The Impact of Illness Perceptions and Coping on the Association Between Back Pain and Health Outcomes in Patients Suspected of Having Axial Spondyloarthritis: Data From the SPonddyloArthritis Caught Early Cohort

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Objective. To investigate whether illness perceptions and coping influence the relationship between back pain and health outcomes in patients suspected of having axial spondyloarthritis (SpA).

Methods. In the SPonddyloArthritis Caught Early cohort, regression models were computed at baseline, with back pain intensity (range 0–10) as the determinant and health-related quality of life, the physical component summary score (PCS) and mental component summary (MCS) of the Short Form 36 (SF-36) health survey, or work productivity loss as outcomes. Subsequently, using Leventhal’s Common-Sense Model of Self-Regulation, illness perceptions and, thereafter, coping were added to the models. Analyses were repeated for patients diagnosed and classified as having axial SpA according to the Assessment of SpondyloArthritis international Society axial SpA criteria (ASAS axial SpA), patients only diagnosed with axial SpA (axial SpA–diagnosed only), and those with chronic back pain.

Results. A total of 424 patients (145 with ASAS axial SpA, 81 with only a diagnosis of axial SpA, and 198 with chronic back pain); 64% of the total group were female, the mean ± SD age was 30.9 ± 8.1 years, and the mean ± SD symptom duration was 13.3 ± 7.1 months) were studied. In all patients, the strength of the associations between back pain and the PCS, back pain and the MCS score, and back pain and loss of work productivity were decreased by adding illness perceptions to the model, but explained variance improved. Adding coping to these models did not change the results. Comparable results were observed in all subgroups.

Conclusion. Illness perception, but not coping, is important in the relationship between back pain and HRQoL and work productivity loss in patients suspected of having axial SpA, irrespective of subgroup. This finding suggests that targeting illness perceptions could improve health outcomes in patients suspected of having axial SpA.

INTRODUCTION

The disease burden in patients with axial spondyloarthritis (SpA) is significant. Treatment aimed at reducing the burden of disease consists of a combination of pharmacologic treatment, education, and exercise (1). Leventhal’s Common-Sense Model of Self-Regulation (“Common-Sense Model” [CSM]) (2) has been shown to be helpful for understanding patients’ responses to various rheumatic diseases and diseases related to axial SpA such as psoriasis or inflammatory bowel disease (3–5). However, the CSM has not yet been studied in patients with axial SpA or in patients with chronic back pain who are suspected of having axial SpA (6).

The CSM is a theoretical framework used to describe and understand a patient’s responses to an illness and its characteristics (e.g., swollen joints) (2) (see Figure 1). According to the CSM, patients perceive an illness and its
characteristics as a health threat and respond to this threat by generating illness perceptions. According to this model, illness perceptions directly influence coping strategies, which in turn influence health outcomes such as health-related quality of life (HRQoL) and loss of work productivity, in order to reestablish a patient’s normal health state. Illness perceptions are ideas formulated by patients that help them make sense of their illness, such as perceived personal control over the disease or the experienced negative emotions that they attribute to the disease. In contrast, coping strategies are cognitive and behavioral strategies used to manage stress associated with having to live with the illness (e.g., actively diverting attention from the illness or adapting the level of physical activity).

Relatively little is known about illness perceptions in patients with axial SpA, especially those with early axial SpA. In a study by Hyphantis and colleagues in patients with longstanding ankylosing spondylitis (AS), “illness concern” (i.e., more concerns about the disease) was found to be associated with worse physical HRQoL (7). Different results concerning illness perceptions in patients with chronic back pain have been reported. Most studies showed that patients with chronic back pain strongly believe in “severe consequences” (e.g., held strong beliefs in “severe consequences,” strong beliefs that the disease is “chronic,” and have “negative emotions” toward their disease (8–10).

Two studies in patients with AS showed that avoidant coping styles “decreasing activities” and “pacing” were associated with more pain and worse physical and mental functioning (11,12). These 2 coping strategies were also strongly related to withdrawal from the workforce (13). In patients with chronic back pain, maladaptive coping strategies such as “avoiding physical activity” were associated with negative health outcomes such as increased pain and disability (14). However, knowledge about illness perceptions and coping in patients with early axial SpA is lacking. Furthermore, little is known about how both illness perceptions and coping impact health outcomes in (early) axial SpA.

Exploring use of the CSM in patients with early axial SpA and chronic back pain is important, because it may enable health care professionals to identify illness perceptions and coping strategies that are susceptible to additional treatment strategies aimed at decreasing the burden of disease in these patients. In the current study, we first investigated the association between back pain and HRQoL or loss of work productivity, and subsequently, we investigated the influence of illness perceptions and coping on these associations in patients suspected of having axial SpA and in subgroups. We used the CSM as the theoretical model. We hypothesized that having severe back pain is associated with lower HRQoL and greater loss of work productivity, and that the strengths of these associations are amplified by negative illness perceptions and maladaptive coping strategies in patients suspected of having axial SpA. We further hypothesized that the relationship between illness perceptions and coping strategies differs across subgroups.

Back pain is a self-reported and subjective symptom that is prevalent among patients with axial SpA. Therefore, we thought it would be interesting to additionally investigate the previously mentioned associations, using an objective sign that is typical for axial SpA. We considered inflammation on magnetic resonance imaging of the sacroiliac joints (MRI-SI) as measured by the Spondyloarthritis Research Consortium of Canada score for the SI joints (SPARCC-SI) to be a good candidate for being the objective sign. We hypothesized that illness perceptions and coping strategies have little influence on these

![Figure 1. Flow diagram representing Leventhal’s Common-Sense Model of Self-Regulation. Adapted, with permission, from ref. 38.](image-url)
associations, because a patient is unaware of his or her SPARCC-SI score. Consequently, all analyses were also performed using the SPARCC-SI score instead of back pain as the independent variable in a group of patients who were diagnosed and classified as having axial SpA.

**PATIENTS AND METHODS**

Baseline data from the SPondyloArthritis Caught Early (SPACE) cohort of patients who were included between January 2009 and February 2017 were used. Briefly, the SPACE cohort is a prospective inception cohort of patients with chronic back pain (≥3 months but ≤2 years, and onset before age 45 years) (15). Dutch, Norwegian, and Italian rheumatology outpatient clinics participated in the SPACE study. Approval by local medical ethics committees (Medical Ethics Committee, Leiden University Medical Center [approval no. P08.105]; regional committee for medical and health research ethics in South-East Norway [approval no./ID 2014/426]; and Azienda Ospedaliera di Padova [approval no. 2438P]) was obtained. Informed consent was obtained from all study participants prior to inclusion.

