Corticosteroids for COVID-19 symptoms and quality of life at 1 year from admission

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Abstract
The long-term evolution of COVID-19 is unknown, making it necessary to study the persistence of symptoms over time and their impact on quality of life in people who have had the disease. We analyzed these aspects 1 year after admission for COVID-19 and explored the influence of treatment with systemic corticosteroids during the acute phase of the illness. This observational cohort study took place in a tertiary hospital in March and April 2021 and included people admitted due to infection with SARS-CoV-2 in March, April, or May 2020. We excluded patients who had died, were unreachable or had substantial cognitive impairment. A telephone survey was undertaken to assess the presence of symptoms related to COVID-19 and to administer the SF-36 quality of life questionnaire. Other variables collected were demographic and clinical data along with the treatment received and the evolution over time. We analyzed 76 patients, including 44 who did not receive corticosteroids and 32 who did. Most symptoms were less frequent in the group that received corticosteroids, with statistically significant differences for headache, dysphagia, chest pain, and depression. These patients also showed significantly better outcomes in the SF-36 domains for “bodily pain” and “mental health.” Corticosteroids administered in the acute phase of COVID-19 could attenuate the presence of long-term symptoms and improve patients’ quality of life.

KEYWORDS
corticosteroids, COVID-19, long COVID, post-acute COVID-19 syndrome, quality of life, SARS-CoV-2

1 | INTRODUCTION

COVID-19 is a disease produced by the SARS-CoV-2 virus (severe acute respiratory syndrome coronavirus 2), which is responsible for the current pandemic1,2 and associated with a wide array of symptoms. About 80% of cases produce mild symptoms like anosmia and ageusia (loss of smell and taste), headache, muscle pain, fever, cough, and diarrhea, among others.3,4 In other cases, the virus triggers a dysregulation of the immune system, which produces an acute respiratory distress syndrome (ARDS) that leads to admission to the intensive care unit (ICU).4-6 In this context of systemic hyperinflammation, corticosteroids have been used as nonselective anti-inflammatory agents, demonstrating significant reductions in mortality.7

The evolution and long-term sequelae of this emerging disease are still unknown, heightening the need to follow patients over time.
Recent contributions to the medical literature have described diverse clinical scenarios during follow-up, including the persistence of symptoms from the acute phase of illness, the onset of new symptoms in the postinfection period, exacerbation of symptoms that were present before the viral infection, and a wide range of post-COVID-19 sequelae.\(^9\) These situations should be detected to ensure treatment and improve patients' quality of life, which may be impaired following the acute episode. However, the best therapeutic approach and pharmacological management are currently unknown.\(^9\) In this context, the health professionals participating in this study had the impression, from anecdotal observation, that patients who received corticosteroids during their admission had a better long-term evolution, with fewer symptoms related to the COVID-19 infection.

This study aims to test that hypothesis, analyzing the impact of corticosteroid treatment on the presence of symptoms and quality of life 1 year after initial admission with COVID-19.

## MATERIALS AND METHODS

### 2.1 Study design and participants

This observational cohort study took place in a tertiary hospital in March and April 2021 and included adults (≥18 years) admitted to the infectious diseases ward due to COVID-19 in March, April, or May 2020. Infection with SARS-CoV-2 was confirmed by real-time polymerase chain reaction (RT-PCR) or serology. We excluded patients who had died, were unreachable or had substantial cognitive impairment.

### 2.2 Variables

A telephone survey was undertaken to assess the presence of symptoms related to COVID-19 (Annex 1). The questionnaire included items on the most frequently reported symptoms according to patient management guidelines in COVID-19.\(^9\) In their response to questions about whether they had specific symptoms, participants could answer yes or no, and if the symptom was present, specify whether it had appeared after the acute phase. There was also an open question about any other symptoms they had experienced, apart from those mentioned in the interview.

