Original research

National registry for patients with inflammatory rheumatic diseases (IRD) infected with SARS-CoV-2 in Germany (ReCoVery): a valuable mean to gain rapid and reliable knowledge of the clinical course of SARS-CoV-2 infections in patients with IRD

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ABSTRACT

Objectives Patients with inflammatory rheumatic diseases (IRD) infected with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) may be at risk to develop a severe course of COVID-19. The influence of immunomodulating drugs on the course of COVID-19 is unknown. To gather knowledge about SARS-CoV-2 infections in patients with IRD, we established a registry shortly after the beginning of the pandemic in Germany.

Methods Using an online questionnaire (www.COVID19-rheuma.de), a nationwide database was launched on 30 March 2020, with appropriate ethical and data protection approval to collect data of patients with IRD infected with SARS-CoV-2. In this registry, key clinical and epidemiological parameters—for example, diagnosis of IRD, antirheumatic therapies, comorbidities and course of the infection—are documented.

Results Until 25 April 2020, data from 104 patients with IRD infected with SARS-CoV-2 were reported (40 males; 63 females; 1 diverse). Most of them (45%) were diagnosed with rheumatoid arthritis, 59% had one or more comorbidities and 42% were treated with biological disease-modifying antirheumatic drugs. Hospitalisation was reported in 32% of the patients. Two-thirds of the patients already recovered. Unfortunately, 6 patients had a fatal course.

Conclusions In a short time, a national registry for SARS-CoV2-infected patients with IRD was established. Within 4 weeks, 104 cases were documented. The registry enables to generate data rapidly in this emerging situation and to gain a better understanding of the course of SARS-CoV2-infection in patients with IRD, with a distinct focus on their immunomodulatory therapies. This knowledge is valuable for timely information of physicians and patients with IRD, and shall also serve for the development of guidance for the management of patients with IRD during this pandemic.

INTRODUCTION

Since 12 March 2020, the WHO has declared the outbreak of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) disease
2019 (COVID-19) a pandemic. The main symptoms of respiratory infection include fever and cough.\textsuperscript{1,2} Eighty-one per cent of the patients show a mild course, 14\% are seriously affected and 5\% of the patients are critically ill.\textsuperscript{2} On 25 April 2020, the Johns Hopkins University reported 2,954,106 confirmed cases worldwide.\textsuperscript{3} The mortality rate at that time point was 6.95\% (n=205,398).\textsuperscript{3} The presence and the number of comorbidities (eg, arterial hypertension (AHT), coronary heart disease), age (immunosenescence) and lifestyle factors (eg, smoking) appear to have deteriorating effects on the course of the infection.\textsuperscript{2} In this situation, patients with inflammatory rheumatic diseases (IRD) may face a particular risk as their disease, especially when clinically active, and their immunomodulatory treatment may impact the course of COVID-19 infection. However, firm knowledge of the course of SARS-CoV-2 infection in patients with IRD is missing, and therefore, evidence-based recommendations for the management of COVID-19 in patients with rheumatic disorders and anti-rheumatic treatments are lacking.

The German Society for Rheumatology (DGRh) developed at an early-stage first concise recommendations for the management of patients with IRD during the COVID-19 pandemic.\textsuperscript{4} Interruption or reduction of immunomodulators is not recommended as this might result in relapses, which would increase the risk of infections, and in addition may lead to the necessity of an even riskier treatment (ie, additional glucocorticoids (GC)).\textsuperscript{5} In a cross-sectional multicentric study, patients with IRD were asked anonymously in the first weeks of the pandemic on their opinion of their immunomodulating therapy. Over 90\% of the patients followed the recommendation of the rheumatologists to continue the antirheumatic therapy, and only a small percentage of patients terminated the therapy on their own.\textsuperscript{6} Whether first recommendations based on responsible care and expert’s opinion is indeed the best option for the management of patients with IRD urgently needs to be validated.

Incidence and course of COVID-19, including lethal outcomes, vary considerably in different cohorts according to pre-existing conditions and healthcare systems. Investigation of special disease groups may contribute to a better understanding of the role of the immune system regarding the risk to get infected or to develop a more severe course of COVID-19. Therefore, patients with IRD, who are treated with different types, combinations and dosages of immunomodulatory therapies represent an interesting population to collect data regarding SARS-CoV-2 infection.

Registries with a large number of case reports are required to answer the questions, whether antirheumatic drugs increase or decrease the risk for a severe course of SARS-CoV-2 infection. As necessary data cannot be extracted from clinical charts or health insurance records, the DGRh and the Justus-Liebig University Gießen decided to establish a web-based registry, which allows a rapid and timely collection of IRD cases with confirmed SARS-CoV-2 infections in Germany to analyse the clinical course of SARS-CoV-2 infections in patients with IRD and to develop guidance for the management of patients with IRD during the COVID-19 pandemic.

