>23–26</sup> Therefore, in a multidisciplinary care approach, a proper evaluation and management of comorbidities could be facilitated in order to improve outcomes in complex patients.

One of the main outcomes of our study is the great level of satisfaction in our patients. In their opinion, for most of patients this multidisciplinary unit compared with their usual care was better or much better. Interestingly, all of them considered that the disease was better controlled in this multidisciplinary care unit. Data from a case series of patients with psoriasis and PsA followed following a multidisciplinary care model was also very high.<sup>14</sup> This is very important because in chronic diseases the adherence and compliance with treatments and visits is vital and is very influenced with the overall level of satisfaction of patients with the care they receive.<sup>27</sup>

Following, this study has some limitations. First, we have no control group. Ideally our results should have been compared with traditional care. However, the rate of diagnosis confirmation and treatment changes suggest that, to some extent, we are making earlier diagnosis of PsA and probably optimizing patient's treatments. On the other hand, we did not used a generic or specific questionnaire to assess patient's satisfaction and this was retrospectively evaluated. But taking into account the very high level of satisfaction reported we are confident that this multidisciplinary care unit really meant a good option for patients with psoriasis or PsA.

In summary, collaboration between expert rheumatologists and dermatologists in a multidisciplinary care unit facilitates a proper and overall evaluation of the skin and musculoskeletal burden, subsequently leading to an earlier diagnosis of PsA, comorbidities and risk factors and to a more comprehensive treatment approach. As a consequence the disease outcomes could improve as well as patient's satisfaction.

## Ethical Disclosures

**Protection of human and animal subjects.** The authors state that for this investigation the procedures followed conformed to the ethical standards of the responsible human experimentation committee (Hospital Can Misses, Ibiza), and in agreement with the World Medical Association and the Declaration of Helsinki.

**Confidentiality of data.** The authors declare that the confidentiality of the data has been guaranteed.

**Right to privacy and informed consent.** The authors state that the patient's informed consent was requested.

### Funding

This manuscript was developed by an unrestricted grant from Pfizer.

### Conflict of Interest

Dr. Loza reports grants from Pfizer, during the conduct of the study; grants from Pfizer, grants from Abbvie, grants from Roche, grants from MSD, grants from BMS, grants from Novartis, grants from UCB, outside the submitted work. The rest of authors refer no conflicts of interest.