The survey on symptomology was followed by the SF-36 quality of life questionnaire.\(^10\) The SF-36 is a general questionnaire that is quick and straightforward to administer, generating a numerical score that enables assessment and comparison of key health domains (physical functioning, physical role limitations, bodily pain, social functioning, mental health, emotional role limitations, energy/vitality, and general health perceptions) in different populations. For each domain, the SF-36 generates a percentage value that is directly and proportionally related to a better quality of life in that domain. Before the administration of both surveys, the patient gave verbal consent, which is reflected in the data collection file.

In addition, the electronic health record (Orion Clinic program) was reviewed to extract demographic variables (age, sex); length of hospital stay; clinical data (relevant medical history, body mass index [BMI], age-adjusted Charlson comorbidity index, development of mild/moderate/severe acute respiratory distress syndrome); treatment received during admission (lopinavir/ritonavir, hydroxychloroquine, tocilizumab, anakinra, systemic corticosteroids); need for noninvasive or invasive mechanical ventilation), and admission to the ICU.

### 2.3 Statistical analysis

We used SPSS statistical software (v.23; IBM) for analyses. In the initial descriptive study, quantitative variables were expressed as the mean (95% confidence interval, CI) or median (interquartile range, IQR), depending on the normality of the distribution. Categorical variables were described as absolute and relative frequencies. Statistical comparisons were undertaken, using the \( \chi^2 \) test for categorical variables and the Student t-test or Mann–Whitney U test, as appropriate, to compare categorical with quantitative variables.

Our center’s clinical research ethics committee approved the study. Patient confidentiality was preserved, and the performance of the study complied with the principles of the Declaration of Helsinki (2013) and good clinical practice rules.

### 3 RESULTS

A total of 124 patients were admitted for COVID-19 during the study period. We excluded 48 of these from the study (24 died, 13 presented significant cognitive impairment, and 11 were unreachable), for a final sample of 76 people, 32 of whom received corticosteroids and 44 of whom did not. Table 1 details their clinical epidemiological characteristics, comorbidities, treatment, and evolution, according to whether they received corticosteroids. The groups were largely similar across different variables, although the steroids group presented a larger proportion of men (75.0% vs. 52.3%, \( p = 0.004 \)) and people with hypertension (71.9% vs. 40.9%, \( p = 0.007 \)).

Regarding the complications experienced over the first year of follow-up, two patients—neither of whom received corticosteroids—were readmitted, one for stroke and the other for a urinary tract infection. The most frequently reported symptoms at 1 year of infection were forgetfulness (45.5% in patients that did not receive corticosteroid treatment vs. 43.8% in those who did), followed by asthenia (43.2% and 34.4% in the respective groups) and arthromyalgia (38.6% and 37.5%).

Patients who had received steroids presented significantly less headache, dysphagia, chest pain, and depression at 1 year. The rest of the symptoms also tended to appear less frequently in this group, with the exception of the loss of taste/smell, odynophagia, diarrhea, and alopecia. The steroids group also presented a lower number of symptoms (median 2, IQR 1–4) compared to the non-steroids group.
| Variables                                      | No steroids (N = 44) | Steroids (N = 32) | p     |
|-----------------------------------------------|----------------------|-------------------|-------|
| **Men, n (%)**                                | 33 (52.3%)           | 24 (75.0%)        | 0.004 |
| **Age, median (IQR), years**                  | 61.5 (52.7–72.5)     | 68.5 (60.2–75.7)  | 0.53  |
| **Charlson comorbidity index, median (IQR)**  | 2 (1–3)              | 3 (2–4)           | 0.70  |
| **BMI, median (IQR), (kg/m²)**                | 29.5 (25.8–31.8)     | 28 (26.3–30.2)    | 0.59  |
| **Length of hospital stay, median (IQR), days**| 12 (8.2–20.7)        | 21 (16–25)        | 0.86  |

