EULAR recommendations for the role of the nurse in the management of chronic inflammatory arthritis

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ABSTRACT

Objectives The authors aim to develop European League Against Rheumatism recommendations for the role of the nurse in the management of patients with chronic inflammatory arthritis, to identify a research agenda and to determine an educational agenda.

Methods A task force made up of a multidisciplinary expert panel including nurses, rheumatologists, occupational therapist, physiotherapist, psychologist, epidemiologist and patient representatives, representing 14 European countries, carried out the development of the recommendations, following the European League Against Rheumatism standardised operating procedures. The task force met twice. In the first meeting, the aims of the task force were defined, and eight research questions were developed. This was followed by a comprehensive, systematic literature search. In the second meeting, the results from the literature review were presented to the task force that subsequently formulated the recommendations, research agenda and educational agenda.

Results In total, 10 recommendations were formulated. Seven recommendations covered the contribution of nurses to care and management: education, satisfaction with care, access to care, disease management, psychosocial support, self-management and efficiency of care. Three recommendations focused on professional support for nurses: availability of guidelines or protocols, access to education and encouragement to undertake extended roles. The strength of the recommendations varied from A to C, dependent on the category of evidence (1A–3), and a high level of agreement was achieved. Additionally, the task force agreed upon 10 topics for future research and an educational agenda.

Conclusion 10 recommendations for the role of the nurse in the management of chronic inflammatory arthritis were developed using a combination of evidence-based and expert consensus approach.

INTRODUCTION

In rheumatology, registered nurses often act as the interface between patients and other members of the multidisciplinary team. As a result of new treatment regimens and organisational developments, the role of the nurse is undergoing great change. However, there are large differences across countries and regions. In several European countries, rheumatology as a nursing specialty does not exist, whereas in other countries, it has developed into a recognised specialty with nurses undertaking advanced and extended roles. These include self-management support, patient education and counselling, intra-articular injections, recommendation for and the prescription of drug treatments, referral to other health professionals, hospital admission of patients, manning telephone advice lines and monitoring disease-modifying and biological treatments. Nurse-led clinics have been established, and their effectiveness has been shown to bring added value to patients’ outcomes at a lower cost. While some countries have accepted that interventions undertaken by nurses are essential to effectively tackle the challenges of chronic illness in an economic and integrated fashion, this concept has not developed everywhere.

The aim of the present study was to evaluate the currently available literature according to the European League Against Rheumatism (EULAR) standardised operating procedures in order to provide recommendations for the role of the nurse in the management of chronic inflammatory arthritis (CIA). This evidence-based approach was complemented by an expert consensus approach.

METHODS

A multidisciplinary task force made up of 15 nurses, a rheumatologist, an occupational therapist, a psychologist, a physiotherapist, two patient representatives and a research fellow, representing several European countries, met twice under the leadership of two conveners and a rheumatologist/clinical epidemiologist.

During the first meeting, the task force formulated eight research questions. These questions served as a guide to the systematic literature review (SLR) and subsequently as the basis for the recommendations. The term ‘CIA’ was confined to rheumatoid arthritis (RA), ankylosing spondylitis and psoriatic arthritis, thereby excluding other systemic inflammatory conditions (eg, systemic lupus erythematosus) and non-inflammatory disorders (eg, fibromyalgia).

The target population for the recommendations was chosen to be healthcare professionals working...
in the field of rheumatology (rheumatologists, nurses and other disciplines), patients and policymakers.

After translation of the research questions into relevant search terms (online supplementary appendix 1), an extensive SLR of MEDLINE, Embase, Cochrane CENTRAL, Cumulative Index to Nursing and Allied Health Literature and PsycINFO was performed in August 2010. Two main search terms—‘inflammatory arthritis’ and ‘nurse’—were searched for in titles, keywords or full texts using MeSH (Medical Subject Headings) headings, subheadings, thesaurus or free text words and truncation symbols. No limitations with regard to publication type, research type, language or time period were applied. Selected titles and abstracts were screened independently for eligibility by AvT and YvE-H. Eligibility criteria were ‘inflammatory arthritis’, ‘interventions undertaken by nurses’ and ‘relevant outcomes to answer the research questions’. The selection was shared with the other members of the task force, and suggestions and comments by the experts complemented the result of the SLR. Additionally, abstracts from American College of Rheumatology and EULAR meetings (2008–2010) were searched (selection procedure is shown in online supplementary appendix 2).

