Considerations for improving quality of care of patients with rheumatoid arthritis and associated comorbidities

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ABSTRACT

Objective Rheumatoid arthritis (RA) is a chronic autoimmune inflammatory disorder with a global prevalence of approximately 0.5–1%. Patients with RA are at an increased risk of developing comorbidities (eg, cardiovascular disease, pulmonary disease, diabetes and depression). Despite this, there are limited recommendations for the management and implementation of associated comorbidities. This study aimed to identify good practice interventions in the care of RA and associated comorbidities.

Methods A combination of primary research (180+ interviews with specialists across 12 European rheumatology centres) and secondary research (literature review of existing publications and guidelines/recommendations) were used to identify challenges in management and corresponding good practice interventions. Findings were prioritised and reviewed by a group of 18 rheumatology experts including rheumatologists, comorbidity experts, a patient representative and a highly specialised nurse.

Results Challenges throughout the patient pathway (including delays in diagnosis and referral, shortage of rheumatologists, limited awareness of primary care professionals) and 18 good practice interventions were identified in the study. The expert group segmented and prioritised interventions according to three distinct stages of the disease: (1) suspected RA, (2) recent diagnosis of RA and (3) established RA. Examples of good practice interventions included enabling self-management (self-monitoring and disease management support, for example, lifestyle adaptations); early arthritis clinic; rapid access to care (online referral, triage, ultrasound-guided diagnosis); dedicated comorbidity specialists; enhanced communication with primary care (hotline, education sessions); and integrating patient registries into daily clinical practice.

Conclusion Learning from implementation of good practice interventions in centres across Europe provides an opportunity to more widely improved care for patients with RA and associated comorbidities.

Key messages

What is already known about this subject
► RA has a global prevalence of about 0.5–1%. The global burden of RA has risen, despite no significant change in prevalence.
► Despite publication of international guidelines and recommendations on RA management, there are limited international recommendations on the management of the several RA-associated comorbidities.
► There are many challenges in the delivery for RA care and associated comorbidities which exist at the level of the healthcare system, healthcare providers and patient, across the patient pathways.

What does this study add
► This initiative highlights good practice interventions which are in place at selected centres across Europe and addresses some of the challenges as well as opportunities in delivery of care.
► An expert multi-disciplinary panel consisting of rheumatologists, specialists in associated comorbidities, a patient representative and a highly specialist nurse prioritised, by consensus, the good practice interventions at three different disease stages; suspected RA, recent diagnosis of RA and established RA.
► The study findings inform European considerations on good practice management of RA and associated comorbidities.

How might this impact on clinical practice
► The interventions highlighted in this report could be potentially implemented by and adapted to varying healthcare settings to improve the quality of care of patients with RA and their associated comorbidities.

INTRODUCTION

Rheumatoid arthritis (RA) is one of the most common autoimmune inflammatory diseases worldwide, affecting between 0.5% and 1% of
the global population. Although life expectancy of patients with RA has increased over the last decades, they are still at increased risk of developing comorbid conditions.

Comorbidities discussed in this paper refer to conditions that may either be directly or indirectly caused by RA, RA treatment or by RA outcomes, such as decreased mobility and functional impairment. Multi-morbidities related to increasing age and lifestyles of patients with RA are also of note and included in this paper’s definition of comorbidities.

When compared to the general population, patients with RA have the following adjusted lifetime HRs: 1.41 for the development of cardiovascular disease (40–70% increased risk), 896 for interstitial lung disease (ILD) (7% increased risk) and 15 for diabetes. Estimates of the prevalence of depressive disorder in patients with RA range between 13% and 20%, which is 2–3 times higher than for the general population while the disability-adjusted life years of patients with RA has increased from 3.3 million in 1990 to 4.8 million in 2010.

Guidelines and recommendations, such as the 2016 EULAR points to consider for reporting, screening for and preventing selected comorbidities, primarily address screening and prevention of comorbidities. However, there is a clear need for recommendations on the management of RA-associated comorbidities.

The objectives of this study were to (1) understand the current challenges in the management of patients living with RA and associated comorbidities, (2) identify and collate approaches undertaken by rheumatology teams to address these challenges and (3) discuss potential strategies to replicate and implement these interventions across healthcare systems and care settings to improve patient outcomes.

MATERIALS AND METHODS

The study followed a three-step approach:

1. A literature review using an integrative approach to identify reported and evidence-based challenges and good practice examples in the management of RA and its associated comorbidities. The review included
   - Academic literature search. A database search for articles in PubMed using a combination of prioritised search terms was conducted. The search terms included rheumatoid arthritis, comorbidities, cardiovascular risk, diabetes, depression, interstitial lung disease, management, early identification, diagnosis, challenges in care, patient outcomes, guidelines, recommendation, quality of care, screening, diagnosis, delay referral, treat to target, disease burden, multidisciplinary approach to care, self-management, personalised care, etc. Additional articles were retrieved through the citation-tracking of original publications and were based on the recommendation from the expert panel.

