# Challenges of Egyptian patients with systemic lupus erythematosus during the COVID-19 pandemic

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#### Abstract

**Objectives**: The coronavirus disease 2019 (COVID-19) pandemic in Egypt is part of the worldwide COVID-19 pandemic that has contributed to substantial deterioration of healthcare systems. The aim of this study was to assess the challenges faced by Egyptian systemic lupus erythematosus (SLE) patients during the COVID-19 pandemic.

**Material and methods**: This questionnaire-based study was carried out on 200 patients with SLE from Egypt. The questionnaire provided covered socioeconomic status, lupus disease data, information about COVID-19 infection, and medical and family history of COVID-19 infection.

**Results**: The mean age of the participants was 30.1 ±8.4 years. 140/200 (70%) of the participants reported difficulty in obtaining medications during the COVID-19 pandemic, particularly antimalarials (60%). The lupus disease condition became worse because of the drug shortage in half of the participants. Wearing protective masks (74%) and using disinfectants of the hands several times per day (67%) were the most reported used measures. Forty patients (20%) had to stop or reduce taking nonsteroidal anti-inflammatory drugs while 10 patients (5%) had to start taking antimalarials as a prophylaxis against COVID-19 infection. Among those who needed hospitalization, the main cause was lupus activity, and most of them (71%) experienced difficulty in hospital admission. Thirty-two patients (16%) had confirmed COVID-19 infection. About half of them had lupus flare and had to change the medications used for treatment of lupus.

**Conclusions**: The current COVID-19 pandemic has a negative impact on the healthcare provided to SLE patients in Egypt. Patients with SLE faced a shortage of their medications, especially antimalarials, and difficulty in hospital admission.

Key words: systemic lupus erythematosus, COVID-19, healthcare.

## Introduction

The COVID-19 infection has rapidly escalated into a global health problem which started in 2019 in China and continues to spread in most countries despite control measures [1]. The COVID-19 pandemic has put an unexpected burden on healthcare systems and even threatens to overwhelm them [2].

The problem of non-COVID patients is more complex in developing countries where most patients are dependent on government hospitals [2]. In an earlier report, Egypt had the highest risk of COVID-19 infection importation among African countries [3].

Patients with autoimmune diseases are at a higher risk of COVID-19 infection [4]. Additionally, there are many challenges faced by these patients during the COVID-19 pandemic [5]. Systemic lupus erythematosus (SLE) is a chronic autoimmune disease characterized by aberrant activity of the immune system with multiple clinical manifestations [6].

Infection is one of the most common causes of morbidity and mortality in SLE [7]. High dose glucocor-

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ticosteroids (GCs) and immunosuppressive agents are well-known risk factors for infections in addition to impaired cellular and humoral immunity among these patients [7].

To the best of our knowledge, no previous studies have evaluated the effect of the COVID-19 pandemic on Egyptian SLE patients' health status from the standpoint of the patients. The aim of this study was to assess the tremendous challenges faced by Egyptian SLE patients during the COVID-19 pandemic in terms of drug shortage, hospitalization, and COVID-19 infection.

## Material and methods

This cross-sectional study was carried out on 200 SLE patients aged > 18 years from Egypt. The study was conducted between June 2020 and December 2020. The patients were recruited from our rheumatology and immunology unit (inpatient and outpatient). Patients who had any other rheumatic, neurological or musculo-skeletal diseases were excluded from the start.

The study protocol was approved by the Institutional Research Board of the Faculty of Medicine, Mansoura University (approval registration number: R.21.03.1257). The study was explained to all participants and informed written consent was obtained from each one of them before starting the study. Sociodemographic, clinical and therapeutic data were collected.

The researchers distributed the questionnaire among participants. The questionnaire was administered by 2 methods: either self-administered for those who could read and write or researcher-administered for those who could not. The questionnaire was originally designed to answer questions about health status, knowledge and information about the COVID-19 pandemic and medical and family history of the COVID-19 infection in rheumatoid arthritis (RA) patients [8].

All items had the same contents with the wording revised to reflect SLE instead of RA. A "yes" or "no" response was provided and all questions were closed (only 2 questions in the "History of the COVID-19 infection" section were open-ended).

#### Statistical analysis

Statistical analysis was performed with SPSS 20.0 for Windows (SPSS Inc., Chicago, IL, USA). Continuous data were expressed as mean ± standard deviation or percentages and frequency; the Shapiro-Wilk test was used to assess the normality of distribution.

#### Results

The sample consisted of 200 SLE patients. The patients' data are described in Table I.