During the second meeting, the formulation of the recommendations was discussed by the entire group until a consensus was reached. The recommendations were graded based on the level of evidence of the literature found (online supplementary appendix 3). This was sent to each participant for final approval and voting on a scale from 0 to 10. Finally, the task force agreed upon the formulation of a research agenda and an educational agenda.

**RESULTS**

In total, 54 studies met the inclusion criteria (online supplementary appendix 4). The selection comprised 1 meta-analysis, 8 randomised controlled trials (RCTs), 2 controlled clinical trials, 9 quasi-experimental studies and 34 descriptive studies. As the majority of the studies were descriptive, it was acknowledged that there was a high risk of bias.

Table 1 shows the 10 recommendations with a level of evidence ranging from 1A to 3. The strength of recommendations varied from A to C, and a high level of expert agreement was achieved.

**Recommendations**

All recommendations relate to care delivered by registered nurses with a specific training in rheumatology. This care includes the monitoring of disease consequences on the level of daily activities, participation and psychosocial consequences and, increasingly, the monitoring of disease activity, drug treatment and drug side effects.

The task force judged that the level of evidence for the role of the nurse is far greater in the management of RA than in the management of ankylosing spondylitis or psoriatic arthritis, and therefore, the recommendations should be regarded as points to consider for patients with these conditions.

The order of the recommendations follows the discussion in the task force. The first three recommendations are formulated from the patients’ perspective, and the other recommendations are formulated from the nurses’ perspective.

1. Patients should have access to a nurse for education to improve knowledge of CIA and its management throughout the course of their disease.

Patient education is defined as a planned process aiming to improve coping strategies and increase self-care abilities. A statistically significant increase in the patient’s knowledge of the disease process, treatment strategies (eg, drug treatment), physiotherapy and self-management strategies (eg, joint protection techniques) was found in patients with RA who were educated during monitoring or who received a specific educational programme from nurses. Moreover, statistically significant greater levels of knowledge were found in patients monitored and educated by a nurse compared to patients monitored by doctors.

Involvement in disease management allows nurses to offer timely education to newly diagnosed and established patients. In addition to information about their disease and treatments, education should address risk factors for comorbidities, such as cardiovascular problems. Nurses can also play an important role in educating patients about the principles of ‘treat to target’ in order to enhance adherence. Overall, the literature demonstrates that education by nurses improves the patients’ knowledge of their disease and disease-related issues.

2. Patients should have access to nurse consultations in order to experience improved communication, continuity and satisfaction with care.

Satisfaction with care is considered an indicator of the quality of care. The majority of the studies showed statistically significant increased satisfaction with information, empathy, technical quality and attitude of the professional, as well as access to care in patients with RA when monitored by a nurse compared to monitoring by doctors or other health professionals. There were no differences in patient satisfaction after intra-articular injections given by either a nurse or a doctor. Patients valued nurses’ communication skills and nursing care in terms of advice to use assistive devices, referral to other members of the multidisciplinary team and education.

Quality and continuity of care in nurse-led clinics were perceived as good. Holistic care and patient-centred information were found to contribute to patient satisfaction.

However, some studies report conflicting evidence. One study did not find statistically significant changes in patients’ satisfaction after monitoring by a nurse practitioner. In another study, patients’ satisfaction with multidisciplinary team care was found to be significantly higher than care coordinated by a clinical nurse specialist.

Nurses tend to be accessible to patients and can facilitate access to services. Continuity of care provides the opportunity to establish a confidential and long-standing patient–professional relationship that is considered to be important by patients. Despite some conflicting evidence, the task force concluded that disease monitoring or follow-up care by nurses enhances patients’ perceptions of care.

