Spinal Radiographic Changes in Ankylosing Spondylitis: Association with Clinical Characteristics and Functional Outcome

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ABSTRACT. Objective. To determine which patients with ankylosing spondylitis (AS) have radiographic spinal damage and to investigate the relation between radiographic spinal changes and limitations in physical function.

Methods. A cross-sectional nationwide study in Belgium of patients with AS under the care of a rheumatologist. The treating physician completed a questionnaire including clinical disease manifestations and laboratory findings (HLA-B27 and C-reactive protein), and classified spinal radiographs into 3 categories: (1) no AS-related spinal abnormalities; (2) syndesmophytes; and (3) spinal ankylosis. Patients completed the Bath AS Disease Activity Index (BASDAI) and the Bath AS Functional Index (BASFI). Ordinal regressions were performed to quantify the relationship between clinical manifestations and spinal radiographic changes. Generalized linear models were computed to quantify relationships among clinical manifestations, radiographic spinal changes, and functioning (BASFI).

Results. A total of 619 patients fulfilled modified New York criteria for definite AS and had evaluable radiographic data; 68% were male and disease duration was 17.5 (SD 12.2) years. Male sex, younger age at symptom onset, and hip involvement were associated with radiographic changes; but HLA-B27, peripheral arthritis, and extraarticular disease status (uveitis, psoriasis, and inflammatory bowel disease) were not. Older age, BASDAI, hip involvement, and spinal change contributed to BASFI; but sex, disease duration, peripheral arthritis, and extraarticular manifestations did not.

Conclusion. Radiographic spinal changes in patients with AS are seen more often in men and those with hip involvement. BASFI status indicates the influence of radiographic changes and hip involvement, but does not reflect the presence of peripheral arthritis and does not differ between men and women. (First Release May 15 2009; J Rheumatol 2009;36:1249–55; doi:10.3899/jrheum.080831)

Key Indexing Terms:
ANKYLOSING SPONDYLITIS  GENDER  RADIOGRAPHY  HEALTH STATUS

Ankylosing spondylitis (AS) is considered prototypical for the group of chronic rheumatological conditions referred to as the spondyloarthropathies. The hallmark of the disease is inflammation of the sacroiliac joints resulting in progressive structural changes that are eventually evaluable by conventional radiography1,2. Apart from sacroiliitis, AS is clinically heterogeneous, which adds to the challenge in understanding the pathophysiology and assessing all aspects of its clinical outcome. Sacroiliitis, spondylitis, peripheral arthritis, enthesitis, and involvement of shoulders or hips (known as root joints) contribute to the articular burden of AS. A number of extraarticular manifestations, including uveitis, psoriasis, and inflammatory bowel disease (IBD) can complicate the clinical presentation of the disease3.

Spinal syndesmophyte formation and ankylosis are the most striking features of AS and can be seen on conventional radiography. Only a few studies have investigated the relation between the clinical characteristics and radiographic spinal changes4-6. The association of uveitis with spinal change has been investigated in 3 studies, but the results are conflicting7-9. Brophy, et al found no association of IBD or psoriasis with spinal changes5. To our knowledge, the relation between the articular manifestations and ankylosis has never been addressed. In addition to the disease-related clin-
ical manifestations, intrinsic patient characteristics (e.g., sex) can be associated with the presence of radiographic changes. In this regard, all but one of 13 studies showed that women had fewer radiographic changes in lumbar and thoracic spine, independent of the scoring method applied.

There are few studies addressing the influence of radiographic spinal changes on patient outcomes. Taylor, et al. found that ankylosis correlated with more pain, stiffness and sleep disturbance; and Lee, et al. and Doran, et al. reported that radiographic status was associated with function in univariate and multivariate analyses. Landewé, et al. reported in a longitudinal analysis over 2 years that disease activity and radiographic damage assessed by the modified Stoke Ankylosing Spondylitis Spine Score (mSASSS) were independent contributors of change in function after adjusting for age and sex.

The aim of our study was to assess (1) the relationship between the broad range of clinical characteristics of patients with AS and radiographic spinal damage; and (2) the influence of spinal changes, in addition to the clinical characteristics, on functional outcome.

MATERIALS AND METHODS
Rheumatologists and patients. Between February 2004 and February 2005, all 204 Belgian rheumatologists were invited to register their patients with AS for an epidemiologic study. A random subsample of patients was included in a cross-sectional clinical study. Those fulfilling the modified New York criteria for definite AS and having a spinal radiographic examination within 2 years were considered for analyses.