All patients underwent the same diagnostic evaluation at baseline, consisting of medical history, physical examination, questionnaires, laboratory assessments (i.e., HLA-B27), and imaging including plain radiographs of the pelvis and coronal oblique MRI-SI (1.5T, 4-mm slice thickness). Patients were unaware of their diagnosis until the full assessment was performed. Treating rheumatologists provided the diagnosis, using clinical findings and local readings of the images. Patients in whom axial SpA was diagnosed were classified according to the Assessment of SpondyloArthritis international Society (ASAS) axial SpA criteria (16), based on central reading of images.

Analyses were performed in all patients as well as the following subgroups: patients diagnosed with axial SpA and classified according to the ASAS axial SpA criteria (ASAS axial SpA), patients diagnosed with axial SpA only (axial SpA–diagnosed only), and patients diagnosed with chronic back pain.

Back pain intensity was assessed by asking patients to report the extent of back pain in the past 7 days on a numeric rating scale (NRS) ranging from 0 (no pain) to 10 (unbearable pain). Inflammation suspected of being axSpA on MRI-SI was quantified by 3 central readers according to the SPARCC-SI scoring method, and the average continuous SPARCC-SI score from 3 readers was calculated. Four quadrants were scored for each SI joint, and additional scores were given to lesions characterized by depth or intensity, resulting in a total score ranging from 0 to 72 (17).

Illness perceptions were assessed with the Revised Illness Perception Questionnaire (IPQ-R), which consists of 3 sections (18,19). The first section is the illness identity dimension, in which patients are asked about their experience with particular symptoms (15 items) and the perceived relationship with back pain. The numbers of symptoms with a perceived relationship is summed. The second section of the IPQ-R consists of 7 dimensions: “consequences” (perceived impact of the disease on the patient’s life), “acute/chronic timeline” (perceived likeliness of chronicity of the disease), “personal control” (perceived personal control over the disease), “treatment control” (perceived efficacy of treatment), “illness coherence” (extent to which patients feel they understand their disease), “cyclical timeline” (the patient’s perceptions of variability of her or her disease), and “emotional representation” (the patient experienced negative emotions due to the disease). The third section (causal attributions) consists of 18 possible causes that patients may attribute to their disease. Five dimensions were calculated: “psychological attributions,” “risk factors,” “immunity,” “accident,” and “chance.” The subscales of the second and third sections used Likert scales to score all items (1 = strongly disagree and 5 = strongly agree). Higher scores indicate stronger beliefs in that dimension (second section) or stronger beliefs in a dimension being a cause of the disease (third section) (19).

Coping strategies were assessed with the Coping with Rheumatic Stressors (CORS) questionnaire. The questionnaire is aimed at dealing with the most important stressors in rheumatic diseases: pain, limitations, and dependence (20,21). “Comforting cognitions” (putting pain in perspective), “decreasing activities,” and “diverting attention” (thinking about/focusing on something else) refer to coping with pain. Coping with limitations is measured by “optimism,” “pacing” (adapting/lowering the level of activity), and “creative solution seeking” (searching for creative solutions to cope with the limitations in daily life). The 2 styles of coping with dependence are “accepting” (making efforts to accept the level of dependence) and “showing consideration” (considering the feelings of others). Higher scores indicate preferential use of a particular coping strategy. The mean scores for each subscale of both the IPQ-R and CORS questionnaires were calculated.

Work productivity was assessed by the Work Productivity and Activity Impairment questionnaire (WPAI: general health, version 1.0). Patients were asked to report, e.g., the number of work hours missed due to their disease, the number of hours that they actually worked, and the impact of their disease on work productivity, scored on an NRS from 0 (health problems had no effect on work) to 10 (health problems completely prevented working) in the past week. The summary measure work productivity loss (i.e., total work impairment due to chronic back pain) on a scale from 0% (no work productivity loss) to 100% (total work productivity loss) was calculated. Greater impairment is indicated by higher percentages (22).

HRQoL was assessed with the Short Form 36 (SF-36) health survey (23), which consists of 8 subscales. After recoding and recalibration were performed, raw scale scores were transformed into scale scores ranging from 0 (worst health) to 100 (best health). These scores were weighted according to sex- and age-matched scores for patients in each country (24,25). Dutch-weighted scores were used for all Italian patients (n = 57; [13%]), because no Italian sex- and age-matched scores were available. Two summary scores, the physical component summary score (PCS) and the mental component summary score (MCS), were calculated and transformed to compare the scores with the general population mean of 50. Higher scores indicate better HRQoL (26).
Categorical variables are presented as the number (frequency) and continuous variables as the mean ± SD. Back pain was used in models as an independent variable in analyses of all patients and subgroup analyses, while the SPARCC-SI score was used only in models that included patients with ASAS axial SpA. Pearson’s correlation coefficients were calculated to determine differences between back pain or SPARCC-SI score, illness perceptions, coping, and outcome measures (PCS, MCS, or work productivity loss) were calculated. All variables that had a significant correlation ($P < 0.05$) with the dependent variables (PCS, MCS, or work productivity loss) were included in a multistep linear regression model for that outcome. Models were adjusted for age and sex, by default. Illness perceptions were added to the basic model with back pain intensity or SPARCC-SI score as an outcome in the first step, and coping strategies were added in the second step. Likelihood ratio tests were used to determine whether the addition of each step independently improved the model. Data analyses were performed using Stata SE version 14.

RESULTS

Baseline data were available for 550 patients included in the SPACE cohort. Patients were excluded from further analyses when a complete questionnaire ($n = 39$) or scales of the questionnaires ($n = 87$) were missing. For the current analysis, 424 patients were used. Compared with all patients included in the analyses, patients who were excluded from the analyses less often had a diagnosis of axial SpA (axial SpA diagnosis in 53% of included patients and 39% of excluded patients; $P = 0.012$) and fewer clinical SpA features (2.6 features in included patients and 2.0 features in excluded patients; $P = 0.002$).