3. Patients should have access to nurse-led telephone services to enhance continuity of care and to provide ongoing support.

The unpredictable, fluctuating nature of rheumatic diseases means that rapid access to advice is of utmost importance for patients. Telephone helplines were found to support accessibility to care. Patients perceived that telephone helplines support enhanced continuity of care and provided them with access to a knowledgeable practitioner who would often be the first point of contact at times of need. Mostly, patients stated that they would call again if needed. The contents of the service that was offered by the helplines were not clearly described in all studies, and its focus varied. Some services provided regular follow-up, while others focused on support by answering disease-related questions or worries that might avoid unnecessary consultations with general practitioners.

There were...
recommendations for rheumatology nursing management of CIA

<table>
<thead>
<tr>
<th>Recommendations</th>
<th>Category of evidence</th>
<th>Strength of recommendation</th>
<th>Agreement (0–10, mean ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Patients should have access to a nurse for education to improve knowledge of CIA and its management throughout the course of their disease</td>
<td>1B</td>
<td>A</td>
<td>9.9 ± 0.2</td>
</tr>
<tr>
<td>2 Patients should have access to nurse consultations in order to experience improved communication and satisfaction with care</td>
<td>1B</td>
<td>A</td>
<td>9.1 ± 0.6</td>
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<tr>
<td>3 Patients should have access to nurse-led telephone services to enhance continuity of care and to provide ongoing support</td>
<td>3</td>
<td>C</td>
<td>9.2 ± 0.8</td>
</tr>
<tr>
<td>4 Nurses should participate in comprehensive disease management to control disease activity, to reduce symptoms and to improve patient-preferred outcomes</td>
<td>1A</td>
<td>A</td>
<td>9.4 ± 0.8</td>
</tr>
<tr>
<td>5 Nurses should identify, assess and address psychosocial issues to minimise the chance of patients’ anxiety and depression</td>
<td>1B</td>
<td>A</td>
<td>9.4 ± 0.7</td>
</tr>
<tr>
<td>6 Nurses should promote self-management skills in order that patients might achieve a greater sense of control, self-efficacy and empowerment</td>
<td>3</td>
<td>C</td>
<td>9.7 ± 0.5</td>
</tr>
<tr>
<td>7 Nurses should provide care that is based on protocols and guidelines according to national and local contexts</td>
<td>3</td>
<td>C</td>
<td>9.4 ± 1.0</td>
</tr>
<tr>
<td>8 Nurses should have access to and undertake continuous education in order to improve and maintain knowledge and skills</td>
<td>3</td>
<td>C</td>
<td>9.7 ± 0.5</td>
</tr>
<tr>
<td>9 Nurses should be encouraged to undertake extended roles after specialised training and according to national regulations</td>
<td>3</td>
<td>C</td>
<td>9.5 ± 0.8</td>
</tr>
<tr>
<td>10 Nurses should carry out interventions and monitoring as part of comprehensive disease management in order to achieve cost savings</td>
<td>1B</td>
<td>C</td>
<td>8.8 ± 1.3</td>
</tr>
</tbody>
</table>

CIA, chronic inflammatory arthritis.

also examples of helplines that focused specifically on triage to identify patients that required fast-track clinical assessment. In countries where telephone services are not available, information and support are sometimes given by email. In general, both means of support contribute to enabling patient accessibility and appropriate care.

4. Nurses should participate in comprehensive disease management to control disease activity, to reduce symptoms and to improve patient-preferred outcomes.

Nurses have increasingly combined providing support to patients in a broad spectrum of disease-related problems, such as psychosocial problems and limitations in participation, with disease monitoring. Several studies showed that nurse-led care results in equivalent suppression of disease activity in patients with RA, in comparison with medical care. Moreover, appropriately trained nurses were able to detect early arthritis, make referrals, determine necessary interventions and change medications.