The study was approved by the national license bureau for noninterventional research, and all patients provided signed informed consent prior to inclusion. A detailed description of the sampling and epidemiology of rheumatologists and patients is described elsewhere.

Clinical characteristics and functional outcome. A questionnaire was completed by the treating rheumatologist based on the patients’ records and included the items of the modified New York criteria, age at symptom onset, the presence of HLA-B27 antigen, the last available value of C-reactive protein (CRP), and the history of past or present peripheral arthritis, enthesis and extraarticular disease. Extraarticular disease manifestations included anterior uveitis, psoriasis, ulcerative colitis, and Crohn’s disease [combined and referred to as irritable bowel disease (IBD)]. To assess disease activity patients completed the Bath AS Disease Activity Index (BASDAI) and to assess limitations in mobility, they underwent a clinical examination by the rheumatologist, who recorded the components of the Bath AS Metrology Index (BASMI). Physical function was assessed through the self-reported Bath AS Functional Index (BASFI).

Radiographic changes. Consistent with the naturalistic design of the study, information on radiographic abnormalities was based on the latest available documents if not performed at the time of the study visit and were scored by the treating rheumatologist. The sacroiliac joints were scored according to the modified New York criteria. Spinal radiographic changes had to be classified into 3 (exclusive) categories: (1) no spinal abnormalities related to AS; (2) presence of syndesmophytes; or (3) presence of spinal ankylosis.

Statistical analyses. Descriptive statistics were used to describe the data by the calculation of means with standard deviation and range for continuous data and frequencies for dichotomous and ordinal data. Differences and frequencies between the 3 subgroups with different spinal changes were tested by computing odds ratios and 95% confidence intervals. Associated p values were calculated by chi-square statistics for dichotomous data and Kruskal-Wallis tests for continuous data.

To explore the effect of one or more variables on radiographic changes, a logit link function (ordinal logistic regression) was used, first correcting for the potential confounders age at onset, disease duration since symptom onset, and sex. Variables were HLA-B27, elevated CRP, extraspinal manifestations (ever hip arthritis, arthritis, or enthesitis) and extraarticular manifestations (uveitis, psoriasis, and IBD). Interactions between the sets of variables were explored in additional analyses that included for the main effect variables and the interaction term for each interaction studied. The exponents of the estimates of those ordinal logistic regression analyses can be interpreted as odds ratios for each level of radiographic change.

RESULTS
Rheumatologists and patient characteristics. Eighty-nine of 204 invited rheumatologists (44%) agreed to participate and represented all the geographical regions of Belgium as well as a mix of academic and nonacademic practices. A total of 1023 patients were included in the study. All patients fulfilled the modified New York criteria, of whom 847 (83%) met the criteria for definite AS and 176 (17%) the criteria for probable AS. Six hundred nineteen (72%) patients with definite AS had radiographs available from the last 2 years. Sixty-seven percent were male, with a mean age of 43.9 (± 12.1) years, and mean disease duration since symptom onset was 17.5 (± 12.2) years. In addition, 83% were HLA-B27-positive.

Radiographic spinal change was recorded in 294 of 619 patients (47%). One hundred twenty-four (20%) had spinal ankylosis, 170 (27%) syndesmophytes, and 325 (53%) no spinal abnormalities related to AS. A cumulative history of peripheral arthritis was recorded in 351 (56.8%) patients, enthesitis in 215 (47.7%), and hip involvement in 166 (26.6%). A hip arthroplasty had been performed in 36 (5.8%) patients. Twenty-seven percent of patients had spinal changes as well as peripheral arthritis and 155 (25%) had neither radiographic change nor arthritis. A cumulative history of extraarticular manifestations was registered for uveitis in 169 patients (27.3%), psoriasis in 68 (11.0%), and IBD in 54 (8.7%). Forty percent of patients had at least one extraarticular manifestation.

Missing data. Patients with a missing radiographic evaluation were older, 46.6 versus 43.9 years (difference 2.7 yrs; 95% CI 0.7–0.4), had longer symptom duration, 13.6 versus...
11.4 years (difference 2.2 yrs; 95% CI –0.15 to 3.8), and had slightly better physical functioning on BASFI, 4.8 versus 5.2 (difference 0.4; 95% CI 0.08–0.73), but did not differ with respect to sex, peripheral arthritis or hip involvement, or HLA-B27 status. From 619 patients with radiographic evaluations, HLA-B27 data were available in 78% and complete BASMI in 58%, the remaining 42% having incomplete BASMI measurements. Patients with missing HLA-B27 were older, 46.9 compared to 42.4 years (difference 4.5 yrs; 95% CI 2.6–6.5), and patients with missing BASMI had higher BASDAI, 5.8 versus 5.1 (difference 0.7; 95% CI 0.4–1.0), and BASFI, 5.7 versus 4.9 (difference 0.8; 95% CI 0.4–1.2).

