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Incidence and Predictive Factors for Orthopedic Surgery in Patients with Psoriatic Arthritis

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ABSTRACT. Objective. To investigate the incidence of orthopedic procedures in patients with psoriatic arthritis (PsA), and how patient characteristics, time of diagnosis, and treatment affect the need for surgery. Methods. We reviewed the medical history of 1432 patients with possible PsA at Haukeland University Hospital in Bergen, Norway. There were 590 patients (mean age 49 yrs, 52% women) who had sufficient journal information and a confirmed diagnosis of PsA, and who were included in the present study. Relevant orthopedic procedures were obtained from the hospital’s administrative patient records. Survival analyses were completed to evaluate the effect of different factors such as year of diagnosis, age, sex, radiographic changes, disease activity, and treatment, on the risk of surgery. Results. There were 171 procedures (25% synovectomies, 15% arthrodesis, and 53% prostheses) performed on 117 patients. These factors all increased the risk of surgery: female sex [relative risk (RR) 1.9, p = 0.001], age ≥ 70 years at diagnosis (RR 2.4, p = 0.001), arthritis in initial radiographs (RR 2.2, p = 0.006), and maximum erythrocyte sedimentation rate 30-59 mm/h (RR 1.6, p = 0.026). Time period of diagnosis had no effect on the outcome. In a subanalysis of surgery exclusive of hip and knee arthroplasty, diagnosis in earlier years (1954–1985 vs 1999–2011) was a risk factor (RR 2.1, p = 0.042). Antirheumatic treatment changed significantly over time. Conclusion. There were 20% of patients with PsA who needed surgery. We found that the prognosis of patients with PsA did not change regarding the risk of orthopedic surgery, despite the change in treatment. A possible explanation is the increase in large joint replacements in the general population. (J Rheumatol First Release September 1 2018; doi:10.3899/jrheum.180203)

Key Indexing Terms: PSORIATIC ARTHRITIS ORTHOPEDIC PROCEDURE PROGNOSIS EPIDEMIOLOGY

Psoriatic arthritis (PsA) develops in up to 30% of patients with psoriasis, and the prevalence worldwide has been estimated between 0.02% and 0.25%. A study of the population of Hordaland county in western Norway, which is the population we address in this study, found a prevalence of 0.2%, and polyarthritis was the most frequent subclass, documented in 68.6%.

PsA with peripheral joint involvement is a progressive disease in most patients, and erosions are seen in 47% within the first 2 years. Individuals with polyarthritis are the most susceptible to bone erosions and deformities. Previous studies have been conflicting regarding whether PsA is as destructive radiologically as rheumatoid arthritis (RA).

Spontaneous remission of PsA is rare, and if nonsteroidal antiinflammatory drugs (NSAID) and intra-articular steroids are not sufficient, a synthetic disease-modifying antirheumatic drug (DMARD) is often prescribed to ease pain and restore function. No synthetic DMARD has, however, been proven to slow or prevent radiographic changes. Biologic treatment is recommended when other agents are not sufficient, and it has been shown to give better control of structural damage.

Orthopedic surgery has been a necessary part of treating patients with PsA when medication fails to prevent inflammation and joint destruction. Surgery can thus be considered a proxy for the degree of inflammatory activity, and studying time trends in orthopedic surgery gives valuable information regarding the prognosis of patients with inflammatory arthritis. In previous studies, there is a large discrepancy in the incidence of orthopedic surgery in patients with PsA. A study of 444 patients published in 1998 found that 7% had a musculoskeletal procedure performed. In a more recent study of 269 patients published in 2016, 48.3% received orthopedic surgery. This total, however, also included arthroscopies, meniscus surgery, carpal tunnel surgery, surgery of the spine, and others (205 of 280 procedures). We have not found any data on changes over time. In patients...
with RA there has been a gradual declining incidence of orthopedic interventions. The change in available medical treatment is believed responsible for this. Because synthetic DMARD may be less efficient in patients with PsA, it is uncertain whether a decline of the same magnitude can be expected among these patients. It is also possible that a change, if present, would occur later, after the introduction of tumor necrosis factor (TNF)-α inhibitors in this patient group.

We investigated the occurrence of orthopedic surgery among patients diagnosed with PsA in western Norway, and whether patient characteristics, treatment, and year of diagnosis affected the need for surgical intervention.