   - Grey literature search. The publicly available grey literature was investigated by applying the same prioritised search terms in search engines. Non-English-language references were excluded unless there was sufficient explanatory text in English. The time allotted for the grey literature search was 1 hour or until saturation was reached, whichever came first. Saturation was defined as not identifying new literature to include in analysis for 40 min or five consecutive search pages, whichever came first.

   - Geography: the centres, each located in a different European country, were selected to represent a multinational approach to RA management.

   - Focus: the centre’s experience with RA was considered; both specialised and more generalist centres were selected to ensure the results of the study would be applicable to the wider clinical community.

   - Centre type: the centres’ models and involvement in the wider healthcare ecosystem (eg, private vs public funding, size, partnerships within the community, etc) were assessed.

   - Participating centres included Diakonhjemmet Hospital (Oslo, Norway), Cochin Hospital (Paris, France), Institute of Rheumatology (Prague, Czech Republic), Hospital Santa Maria (Lisbon, Portugal), University Hospital La Paz (Madrid, Spain), Chapel Allerton Hospital and University of Leeds (Leeds, UK), University Clinic Saint-Luc (Brussels, Belgium), Geneva University Hospital (Geneva, Switzerland), Sint Maartenskliniek (Nijmegen, Netherlands), Karolinska University Hospital and the Centre for Rheumatology (Stockholm, Sweden), Rigshospitalet (Copenhagen, Denmark) and ASST Gaetano Pini-CTO Institute (Milan, Italy).

   - Approximately 180 qualitative semi-structured interviews were conducted to gain a robust understanding of the management of RA and associated comorbidities through the visits. An interview guide was created to facilitate the structured collection of quantitative and qualitative insights. The questions were designed to capture the RA service set-up and patient pathway design within each centre and assessing the key challenges, which were categorised as ‘gaps in care’ and ‘drivers’ (ie, reasons for gaps in care).

   - All the interviews were conducted on an individual basis over a duration of approximately 1 hour. Before the start of the interview, all participants were given an overview of the project objectives.

   - The individuals interviewed included a wide range of healthcare professionals (HCPs) such as rheumatologists, nurses, physiotherapists, occupational therapists and RA-associated comorbidity specialists, pharmacists, practice management.
Interview responses were systematically analysed for insights and patterns. Thematic analysis was undertaken to identify all challenges and interventions, which were subsequently reviewed by the expert panel.

3. The findings from the above secondary and primary research were tested with an expert panel. This panel of European experts was set up to ensure relevance in a range of healthcare systems. It consisted of 18 members including 16 clinicians of which there were 12 rheumatologists (one from each centre visited), 1 cardiologist, 1 pulmonologist, 1 diabetologist and 1 psychologist (ie, one representative for the four preselected common comorbidities). The panel also included one rheumatology specialist nurse, acknowledging evidence that highlights the importance of the role of nurses in practice and one patient representative to capture the views and perspectives of patients. The expert panel was involved throughout the study and met on three occasions to guide the project objectives and discuss project findings. Ethical approval was not required for this study, since it did not include data collection from patients or other individuals.

RESULTS

Several challenges in the management of RA and associated comorbidities were identified. Gaps in the care of patients with RA included the following: delays in securing a diagnosis consisting of seeking medical advice (>12 weeks\textsuperscript{14}); referral delays (with a median of 4 visits to the general practitioner before patients are referred to a rheumatologist\textsuperscript{15}) leading to an overall delay in diagnosis (in several countries, it can take up to a year from symptom onset\textsuperscript{16}); and upon diagnosis and treatment initiation, poor patient adherence to therapy (varying between 30\% and 80\%\textsuperscript{17}) and lack of monitoring (up to 50\% of patients do not have access to an regular review in a secondary care setting\textsuperscript{18}). Drivers of these gaps in care were evident at a healthcare system, patient and HCP level (table 1).

Challenges were also evident throughout the patient pathway for the care of RA-associated comorbidities. One pertinent challenge was the limited guidance for the care of comorbidities in patients with RA (table 2).

Table 1  Gaps in care and their associated drivers evident across the journey of patients with RA