**Characteristics of patients according to radiographic subgroup.** Table 1 describes the differences in patients’ characteristics in terms of their radiological involvement (no AS-related spinal abnormalities, syndesmophytes, or spinal ankylosis). No differences in age at onset of axial symptoms or BASDAI or elevated CRP could be found. In contrast, more radiographic spinal changes were significantly associated with longer disease duration and worse BASFI and BASMI scores. Twenty-seven percent of patients with symptom duration < 10 years showed AS-related radiographic spinal changes (syndesmophytes or ankylosis). This proportion increased to 77% in patients with more than 30 years’ symptom duration.

**Factors associated with radiographic spinal changes.** The ordinal regression models first confirmed that age at onset, symptom duration, and sex were confounders (Table 2; first section). A significant interaction between disease duration and sex was found, indicating that male patients not only had more frequent spinal changes but were also more likely to have an increase in spinal changes when disease duration increased. Figure 1 shows the proportion of men and women with spinal ankylosis for cohorts with disease duration < 10 years, 10–30 years, and ≥ 30 years.

**Variables associated with BASFI.** General linear regression analyses showed that age, BASDAI, radiographic changes, and hip involvement, but not sex, disease duration, peripheral arthritis, enthesitis, uveitis, psoriasis, or IBD, contributed to physical function measured by BASFI (Table 3). Further analysis of interaction terms showed significant interaction between symptom duration and hip involvement, indicating that patients with hip involvement were more likely to have an increase in spinal changes when disease duration increased.

**Factors associated with BASMI.** General linear regression analyses showed that age, BASDAI, radiographic changes, and hip involvement, but not sex, disease duration, peripheral arthritis, enthesitis, uveitis, psoriasis, or IBD, contributed to physical function measured by BASMI (Table 3). First, the initial model (described above) was confirmed as the estimates for beta were comparable, indicating the MCAR assumption could be considered valid. Adding the BASMI

<table>
<thead>
<tr>
<th>Table 1. Characteristics of patients with definite AS (modified New York criteria) and available radiographic examination according to the degree of radiographic change.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>No AS Abnormalities, n = 325</strong></td>
</tr>
<tr>
<td>Age at onset, yrs, mean (SD) [range]</td>
</tr>
<tr>
<td>Symptom duration, yrs, mean (SD) [range]</td>
</tr>
<tr>
<td>Male, %</td>
</tr>
<tr>
<td>HLA-B27 (n = 489), %</td>
</tr>
<tr>
<td>Elevated CRP (n = 584), %</td>
</tr>
<tr>
<td>BASDAI, mean (SD)</td>
</tr>
<tr>
<td>BASFI, mean (SD)</td>
</tr>
<tr>
<td>BASMI (n = 360), mean (SD)</td>
</tr>
</tbody>
</table>

CRP: C-reactive protein; BASDAI: Bath AS Disease Index; BASFI: Bath AS Functional Index; BASMI: Bath AS Metrology Index.

In the second step, the different extraspinal and extraarticular manifestations (hip changes, peripheral arthritis, enthesitis, uveitis, psoriasis, IBD) were added to this model. Hip involvement was the only variable associated with radiographic spinal changes (Table 2; section 2). Further analysis of interaction terms showed significant interaction between symptom duration and hip involvement, indicating that patients with hip involvement were more likely to have an increase in spinal changes when disease duration increased.