| Comorbidities, n (%)                          |                      |                   |       |
|-----------------------------------------------|----------------------|-------------------|-------|
| Tobacco use                                   |                      |                   |       |
| Ex-smoker                                     | 7 (15.9%)            | 5 (15.6)          | 0.21  |
| Current smoker                                | 4 (9.1%)             | 0                 |       |
| Hypertension                                  | 18 (40.9%)           | 23 (71.9%)        | 0.007 |
| Dyslipidaemia                                 | 18 (40.9%)           | 12 (37.5%)        | 0.76  |
| Atrial fibrillation                           | 2 (4.5%)             | 1 (3.1%)          | 0.75  |
| COPD                                          | 1 (2.3%)             | 0                 | 0.39  |
| Chronic bronchitis                            | 1 (2.3%)             | 0                 | 0.39  |
| Asthma                                        | 2 (4.5%)             | 0                 | 0.22  |
| Diabetes mellitus without target organ damage | 4 (9.1%)             | 4 (12.5%)         | 0.63  |
| Diabetes mellitus with target organ damage    | 2 (4.5%)             | 2 (6.3%)          | 0.74  |
| Obesity                                       | 17 (38.6%)           | 11 (34.4%)        | 0.70  |
| Anxiety                                       | 3 (6.8%)             | 1 (3.1%)          | 0.48  |
| Depression                                    | 2 (4.5%)             | 1 (3.1%)          | 0.75  |

| Development of acute respiratory distress syndrome, n (%) |       |
|----------------------------------------------------------|-------|
| Mild                                                      | 1 (2.3%) | 2 (6.3%) | 0.10  |
| Moderate                                                  | 1 (2.3%) | 2 (6.3%) |       |
| Severe                                                    | 8 (18.2%) | 12 (37.5%) |      |

| Treatment received, n (%)                                |       |
|----------------------------------------------------------|-------|
| Lopinavir/ritonavir                                       | 31 (70.5%) | 25 (78.1%) | 0.45  |
| Hydroxychloroquine                                        | 39 (88.6%) | 30 (93.8%) | 0.45  |
| Tocilizumab                                               | 1 (2.3%) | 1 (3.1%) | 0.82  |
| Anakinra                                                  | 0      | 0      |       |

| Need for ventilation and admission to ICU, n (%)         |       |
|----------------------------------------------------------|-------|
| Noninvasive mechanical ventilation                       | 4 (9.1%) | 7 (21.9%) | 0.12  |
| Invasive mechanical ventilation                          | 7 (15.9%) | 5 (15.6%) | 0.97  |
| ICU admission                                             | 7 (15.9%) | 6 (18.8%) | 0.95  |

Abbreviations: BMI, body mass index; COPD, chronic obstructive pulmonary disease; ICU, intensive care unit; IQR, interquartile range.

Table 1: Participant characteristics at 1 year from admission for COVID-19, according to the administration of corticosteroids.

Our results suggest that patients who receive corticosteroids during their admission for COVID-19 are less likely to experience persistent symptoms and more likely to have a better quality of life at 1 year of admission.

The long-term persistence of COVID-19 symptoms is currently among the topics attracting the most scientific interest. However, our exhaustive literature search yielded few publications analyzing this aspect in patients who were admitted for this disease, and the studies that do exist are heterogeneous, have a limited follow-up, and do not examine the association with corticosteroids—the only treatment that has been proven to increase survival in the short term and whose use has been widely adopted globally.7

Chopra et al. performed telephone interviews in 488 patients in Michigan (USA) 60 days after their admission due to COVID-19, and 32.6% reported persistent symptoms, including 18.9% who described new and sometimes worsening symptoms.1 The most common were dyspnea, cough, and the persisting loss of smell and taste. In Wuhan (China), Wang et al. assessed 131 patients (52.6% with severe COVID-19 symptoms) 40.4% had symptoms (especially cough, dyspnea, and expectoration), but at four weeks these persisted in just 9.1%, with no relation to the severity of the acute illness.12