<table>
<thead>
<tr>
<th>Stage of patient journey</th>
<th>Gaps in care</th>
<th>Drivers*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Awareness and prevention</td>
<td>Delays in patients seeking medical advice</td>
<td>Low public awareness\textsuperscript{14}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Lack of patient education\textsuperscript{19}</td>
</tr>
<tr>
<td>Referral</td>
<td>Delayed referrals from PCPs to rheumatologist</td>
<td>Limited awareness of PCPs on signs and symptoms of RA\textsuperscript{20}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Long waiting times for specialists\textsuperscript{21}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Joint stiffness and inflammation are common for many other conditions</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Lack of knowledge regarding the referral pathway or the importance of rapid referral\textsuperscript{21}</td>
</tr>
<tr>
<td>Diagnosis</td>
<td>Delay in diagnosis</td>
<td>Complex diagnostic requirements\textsuperscript{14}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Limited availability of imaging facilities\textsuperscript{51,22}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Shortage of rheumatologists\textsuperscript{22}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Limited number of HCPs trained to interpret diagnostic results (including imaging)\textsuperscript{14}</td>
</tr>
<tr>
<td>Treatment and management (pharmacological and non-pharmacological)</td>
<td>Delayed treatment initiation</td>
<td>Budgetary cap and restrictive clinical recommendations regarding choice of therapy\textsuperscript{22}</td>
</tr>
<tr>
<td></td>
<td>Poor patient adherence to therapy</td>
<td>Disease management not aimed at optimising quality of life\textsuperscript{23}</td>
</tr>
<tr>
<td></td>
<td>Lack of coordinated treatment</td>
<td>Deviation from recommendations, or difficulty in implementing them into clinical practice\textsuperscript{22}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Lack of regular review and engagement with patients’ needs\textsuperscript{24}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Treatment adherence\textsuperscript{23}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Competing lifestyle factors\textsuperscript{23}</td>
</tr>
<tr>
<td>Follow-up</td>
<td>Lack of monitoring</td>
<td>Capacity constraint—leading to long waiting time for examinations with imaging modalities\textsuperscript{22}</td>
</tr>
<tr>
<td></td>
<td>Suboptimal patient outcomes</td>
<td>Lack of regularity in follow-up care\textsuperscript{22}</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Access to rheumatologists\textsuperscript{22}</td>
</tr>
</tbody>
</table>

*List not exhaustive.

HCP, healthcare professional; PCP, primary care professional; RA, rheumatoid arthritis.
Through secondary research, several guidelines and recommendations were identified for some comorbidities, but the presence of guidelines differed depending on the comorbidity (table 3).

The expert panel recommended to frame the findings of the study against three different stages of disease (table 4). These disease stages were used to segment and prioritise good practice interventions. In total, 18 good practice interventions were identified, through the centre visits, which span across the whole patient pathway. Interventions were highlighted as applicable to one, multiple or all disease stages (table 5).

All 18 interventions were reviewed by the expert panel and prioritised by ranking the highest impact on patient outcomes, against each stage of the disease: suspicion of RA; recent diagnosis of RA; established disease (while recognising that for each one of them, patients have distinct needs and requirements). The top three interventions for each disease stage are shown in table 6 and are described in the following sections.

**Interventions: diagnosing RA and early intervention**

**Rapid access to specialised care**

Patients with suspected RA must be seen rapidly to facilitate the effective prevention and management of disease progression and joint damage. Delays in diagnosis and referral from primary care mean patients can arrive at secondary centres with irreversible damage to their joints and other organs. It is vital that these patients are triaged rapidly into care and start treatment straight away. Rapid access to care may include online referral systems (patient self-enrollment) and access to diagnostic services (e.g., blood tests and ultrasound). Hospital Universitario La Paz has implemented an online direct communication system between the rheumatology department and primary care practitioners to allow two-way messaging system for rapid patient referrals within 24 hours. Several centres run rapid access clinics, such as the ‘clinically suspect arthralgia’ clinic for undifferentiated arthralgia at the Institute of Rheumatology in the Czech Republic, the pre-RA CCP clinic Leeds Hospital in the UK and the sophisticated triaging system at Hospital de Santa Maria which aims to increase accessibility for new patients and rapidly assess the presence and severity of RA. These services have reduced waiting times, improved the experience for patients and facilitated two-way communication between primary and secondary care.

**Early arthritis clinic**

Early treatment reduces overall disease progression and therefore prevents the accumulation of damage that causes irreversible joint deformities. Clinics are dedicated to ensuring timely clinical assessment and diagnosis of patients with suspected RA, including initial triage, to which patients can be referred by
primary care physicians (PCPs) and other specialists. The Institute of Rheumatology, Hospital de Santa Maria, Cliniques Universitaires Saint-Luc, Sint Maartenskliniek, Diakonhjemmet Hospital, Leeds Teaching Hospitals, Karolinska University Hospital, Hospital Universitario La Paz, ASST Gaetano Pini-CTO Institute and Rigshospitalet have all implemented early arthritis clinics leading to more patients being diagnosed quickly and starting early disease-modifying therapy combined with a treat-to-target strategy; with follow-up maintained in the early arthritis clinic until target is reached and/or directed to escalated treatment pathways and service as indicated.

### Enhanced communication with primary care

Often patients are not transitioned between different levels of the healthcare system in a timely manner, which can be due to several factors including lack of awareness of referral pathways and systems, suboptimal communication between specialists and PCPs, and sharing of up-to-date patient information. Enhanced communication strategies identified include a hotline for PCPs to access specialist advice within 48 hours.