**PATIENTS AND METHODS**

On 18 March 2020 the COVID-19 Registry Task Force was founded by the DGRh consisting of rheumatologists, epidemiologists and information technology specialists. In cooperation with biostatisticians and data-protection specialists to ensure mutual understanding of research objectives and scientifically and legally appropriate data collection, a database-driven online questionnaire was developed and has been launched on 30 March 2020.

Following are the key questions of the registry:

- What is the course of COVID-19 in patients with IRD?
- Does geographical location/different access to healthcare within Germany affect the course?
- Does a specific immunomodulatory treatment ameliorate or deteriorate the course of COVID-19?
- Which other factors are associated with a poor outcome of COVID-19 in patients with IRD?

Patients with a known IRD and positive testing for SARS-CoV-2 were included by their treating rheumatologists after registration. Patients had to have positive PCR-swabs for SARS-CoV-2 for inclusion in our registry. In contrast to other registries, presumptive diagnosis of COVID-19 based on suggestive symptoms, X-ray or CT findings or on other ‘unknown’ findings was not included.

The database includes, for example, federal state, age, weight, height, detailed rheumatological diagnosis, comorbidities, global disease activity antirheumatic medication at time of study and changes due to the infection. In addition, the course and outcome of the SARS-CoV-2 infection are also key parameters. Missing data on diagnosis, outcome and therapies can be queried by directly contacting the participating physicians. Periodic critical evaluation of the registry is carried out by the task force to ensure that the objectives are being met.

The disease activity from the last rheumatological visit was reported and divided into four groups: remission, low disease activity, moderate disease activity and high disease activity. This classification and approach are concordant to the EULAR/Global Rheumatology Alliance (GRA) database. Specific disease activity scores, for example, DAS-28, are not included in the survey.

Data entered in an electronic case report form with the URL https://www.COVID19-rheuma.de/ are directly stored into an SQL-database on a dedicated server located in Germany and certified according to DIN ISO/IEC 27001 using encryption and secure communication protocols (SSL/TLS and HTTPS). Data entered in these forms are checked for plausibility immediately. Web-forms use dynamic menus and subquestions. Data allowing for identification of individual patients are
omitted, and reidentification is only possible via local files in the respective rheumatological unit.

In addition, patients with IRD affected by SARS-CoV-2 can contact a rheumatologist at the coordination site of the project, who collects the respective data sets via a telephone-interview (after informed consent of the patient) entering the pseudonymised data also directly into the same web-form/database. The lockdown appointments of in- and outpatients were reduced remarkably to spare capacity for the treatment of patients with severe COVID-19 infections and to minimise patients and physician’s infection risks during outpatients’ appointment. Especially, visits of patients which showed controlled disease activity were skipped or postponed. However, to give these patients the opportunity to report their case if they were tested positive for SARS-CoV-2, the physician telephone interview was set up. However, these data entries will be flagged and can be analysed separately from those directly entered by their treating physician.

After the study was approved by the ethics committee of the Justus-Liebig-University Giessen (#52-50) and registered (EuDRAC 2020-001958-21), the web-based questionnaire was announced to German rheumatologists on 30 March 2020. Later, the German database was, in part, adjusted to the EULAR COVID-19 database when the latter was launched (https://www.eular.org/eular_COVID-19_database.cfm) to facilitate the transfer of German anonymised data into the European database in the future.

Participating centres consist of academic and non-academic rheumatology clinics, and private practices in Germany. Most of them have been informed directly using established dissemination channels of the DGRh. Announcements of the project have been posted and are being regularly updated on the website of the DGRh, the homepage of the ‘www.COVID19-rheuma.de’ registry, via social media channels, and print media. The registry is also fully supported by the Professional Association of German Rheumatologists ‘Berufsverband Deutscher Rheumatologen’, and by the national patient organisation ‘Deutsche Rheuma-Liga’, its state representatives, the Association of Rheumatology Clinics ‘Verband der Rheumatologischen Akutkliniken e.V’ and several other disease-specific patient associations.

The completed data were reviewed and queried in case of uncertainties. Analysis was performed descriptively using SPSS Statistics. Median was calculated for age and body mass index (BMI) of the fatal courses. Data in figures are shown in percentages using GraphPad Prism 6 (GraphPad Software).

RESULTS

Between 30 March (first patient in) and 25 April, 104 patients with IRD and a SARS-CoV-2 infection were documented in the database by rheumatologists. The development of recruitment during the first 4 weeks is depicted in figure 1. To date, 138 rheumatologists have registered to take part in the project and 71 have already documented at least one patient. Entering a patient case took on average 5 min. Each data entry is logged by a time stamp allowing calculations of the time used for the documentation procedure up to the time required for single items in the survey.