In total, 145 of 424 patients were categorized as having ASAS axial SpA (diagnosed by the rheumatologist and classified as axial SpA), 81 of 424 were categorized as having a diagnosis of axial SpA only (diagnosed by a rheumatologist as axial SpA only), and 198 of 424 were categorized as having chronic back pain. The mean ± SD age of all patients was 30.9 ± 8.1 years, and the mean ± SD symptom duration was 13.3 ± 7.1 months; these values were comparable with those in the different subgroups (Table 1). The

| Table 1. Baseline characteristics of and assessment results for 424 patients with chronic back pain in the SPACE cohort, according to subgroups* |
|-----------------|-----------------|-----------------|-----------------|
| **Baseline characteristics** |
| | All patients (n = 424) | ASAS axial SpA (n = 145) | Axial SpA diagnosis only (n = 81) | Chronic back pain (n = 198) |
| Age, mean ± SD years | 30.9 ± 8.1 | 30.1 ± 7.8 | 32.5 ± 7.8 | 30.8 ± 8.5 |
| Female sex, no. (%) | 272 (64) | 73 (50) | 53 (65) | 146 (74) |
| Symptom duration, mean ± SD months | 13.3 ± 7.1 | 13.7 ± 7.2 | 12.2 ± 6.3 | 13.4 ± 7.4 |
| Inflammatory back pain | 295 (70) | 120 (83) | 64 (79) | 111 (56) |
| Good response to NSAIDs† | 190 (45) | 87 (60) | 46 (57) | 57 (29) |
| Uveitis | 36 (9) | 26 (18) | 4 (5) | 6 (3) |
| Psoriasis | 51 (12) | 22 (15) | 18 (22) | 11 (6) |
| Inflammatory bowel disease | 32 (8) | 7 (5) | 13 (16) | 12 (6) |
| Enthesitis | 188 (44) | 76 (52) | 33 (41) | 79 (40) |
| Dactylitis | 91 (22) | 34 (24) | 43 (53) | 14 (7) |
| Peripheral arthritis | 28 (7) | 13 (9) | 11 (14) | 4 (2) |
| Elevated ESR/CRP level | 69 (16) | 28 (19) | 24 (30) | 17 (9) |
| X-SI-positive | 178 (42) | 130 (90) | 5 (6) | 43 (22) |
| MRI-SI-positive | 177 (42) | 61 (42) | 25 (32) | 31 (16) |
| Use of NSAIDs | 32 (8) | 28 (19) | 0 (0) | 4 (2) |
| No. of SpA features, mean ± SD‡ | 281 (66) | 112 (77) | 54 (67) | 115 (58) |
| Assessment results | | | | |
| Back pain, mean ± SD (0–10 scale) | 4.8 ± 2.4 | 4.4 ± 2.3 | 4.4 ± 2.6 | 5.4 ± 2.3 |
| SPARCC-SI score, mean ± SD (range 0–72) | 1.8 ± 4.9 | 4.8 ± 7.4 | 0.4 ± 1.7 | 0.1 ± 0.6 |
| PCS, mean ± SD (range 0–100) | 26.9 ± 14.8 | 28.2 ± 15.0 | 29.1 ± 13.8 | 25.1 ± 14.9 |
| MCS, mean ± SD (range 0–100) | 47.2 ± 12.7 | 48.2 ± 13.9 | 44.8 ± 12.2 | 47.5 ± 12.0 |
| WPL, mean ± SD % (range 0–100) | 42.5 ± 32.1 | 37.8 ± 31.5 | 35.1 ± 30.4 | 49.5 ± 32.0** |

* Values are the number (%) unless specified otherwise. SPACE = SpondyloArthritis Caught Early; ASAS = Assessment of SpondyloArthritis international Society; SpA = spondyloarthritis; ESR = erythrocyte sedimentation rate; CRP = C-reactive protein; X-SI = radiography of the sacroiliac (SI) joints; MRI-SI = magnetic resonance imaging of the SI joints; SPARCC-SI = Spondyloarthritis Research Consortium of Canada score for the SI joints; MCS = mental component summary score; PCS = physical component summary score; WPL = work productivity loss.

† Back pain no longer present or is much better 24–48 hours after administration of a full dose of a nonsteroidal antiinflammatory drug (NSAID).

‡ Only 329 patients were evaluated.

§ Only 110 patients were evaluated.

¶ Only 65 patients were evaluated.

** Only 144 patients were evaluated.
majority of patients were female (50% of patients with ASAS axial SpA, 65% of patients with axial SpA–diagnosed only, and 74% of patients with chronic back pain). Patients with ASAS axial SpA and those with only axial SpA diagnosed had more SpA features (excluding HLA–B27 and imaging) (mean 3.3 and mean 3.5 features, respectively) compared with patients with chronic back pain (mean 1.7 features). In patients with chronic back pain, back pain was more severe than that in the ASAS axial SpA and the axial SpA–diagnosed–only group (Table 1). A greater percentage of patients in the ASAS axial SpA group had inflammation on MRI-SI compared with patients with chronic back pain (mean 1.7 features). In patients with chronic back pain, back pain was more severe than that in the ASAS axial SpA and the axial SpA–diagnosed–only group (Table 1). A greater percentage of patients in the ASAS axial SpA group had inflammation on MRI-SI compared with the percentage in the axial SpA–diagnosed only group and the chronic back pain group. The mean PCS was decreased in all groups compared with the general population (mean of 50), but the mean MSC was comparable with that in the general population patients, but the mean MCS was comparable with that in the general population (Table 1). Seventy–seven percent (n = 326) of all patients were in the work force at baseline. In 7 patients, loss of work productivity could not be calculated. Work productivity loss was comparable in the ASAS axial SpA group and the axial SpA–diagnosed only group but was higher in patients with chronic back pain (see Table 1). Statistically significant differences in mean scores were observed for the illness perceptions “personal control” (mean scores 3.3 in the axial SpA group, 3.2 in the axial SpA–diagnosed only group, and 3.0 in the group with chronic back pain), “treatment control” (mean scores 3.5 in the ASAS axial SpA group, 3.3 in the axial SpA–diagnosed

Table 2. Multiple-step linear regression model with back pain, illness perception, and coping explaining HRQoL and work productivity loss among all patients (n = 424)*