The mean age of the participants was 30.1 years (with SD  $\pm$ 8.4). Among the participants, 180 (90%) were female and 172 (86%) were from urban areas. About three quarters of the studied patients were highly educated and 80% belonged to the medium economic class.

Lupus activity scores showed inactivity in 65%, mild in 17%, moderate in 16% and severe disease activity in 3% of the study patients. Corticosteroids and mycophenolate mofetil (MMF) and antimalarials were the most used medications (87%, 87% and 70% respectively).

Most patients (90%) were compliant with therapy. However, 140/200 (70%) of the participants reported difficulty in obtaining medications during the COVID-19 pandemic. It was particularly hard to obtain antimalarials (60%). Lupus disease activity was affected by the drug shortage in about half of the study participants.

The protective measures taken for COVID-19 infection, sources of information about the COVID-19 virus and the impact of the pandemic on the participants are shown in Table II. Twenty-eight (14%) patients reported that they had lost their jobs while 26 patients (13%) reported that their work was severely impacted by the crisis.

Among the precautions taken for prophylaxis against infection, wearing protective masks (74%) and using disinfectants of the hands several times per day (67%) were the most reported measures.

Forty patients (20%) had to stop or reduce taking NSAIDs to guard against COVID-19 infection while 10 patients (5%) had to start taking antimalarials as a prophylaxis against COVID-19 infection.

Among 34 patients (17%) who needed hospitalization, 26 (76%) were mainly due to lupus activity and 24 (71%) of them experienced difficulty in hospital admission because of the COVID-19 pandemic.

About 40% of the participants reported that they had at least one close relative with confirmed COVID-19 infection. In addition, thirty-two patients (16%) had confirmed COVID-19 infection. Most of them became clinically free after treatment.

However, about half of them had lupus flare and had to change the medications used for treatment of SLE. Cutaneous and musculoskeletal manifestations were the most frequently reported presentation of SLE flare following COVID-19 infection. Figure 1 illustrates the disparities in questionnaire responses between males and females.

Among studied patients 58.6% of females vs. 20% of males would use telemedicine for remote contact with the rheumatologist. Most female patients were keener than males to wear masks (76.7% vs. 50% respectively).

Figure 2 illustrates the difference in response to the questionnaire between the patients from urban and rural areas.

Parameters	SLE patients ( <i>n</i> = 200) <i>n</i> (%)	Parameters	SLE patients ( <i>n</i> = 200) <i>n</i> (%)
Sociodemographic parameter	S	Methotrexate	2 (1)
Age [years]	30.1 ±8.4	Leflunomide	6 (3)
Gender (female)	180 (90)	Intravenous immunoglobulin	4 (2)
Residence		Rituximab	4 (2)
Urban	172 (86)	Mode of obtaining medications	
Rural	28 (14)	Self-payment	152 (76)
Education level		Medical insurance	42 (21)
Low	14 (7)	Governmental supply	12 (6)
Moderate	38 (19)	Regularity of drug intake	180 (90)
High	148 (74)	Difficulty to obtain the drug	140 (70)
Economic class		Corticosteroids	8 (4)
Low	18 (9)	Azathioprine	12 (6)
Medium	160 (80)	Mycophenolate mofetil	8 (4)
High	22 (11)	CQ/HCQ	120 (60)
Special habits		Disease affected by drug shortage	96 (48)
Tea/coffee	120 (60)	Regular contact with	174 (87)
Smoking	16 (8)	the rheumatologist	
Alcohol	4 (2)	Modes of remote contact	110 (55)
Addiction	2 (1)	Phone	76 (38)
Lupus activity (SLEDAI)		WhatsApp	94 (47)
Inactive	130 (65)	Messenger	24 (12)
Mild	34 (17)	Telegram	2 (1)
Moderate	31 (16)	Physician's website/page	6 (3)
Severe	5 (3)	Associated diseases/comorbidities	
Health status during COVID-19 pandemic parameters		Diabetes mellitus	4 (2)
Medications used		Hypertension	26 (13)
Corticosteroids	174 (87)	Cardiovascular	6 (3)
CQ/HCQ	140 (70)	Chest problems	12 (6)
Azathioprine	10 (5)	Renal problems	22 (11)
Cyclosporine	62 (31)	Hepatic problems	2 (1)
Mycophenolate mofetil	174 (87)	Allergy problems	10 (5)

**Table I.** Sociodemographic and health status of the patients including the medications received, availability and their contact with the rheumatologist

*CQ/HCQ – hydroxychloroquine, SLE – systemic lupus erythematosus, SLEDAI – Systemic Lupus Disease Activity Index. All parameters are described as number (percent), except age, which is described as mean ± SD.* 

### Discussion

Systemic lupus erythematosus is a chronic autoimmune multisystem disorder with a broad-spectrum clinical presentation. The incidence and prevalence of SLE are significantly increasing worldwide [9].