### Table 3

<table>
<thead>
<tr>
<th>Comorbidity</th>
<th>Authors</th>
<th>Guidelines/recommendations</th>
</tr>
</thead>
</table>
| Cardiovascular disease | Regulatory bodies or disease associations | 1. 2016 Update EULAR recommendations for CVD risk management \cite{30}
2. 2016 EULAR points to consider for reporting, screening for and preventing selected comorbidities \cite{12}
3. 2019 European Society of Cardiology/European Atherosclerosis Society guidelines for the management of dyslipidaemias \cite{31}
4. Spanish Society of Rheumatology (2011). Update of the clinical practice guideline for the management of rheumatoid arthritis in Spain \cite{32}
5. Haute Autorite De Sante (2017): Clinical practice guidelines: Rheumatoid Arthritis \cite{33} |
| Pulmonary disease      | Academic literature | 1. Diagnosis of Idiopathic Pulmonary Fibrosis. An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline \cite{34} (associated rheumatological diseases are discussed from a pulmonary perspective)
2. 2019 Swedish Respiratory Society Guidelines for Idiopathic Pulmonary Fibrosis \cite{35} |
| Diabetes               | No existing guidelines or recommendations were found at a regional or national level for diabetes |
| Depression             | Existing recommendations/points to consider published by regulatory bodies or disease associations | 1. 2016 EULAR points to consider for reporting, screening for and preventing selected comorbidities \cite{12}
2. NICE (2009): Depression in adults with a chronic physical health problem: recognition and management \cite{36} |

ALAT, Latin American Thoracic Association; ATS, American Thoracic Society; CVD, cardiovascular disease; ERS, European Respiratory Society; JRS, Japanese Respiratory Society; NICE, National Institute for Health and Care Excellence; RA, rheumatoid arthritis.

### Table 4

<table>
<thead>
<tr>
<th>Disease stage</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suspicion of RA</td>
<td>Patients who have not yet been given a diagnosis but may have interacted with primary care or had a first contact with a rheumatologist, and who may have suspected RA (inflammatory manifestations not diagnosed)</td>
</tr>
<tr>
<td>Recent diagnosis of RA</td>
<td>Patients have been given a diagnosis of RA and usually have been started on treatment with regular follow-up</td>
</tr>
<tr>
<td>Established disease/structural damage</td>
<td>Patients have had a diagnosis of RA for several years or may have presented late (most likely from less developed healthcare economies). At this stage, treatments may have become progressively less effective at reducing inflammation and preventing further joint damage</td>
</tr>
</tbody>
</table>

RA, rheumatoid arthritis.
Table 5  Good practice interventions relevant to the three disease stages of patients with RA

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Definition</th>
<th>Suspected RA</th>
<th>Recently diagnosed with RA</th>
<th>Established RA/structural damage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rapid access to care</td>
<td>Fast-track access to care for patients with RA done via online referral form reviewed every 24 hours, hotline leading to appointments within 48 hours for diagnostic services including blood tests and joint imaging</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
<tr>
<td>Enhanced communication across wider care team</td>
<td>Availability of reliable communication channels (eg, emails, online forms) enabling easy dialogue between specialists and PCPs; and providing and coordinating education programmes to ensure the wider care team are kept up-to-date with developments in best practice care</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
<tr>
<td>Early arthritic clinic</td>
<td>Clinic dedicated to ensuring timely clinical assessment and diagnosis of patients with suspected RA</td>
<td>X</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Comprehensive comorbidity assessment</td>
<td>Comorbidity assessment in patient baseline assessment and follow-up of newly diagnosed patients</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
<tr>
<td>Tailored education to patients and family members</td>
<td>Programmes to increase understanding of diagnosis, treatment plans, and how to live with the disease that is sensitive to individual patient needs</td>
<td>X</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Role of the care coordinator</td>
<td>Care coordinator role to help to manage the burden of navigating contacts across multiple HCPs</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Dedicated comorbidity specialist HCP</td>
<td>Specific role or clinic to support the management of comorbidities in the context of RA</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Enabling self-management</td>
<td>Provision of tools and resources to patients to monitor and manage their RA and reduce dependence on healthcare services</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Enhanced therapy services</td>
<td>Additional care centred around non-physician-led management including care led by therapists to promote rehabilitation and enablement</td>
<td></td>
<td>X</td>
<td></td>
</tr>
<tr>
<td>Day clinic services</td>
<td>The coordination of services enabling provision of stacked outpatient appointments across specialities and disciplines over 1 day or session</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Virtual engagement with patients</td>
<td>Digital enablement of autonomy, self-management and empowerment by providing a channel of direct communication with attending physicians in addition to online access to education around the disease, networks and peer support</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Integrating patient registries into daily clinical practice</td>
<td>Employment of evidence-based practice, informed by clinical research and supported by systematic capture and monitoring of data, in order to improve the quality of clinical care delivery and promote evolution of care models</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Patient-centred care journey</td>
<td>Curation of care processes and physical environment that enables the patient to feel empowered and supported during their journey through care</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Effectively using the skill mix of the multidisciplinary care team</td>
<td>Enablement of non-physician HCPs to taking on greater responsibility in the assessment and management of patients</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Integrative and shared care solutions</td>
<td>Ensuring regular communication between all attending physicians and other HCPs as part of a holistic and integrated approach to care</td>
<td>X</td>
<td></td>
<td>X</td>
</tr>
</tbody>
</table>