The 104 patients reported so far included 40 males, 63 females and 1 diverse, the age range was between 23 and 87 years (median age 56 years; figure 2A). Most patients...
with COVID-19 (24%) were between 50 and 59 years old. About two-thirds of the patients were reported from private practices, 32% from hospitals and 6% of the patients had a telephone interview by a rheumatologist at the coordination site. In relation to population numbers, most cases have been notified from Hamburg, Baden-Württemberg, Berlin, North Rhine-Westphalia and Rhineland Palatinate (table 1). The presence of antibodies was not included in our registry, which is comparable to the EULAR and GRA COVID-19 database.

Regarding diagnosis, 45% of the patients with IRD had rheumatoid arthritis, 18% were diagnosed with psoriatic arthritis, 10% with ankylosing spondylitis and 5% systemic sclerosis. The proportion of other IRD was below 5% (figure 2B).

Prior to the SARS-CoV-infection, 34% of the patients were receiving conventional synthetic disease-modifying antirheumatic drugs (csDMARDs), 42% were on bDMARDs, 24% in monotherapy and 18% in combination with csDMARDs. Forty-three per cent were treated with GC, 7% without any other immunomodulation and 8% of the patients did not receive DMARDs or GC (figure 2C).

In 59% of the cases, relevant comorbidities were reported. This included AHT in 37% of the patients, other relevant comorbidities in 15%, cardiovascular...
One-third (32%) of the patients was hospitalised, in 70% (23/33) of these cases, oxygen supply was necessary, 15% (5/33) received non-invasive and 24% (8/33) invasive ventilation. Unfortunately, six patients (6%) died in the context of COVID-19 (figure 3B), three male and three female patients (table 2). The median age in this group was 71 years (range 59–80 years), the median BMI 27.8 kg/m² (range 23.9–40.6 kg/m²). All patients had at least AHT and/or other CVD (table 2). All deceased patients needed to be ventilated invasively. Five of the six patients were treated with low-dose GC (≤7.5 mg/day), which was not interrupted due to the infection in four cases. Only two of eight patients recovered after invasive ventilation.

Hospitalised patients were older than non-hospitalised patients (median age 69 vs 52 years). Even though women represented 62% of all registered patients, the hospitalisation rate was equal in both genders (48% male vs 52% female). More hospitalised patients were on GC (64% vs 34%), but less were treated with bDMARD (33% vs 48%). In addition, more comorbidities (>2) were documented in hospitalised patients (48% vs 21%, figure 4).

At the last rheumatological visit, 37% (38/104) of the patients were in remission, 17% (18/104) patients were reported to have low disease activity, 13% (13/104) moderate and 2% (2/104) high disease activity. In 29% (30/104), disease activity was not reported. Interestingly, the two cases with reported high disease activity developed a severe course of COVID-19 leading to hospitalisation. Comparing remission (33% vs 38%) and moderate (9% vs 14%) disease activity, the proportion was similar in the hospitalised versus non-hospitalised groups, and different in patients with low disease activity (27% vs 13%).

### DISCUSSION

Patients affected by IRD are at an increased overall risk of infection compared to the general population. However, the infection risk varies and is highly dependent on the type and activity of the autoimmune disease, on comorbidities and on the intensity of the immunosuppressive/immunomodulatory treatment. Most patients with IRD are treated with GC, csDMARDs, bDMARDs and tsDMARDs on a regular basis. Especially, GC increase the risk of serious infection in a dose-dependent manner. Moreover, treatment with DMARDs can be associated with infectious complications. Most frequently, these are of bacterial origin, but also certain viral infections like Herpes zoster may complicate the course of many anti-rheumatic therapies. Treatment with tumour necrosis factor (TNF) inhibitors is associated with an increased risk of serious infections at the beginning of treatment, but when effective, the risk decreases due to better functional capacity and decreased use of GC.

Currently, there is no evidence, whether and to what extent patients with IRD are at an increased risk for COVID-19 and if antirheumatic treatment, especially GC and DMARDs, are harmful to patients in the context of COVID-19 and if antirheumatic treatment, especially GC and DMARDs, are harmful to patients in the context of COVID-19 and if antirheumatic treatment, especially GC and DMARDs, are harmful to patients in the context.
Figure 3  Symptoms and courses of COVID-19 infection (in %). (A) Distribution of the symptoms of COVID-19 infection (in %): 86% of the affected patients had more than two symptoms, 69% reported cough, 59% fever, 42% fatigue, 36% headache, 33% myalgia, 32% dyspnea, 26% loss of odour, 25% loss of taste, 23% rhinitis, 16% loss of appetite, 15% vertigo, 15% diarrhoea, 13% expectoration, 13% other symptoms, 7% had no symptoms, 3% abdominal pain and 2% vomit. (B) Distribution of the course of COVID-19 infection (in %): 60% of the patients already recovered, 32% of the patients needed to be hospitalised, 22% of the patients were treated with oxygen (5% non-invasive ventilation (NIV), 8% invasive ventilation (IV)). Six deadly courses were already reported.
### Table 2  Characteristics of fatalities