|                      | PCS |       |       |       |       |       |
|----------------------|-----|-------|-------|-------|-------|-------|
|                      | Range | B   | 95% CI | B     | 95% CI | B     | 95% CI |
| Model 1 (basic model) |       |     |       |       |       |       |
| Back pain            | 0–10 | −3.5 | −3.9, −3.0† | −0.9 | −1.4, −0.4‡ | 7.7 | 6.5, 8.9† |
| Age (years)          | 0.4  | 0.3, 0.6† | −0.1 | −0.3, 0.04 | −0.1 | −0.5, 0.3 |
| Female sex           | 2.7  | 0.3, 5.1† | 0.7 | −1.9, 3.3 | 5.7 | −0.3, 11.6 |
| Model 2 (basic model plus illness perceptions) |       |     |       |       |       |       |
| Back pain            | 0–10 | −2.7 | −3.2, −2.2† | −0.1 | −0.5, 0.4 | 6.3 | 5.0, 7.7† |
| Age (years)          | 0.5  | 0.4, 0.6† | −0.1 | −0.2, 0.1 | −0.2 | −0.5, 0.2 |
| Female sex           | 3.5  | 1.2, 5.8§ | 1.7 | −0.5, 4.0 | 4.7 | −1.3, 10.7 |
| Identity             | 0–15 | −0.3 | −0.8, 0.1 | −0.6 | −1.1, −0.2§ | 0.3 | −0.9, 1.5 |
| Consequences         | 1–5  | −6.9 | −8.6, −5.1† | −0.2 | −1.8, 1.5 | 8.6 | 3.9, 13.2† |
| Timeline (acute/chronic) | 1–5 | 0.5 | −1.1, 2.2 |     |       |     |       |
| Personal control     | 1–5  | 0.9 | −1.1, 2.8 | 0.8 | −1.0, 2.6 | −2.3 | −7.2, 6.2 |
| Treatment control    | 1–5  | 0.7 | −1.7, 3.1 |     |       |     |       |
| Illness coherence    | 1–5  | 0.03 | −1.4, 1.5 | 0.1 | −1.3, 1.4 | −2.4 | −6.2, 1.4 |
| Emotional representation | 1–5 | 2.4 | 0.8, 4.1§ | −5.0 | −6.6, −3.4† | −0.2 | −4.6, 4.2 |
| Psychological attributions | 1–5 | 1.4 | −0.3, 3.1 | −5.4 | −7.2, −3.7† | −0.4 | −4.3, 3.4 |
| Risk factors         | 1–5  |     |       | 2.0 | −0.5, 4.5 |     |       |
| Immunity             | 1–5  | −1.3 | −3.3, 0.4 | −1.0 | −2.0, 0.7 |     |       |
| Accident             | 1–5  | −0.9 | −2.0, 0.3 | 0.5 | −0.7, 1.7 |     |       |
| Chance               | 1–5  | −1.2 | −2.1, −0.2‡ |     |       |     |       |
| Model 3 (basic model plus illness perceptions and coping) |       |     |       |       |       |       |
| Back pain            | 0–10 | −2.3 | −2.8, −1.9† | −0.1 | −0.5, 0.4 | 5.9 | 4.7, 7.2† |
| Age (years)          | 0.5  | 0.4, 0.6† | −0.1 | −0.2, 0.1 | −0.2 | −0.6, 0.2 |
| Female sex           | 3.7  | 1.6, 5.8§ | 1.1 | −1.1, 3.3 | 4.3 | −1.4, 10.0 |
| Identity             | 0–15 |     |       | −0.6 | 1.0, −0.2§ |     |       |
| Consequences         | 1–5  | −4.7 | −6.4, −3.1† |     |       | 6.4 | 2.2, 10.6§ |
| Emotional representation | 1–5 | 2.2 | 0.8, 3.6§ | −5.1 | −6.5, −3.7† |     |       |
| Psychological attributions | 1–5 | −2.0 |       | −4.4 | −5.8, −3.1† |     |       |
| Chance               | 1–5  | −1.0 | −1.9, −0.1‡ |     |       |     |       |
| Comforting cognitions | 1–4 |     |       | 2.1 | −1.0, 5.2 |     |       |
| Decreasing activities | 1–4 | −4.0 | −6.7, −1.3§ | −0.6 | −2.7, 1.4 | 7.3 | −0.2, 14.8 |
| Diverting attention  | 1–4  |     |       | −0.7 | −3.2, 1.9 |     |       |
| Optimism             | 1–4  | 1.7 | −0.2, 3.6 | 0.3 | −2.1, 2.8 |     |       |
| Pacing               | 1–4  | −3.3 | −6.3, −0.3‡ |     |       | 5.1 | −3.2, 13.5 |
| Creative solution-seeking | 1–4 | −1.0 | −3.4, 1.3 |     |       | −0.6 | −7.1, 5.9 |
| Accepting            | 1–4  | −0.8 | −2.6, 1.1 |     |       | 0.1 | −5.2, 5.3 |
| Consideration        | 1–4  | −1.6 | −3.8, 0.7 |     |       | 1.9 | −3.9, 7.7 |

* Work productivity loss (WPL) was assessed in only 319 patients. P values indicate a statistically significant association with the outcome. HRQoL = health-related quality of life; PCS = physical component summary score; MCS = mental component summary score; 95% CI = 95% confidence interval.
† P < 0.001.
‡ P < 0.05.
§ P < 0.01.
only group, and 3.3 in the group diagnosed as having chronic back pain), “illness coherence” (mean scores 3.3 in the ASAS axial SpA group, 3.1 in the axial SpA–diagnosed only group, and 2.8 in the group with chronic back pain), “emotional representation” (mean scores 2.7 in the ASAS axial SpA group, 2.9 in the axial SpA–diagnosed only group, and 2.9 in the group with chronic back pain), and “psychological attributions” (mean scores 1.9 in the ASAS axial SpA group, 2.2 in the axial SpA–diagnosed only group, and 2.1 in the group with chronic back pain). No differences in other illness perceptions or coping strategies between the 3 subgroups were observed (data not shown).

**Analysis in all patients.** In the basic model including all patients, the PCS decreased by 3.5 points (R² = 0.37, P < 0.001), the MCS decreased by 0.9 points (R² = 0.03, P < 0.001), and work productivity loss increased by 7.7% (R² = 0.36, P < 0.001) per point increase in the severity of back pain (Tables 2 and 3). After adding illness perceptions to the model, the impact of a 1-point increase in back pain on the PCS and work productivity loss became smaller and resulted in a decrease of 2.7 points (P < 0.001) and an increase of 6.3% (P < 0.001), respectively, and the association between back pain and the MCS was no longer statistically significant (B = –0.1, P = 0.838) (Tables 2 and 3). The model performance improved by adding illness perceptions: more variance was explained in the PCS (R² = 0.47), the MCS (R² = 0.32), and work productivity loss (R² = 0.40) compared with the basic model, and these differences were statistically significant. After further adding coping strategies to the model, the associations between back pain and the PCS or work productivity loss changed only slightly (–2.3 points [P < 0.001] and 5.9% [P < 0.001], respectively, per point increase in back pain) while the association with the MCS score remained the same (–0.01 points per point back pain; P = 0.762) compared with the model with illness perceptions only (Tables 2 and 3). Explained variance did not further improve statistically significantly (PCS, R² = 0.53; MCS, R² = 0.32; work productivity loss, R² = 0.42) by adding coping strategies.