Continued
linked information systems leading to collaboration between primary and secondary care, educational and training sessions. Hôpital Cochin implemented the ‘Hospital and City Rheumatology Network’ and Diakonhjemmet Hospital have a dedicated primary care coordinator. ASST Gaetano Pini-CTO Institute delivers education and training for PCPs in Italy regarding RA, developing a network with primary care. Enhanced communication facilitates improved integration of services and patients accessing specialist treatment in a timely manner.

**Interventions: management of comorbidities**

**Comprehensive comorbidity assessment**

Certain patients with RA are at increased risk of morbidity and mortality due to existence of comorbidities such as cardiovascular disease, ILD, diabetes and depression. Systems for cross-specialty care and follow-up can be fragmented. Centres may implement regular (typically 6-monthly) full comorbidity assessment as a standalone service or in conjunction with RA appointments. Hôpital Cochin has a Comorbidities Education in Rheumatoid Arthritis (COMEDRA) service which is a doctor-led programme on RA comorbidity management and supported by a programme coordinator, and Hospital Universitario La Paz has a systematic inflammatory osteoporosis screening. Comorbidity assessments may lead to reduced patient mortality due to comorbidity complications in patients with RA (eg, cardiovascular disease and ILD).

**Dedicated comorbidity specialist**

Comorbid diseases may not always be given appropriate focus and attention because rheumatologists are not specialists in these areas. To overcome this, centres may deploy an in-house comorbidity specialist for example, a cardiologist dedicated to the cardiovascular manifestations of rheumatic disorders. Diakonhjemmet Hospital has a preventive cardio-rheuma clinic run by a cardiologist employed in the Department of Rheumatology, and Hospital de Santa Maria has a dedicated psychologist for patients with RA. The input of comorbidity specialists may raise awareness regarding comorbidities among rheumatology HCPs. At the Karolinska University Hospital, for instance, pulmonary complications can be presented during discussion rounds with the participation of pulmonologists, rheumatologists and radiologists.

**Combined clinics**

To effectively and more efficiently manage comorbidities associated with RA, with minimum burden on patient hospital visits, centres may implement combined clinics with both rheumatologists and the respective comorbidity specialists. Hospital de Santa Maria, ASST Gaetano Pini-CTO Institute and Hospital Universitario La Paz have combined clinics with rheumatologists and pulmonologists to manage patients with RA with or at risk of ILD. Hospital de Santa Maria has a joint rheumatobstetrics clinic to manage the potential problems that

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**Table 5** Continued

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Definition</th>
<th>Suspected RA</th>
<th>Recently diagnosed with RA</th>
<th>Established RA/structural damage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Collaborating with PAGs</td>
<td>Increasing communication with local and regional patient advocacy groups through working group sessions, conference attendance, newsletters and patient liaisons</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
<tr>
<td>Developing care networks</td>
<td>Developing networks with community-based RA services to maintain high-quality ‘joined-up’ care past the point of discharge</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
<tr>
<td>Quality management programmes</td>
<td>A coordinated approach and a robust system for measuring, tracking and improving the quality of care</td>
<td>X</td>
<td>X</td>
<td>X</td>
</tr>
</tbody>
</table>

HCPs, healthcare professionals; PAGs, patient advocacy groups; PCPs, primary care professionals; RA, rheumatoid arthritis.

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**Table 6** Prioritised interventions per disease stage

<table>
<thead>
<tr>
<th>Disease stage</th>
<th>Prioritised interventions*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suspicion of RA</td>
<td>1. Rapid access to care.</td>
</tr>
<tr>
<td></td>
<td>2. Enhanced communication with primary care.</td>
</tr>
<tr>
<td></td>
<td>3. Early arthritis clinic.</td>
</tr>
<tr>
<td>Recent diagnosis of RA</td>
<td>1. Enabling self-management.</td>
</tr>
<tr>
<td></td>
<td>2. Early arthritis clinic.</td>
</tr>
<tr>
<td>Established disease/structural damage</td>
<td>1. Dedicated comorbidity specialist.</td>
</tr>
<tr>
<td></td>
<td>2. Integrating patient registries into daily clinical practice.</td>
</tr>
<tr>
<td></td>
<td>3. Enabling self-management.</td>
</tr>
</tbody>
</table>

*The top three priority interventions are listed.