<table>
<thead>
<tr>
<th>IRD</th>
<th>PsA</th>
<th>RA</th>
<th>RA</th>
<th>RA</th>
<th>PsA</th>
<th>RA</th>
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</thead>
<tbody>
<tr>
<td>Age (years)</td>
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<td>70</td>
<td>80</td>
<td>80</td>
<td>59</td>
<td>72</td>
</tr>
<tr>
<td>Gender</td>
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<td>M</td>
<td>F</td>
<td>F</td>
<td>M</td>
<td>F</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>40.6</td>
<td>27.8</td>
<td>28.7</td>
<td>23.9</td>
<td>27.8</td>
<td>26.6</td>
</tr>
<tr>
<td>Antirheumatic therapy</td>
<td>GC</td>
<td>SSZ</td>
<td>GC, MTX</td>
<td>GC, MTX, RTX</td>
<td>GC, SSZ</td>
<td>GC, MTX, ABC</td>
</tr>
<tr>
<td>Comorbidities</td>
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<td>AHT</td>
<td>CVD, AHT, COPD</td>
<td>AHT, Osteoporosis</td>
<td>CVD, AHT, Other</td>
<td>AHT, COPD</td>
</tr>
<tr>
<td>Disease duration (COVID-19)</td>
<td>6 days</td>
<td>8 days</td>
<td>14 days</td>
<td>21 days</td>
<td>18 days</td>
<td>20 days</td>
</tr>
<tr>
<td>Symptoms (COVID-19)</td>
<td>Fever</td>
<td>Dyspnea</td>
<td>Fever</td>
<td>Dyspnea</td>
<td>Fever</td>
<td>Cough</td>
</tr>
<tr>
<td>Chronic obstructive pulmonary disease (COPD)</td>
<td>Chest pain</td>
<td>Fever</td>
<td>Cough</td>
<td>Dyspnea</td>
<td>Vertigo</td>
<td>Fatigue</td>
</tr>
<tr>
<td>Invasive ventilation</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>Interruption DMARD</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes, MTX</td>
<td>Yes, MTX and RTX</td>
<td>No</td>
<td>Yes, ABC and MTX</td>
</tr>
</tbody>
</table>

ABC, abatacept; AHT, arterial hypertension; COPD, chronic obstructive pulmonary disease; CVD, cardiovascular diseases; DMARD, disease-modifying antirheumatic drugs; F, female; GC, glucocorticosteroids; IRD, inflammatory rheumatic diseases; M, male; MTX, methotrexate; PsA, psoriatic arthritis; RA, rheumatoid arthritis; RTX, rituximab; SSZ, sulphasalazine.

**Figure 4** Distribution between hospitalised and non-hospitalised patients. Median age in group of hospitalised patients (HOSP) was 69 years and in non-hospitalised (NON-HOSP) was 52 years. More male patients were hospitalised (48%) compared to NON-HOSP (34%). HOSP were treated in 64% of the cases with GC, NON-HOSP in 34%. Thirty-three per cent of HOSP received bDMARDs compared to 48% of NON-HOSP. In HOSP, more comorbidities were observed (48%) compared to NON-HOSP (21%). bDMARD, biological disease-modifying antirheumatic drugs; Comorb., comorbidities; GC, glucocorticosteroids.
Taken together, as a first result of our project, it could be shown that establishing an online-registry of patients with IRD and COVID-19 is feasible in short time and allows for rapid collection of possibly relevant data in the context of the SARS-CoV-2 pandemic. As in the general population, also in our study, male gender might be a potential risk factor for COVID-19 in patients with IRD, and as in the global register, treatment with GC seemed to be a disease-related risk factor in patients with IRD. With recruitment of more patients in the near future the trend of lower hospitalisation rates in patients with IRD treated with bDMARDs compared to those treated with csDMARDs will be further investigated. It is also worthwhile to compare data from Germany with those from other countries in and outside the European Union.

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Contributors RH, UM-L, TS, HS-K and CS performed the study design. RH performed the research, analysed, and interpreted the data. RH, UM-L, TS, HS-K and CS wrote the manuscript. RH, UM-L, TS, BFH, AK, H-ML, ACR, JGR, AS, REV, HS-K and CS performed physician recruitment. All authors contributed to preparation of the project, and read and approved the final manuscript.

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