In the third model, having stronger beliefs in severe consequences (illness perception “consequences”; B = –4.7) or chance as a cause for the disease (illness perception “chance”; B = –1.0), and more use of the coping strategies “decreasing activities” (B = –4.0) and “pacing” (B = –3.3) were statistically significantly associated with a lower PCS. The illness perception “emotional representation” (having more negative emotions toward the disease) was associated with a better PCS (B = 2.2). Attributing more symptoms to the disease (illness perception “identity”; B = –0.6), having more negative emotions toward the disease (B = –5.1), and having stronger beliefs in psychological attributions as a cause (illness perception “psychological attributions”; B = –4.4) were statistically significant associated with a lower MCS. Having stronger beliefs in severe consequences was statistically significantly associated with more work productivity loss (B = 6.4).

**Subgroup analyses.** Similar results were observed in the ASAS axial SpA group (Tables 3 and 4), the axial SpA–diagnosed only group (Tables 3 and 5), and the chronic back pain group (Tables 3 and 6) separately. The negative association between back pain and the MCS was observed only in the basic model. The strength of the associations between back pain and the PCS or work productivity loss decreased only slightly.

| Table 3. Adjusted R² and –2 log likelihood ratios of the multiple-step linear regression models for each group of patients* |
|-----------------------------------------------|---------------|--------------|--------------|
| | PCS | | | | MCS | | | | | | WPL | | | |
| | Adjusted | –2 log likelihood | Adjusted R² | –2 log likelihood | Adjusted R² | –2 log likelihood |
| All patients | | | | | | | | | | | | | |
| Basic model | 0.37 | –1,645.4 | 0.03 | –1,672.7 | 0.36 | –1,486.3 |
| Basic model + illness perceptions | 0.47 | –1,601.3‡ | 0.32 | –1,593.1‡ | 0.40 | –1,473.4 |
| Basic model + illness perceptions and coping | 0.53 | –1,578.8 | 0.32 | –1,593.3 | 0.42 | –1,467.0 |
| ASAS axial SpA | | | | | | | | | | | | | |
| Basic model | 0.28 | –573.5 | 0.04 | –581.7 | 0.33 | –511.8 |
| Basic model + illness perceptions | 0.42 | –553.9‡ | 0.36 | –548.3‡ | 0.40 | –503.3‡ |
| Basic model + illness perceptions and coping | 0.45 | –552.7 | 0.38 | –547.0 | 0.43 | –499.9 |
| Axial SpA–diagnosed only | | | | | | | | | | | | | |
| Basic model | 0.37 | –306.6 | 0.02 | –314.5 | 0.29 | –300.8 |
| Basic model + illness perceptions | 0.48 | –296.6‡ | 0.26 | –299.9‡ | 0.35 | –297.0‡ |
| Basic model + illness perceptions and coping | 0.49 | –295.7 | 0.25 | –303.3‡ | 0.36 | –297.1‡ |
| Chronic back pain | | | | | | | | | | | | | |
| Basic model | 0.42 | –760.8 | 0.03 | –768.6 | 0.37 | –668.6 |
| Basic model + illness perceptions | 0.48 | –747.0‡ | 0.29 | –731.5‡ | 0.39 | –664.3‡ |
| Basic model + illness perceptions and coping | 0.50 | –722.1‡ | 0.29 | –734.1 | 0.47 | –654.3‡ |

* HRQoL = health-related quality of life; PCS = physical component summary score; MCS = mental component summary score; WPL = work productivity loss; ASAS = Assessment of SpondyloArthritis international Society; SpA = spondyloarthritis.
‡ Statistically significant (P < 0.05) for the model compared with previous model.
† If no coping dimension could be added to model 2 (basic model + illness perceptions), all nonsignificant illness perceptions were removed from model 3 (basic model + illness perceptions and coping).
Table 4. Multiple-step linear regression model with back pain, illness perceptions, and coping explaining variance in HRQoL and work productivity loss among patients with ASAS axial SpA (n = 145)*

|                     | Range | PCS     |         |     |         |     |                 | MCV | B 95% CI |         |     |         |     |                 |
|---------------------|-------|---------|---------|-----|---------|-----|-----------------|-----|---------|---------|-----|---------|-----|-----------------|
|                     |       | B       |         | B   | 95% CI  | B   | 95% CI          | B   | 95% CI  |
| **Model 1 (basic model)**                   |       |         |         |     |         |     |                 |     |         |         |     |         |     |                 |
| Back pain           | 0–10  | −3.4    | −4.3, −2.4† | −1.0 | −2.0, −0.02‡ | 7.7 | 5.5, 9.9†       |     |         |         |     |         |     |                 |
| Age (years)         | 0.4   | 0.1    | 0.7§    | −0.2 | −0.5, 0.1 | 0.3 | −0.4, 0.9       |     |         |         |     |         |     |                 |
| Female sex          | 3.3   | −1.0   | 7.6     | 4.2  | −0.3, 8.7 | 5.6 | −4.3, 15.5      |     |         |         |     |         |     |                 |
| **Model 2 (basic model plus illness perceptions)**     |       |         |         |     |         |     |                 |     |         |         |     |         |     |                 |
| Back pain           | 0–10  | −2.3    | −3.2, −1.4† | −0.3 | −1.2, 0.6 | 5.7 | 3.3, 8.0†       |     |         |         |     |         |     |                 |
| Age (years)         | 0.6   | 0.3    | 0.8‡    | −0.1 | −0.4, 0.1 | 0.03 | −0.6, 0.7       |     |         |         |     |         |     |                 |
| Female sex          | 3.5   | −0.5   | 7.6     | 2.9  | −1.0, 6.8 | 2.5  | −7.6, 12.6      |     |         |         |     |         |     |                 |
| Identity            | 0–15  | −0.7    | −1.5, 0.1 | −0.4 | −1.2, 0.4 | 1.7  | −0.2, 3.6       |     |         |         |     |         |     |                 |
| Consequences        | 1–5   | −8.4    | 11.9, −4.9† | 1.6  | −1.8, 4.9 | 9.3  | 0.8, 17.8‡      |     |         |         |     |         |     |                 |
| Timeline (acute/chronic) | 1–5 | −0.1    | −2.8, 2.7 |     | −       | 0.8  | −5.8, 7.5       |     |         |         |     |         |     |                 |
| Personal control    | 1–5   | −       | −       | −    | −       | −2.9 | −10.4, 4.6      |     |         |         |     |         |     |                 |
| Illness coherence   | 1–5   | −       | −       | −    | −0.5   | −3.0 | −2.0           |     |         |         |     |         |     |                 |
| Emotional representation | 1–5 | 1.6    | −1.3, 4.5 | −6.8 | −10.0, −3.7† | 3.5  | −4.0, 11.0      |     |         |         |     |         |     |                 |
| Psychological attributions | 1–5 | −       | −       | −    | −6.6   | −7.6 | −11.1, −7.8†   |     |         |         |     |         |     |                 |
| Risk factors        | 1–5   | −       | −       | −    | 2.9    | −1.7 | −7.5           |     |         |         |     |         |     |                 |
| Immunity            | 1–5   | 0.7     | −2.1, 3.5 | −1.0 | −4.2, 2.3 | −     | −              |     |         |         |     |         |     |                 |
| Accident            | 1–5   | −2.0    | −4.0, 0.1 | −    | −       | −     | −              |     |         |         |     |         |     |                 |
| **Model 3 (basic model plus illness perceptions and coping)** |       |         |         |     |         |     |                 |     |         |         |     |         |     |                 |
| Back pain           | 0–10  | −2.1    | −3.0, −1.2† | −0.2 | −1.1, 0.7 | 5.5  | 3.2, 7.8†       |     |         |         |     |         |     |                 |
| Age (years)         | 0.6   | 0.3    | 0.8‡    | −0.1 | 0.3, 0.2 | 0.1  | −0.5, 0.8       |     |         |         |     |         |     |                 |
| Female sex          | 2.9   | −1.1   | 6.8     | 2.6  | −1.2, 6.4 | 7.6  | −2.3, 17.1      |     |         |         |     |         |     |                 |
| Consequences        | 1–5   | −7.1    | −10.2, −3.9† | −    | −       | 8.9  | 1.3, 16.4†     |     |         |         |     |         |     |                 |
| Emotional representation | 1–5 | −       | −       | −    | −5.6   | −8.3 | −2.8†          | −     |         |         |     |         |     |                 |
| Psychological attributions | 1–5 | −       | −       | −    | −5.6   | −8.2, −3.1† | −     | −         |         |     |         |     |                 |
| Comforting cognitions | 1–4 | −       | −       | −    | 3.8    | −1.4 | −9.0           | −     |         |         |     |         |     |                 |
| Decreasing activities | 1–4 | −2.7   | −8.4, 3.0 | −2.7 | −6.4, 1.1 | 2.6  | −11.2, 16.4     | −     |         |         |     |         |     |                 |
| Optimism            | 1–4   | −       | −       | −    | −1.3   | −5.6 | 3.0            | −     |         |         |     |         |     |                 |
| Pacing              | 1–4   | −2.8    | −9.0, 3.4 | −    | −       | 13.4 | −2.2, 28.9      | −     |         |         |     |         |     |                 |
| Creative solution-seeking | 1–4 | −0.3   | −4.3, 3.8 | −    | −       | −4.4 | −14.3, 5.5      | −     |         |         |     |         |     |                 |
| Accepting           | 1–4   | −1.8    | −5.2, 1.6 | 2.6  | −0.6, 5.8 | 5.5  | −2.8, 13.7      | −     |         |         |     |         |     |                 |