RA, rheumatoid arthritis.
can arise during pregnancy in rheumatological patients. Leeds Teaching Hospitals has several joint comorbidity services with dermatology, gastroenterology, immunology, neurology, respiratory, renal and obstetrics. These combined or stand-alone clinics may cover the wider aspects of RA management, including lifestyle factors. For example, Rigshospitalet in Copenhagen offers a sleep clinic and smoking cessation clinic for patients with RA helping patients to live healthier lives. Joint clinics may lead to a greater focus on comorbidities and involvement of specialist comorbidity expertise leading to more effective decisions and better patient outcomes.

Interventions: encouraging patient self-management and empowering health professionals

Effectively using the skill mix of the multidisciplinary care team

The findings of this study indicated an increasing number of patients with RA requiring comprehensive care and management. Rheumatologists may not have enough time, capacity or specific skills to address these important problems outside diagnosis, inflammatory assessment and pharmacological treatment. Other health professionals (eg, nurses, physio- and occupational therapists, care coordinators and administrative staff) often have the competencies that are well suited for some of the tasks required to ensure best practice. By facilitating a more active role of these HCPs in the management of patients, they are able to bring a different dimension which adds to a holistic and patient-centred approach to care. For example, at Sint Maartenskliniek, the pharmacist is a key member of the care team, delivering medication support for patients. At Diakonhjemmet Hospital, Cliniques Universitaires Saint-Luc, Rigshospitalet and Leeds Teaching Hospitals, the enhanced role of the nurse helps support rheumatologists in care delivery (eg, in Leeds the nurse-delivered services at the centre include outpatient clinics and a day care unit where rheumatology nurses manage the delivery of biologics to patients). Such activities can be done in various ways between countries and health systems as shown in online supplementary table S1.

Enabling self-management

Patients may not always have long enough consultation sessions with rheumatologists and rheumatology nurses and other HCPs to fully educate themselves about the impact of the disease and how to manage it. As part of the COMEDRA initiative at Hôpital Cochin, patients are taught to self-assess and self-monitor their disease activity and more importantly to feel comfortable to undertake this. At Diakonhjemmet Hospital a ‘learning and coping’ centre is run by a multidisciplinary team led by an experienced nurse. Cliniques Universitaires Saint-Luc enables self-management through personalising care delivered to patients. All these interventions have been put in place not only to improve adherence to treatment but also to help patients manage their lives and not just their disease activity; it is about empowerment and feeling in control.

Interventions: care delivery process optimisation

Integrating patient registries into daily clinical practice

Several registries have been implemented at different centres to collect longitudinal real-life data, for example, at the Institute of Rheumatology, Swiss Clinical Quality Management in Rheumatic Diseases at the Hôpitaux Universitaires de Genève; Swedish Rheumatology Quality Registry at the Karolinska University Hospital and the Centre for Rheumatology in Stockholm; Danish Database for Biological Therapies at Rigshospitalet; and the Norwegian Disease-Modifying Antirheumatic Drug study at Diakonhjemmet Hospital. Hospital de Santa Maria uses the Portuguese Reuma.pt registry to support research and clinical practice and Leeds Teaching Hospitals has a focus on research to support clinical practice. Registries may inform real-life results of clinical management from different participating centres and thereby also provide information which may stimulate the different centres to improve their quality of care. Further, linkages between registries may help the understanding of the relationships between RA and comorbidities. Karolinska University Hospital has had a strong focus on this type of research.

Patient-centred care journey

Patients need to be satisfied and confident with their care to be fully engaged with their care. Patient-centred care can be defined as when patients are empowered throughout the duration of their journey whether it relates to the physical environment or the experience of the services received. At the Sint Maartenskliniek, patient-centred design reduces time spent travelling between each care station, and provides more time with the HCPs; all of which improves the overall patient experience. This intervention enables HCPs to improve patient engagement and helps empower and support patients throughout their care journey.

Key considerations in selected comorbidity care for patients with RA

The visits to study centres (primary research) and the input from the expert panel provided several key considerations for the care of comorbidities (cardiology, pulmonology, diabetes and depression) in patients with RA.

Cardiovascular disease (CVD): Key elements to consider for improvement of the quality of care of CVD in patients with RA are listed as follows:

- Communication about CVD risk factors between medical professionals as well as screening of CVD risk factors. It was noted that CVD risk prediction in patients with RA using risk calculators developed for the general population is generally inaccurate in the prediction of future CVD. Modifying the predicted
may also not necessarily reclassify patients to their appropriate risk class. However, adding information on carotid plaque/atherosclerosis may reclassify patients to a more appropriate risk class in up to 30–60% of cases.47

**Organisation and responsibility of CVD risk management through cardiologists working alongside or within rheumatology departments, for example, preventive cardio-rheumatology clinics. It was noted that lipid-lowering, antihypertensive therapies and non-pharmacological recommendation (eg, focusing on lifestyle factors) can be implemented safely and effectively with recommended goal achievement over approximately three consultations in 80–90% of the patients.48 Systemic inflammation or lipid levels at baseline or antirheumatic medication do not have an impact on statin dose needed to obtain low-density lipoprotein goals.49**