Not much is known about the impact of the COVID-19 pandemic on SLE patients. In this questionnaire-based study, we studied the tremendous challenges faced by

Egyptian SLE patients during the COVID-19 pandemic in terms of drug shortage, hospitalization, and COVID-19 infection.

A significant number of our participants n = 140 (70%) reported difficulty to obtain drugs during the COVID-19 pandemic. Medication shortages were of particular concern during the COVID-19 outbreak [10]. In the same context, antimalarial drug shortage due to the COVID-19 outbreak affected more than half of our study cohort (60%).

Parameters (during COVID-19 pandemic)	SLE patients (n = 200) n (%)
Work affected	
Mild	54 (27)
Moderate	38 (19)
Severe	26 (13)
Lost	28 (14)
Protective measures taken	
Staying at home	116 (58)
Wearing a mask	148 (74)
Social distancing	122 (62)
Using disinfectants/alcohol	134 (67)
Source of information on COVID-19	
Physicians	56 (28)
Friends	26 (13)
Media	116 (58)
Social media	150 (75)
Others (relatives/MOH/at work)	88 (44)
Consider their knowledge sufficient	120 (60)
Stopped or reduced taking NSAIDs	40 (20)
Considered complementary therapy	46 (23)
Received hydroxychloroquine for prevention	10 (5)
Needed hospitalization	34 (17)
Activity	26 (13)
Pulmonary embolism	6 (3)
Premature labor	2 (1)
Found difficulty in hospitalization	24 (12)
History of COVID-19 infection	
Close relatives affected	76 (38)
Protective measures considered	68 (34)
Patients with confirmed COVID-19	32 (16)
Free after treatment	24 (12)
Carrier	2 (1)
Still infected	6 (3)
The disease was affected (flare)	18 (9)
Cutaneous manifestations	7
Arthritis	5
Lupus nephritis	2
Livedoid vasculopathy	1
Lupus vasculitis	1
Unknown	2
Change in the lupus medications	16 (8)

**Table II.** Protective measures taken for COVID-19, sourceof information and influence of the pandemic

MOH – Ministry of Health, NSAIDs – nonsteroidal anti-inflammatory drugs. Antimalarials are commonly used as the standard of care for management of SLE [11]. The use of antimalarial agents in SLE is very beneficial [12]. Their use is associated with lower rates of new organ damage, disease activity and thromboembolic events [13].

Also, antimalarials have recently been touted as one of the weapons against COVID-19 infection [14]. Additionally, US President Donald Trump stated that antimalarials showed great promise and "could be a gamechanger" in the treatment of COVID-19 infection [15].

However, a retrospective multicenter study published in the *Journal of the American Medical Association* [16] showed that use of hydroxychloroquine (HCQ) to treat COVID-19 infected patients was associated with higher mortality.

Nevertheless, there is widespread off-label use of antimalarials for both prophylaxis and treatment of COVID-19 infection, especially in African countries [17], which may be the main cause of shortage of these drugs [18].

In a national survey conducted on 531 Canadian rheumatologists, about three quarters of the responders were concerned about the antimalarial drug shortage, while 60% of them had been contacted by the patients regarding difficulties accessing HCQ [19].

Shortage of these drugs may cause significant problems for SLE patients who are currently being treated with these drugs [20]. In a retrospective chart review of 509 SLE patients, patients who discontinue antimalarials were at higher risk of disease flares [21].

Unfortunately, no other drug provides similar benefits as HCQ [22]. Therefore, SLE patients are facing challenges with the improper use of their essential medications, and health care policies should take this point into account [23].

To solve this problem in Italy, SLE patients are advised to reduce their doses of antimalarial agents in order to overcome the widespread shortage of these medications [15]. In this context, the German Federal Institute for Drugs and Medical Devices reiterates that any prescription of HCQ in Germany should include an in-label diagnosis justifying its use and prohibits off-label use of that drug except for clinical trials [24].

Our results showed that patients who reported drug shortage also reported more exacerbation of lupus manifestations. It is well known that stoppage of medications of SLE is associated with disease flare [25].