* Work productivity loss (WPL) was assessed in only 110 patients. HRQoL = health-related quality of life; ASAS = Assessment of SpondyloArthri-
tis international Society; SpA = spondyloarthritis; PCS = physical component summary score; MCS = mental component summary score; 95% CI = 95% confidence interval.
‡ P < 0.001.
§ P < 0.05.
¢ P < 0.01.

Productivity loss decreased after adding illness perceptions to all basic models, although the model performance improved. Results did not change when coping strategies were added to illness perceptions. The same illness perceptions and coping strategies that were associated with PCS, MCS, and work productivity loss in all patients were also associated with these outcomes in each subgroup of patients. Only small differences were found (see Tables 3–6).

SPARCC-SI score in patients with ASAS axial SpA. All analyses were repeated using the SPARCC-SI score instead of back pain in patients with ASAS axial SpA to investigate whether an objective disease measure would yield results similar to those obtained using back pain intensity. In the basic model, the PCS decreased by 0.8 point (P < 0.001), the MCS increased by 0.6 point (P < 0.001), and work productivity loss increased by 0.9% (P = 0.035) per point increase in the SPARCC-SI score (see Supplementary Table 1, available on the Arthritis Care & Research web site at http://onlinelibrary.wiley.com/doi/10.1002/acr.23566/abstract). After illness perceptions and coping strategies were added, the PCS decreased by 0.8 point and 0.7 point, respectively, the MCS increased by 0.5 point and 0.5 point, respectively, and work productivity loss 1.1% and 0.9%, respectively. These results are different from those using models with back pain, because the strength of the associations was not influenced by adding illness perceptions and coping strategies.

DISCUSSION

To our knowledge, this is the first study that used the CSM as a theoretical framework to investigate patients' responses to 1) back pain in patients with chronic back
Therefore, be actively explored and taken into consideration outcomes in these patients. Illness perceptions should, perceptions play an important role in determining medical health care professionals should be aware that illness per-

axial SpA and chronic back pain. Rheumatologists and work productivity, the focus should also be on tar-

In patients with chronic back pain, no associations between illness perceptions and HRQoL or work productivity loss have been investigated, as far as we know. Only one previous study investigated the association between illness

**Table 5. Multiple-step linear regression model with back pain, illness perceptions, and coping explaining variance in HRQoL and work productivity loss among axial SpA–diagnosed only patients (n = 81)**

| Model 1 (basic model) | PCS | MCS | WPL |
|-----------------------|-----|-----|-----|
| Back pain             | B   | 95% CI | B    | 95% CI | B    | 95% CI |
| 0–10                  | -3.2 | -4.2, -2.3† | -0.6 | -1.7, 0.4 | 6.9   | 4.3, 9.6† |
| Age (years)           | 0.2  | -0.1, 0.5 | 0.1  | -0.3, 0.4 | 0.2   | 0.2, -0.7, 1.1 |
| Female sex            | 5.5  | 0.3, 10.7‡ | -4.6 | -10.4, 1.2 | -0.3  | -14.0, 13.4 |

Model 2 (basic model plus illness perceptions)

| Back pain             | B   | 95% CI | B    | 95% CI |
| 0–10                  | -2.6 | -3.5, -1.7† | 0.1  | -0.9, 1.1 | 6.2   | 3.5, 8.8† |
| Age (years)           | 0.3  | 0.03, 0.6‡ | 0.1  | -0.2, 0.4 | 0.1   | -0.6, 1.0 |
| Female sex            | 5.9  | 1.1, 10.6‡ | -4.0 | -9.3, 1.3 | 3.4   | -10.3, 17.1 |
| Identity              | 0–15 | -0.6 | -1.6, 0.5 | -0.8 | -2.0, 0.3 | -2.2  | 0.5, 4.6 |
| Consequences          | 1–5  | -5.1 | -8.2, -2.0§ | 1.3  | -2.1, 4.8 | 1.4   | -6.8, 9.7 |
| Illness coherence     | 1–5  | -2.3 | -5.5, 0.9 | -1.2 | -5.0, 2.6 | 1.5   | -0.6, 3.9 |
| Emotional representation | 1–5  | 2.3  | -1.0, 5.6 | -4.5 | -8.3, -0.7† | -5.3  | -3.3, -7.3 |
| Psychological attributions | 1–5  | -2.0 | -5.7, 1.8 | -2.1 | -5.8, 1.6 | 1.7   | -0.7, 4.1 |
| Immunity              | 1–5  | -3.7 | -7.2, -0.2§ | 1.3  | -2.1, 4.8 | 1.4   | -6.8, 9.7 |