Patients with RA also perceived statistically significant less pain and fatigue when monitored by nurses, compared to doctors. Research showed that these symptoms contribute to patients’ perceptions of disease impact, and therefore, patients consider them as preferred outcomes in clinical trials.

There is evidence that nurses can manage CIA appropriately, that they can contribute to symptom control and that they can play a role in the early detection of arthritis.

5. Nurses should identify, assess and address psychosocial issues to minimise the chance of patients’ anxiety and depression.

The psychosocial impact of CIA is considerable, and anxiety and depression are both well-known comorbidities in RA. One study found a statistically significant reduction in anxiety and depression in patients with RA after monitoring by a nurse. Others showed equivalent reductions in anxiety and depression in patients receiving nurse-led or medical care. It has been shown that cognitive–behavioural interventions provided by a nurse can statistically significantly improve the emotional well-being in patients with RA. Moreover, quality of life was comparable between patients receiving either nurse-led or multidisciplinary team care. Psychosocial adjustment, which is considered as overall adaptation to the impact of RA, was comparable in patients monitored by either a nurse or a doctor, but patients monitored by a nurse reported supplementary increased social activities. Patients valued the opportunity to discuss the wider implications of their condition with a nurse. Counselling in psychological issues is considered important, but undertaking this role depends strongly on the level of the problems, the skills of the nurse and the ability to access other sources of support when required—for example, the availability of psychologists in assisting with these tasks. Therefore, the task force considered identifying problems and referring to other professionals when needed as key components of nursing care.

6. Nurses should promote self-management skills in order that patients might achieve a greater sense of control, self-efficacy and empowerment.

Self-efficacy refers to the belief that one can successfully execute the behaviour required to attain certain goals and, thus, to feel in control. Research shows that nurses are likely to contribute to increased perceived control, levels of self-efficacy and empowerment. Patients reviewed by a clinical nurse specialist in a drug-monitoring clinic perceived clinically relevant increased perception of control. After a cognitive–behavioural intervention provided by nurses, patients demonstrated a statistically significant increase in personal coping resources, such as competency beliefs and decrease of helplessness. Nurse-led management, information and support have been shown to increase self-efficacy beliefs and self-reliance of patients as well as to contribute to patient empowerment.

However, some studies were unable to demonstrate any changes in lifestyle, self-management behaviour or self-efficacy after interventions undertaken by nurses. Support with regard to self-management comprises all actions that encourage patients to manage their own disease, and this is a task for all members of the multidisciplinary team. As self-management support is multifaceted, the task force considered this as an important role of the nurse, rather than a single intervention undertaken by nurses.

7. Nurses should provide care that is based on protocols and guidelines according to national and local contexts.

Guidelines provide research-based options for decisions, whereas protocols describe steps to be taken to reduce variation in the treatment of patients. Guidelines and protocols are essential for all healthcare professionals to ensure safe and...
Recommendation

high-quality care. Often, these guidelines and protocols will be adapted to a national or local context.56

It has been demonstrated that structured implementation material supported nurses in the guidance of patients with a complex treatment regimen. Furthermore, nurses’ perceived capability for guidance statistically significantly increased after an educational session.57 Guidelines have been found to support nurses’ clinical decision-making skills with regard to assessment and treatment.56 referral,42 supplementary prescription,41 and therefore contribute to evidence-based nursing59 and holistic care.41 58 Protocols have been found to support continuity and safety of care in terms of immediate and appropriate adjustments of treatment41 42 60 and enabled nurses to discriminate between early arthritis and other conditions.61

However, standardised care should be implemented alongside national and regional regulations yet enable an individualised, patient-centred approach. Overall, guidelines and protocols are likely to support nurses in providing evidence-based care.8 Nurses should have access to and undertake continuous education in order to improve and maintain knowledge and skills.