Systemic lupus erythematosus patients have been overwhelmed with warnings of their higher risk of COVID-19 infection and severe illness [15]. Shortage of essential medications for SLE patients may be associated with high emotional distress and anxiety, which may also be associated with a worse disease outcome [26].

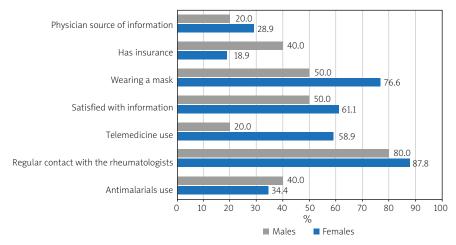
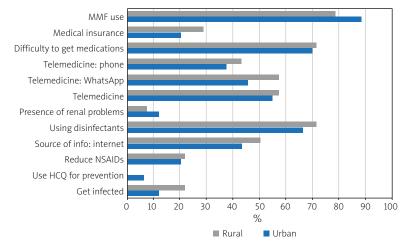


Fig. 1. Frequency of responses to questionnaire in Egyptian lupus patients by gender.



HCQ – hydroxychloroquine, MMF – mycophenolate mofetil, NSAIDs – nonsteroidal anti-inflammatory drugs.

Fig. 2. Frequency of responses to questionnaire in Egyptian lupus patients by residence.

Most of the study patients stated that they were in regular contact with the rheumatologist during the COVID-19 pandemic. In point of fact, SLE patients face challenges in disease followup during the COVID-19 pandemic. Those who keep visiting the doctor are at risk of COVID-19 infection, while at the same time, others are at risk of disease flare [27].

Therefore, it may be better to replace physical consultations of patients with SLE by telemedicine consultations in order to prevent the risk of contagion [28]. In Germany for example, the COVID-19 pandemic led to acute restriction of outpatient care options for patients with rheumatic diseases, which led to a temporary reduction of patients' visits to doctors' offices by about 50% [29]. As a result, medical care was widely switched to telemedicine [29].

A substantial number of the participants used telemedicine as a mode of contact with their physician. In a survey conducted among members of the Dutch Rheumatology Society, the results revealed that more than 80% of outpatients' visits were performed via telephone during the pandemic [30]. However, it is not easy to judge which patients may be suitable for telemedicine consultation [31].

A significant proportion of our cohort (17%) reported that they were in need of hospitalization during the COVID-19 pandemic. The most common reason for hospitalization was lupus flare. COVID-19 related distress in rheumatic patients is associated with worse health scores [32]. Disease flare and medication access difficulties were common among rheumatic patients during the COVID-19 pandemic [32].

Regrettably, most of those who needed hospitalization experienced difficulty in hospital admission. The COVID-19 pandemic is associated with "collateral damage" caused by the decrease of emergency room visits for non-COVID-19 conditions [33].

There is a decrease (65.4%) in hospitalization of SLE patients during the COVID-19 pandemic [34]. The pan-

demic has promoted crowding of hospital places with COVID-19 patients. Some hospitals had to suspend other services during the pandemic [35].

Systemic lupus erythematosus patients are at increased risk of infection either due to lupus itself or immunosuppressive medications [36]. The initial results of the COVID-19 Global Rheumatology Alliance Registry show that 19 (17%) of 110 rheumatic patients diagnosed with COVID-19 were patients with lupus [37].

According to our results, 16% of the studied patients reported confirmed infection with COVID-19. The clinical course of our infected cohort seems to be favorable, as most of them (75%) reported complete resolution of infection without significant complications. This is compatible with the results of a survey conducted among 126 lupus patients in Italy which revealed 12 patients with confirmed or suspected COVID-19 infection [38].

The most recent studies showed controversial results about COVID-19 sequelae in SLE patients [39]. It seems that SLE patients may be at higher risk of an unfavorable outcome during the COVID-19 pandemic [40].

## Conclusions

This study had some limitations. The method of data collection was based on self-reporting. The information provided by the participants could not be verified for clinical accuracy. Additionally, we could not get information from patients with severe COVID-19 infection. The small number of the participants may be another limitation. Finally, the COVID-19 outbreak is still ongoing, and possible new infections in our cohort cannot be excluded.

However, this study had several strengths; it targeted an Egyptian cohort. To the best of our knowledge, this is the first paper that addresses the impact of the COVID-19 pandemic in Egyptian lupus patients.

Further studies on a large number of patients and from different centers in Egypt are warranted to confirm that these data help improve management of SLE patients during the COVID-19 pandemic. Multinational studies may also shed light on this topic.

The authors declare no conflict of interest.

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