**Table 2. Association of demographic, extraspinal, and extraarticular manifestations with radiographic spinal changes.**

<table>
<thead>
<tr>
<th></th>
<th>exp B (OR)</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at onset</td>
<td>0.03</td>
<td>0.02–0.05</td>
</tr>
<tr>
<td>Symptom duration</td>
<td>0.09</td>
<td>0.07–1.01</td>
</tr>
<tr>
<td>Sex</td>
<td>1.67</td>
<td>1.27–2.08</td>
</tr>
<tr>
<td>Estimate when controlling for age at onset, sex, and disease duration</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hip involvement, disease duration</td>
<td>0.61</td>
<td>0.24–0.98</td>
</tr>
<tr>
<td>Peripheral arthritis</td>
<td>0.30</td>
<td>–0.04–0.64</td>
</tr>
<tr>
<td>Enthesitis</td>
<td>0.08</td>
<td>–0.14–0.24</td>
</tr>
<tr>
<td>Uveitis</td>
<td>0.31</td>
<td>–0.06–0.72</td>
</tr>
<tr>
<td>Psoriasis</td>
<td>0.10</td>
<td>–0.43–0.62</td>
</tr>
<tr>
<td>Inflammatory bowel disease</td>
<td>–0.37</td>
<td>–0.96–0.21</td>
</tr>
</tbody>
</table>
to the explanatory variables revealed a strong association between BASMI and BASFI (beta 0.33, 95% CI 0.24–0.41). However, the contribution of radiographic changes disappeared and the strength of the association with hip involvement decreased, indicating BASMI was a mediator for these variables. The model including BASMI (confounders, BASMI, and hip involvement) had a better goodness of fit compared to the model without BASMI (confounders, 

Figure 1. Cumulative prevalence of radiographic syndesmophytes and ankylosis according to disease duration for male (n = 417) and female (n = 202) patients.
Table 3. Association of demographic, clinical, and disease characteristics with physical functioning (Bath AS Functional Index).

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Beta</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intercept</td>
<td>-1.52</td>
<td>-2.1 to -0.9</td>
</tr>
<tr>
<td>Age</td>
<td>0.04</td>
<td>0.02-0.05</td>
</tr>
<tr>
<td>Female sex</td>
<td>0.13</td>
<td>-0.17-0.40</td>
</tr>
<tr>
<td>Symptom duration</td>
<td>-0.003</td>
<td>-0.02-0.11</td>
</tr>
<tr>
<td>BASDAI</td>
<td>0.86</td>
<td>0.92-0.19</td>
</tr>
<tr>
<td>Radiographic spinal change</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ankylosis</td>
<td>1.13</td>
<td>0.74-1.52</td>
</tr>
<tr>
<td>Syndesmophytes</td>
<td>0.42</td>
<td>0.11-0.74</td>
</tr>
<tr>
<td>No abnormalities</td>
<td>Reference</td>
<td></td>
</tr>
<tr>
<td>Hip involvement</td>
<td>0.64</td>
<td>0.35-0.92</td>
</tr>
</tbody>
</table>

DISCUSSION

We evaluated characteristics of patients with AS who had radiographic spinal changes. There were no radiographic spinal changes in 53% of the total group; however, in the subgroup with disease duration longer than 30 years, only 23% remained without spinal changes. This suggests that a large majority of patients under care of a rheumatologist have spinal involvement that will eventually lead to changes visible on conventional radiographs. The likelihood of spinal ankylosis was also higher in men, in those with symptom onset at an earlier age, and in patients who had a history of hip involvement. There was no association of HLA-B27 or extraspinal articular disease or extraarticular manifestations with spinal ankylosis.

While it does not seem surprising that disease duration is associated with radiographic changes, there are few prospective longterm studies with data that quantify this observation. The Outcome Assessments in Ankylosing Spondylitis International Study evaluated a cohort of patients under the care of a rheumatologist, and in radiographs scored using the mSASSS (score range 0–72)17, it was shown that over a period of 4 years the average change in score was low, but on the patient level and depending on the criterion, between 37% and 57% of patients had an increase in the spinal scoring system18. Dawes19 reported an average increase of 23 points in mSASSS over 9 years in a group of patients with at least 2 radiographic assessments available. As in our study, Dawes also noted that earlier onset of symptoms was associated with more AS-related radiographic changes.

The increased prevalence and severity of radiographic spinal changes in men and in those with hip involvement confirms previous reports and adds to the evidence4,7,9,20,27. The explanation for the gender difference remains unclear. Although there is no linkage between X-chromosome markers and susceptibility for AS28, it cannot be excluded that severity of AS is X-link determined. Polymorphisms of genes that control tissue calcification and arthritis have been found to be associated with AS, and their relationship with sex is under investigation29,31. The previously observed association between spinal involvement and hip arthritis resulted in the classification of hips as “root” or “axial” joints. No pathophysiological model investigated whether the disease processes in the spine are similar to that in the hips. This remains an interesting area for future research.