The literature demonstrates that nurses undertake a wide variety of interventions, including self-management support, education, counselling, drug monitoring, drug prescription and administration of intra-articular injections.62–69 In a survey undertaken in the UK, the majority of the nurses felt confident in their abilities to undertake these tasks.60 The self-confidence of nurses is supported by knowledge of rheumatic diseases and their treatment, skills in relation to education, counselling and training, collaboration with other health professionals and manual skills.70 Furthermore, nurse education needs to be tailored to reflect research findings if nurses are to maintain and enhance their knowledge and skills appropriately.62 70 Studies have demonstrated that the contents of consultations and issues for patient education and counselling are dependent on the educational level of nurses.71–75 However, it has been reported that knowledge and skills do not appear to be sufficiently covered by basic and advanced training at present.66

Educational opportunities in nursing vary to a large extent if an educational curriculum is not defined accurately. Given the complexity of the tasks and activities that are performed by nurses, ongoing access to well-defined education on a basic, advanced and extended level is needed.

9. Nurses should be encouraged to undertake extended roles after specialised training and according to national regulations.

Increasingly, nursing care expands from a more basic level to an advanced or an extended level. The rationale for these developments comes from several perspectives. From the patient’s perspective, holistic care, patient-centred information and communication skills of nurses are appreciated, and improved outcomes such as knowledge, satisfaction and physical and psychological symptoms were found.9–11 26 27 29 42 From the professional’s perspective, job satisfaction is enhanced by greater autonomy and by optimal use of nurses’ qualities and skills.42 62 From an organisational perspective, advanced and extended roles may prevent fragmentation of care and promote efficiency and accessibility.42 62 Key components for achieving extended roles include performing outpatient procedures, prescribing drugs and treatment and taking a lead in the organisation of local health services.82

However, some concerns about extended nursing roles have been raised among members of the medical and the nursing professions about being ‘placebo-doctors’ or ‘second-rate doctors’.26 62 If role development is patient-focused, aiming to meet patients’ identified needs and to improve overall patient care, the extended role of the nurse should be regarded as a complement to the medical role and not only as a substitution of medical tasks.

The competencies and skills of the nurse should be considered and optimised to further improve patient care, to enhance and mobilise nursing competency and to improve efficiency of care.10 Nurses should carry out interventions and monitoring as part of comprehensive disease management in order to achieve cost savings.

Innovative nurse-led care has advanced to increase efficiency of care. Coordinated care by a clinical nurse specialist was shown to be statistically significantly cost-effective, compared with multidisciplinary inpatient or day-patient care.12 Additional costs for a nurse practitioner to a medical team were found to be partially compensated by taking over tasks from other team members.30 Moreover, nurse-led monitoring led to decreased medical referrals42 or decreased the length of stay in a rehabilitation programme for patients with RA,49 all pointing to cost savings. Furthermore, rheumatology telephone helplines are likely to contribute to cost savings by decreasing the number of unnecessary doctor consultations32 34 36 and by reducing follow-up waiting time.33 Accessibility of care also improved with the nurse being able to discriminate different categories of early arthritis.61 Recent preliminary data suggest a positive cost–benefit of the role of nurse specialists by preventing unscheduled care and hospital admissions.69

However, compared to usual care from a rheumatologist, patients seen by a nurse were more frequently referred to occupational therapists, and this difference was statistically significant.9 54 It is arguable whether this phenomenon is a consequence of the greater emphasis that nurses placed on joint protection and improving functioning in daily living9 and, therefore, could be considered as quality improvement of care rather than increase in costs.54 There is a need for high-quality economic analyses in future research.

Research agenda

In addition to the developed recommendations, a research agenda and an educational agenda were formulated. The research agenda highlights where there is weak or lacking evidence to further optimise the role of the nurse in the management of CIA and is shown in box 1.

Educational agenda

The educational agenda was defined to support educational opportunities for nurses and to guarantee quality in nursing care (shown in box 2).

DISCUSSION

Nurses are the largest group of healthcare professionals, and their role development in the provision of team care for patients with rheumatic diseases follows a worldwide tendency to more proactive, evidence-based care for patients with chronic disorders with nurses fulfilling extended roles.74 75 The role of nurses in the management of CIA appeared to differ greatly between countries and across regions due to their educational level, training and expertise, as well as to national and regional regulations and contexts and funding issues related to overall healthcare provision.