Model 3 (basic model plus illness perceptions and coping)

| Back pain             | B   | 95% CI | B    | 95% CI | B    | 95% CI |
| 0–10                  | -2.4 | -3.4, -1.5† | -0.1 | -1.1, 0.8 | 6.2   | 3.7, 8.9† |
| Age (years)           | 0.3  | 0.1, 0.6‡ | 0.1  | -0.2, 0.4 | 0.1   | -0.7, 1.0 |
| Female sex            | 6.4  | 1.4, 11.3‡ | -4.3 | -9.4, 0.8 | 3.8   | -9.6, 17.2 |
| Consequences          | 1–5  | -3.7 | -7.2, -0.2§ | 1.3  | -2.1, 4.8 | 1.4   | -6.8, 9.7 |
| Illness coherence     | 1–5  | -2.5 | -8.2, 3.2 | -6.8 | -9.6, -4.1† | -5.7  | -3.7, -7.7 |
| Emotional representation | 1–5  | 2.3  | -1.0, 5.6 | -4.5 | -8.3, -0.7† | -5.3  | -3.3, -7.3 |
| Decreasing activities | 1–4  | 3.7  | 10.2, 0.6 | 0.1  | -0.2, 0.4 | 0.1   | -0.7, 1.0 |
| Pacing                | 1–4  | -2.0 | -7.2, 3.2 | -2.1 | -5.8, 1.6 | 1.7   | -0.7, 4.1 |

* Work productivity loss (WPL) was assessed in only 65 patients. HRQoL = health-related quality of life; SpA = spondyloarthritis; PCS = physical component summary score; MCS = mental component summary score; 95% CI = 95% confidence interval.
† P < 0.001.
‡ P < 0.05.
§ P < 0.01.

These findings are important for managing patients with axial SpA and chronic back pain. Rheumatologists and health care professionals should be aware that illness perceptions play an important role in determining medical outcomes in these patients. Illness perceptions should, therefore, be actively explored and taken into consideration in the management plan. To maximally improve health outcomes in patients with axial SpA, psychological support could be given in addition to targeting back pain using drug treatment and physiotherapy. Several studies in other diseases have shown that psychological interventions could potentially change illness perceptions (27–30).

The main aim of our study was to investigate the clinical question of how rheumatologists and health care professionals can maximally improve health outcomes and whether and which illness perceptions and coping strategies are important for disease management in patients with early onset of axial SpA. Therefore, we performed a stepwise regression analysis rather than a mediation analysis. In the regression analysis, the effect of illness perceptions and coping strategies on the relationship between back pain and outcomes can be clearly seen. In a mediation analysis, back pain would be included as a control variable, and therefore this effect would no longer be apparent. The advantage of a mediation analysis would be that all direct and indirect effects of illness perceptions and coping could be evaluated, but the clinical interpretation of the various coefficients in the model is unclear.

In patients with chronic back pain, no associations between illness perceptions and HRQoL or work productivity loss have been investigated, as far as we know. Only one previous study investigated the association between illness
perceptions and HRQoL in patients with longstanding AS. In that study, the Brief Illness Perception Questionnaire (31) was used and showed that higher scores on the illness perception “concern,” part of the “emotional representation” of the disease, were associated with worse physical HRQoL (7). Those findings contrast with the findings in our cohort, in which it was shown that “consequences” and “chance” were associated with decreased physical HRQoL, and that “identity,” “emotional representation,” and “psychological attributions” were associated with decreased mental HRQoL. “Consequences” was also associated with increased loss of work productivity. Differences between our study and the study including AS patients might be explained by the use of different questionnaires (Brief IPQ versus IPQ-R in our cohort) and different patient populations (longstanding AS versus early axial SpA or suspected axial SpA in our cohort). It is possible that other illness perceptions become more important when the disease is longstanding.

### Table 6. Multiple-step linear regression model with back pain, illness perceptions, and coping explaining variance in HRQoL and work productivity loss in patients with chronic back pain (n = 198)*