**Availability of data regarding CVD risk management.**

**Pulmonary disease:** Interventions and results from the study highlight the importance of early screening and detection. This can include chest X-ray, CT and functional assessment with spirometry and six-minute walking test (6MWT). Regular follow-up in outpatient clinics are also important. Management of adverse respiratory events, side effects from medications and smoking cessation advice should be included. The expert panel emphasised the importance of the fieldwork findings from the visits of study centres and especially the collaboration between rheumatologists and pulmonologists. ILD should be diagnosed in multidisciplinary discussions where the presence of rheumatologists is encouraged. These conferences can also enable pulmonologists to consider the possibility of an underlying systemic disease including RA in patients with ILD.

**Diabetes:** Study results highlight the importance of screening for and detecting diabetes, coordination of care between RA and diabetologists and regular follow-ups. The expert panel advised that the effective treatment of RA with targeted therapies can improve long-term glycaemic control in patients with diabetes and RA.50

**Depression:** The results of the study highlight the importance of screening and detection, periodic follow-up, guided self-help and targeted depression history. These can be achieved through dedicated psychological and psychiatric services as well as focusing on impacts of depression for example, sleep disturbance and socioeconomic consequences. The expert panel recommended a strong focus on prevention through paying attention to individual factors (eg, sleep, physical activity, interest in undertaking any activities), encouraging a healthy lifestyle and treatment of depression (moderate–severe) if guided self-help and exercise fail (eg, treatment with cognitive-behavioural treatment or psychotherapy). The expert panel noted that the ideal model to achieve the aforementioned activities would be the integration of psychology resources as part of the wider multidisciplinary team although the experts also recognised local constrains.

**DISCUSSION**

Challenges in the care of RA and associated comorbidities are well documented, and the results of the interviews confirmed the findings from the literature review. However, the study highlighted the different types of challenges experienced by the centres. In order to improve patient care, it is recommended that centres identify and prioritise interventions most suited to their population, patient pathways and current challenges in delivery of care. The need for comprehensive and consensus-approved guidelines and recommendations to improve the care of patients with RA and the several associated comorbidities was clear in the findings. The experts acknowledged that this requires further exploration and expert input with more robust methodology in order to develop guidelines and recommendations.

It was recognised that there were limitations to the study methodology and several actions were taken to mitigate these. One drawback was the lack of patient interviews at each individual centre which limited the patient perspective on challenges in the patient pathways and intervention benefits (eg, benefits of self-management). The role of the very experienced patient representative on the expert committee (NB) helped to overcome this by providing a collective patient perspective, on behalf of patients. Second, the inclusion of one centre per country limited the findings with respect to the representativeness of the wider healthcare system in which they reside. This was taken into consideration when conducting research which did not solely focus on centres themselves but rather their entire ecosystems; including the various institutions they collaborate with, whether formally or informally. However, the goal of this study was rather to find centres that had developed good quality of care practices which could serve as models if implemented in other centres.

Third, centre selection was biased towards larger teaching centres. This was mitigated by gathering and documenting interviewees’ perspectives on how to replicate interventions in smaller and less resource-rich centres, although it was accepted that this was incomplete.

The project group decided to focus on four main comorbidities; CVD, ILD, diabetes and depression. The literature review and also centre visits highlighted other comorbidities of importance. For example, it has been shown that osteoporosis is twice as common in patients with RA compared with controls of the same age51 52 and that fractures are also more common.53 Services to support primary and secondary prevention are of value in improving patient morbidity.

Infections are other important comorbidities which can partly be related to the disease, partly to treatment. Several studies have described treatment-related...
occurrence of infections. Importantly, EULAR has recently published updated recommendations on vaccinations, and editorials have addressed the important aspects of hepatitis and tuberculosis in this population.

Guidelines and/or recommendations on periodontitis, malignancy and gastrointestinal disorders were not assessed, but it was noted that these conditions lead to poor outcomes for patients with RA.

We believe that the most important message from this paper is the description of examples of good clinical practices and care models. One of the main findings was the success of early arthritis clinics. Impressively, many clinics reported to see patients within 1–2 days since they focused on the importance of early diagnosis and treatment. This may appear more aspirational for the majority of departments with limitations in resource and funding to deliver such a service. Nevertheless, these examples illustrate the importance of prioritising early RA pathways of care within rheumatology services to optimise efficient diagnosis and management. However, we also recognise that not all recommendations are feasible in every setting, since different clinical structures and workforce challenges may limit the implementation of for example multidisciplinary care approaches. However, the recommendation may hopefully serve as a benchmark to support efforts to obtain more resources to enhance quality of care.

The strength and impact of this paper would have been improved if data had been available on longitudinal clinical outcomes to demonstrate the benefit of the good practice recommendations shown in table 5. Long-term clinical data are available for some of the recommendations and are included in the reference list.