Evidence for the additional value of nurses was most obvious when disease monitoring by nurses was combined with support
for patients in a broad spectrum of disease-related problems. However, roles, tasks and qualifications should be clearly described in frameworks to practice that include protocols and guidelines.

Nurses tend to be accessible for patients. Given their qualities and skills with regard to coordination of care, they may facilitate increased access to multidisciplinary care. In practice, however, some tasks may be provided by other health professionals, depending on local accessibility and competency. It is arguable whether these tasks have the same quality when provided by different health professionals. This study explicitly deals with nurses, because this profession is often not clearly visible in multidisciplinary team care, but their role should be considered in the context of care delivery of other healthcare providers and an active role of patients. By their continuous presence, nurses can identify and communicate specific areas that can be addressed by other members of the multidisciplinary team.

Ten recommendations for the role of the nurse in the management of patients with CIA were developed. There are some limitations to these recommendations that need to be addressed.

First, the task force decided to include all types of studies that could give insight in nursing care, as only a limited number of RCTs exist, and RCTs alone may not adequately cover the topic. Qualitative studies provide important insight in patients’ individual needs, functional limitations and the extent to which different types of care meet those needs, all of which may improve quality of care. However, the high level of agreement within the task force supported the method used. Agreement with the recommendations by nurses, rheumatologists, healthcare professionals in daily practice and patients will soon be validated simultaneously with dissemination of the recommendations.

The research agenda will support future directions of nursing research, and it is important that high-quality studies, with clear descriptions of nursing roles and interventions, will be conducted as the role of the nurse in care is currently often not clearly stated in studies. Given the research issues and themes of importance, qualitative and quantitative insights into nursing care are needed.

The educational agenda will support access to high-quality education for nurses; will enhance clarity about knowledge, skills and competencies required by nurses, and will reduce diversity of the nursing role within and between countries.

In conclusion, this study provides 10 evidence-based and expert opinion-based recommendations on the role of the nurse in the management of CIA. These recommendations provide a basis for emphasising and optimising rheumatology nursing care in order to contribute to a more standardised level of professional nursing across Europe.

**Recommendation**

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<th>Box 1 Research agenda</th>
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<tbody>
<tr>
<td>To study the contribution of the nurse in improving access to care and in facilitating the effective utilisation of care provided by members of the multidisciplinary team</td>
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<td>To study the role of nurses in optimising ‘treat to target’ in early disease</td>
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<tr>
<td>To study the contribution of the nurse in improving patient-preferred outcomes</td>
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<td>To compare the different components of nursing care in each European country in relation to knowledge and competencies</td>
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<td>To perform cost-effectiveness studies across different European countries, on the role of the nurse in basic and advanced practice</td>
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<td>To study the long-term effects of interventions by a nurse on quality of life, psychosocial and general well-being</td>
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<tr>
<td>To study the contribution of the nurse in improving self-management and self-efficacy</td>
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<tr>
<td>To study the impact of interventions by a nurse on the patient’s employment status and social participation</td>
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<tr>
<td>To define the contribution of the nurse in the prevention of comorbidities</td>
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<tr>
<td>To study the recommendations in different patient populations including ankylosing spondylitis and psoriatic arthritis</td>
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<th>Box 2 Educational agenda</th>
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<tr>
<td>To develop a competency framework for nurses</td>
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<tr>
<td>To develop educational nursing programmes at the basic and advanced level</td>
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To study the contribution of the nurse in improving access to care and in facilitating the effective utilisation of care provided by members of the multidisciplinary team. 21

However, roles, tasks and qualifications should be clearly described in frameworks to practice that include protocols and guidelines.

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**Contributors** During the development of the recommendations, the steering group discussed authorship, and the authors decided to include all participants who attended at least one meeting of the task force. This was also discussed with the members of the task force. All 26 authors contributed to the development process of the recommendations, drafting of the article and final approval of the version of the study to be published.

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