**MATERIALS AND METHODS**

Our data originate from Haukeland University Hospital, which delivers specialist care to about 500,000 inhabitants in western Norway. Because only 2 private practicing rheumatologists operate in the area, the vast majority of patients with rheumatic disease are cared for by the hospital’s department of rheumatology. In general, patients are referred to specialist care at the time of suspected inflammatory rheumatic disease. Some patients with stable disease are later managed by their general practitioner, but most continue to be followed until death or inactive disease.

From the hospital’s administrative patient records, we have data available from 1972 until the present, and some paper files go back even further. Using the International Classification of Diseases (ICD) 8th, 9th, and 10th revisions, we searched for disease codes for PsA (ICD-10 L40.5, ICD-9 7133), and the combination of arthritis or spondylarthropathy (SpA; ICD-10 M06, M46; ICD-8 712, 715; ICD-9 714 and psoriasis (ICD-10 L40, ICD-9 712, ICD-9 696). Because orthopedic surgery is a late outcome, we wished to give all patients the possibility of at least 5 years observation after the date of diagnosis and excluded patients with first hospital contact later than 2011. The number of patients with first contact in the pre-methotrexate (MTX) era (before 1985) and in the era after the introduction of MTX, but before the use of TNF-α inhibitors (1986–1998), was much lower than the number of patients with first contact from 1999 to 2011. To ensure a sufficient number of patients from the first 2 periods, we chose to evaluate all of these, to find possible subjects for inclusion. From the third period, we evaluated patients with at least 2 hospital contacts, to increase the possibility that cases evaluated had a confirmed diagnosis of PsA and could be included in the study. However, the inclusion criteria were the same for all evaluated patients. We also searched the local patient system GoTreatIt and the National Arthritis Registry (NorArthritis; helse-bergen.no/seksjon-engelsk/sekjon-avdeling/Silder/Norwegian-Arthritis-Registry-NorArthritis.aspx) for patients registered under the diagnosis psoriatic arthritis.

Using these sources, 2251 patients remained for evaluation. All of these had contacts with Haukeland University Hospital from 1972 to 2011 coded in accordance with our search, but some had no contact with the department of rheumatology prior to 2012, and others did not consent to participation. There were 819 patients excluded because of this, leaving 1432 patients available for journal review. Of these, 951 had given consent to participate and 481 had died. Following this process, we were able to include 590 patients. The selection process is described in Figure 1. The search performed was very broad, to account for the possibility that in previous years, patients with PsA were not identified and coded that way, but rather as other arthritis conditions. This proved not to be true, and the vast majority of patients included had been identified and coded as PsA.

Each medical record was reviewed for the following information: weight, height, year of diagnosis, number of joints with arthritis (evaluated by the treating rheumatologist) during disease course, whether in fulfillment of the CASPAR criteria for PsA, and the presence of psoriatic dermatitis, and erythrocyte sedimentation rate (ESR) at diagnosis and maximal levels. The medical records did not contain radiographic images, so only the radiologist’s interpretation of these as normal or consistent with arthritis or osteoarthritis (OA) was recorded, and for images of the columnar and sacroiliac joints (SIJ), whether there were signs of SpA. Medications used in the first year and during the course of the disease were also registered. Supplementary data were obtained from the Norwegian Arthritis Registry for patients herein. The study was approved by the Regional Committee for Medical and Health Research Ethics (2016/2207/REK West).

Because MTX was introduced to our patient group around 1986, and biologic treatment in the form of TNF-α inhibitors in 1999, we split the group into 3 depending on diagnosis in different treatment eras: time period 1 included patients diagnosed prior to 1986, time period 2 from 1986 to 1998, and time period 3 from 1999 to 2011. Patient characteristics for each group and in total are described in Table 1. Mean body mass index (BMI) was significantly lower in time period 1 versus time period 3 (p = 0.033).

Statistical analyses were performed in SPSS versions 23 and 24. The level for statistical significance was set to p < 0.05.
Figure 1. Selection process.
RESULTS

Treatment, presented as percentage of patients given each medication in the first year and during the course of the disease is presented in Figure 2. The proportion of patients using NSAID was high in all 3 time periods, but significantly higher in time period 1, both in the first year and during the course of the disease. Within 2 years of disease duration, 37% of all patients had been prescribed MTX. There were 56% who were prescribed MTX during the course of the disease. The proportion of patients receiving MTX within the first year since diagnosis increased from 17% in time period 2 to 43% in time period 3. When looking only at patients in time period 3, 53% were prescribed MTX, and 19% biologics within 2 years of disease onset. The effects of time since diagnosis increased from 17% in time period 2 to 43% in time period 3. When looking only at patients in time period 3, 53% were prescribed MTX, and 19% biologics within 2 years of disease onset. Further details are described in Table 1.