It is envisaged that rheumatology departments could use this study as an initial platform to draw upon models appropriate to local services and pathways for adaptation and implementation. Interested centres are welcome to contact the centres who took part in this study for further knowledge transfer.

The panel of experts recognise that data capture should also be done in the community. Community-based physicians could propose a list of items which, if accepted by rheumatologists, would lead to harmonisation of standards of care across all care settings along the patient journey.

The next steps for the project group are to do additional work on dissemination of our findings. A comprehensive study report has been published on the website of the project accompanied by an appendix with single-centre reports (see Previous publications at the end). More importantly, we also plan to support the implementation of the examples of models of good clinical care into other settings.

In conclusion, we identified 18 models of good clinical practice which are linked to three different disease stages. Further work is needed to explore the ability to implement each of the interventions (eg, the development of tools and manuals detailing their implementation) and the results achieved.

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Acknowledgements The authors thank Sanofi Genzyme who funded this study. The authors are also grateful to Olivia Bryan, Emma Hanson, Jessica O’Neil, Zoe Phillips and Thishi Surendranathan from the KPMG team for their input and contribution to the work.

Contributors All authors contributed to the data collection, to the interpretation and analyses of the results and to the writing of the manuscript. All authors have read and approved the final version of the manuscript submitted on 29 May 2020.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests TKK has received fees for speaking and/or consulting from AbbVie, Biogen, Celltrion, Egsis, Eli Lilly, Histma, MSD, Mylan, Novartis, Oktal, Orion Pharma, Pfizer, Roche, Sandoz, Sanofi and UCB and received research funding to Diakonhjemmet Hospital from AbbVie, BMS, MSD, Pfizer, Roche and UCB. AB has received Grant/research support, fees for consultancies or as a speaker for AbbVie, Pfizer, Novartis, BMS, Nordic, Sanofi-Genzyme, Sandoz, Lilly, UCB and Roche. NB has received fees for speaking and/or consulting from Amgen, Eli Lilly, Grunenthal, GSK, Heart Valve Voice, Janssen, Roche, Sanofi Genzyme and Sanofi Regeron. MHB has received fees for speaking and/or consulting from AbbVie, AstraZeneca, Bristol-Myers-Squibb, Chugai, Eli Lilly, Merck-Serono, Pfizer, Roche, Sandoz and Sanofi, and research funding to University of Leeds from Pfizer, Roche and UCB. PD has received fees for speaking and/or consulting from Bristol-Myers-Squibb, Celltrion, Lilly and Sanofi Genzyme. EGF has received fees for speaking and/or consulting from AbbVie, BMS, Eli Lilly, Gilead, MSD, Novartis, Pfizer, Roche, Sanofi Genzyme and UCB. IG-B has received fees for speaking and/or consulting from Amgen, Akcea, Sanofi Genzyme and Regeneron. FvdH has received fees for consulting from AbbVie, Biogen, Celltrion, Roche, Sanofi Genzyme, Pfizer and Mundipharma. AK has received fees for speaking and/or consulting from UCB, Bristol-Myers-Squibb, MSD, Amgen, Abbvie, Pfizer, Novartis and Sanofi. IG-B and FvdH have received fees from the National Institute for Health Research (NIHR) as part of the NIHR Biomedical Research Centre funding. PdH has received fees for speaking and/or consulting from AbbVie, BMS, GSK and UCB. MØ has received fees for speaking and/or consulting from AbbVie, Bristol-Myers-Squibb, Boehringer-Ingelheim, Celgene, Eli Lilly, Hospira, Janssen, Merk, Novartis, Novo, Orion, Pfizer, Regeneron, Roche, Sanofi and UCB. KP has received fees for speaking and/or consulting from AbbVie.
Rheumatoid arthritis

Amgen, Biogen, Bristol-Myers-Squibb, Egis, MSD and UCB. JP-P has received fees for speaking and/or consulting from AbbVie, MSD, Pfizer, Roche and Tecnome.de. AGS has received fees for speaking and/or consulting from AbbVie, Novartis, Sanofi and Bayer and have an unrestricted research collaboration with Eli Lilly which includes transfer of funds to Diakonhemmet Hospital from Eli Lilly. MS has received research grants from Boehringer Ingelheim and Roche, speaker fees/consultancy from AstraZeneca, Boehringer Ingelheim, GlaxoSmithKline, Novartis, Mundipharma, Sandzo and Roche. MD has received fees for speaking and/or consulting from AbbVie, Biogen, Eli Lilly, BMS, MSD, Novartis, Pfizer, Roche, Sandzo, Sanofi and UCB and his department has received research grants from AbbVie, BMS, MSD, Pfizer, Roche, Lilly, Janssen, Novartis and UCB.

Patient consent for publication Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement Data are available in a public, open-access repository. All data relevant to the study are included in the article or uploaded as online supplementary information.


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