### References

- Moll JM, Wright V. Psoriatic arthritis. *Semin Arthritis Rheum*. 1973;3:55–78.
- Cats A. Psoriasis and arthritis. *Cutis*. 1990;46:323–9.
- Torre Alonso JC, Rodriguez Perez A, Arribas Castrillo JM, Ballina Garcia J, Riestra Noriega JL, Lopez Larrea C. Psoriatic arthritis (PA): a clinical, immunological and radiological study of 180 patients. *Br J Rheumatol*. 1991;30:245–50.
- Haddad A, Li S, Thavaneswaran A, Cook RJ, Chandran V, Gladman DD. The incidence and predictors of infection in psoriasis and psoriatic arthritis: results from longitudinal observational cohorts. *J Rheumatol*. 2016;43:362–6.
- Husni ME. Comorbidities in Psoriatic Arthritis. *Rheum Dis Clin N Am*. 2015;41:677–98.
- Ogdie A, Schwartzman S, Husni ME. Recognizing and managing comorbidities in psoriatic arthritis. *Curr Opin Rheumatol*. 2015;27:118–26.
- Rosen CF, Mussani F, Chandran V, Eder L, Thavaneswaran A, Gladman DD. Patients with psoriatic arthritis have worse quality of life than those with psoriasis alone. *Rheumatology (Oxford)*. 2012;51:571–6.
- Truong B, Rich-Garg N, Ehst BD, Deodhar AA, Ku JH, Vakil-Gilani K, et al. Demographics, clinical disease characteristics, and quality of life in a large cohort of psoriasis patients with and without psoriatic arthritis. *Clin Cosmet Invest Dermatol*. 2015;8:563–9.
- Buder K, Wozel G. Psoriatic arthritis: a diagnostic challenge? *G Ital Dermatol Venereol*. 2010;145:407–14.
- Haroon M, Gallagher P, FitzGerald O. Diagnostic delay of more than 6 months contributes to poor radiographic and functional outcome in psoriatic arthritis. *Ann Rheum Dis*. 2015;74:1045–50.
- Ng CY, Yang YW, Liu SH, Lu JF, Yang LC, Yang CH, et al. SF-36 healthy survey on psoriasis quality-of-life: a study of 414 Taiwanese patients. *J Rheumatol*. 2015;42:159–65.
- Valenzuela F, Silva P, Valdes MP, Papp K. Epidemiology and quality of life of patients with psoriasis in Chile. *Actas Dermosifiliogr*. 2011;102:810–6.
- Gossec L, Smolen JS, Ramiro S, de Wit M, Cutolo M, Dougados M, et al. European League Against Rheumatism (EULAR) recommendations for the management of psoriatic arthritis with pharmacological therapies: 2015 update. *Ann Rheum Dis*. 2016;75:499–510.
- Cobo-Ibanez T, Villaverde V, Seoane-Mato D, Munoz-Fernandez S, Guerra M, Del Campo PD, et al. Multidisciplinary dermatology-rheumatology management for patients with moderate-to-severe psoriasis and psoriatic arthritis: a systematic review. *Rheumatol Int*. 2016;36:221–9.
- Luelmo J, Gratacos J, Moreno Martinez-Losa M, Ribera M, Romani J, Calvet J, et al. A report of 4 years of experience of a multidisciplinary unit of psoriasis and psoriatic arthritis. *Reumatol Clin*. 2014;10:141–6.
- Velez NF, Wei-Passanese EX, Husni ME, Mody EA, Qureshi AA. Management of psoriasis and psoriatic arthritis in a combined dermatology and rheumatology clinic. *Arch Dermatol Res*. 2012;304:7–13.
- Geurts MM, van der Flier M, de Vries-Bots AM, Brink-van der Wal TI, de Gier JJ. Medication reconciliation to solve discrepancies in discharge documents after discharge from the hospital. *Int J Clin Pharm*. 2013;35:600–7.
- Heiner MM. Key barriers to optimal management of adult asthma in Australia: physician and patient perspectives. *Curr Med Res Opin*. 2007;23:1799–807.
- Husted JA, Thavaneswaran A, Chandran V, Eder L, Rosen CF, Cook RJ, et al. Cardiovascular and other comorbidities in patients with psoriatic arthritis: a comparison with patients with psoriasis. *Arthritis Care Res (Hoboken)*. 2011;63:1729–35.
- Jamnitski A, Symmons D, Peters MJ, Sattar N, McInnes I, Nurmohamed MT. Cardiovascular comorbidities in patients with psoriatic arthritis: a systematic review. *Ann Rheum Dis*. 2013;72:211–6.
- Khraishi M, MacDonald D, Rampakakis E, Vaillancourt J, Sampalis JS. Prevalence of patient-reported comorbidities in early and established psoriatic arthritis cohorts. *Clin Rheumatol*. 2011;30:877–85.
- Oliviero F, Scanu A, Galozzi P, Gava A, Frallonardo P, Ramonda R, et al. Prevalence of calcium pyrophosphate and monosodium urate crystals in synovial fluid of patients with previously diagnosed joint diseases. *Joint Bone Spine*. 2013;80:287–90.
- Palomo L, Rubio C, Gervas J. The comorbidity in primary care. *Gac Sanit*. 2006;20 Suppl. 1:182–91.
- Long JA, Husted JA, Gladman DD, Farewell VT. The relationship between patient satisfaction with health and clinical measures of function and disease status in patients with psoriatic arthritis. *J Rheumatol*. 2000;27:958–66.
- Lee S, Mendelsohn A, Sarnes E. The burden of psoriatic arthritis: a literature review from a global health systems perspective. *Pharmacy Ther*. 2010;35:680–9.
- Husni ME, Meyer KH, Cohen DS, Mody E, Qureshi AA. The PASE questionnaire: pilot-testing a psoriatic arthritis screening and evaluation tool. *J Am Acad Dermatol*. 2007;57:581–7.
- Kumar K, Raza K, Nightingale P, Horne R, Chapman S, Greenfield S, et al. Determinants of adherence to disease modifying anti-rheumatic drugs in White British and South Asian patients with rheumatoid arthritis: a cross sectional study. *BMC Musculoskelet Disord*. 2015;16:396.