The department of dermatology seemed to start using MTX earlier than the department of rheumatology in our county, and the 3 patients in time period 1 using MTX in the first year of disease had all been prescribed this from their dermatologist.

There was no significant difference in the prescription of effective medication in patients aged ≥ 70 years versus younger patients.

Surgical procedures. There were 1.4 events per 100 patient-years over the whole study period. When excluding surgery conducted earlier than 1 year prior to diagnosis, we found 171 procedures performed in 117 patients (20%). The procedures performed within 1 year prior to diagnosis were counted as performed at diagnosis, because we had reason to assume an association between surgery and diagnosis. A single procedure was performed on 68%, and 24% who had 2 procedures performed, whereas 9.5% underwent 3–6 surgeries. Of the types of procedures, 53% of the 171 procedures were prosthetic, 25% were synovectomies, and 15% arthrodesis procedures (triple arthrodeses and forefoot procedures included). The distribution of different procedures in diverse joints is described in Table 2. The most frequently operated area was the knee, in which 38% of procedures were performed, followed by the hip (27%). Eight percent of the 590 patients had knee surgery performed, 6.4% who had hip surgery, and 6.1% who had surgery of the hands or feet. Mean time to first procedure was 8.5 years (0–52 ± SD 8.5 yrs). Mean age at surgery was 62 years. Patients were youngest at time of synovectomy (mean age 45 yrs), older at first arthrodesis procedure (mean age 55 yrs), and oldest at arthroplasty surgery (mean age 70 yrs).

Survival analyses. Results are presented in Table 3 and Figure 3. Factors found to affect the risk of a surgical procedure during the course of the disease were older age at diagnosis, female sex, arthritis on initial radiographs, and highest ESR between 30 and 59 mm/h. Time period of diagnosis did not influence the risk. No significant effects were found involving fulfillment of the CASPAR criteria, biological treatment within 2 years of disease onset, MTX treatment within 2 years of disease onset, ESR at disease onset, ≥ 5 joints affected, or BMI ≥ 30 kg/m². OA in initial radiographs was borderline significant, but not significant when analyzing 100 files with imputed values [relative risk (RR) 1.4, 95% CI 0.72–2.64, p = 0.33]. For the other exposure variables, analysis of the files with imputed values did not change the significance of the above-described results.

When performing subanalyses of procedures exclusive of...
Figure 2. Medical treatment in the first year (A) and during the course of the disease (B) for patients diagnosed in 3 different time periods. COX-2: cyclooxygenase 2 inhibitors; NSAID: nonsteroidal antiinflammatory drugs.
DISCUSSION

Our study’s main findings were that 20% of patients with PsA required orthopedic surgery, and that time period of diagnosis did not affect the outcome.

Observational studies have disadvantages, compared to randomized controlled trials (RCT), but describe to a greater extent the prognosis of patients in real life instead of patients treated under ideal conditions. In this study we observed the patients for a mean time of 13.8 (0–63) years, which would be impossible in an RCT because it is less suitable for the investigation of late outcomes.

Female sex, older age, elevated ESR, and arthritis in initial radiographs increased the risk of orthopedic surgery. Others have also found that female sex predicts an unfavorable outcome23,24, whereas in a different study, male sex was a risk factor for early radiographic damage25. For orthopedic surgery exclusive of hip and knee prostheses, patients diagnosed in time period 1 (1954–1985) had an increased risk of surgery compared to those with a diagnosis in time period 3 (1999–2011).

We divided patients into 3 time periods based on the availability of medical treatment at diagnosis. A limitation to this approach is that the change in treatment came gradually, and that new treatment was also made available to patients diagnosed in previous years, although later in the disease course.

Among patients diagnosed in the 3 time periods, we did not find any significant changes in the disease activity as measured by inflammatory variables or radiographic changes.
at disease onset. On the other hand, maximum ESR during the disease course decreased significantly from 43 to 30, from time period 1 to time period 3, and the share with ESR ≥ 60 mm/h during the course of the disease fell from 30% in patients diagnosed before 1986 to 14% in patients diagnosed 1999–2011 (p < 0.001). Significantly more patients among those diagnosed in earlier years also developed radiographic changes. This suggests that the burden of inflammation has become lower in recent years. In the articles by Zangger, et al and Haque, et al, survival analyses were not performed, but patients were divided into 2 groups according to whether they had undergone orthopedic surgery, and the groups were then compared. Zangger, et al found that patients who had an operation had significantly more radiological damage and more actively inflamed joints at first assessment. Haque, et al found that asymmetric mono-/oligoarticular arthritis and the combination of peripheral and axial disease were more frequent among patients with surgeries, whereas there were no differences in treatment between the groups.