In our study radiographic spinal change was a relevant contributor to physical function, together with age and disease activity (BASDAI) and hip involvement. Interestingly, gender had no influence on function, and also did not influence the univariate analyses, despite the higher frequency of radiographic changes among men. In the subgroup of patients with BASMI data, the effect of radiographic change on BASFI disappeared and the influence of hip involvement decreased when taking spinal and hip mobility (BASMI) into account, and improved the goodness of fit of the model. This suggests that an important part of the effect of the radiographic changes on the BASFI can be explained by its effect on the BASMI. The contribution of radiographic changes over time to the BASFI or the Dougados Functional Index was confirmed by Landewé, et al in a study that applied mSASSS to score the extent of radiographic changes11. Similarly to our study, the BASDAI also was a determinant of physical function. Of note, the relation between radiographic changes was shown not only for the syndesmophyte score, but also for the non-syndesmophyte score (squaring of sclerosis erosions). The independent contribution of BASMI data was not explored in this study. Maksymowycz, et al have described strong univariate associations between spinal mobility (using the Edmonton AS Metrology Index; EDASMI) and function (BASFI) or radiographic changes (mSASSS)32. Again, the combined contribution of spinal and radiographic change to function was not investigated. It should be emphasized that although BASMI relates to radiographic damage on the group level, this does not necessarily imply that radiographic staging on the individual level can be replaced by the BASMI33. It was surprising in our study that enthesitis or peripheral arthritis had a low impact on function. It is possible that BASFI is not sensitive for the functional influence of peripheral joint involvement, since this instrument primarily focuses on spinal functioning. Also, physician-reported enthesitis might not be accurate. This might explain why the Spondyloarthritis Research Consortium of Canada enthesitis index showed a significant correlation with BASFI in the validation study34.

Some limitations of our study should be noted. First, the data were collected cross-sectionally and therefore do not allow the conclusion that variables associated with spinal changes are “prognostic” factors. Also, left censorship may have occurred, referring to the phenomenon that patients with less severe disease refrained from further control or had...
no radiographic assessment and therefore the group with longer disease is a selection of patients with worse disease. It is unlikely that this influences the observed relation with sex, since in the cohorts with different disease duration, no differences in prevalence of both sexes were seen. Second, these results pertain to patients with AS under care of a rheumatologist, which will certainly bias toward more severe disease, potentially explaining the large prevalence of ankylosis and extraspinal articular manifestations, especially peripheral arthritis. On the other hand, the study population is relevant, since it represents the population where more intensive treatment will be considered. Nevertheless, due to this source of sampling and the left censorship, the absolute figures on prevalence of radiographic spinal changes should be interpreted with caution. Finally, the design of the study did not allow standardizing the instruments to assess articular and extraspinal disease and radiographic changes. It is difficult to hypothesize whether physician-reported enthesisitis, arthritis, or hip involvement will result in under- or over-reporting. As noted, the absence of a standardized method to report enthesisitis could explain why our study did not find an association with function. When interpreting the relevance of hip involvement, an important factor to explain spinal changes as well as function, the issue of standardization is pertinent. It should be emphasized, however, that similar strong relationships remained when replacing hip involvement with hip arthroplasty. The classification of radiographic changes into the 3 groups by physicians could have led to overlap between the groups with “syndesmophytes” and those with “ankylosis,” since the threshold at which consecutive bridging syndesmophytes are considered “spinal ankylosis” is likely different for each rheumatologist. However, the data suggest that the group classified as “spinal ankylosis” show more extensive radiographic effects than the group classified as “syndesmophytes.” The terminology “radiographic spinal changes” was chosen carefully. Patients can have spinal involvement (pain and stiffness) without changes seen on conventional radiology. There is increasing insight that the structural spinal changes in AS reflect the pathophysiologic remodeling as a consequence of either inflammation or other stress on the joint and therefore are “reparative” and not just “destructive.” A timeframe of 2 years was allowed to include radiographs. It is unlikely this influenced our results since it has been shown that on the group level the radiographic progression over 2 years in an unselected cohort was low, although clear changes were seen at the patient level after 2 years.

The major strength of our study is the large number of patients with AS that were recorded and assessed. This allowed sufficient power to study the relationships of disease manifestations that are present in only a minority of patients.

Almost 47% of patients with AS under care of a rheumatologist had radiographic spinal changes. Patient’s sex and hip involvement, but not uveitis, IBD, peripheral arthritis or psoriasis, were associated with spinal changes. Radiological change contributed to worse physical functioning, but on the group level the presence of radiographic changes seemed to be interchangeable with assessment of spinal mobility assessed by BASMI.

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