| Model | Range | PCS | 95% CI | MCS | 95% CI | WPL | 95% CI |
|-------|-------|-----|--------|-----|--------|-----|--------|
| Model 1 (basic model) | | | | | | | |
| Back pain | 0–10 | -3.7 | -4.4, -3.0 | -1.0 | -1.8, -0.3 | 7.8 | 5.9, 9.6 |
| Age (years) | 0.5 | 0.3, 0.7 | -0.03 | -0.2, 0.2 | -0.4 | -0.9, 0.1 |
| Female | 1.0 | -2.7, 4.7 | -0.2 | -4.0, 3.7 | 8.4 | -1.1, 17.9 |
| Model 2 (basic model plus illness perceptions) | | | | | | | |
| Back pain | 0–10 | -3.0 | -3.8, -2.2 | -0.1 | -0.8, 0.7 | 6.5 | 4.5, 8.6 |
| Age (years) | 0.5 | 0.3, 0.7 | -0.01 | -0.2, 0.2 | -0.4 | -0.9, 0.1 |
| Female sex | 2.5 | -1.1, 6.1 | 1.0 | -2.4, 4.4 | 7.2 | -2.3, 16.7 |
| Identity | 0–15 | -0.6 | -1.2, -0.01 | -0.6 | -1.2, -0.01 | -0.6 | -1.2, -0.01 |
| Consequences | 1–5 | -6.4 | -8.9, -3.9 | -0.8 | -3.2, 1.7 | 10.1 | 2.8, 17.4 |
| Timeline (acute/chronic) | 1–5 | -1.6 | -4.0, 0.9 | -1.6 | -4.0, 0.9 | -1.6 | -4.0, 0.9 |
| Personal control | 1–5 | 0.5 | -2.6, 3.5 | 1.4 | -1.6, 4.3 | 0.3 | -7.8, 8.4 |
| Treatment control | 1–5 | 1.6 | -1.7, 4.9 | 1.8 | -1.7, 5.2 | -0.8 | -2.4, 1.8 |
| Illness coherence | 1–5 | -0.2 | -2.3, 2.0 | 1.0 | -1.1, 3.1 | -0.2 | -2.3, 2.0 |
| Emotional representation | 1–5 | 3.0 | 0.7, 5.4 | -4.1 | -6.4, -1.8 | -1.1 | -7.2, 5.0 |
| Psychological attributions | 1–5 | -4.2 | -6.7, -1.8 | -4.2 | -6.7, -1.8 | -4.2 | -6.7, -1.8 |
| Immunity | 1–5 | 0.2 | -2.2, 2.6 | 0.2 | -2.2, 2.6 | 0.2 | -2.2, 2.6 |
| Accident | 1–5 | -2.2 | -2.9, -1.5 | -0.2 | -0.9, 0.5 | 5.1 | 3.2, 7.1 |
| Model 3 (basic model plus illness perceptions and coping) | | | | | | | |
| Back pain | 0–10 | -2.2 | -2.9, -1.5 | -0.2 | -0.9, 0.5 | 5.1 | 3.2, 7.1 |
| Age (years) | 0.5 | 0.3, 0.7 | -0.04 | -0.2, 0.2 | -0.5 | -1.0, -0.03 |
| Female sex | 3.6 | 0.4, 6.8 | 0.8 | -2.6, 4.2 | 5.2 | -3.7, 14.2 |
| Identity | 0–15 | -0.7 | -1.3, -0.1 | -0.7 | -1.3, -0.1 | -0.7 | -1.3, -0.1 |
| Consequences | 1–5 | -3.4 | -5.7, -1.0 | -5.0 | -6.9, -3.1 | -5.0 | -6.9, -3.1 |
| Emotional representation | 1–5 | 2.6 | 0.7, 4.6 | -3.4 | -5.6, -1.3 | -3.4 | -5.6, -1.3 |
| Psychological attributions | 1–5 | -3.4 | -5.6, -1.3 | -3.4 | -5.6, -1.3 | -3.4 | -5.6, -1.3 |
| Comforting cognitions | 1–4 | -0.2 | -4.4, 4.0 | -0.2 | -4.4, 4.0 | -0.2 | -4.4, 4.0 |
| Decreasing activities | 1–4 | -5.9 | -9.7, -2.2 | -0.5 | -3.6, 2.5 | 10.2 | -1.1, 21.5 |
| Optimism | 1–4 | 2.8 | 0.2, 5.4 | 2.5 | -1.3, 6.4 | 2.5 | -1.3, 6.4 |
| Pacing | 1–4 | -4.5 | -8.5, -0.5 | -4.5 | -8.5, -0.5 | -4.5 | -8.5, -0.5 |
| Creative solution-seeking | 1–4 | -2.5 | -5.6, 0.6 | -2.5 | -5.6, 0.6 | -2.5 | -5.6, 0.6 |
| Accepting | 1–4 | -1.3 | -4.0, 1.5 | -1.3 | -4.0, 1.5 | -1.3 | -4.0, 1.5 |

* Work productivity loss (WPL) was assessed in only 144 patients. HRQoL = health-related quality of life; PCS = physical component summary score; MCS = mental component summary score; 95% CI = 95% confidence interval.

† P < 0.001.
‡ P < 0.01.
§ P < 0.05.

Further, several studies showed that the maladaptive coping strategies “decreasing activities” and “pacing” were associated with worse HRQoL and withdrawal from the workforce in patients with AS (11–13), and that “avoiding physical activity” was associated with increased pain and disability in patients with chronic back pain (14). In our study, increased use of the “decreasing activities” and “pacing” strategies were associated with lower physical HRQoL. These coping strategies were not related to work productivity loss, which could be explained by the fact that work productivity loss and withdrawal from the workforce are different concepts. Moreover, this difference could also reflect the difference between early versus longstanding disease.

In contrast to our expectations, a positive association between illness perception “emotional representation” (having more negative emotions toward the disease) and physical HRQoL was observed in our study. When
examining correlations between “emotional representation” and the PCS, we observed negative associations in all patients and in each subgroup, as expected. Therefore, this effect appears only in the multivariable model, in the context of the other illness perceptions that play a significant role. The context of other illness perceptions might explain why the association between “emotional representation” and HRQoL reversed. Additionally, in our study coping strategies did not have an additional influence over illness perceptions for the association between back pain and HRQoL or work productivity loss, in contrast to our hypothesis. These unexpected findings could be explained by the CSM itself, because the CSM is a self-regulatory model. The CSM implies that individuals use coping strategies based on his or her illness perceptions, and illness perceptions are adapted based on coping strategies by a feedback loop from HRQoL to these factors. Using maladaptive coping strategies decreases HRQoL, and according to the CSM, illness perceptions will be adapted in a manner such that worsening in HRQoL is reduced. Furthermore, the effect of illness perceptions may be balanced by coping strategies. Having more negative emotions associated with a disease could lead to a change in coping strategies from maladaptive toward adaptive in order to decrease worsening of HRQoL. Future studies are needed to investigate this notion further. The effect of coping strategies could have been disadvantaged or changed or different due to the fact that coping strategies cannot be added to the model before illness perceptions are added. However, testing the coping strategies first would violate the CSM.

Our main analysis was performed in all patients suspected of having axial SpA. Remarkably, comparable results were observed in all analyses in all subgroups. This finding may be explained by the fact that patients were unaware of the results of laboratory and imaging tests and diagnosis when they filled out the questionnaires. Therefore, it would be interesting to study the impact of receiving a diagnosis on illness perceptions and coping. Unfortunately, in our cohort we currently have no data on this subject.

Our results suggest that HRQoL and work productivity can be further improved by interventions targeting patients’ cognitions and behavior along with treatment that suppresses pain and inflammation. Targeting patients’ cognitions and behavior along with treatment that suppresses pain is more established not only in studies but also in use of existing treatment options to decrease the burden of disease in patients suspected of having axial SpA. Future research is needed to investigate whether the impact of illness perceptions and coping strategies vary over time, the differences between these factors in early and longstanding disease, as well as the impact of targeting illness perceptions on back pain and physical HRQoL.

In conclusion, in patients suspected of having axial SpA, high intensity of back pain is associated with worsening of physical HRQoL and increasing loss of work productivity. Our results suggest that, in addition to treating back pain, targeting negative illness perceptions could improve HRQoL and work productivity. Our study supports the development of interventions targeting patients’ cognitions in addition to use of existing treatment options to decrease the burden of disease in patients suspected of having axial SpA. Future research is needed to investigate whether the impact of illness perceptions and coping strategies vary over time, the differences between these factors in early and longstanding disease, as well as the impact of targeting illness perceptions on back pain and physical HRQoL.

**AUTHOR CONTRIBUTIONS**

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be published. Ms van Lunteren had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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