There has been a significant change in the prescription of medication. The share of patients receiving potent medication within 2 years of disease onset more than doubled when comparing patients diagnosed 1999–2011 to patients diagnosed 1986–1998.

When investigating surgery among patients with inflammatory arthritis in our region, we found that 31% of patients with RA underwent an orthopedic procedure, whereas for patients with PsA, 20% had joint surgery performed. For RA, diagnosis in earlier years was a significant risk factor for orthopedic surgery. In PsA, however, this did not affect the outcome.

When analyzing only hip and knee arthroplasty, the time period of diagnosis was not a significant risk factor for surgery in patients with RA. This supports the findings from other studies, that while hand and foot surgery in RA has declined, large joint replacements remain unchanged. A possible explanation for this might be the general increase seen in joint replacement surgery in later years, but it has also been discussed whether the inflammation process is different in small joints compared to larger joints, and that the latter could be less affected by antirheumatic therapy. Whereas arthroplasty in joints other than hip and knee were found to be frequent in RA (18% of prosthesis procedures), this was seldom performed in patients with PsA, where 96% of prosthesis surgery were hip and knee procedures. Because large joint replacements account for a greater proportion of surgery in patients with PsA than those with RA (51% vs 33%), this would be expected to weaken the effect of time period of diagnosis. After excluding prosthesis surgery of the hip and knee from the analysis, we find that diagnosis in 1954–1985 increased the risk of surgery compared to diagnosis later than 1998.
A possible confounding factor is that hip and knee arthroplasties may have been conducted on the basis of coexisting OA in patients with PsA. The etiology of joint destruction and the subsequent need for joint replacement surgery in patients with inflammatory arthritis may be hard to detect. This is especially the case for the hip joint, particularly before the use of ultrasound. We found that 57% of patients with knee prosthesis surgery had had arthritis of the knee during disease course, and that 21% of patients with hip prosthesis surgery had had hip joint arthritides, as detected by the treating rheumatologist. One must thus suspect that inflammatory disease was a contributing factor also for large-joint destruction.

Although not to the same extent as for RA, patients with PsA have, to an increasing degree, been treated with synthetic, and in later years, biologic DMARDs. In our study, this change coincides with patients having a lower burden of inflammation, as measured by ESR. Others have found that early referral to an arthritis clinic and tight disease control give a better outcome. However, it has been shown that clinical signs of inflammation and progression of joint destruction might be dissociated, and we could not find a decrease in joint surgery, suggesting that this outcome has not been affected by the change in medication, contrary to what is found for patients with RA. This is in concordance with the knowledge that contrary to the effect of synthetic DMARD on structural damage in patients with RA, the same has not been shown for PsA.

Because biologic treatment has been shown to prevent joint destruction in PsA, it is possible that increasing use of TNF-α inhibitors would lessen the risk of an orthopedic procedure during the disease course. When considering all procedures, this was not the case in our material. Among the 25% of patients prescribed biologic treatment during the course of the disease, only 42% started treatment within 2 years of diagnosis. It might thus be that damage, predicting later orthopedic surgery, had already occurred when treatment was initiated.

We found that 20% of patients with PsA underwent orthopedic surgery. The maximum disease activity seems to have decreased over time, but the risk of surgery was not affected by year of diagnosis. The unchanged prognosis regarding joint surgery in patients diagnosed from 1985 to 1998, compared to those with disease onset in the pre-MTX era, might indicate that MTX and other synthetic DMARD do not prevent joint damage. However, because biologic treatment has been shown to prevent joint damage, one would expect the risk of orthopedic surgery to decline following its introduction, in patients diagnosed from 1999 onward. It is possible that the improved treatment is too recent for a change in prognosis to have become evident. But one must also consider whether the use of biologic treatment might be too infrequent and initiated too late if we seek optimal treatment of patients with PsA.

**ONLINE SUPPLEMENT**

Supplementary material accompanies the online version of this article.

**REFERENCES**