As for European studies, Halpin et al. (UK) analyzed 100 patients (nearly a third of whom had been admitted to the ICU) at 48 days from discharge; 72% reported fatigue; 65.6%, dyspnea; and 46.9%, anxiety.13 In Italy, Carfi et al. reported similar results in 143 patients admitted for COVID-19 and interviewed at 60 days from symptoms onset. Fatigue (53.1%), dyspnea (43.4%), arthralgia (37.3%), and chest pain (21.7%) were the most frequently reported symptoms.14

4 Discussion

Our results suggest that patients who receive corticosteroids during their admission for COVID-19 are less likely to experience persistent symptoms and more likely to have a better quality of life at 1 year of admission.
Moreover, 55% reported having at least three symptoms, and 44.1% saw their quality of life decline, according to the EuroQoL visual analog scale.

In our country, Moreno-Pérez et al. (Alicante) assessed symptoms persistence face-to-face at 2–3 months of infection in 277 people, not all of whom required hospitalization; their median age was 56 years, and 8.7% were admitted to the ICU. In line with the literature, the most frequently reported symptoms were fatigue (34.8%), dyspnoea (34.4%), loss of taste/smell (21.4%), and cough (21.3%).

All of these studies have a quite limited follow-up compared to our study, which could explain the predominance of respiratory symptoms, as the SARS-CoV-2 pneumonia was still quite recent. The most prominent symptoms in our patients had a different nature and included forgetfulness, asthenia, and arthromyalgia.

Treatment with corticosteroids during admission for COVID-19 seemed to have a positive influence in practically all aspects of both symptoms and quality of life at 1 year. This tendency is clear, and the fact that statistically significant differences were not observed may be due in part to the small sample size. Furthermore, regarding the SF-36 quality of life survey, if we analyze the distribution of the data and the IQRs in addition to the median scores, better scores can be inferred in the steroids group for all eight domains.

One possible explanation for these results could be that the symptoms or sequelae following the viral infection are consequences of the organ and tissue damage caused by the intense inflammation during the acute phase of the illness. By lessening that inflammation, its consequences would also be attenuated.

The strengths of this study reside in the direct form of collecting data on symptoms at 1 year, via a personal interview that specifically asked about each symptom and allowed participants to volunteer others. Moreover, our exhaustive literature search did not show any studies with such a long follow-up in patients who had been admitted for COVID-19 or studies that assessed the long-term impact of treatment with corticosteroids.

In contrast, limitations include its small sample size, the performance of non-parametric statistical analyses due to the non-normal distribution of the variables (as also occurs in the cited literature), and the lack of a control group.

| Symptoms    | Presence of symptoms, n (%) | Onset of symptoms following acute phase, n (%) |
|-------------|-----------------------------|-----------------------------------------------|
|             | No steroids (N = 44)        | Steroids (N = 32) | p     | No steroids | Steroids | p     |
| Forgetfulness | 20 (45.5%)                  | 14 (43.8%) | 0.88 | 8 (18.2%)  | 8 (25%)  | 0.47  |
| Asthenia    | 19 (43.2%)                  | 11 (34.4%) | 0.44 | 4 (9.1%)   | 1 (3.1%) | 0.30  |
| Arthromyalgia | 17 (38.6%)                 | 12 (37.5%) | 0.92 | 2 (4.5%)   | 2 (6.3%) | 0.74  |
| Dyspnea     | 13 (29.5%)                  | 6 (18.8%)  | 0.28 | 4 (9.1%)   | 0 (     )| 0.080 |
| Anxiety     | 12 (27.3%)                  | 5 (15.6%)  | 0.23 | 2 (4.5%)   | 3 (9.4%) | 0.40  |
| Headache    | 11 (25%)                    | 2 (6.3%)   | 0.032| 4 (9.1%)   | 1 (3.1%) | 0.30  |
| Depression  | 10 (22.7%)                  | 1 (3.1%)   | 0.016| 3 (6.8%)   | 1 (3.1%) | 0.48  |
| Arthritis   | 10 (22.7%)                  | 2 (6.3%)   | 0.052| 4 (9.1%)   | 1 (3.1%) | 0.30  |
| Cough       | 10 (22.7%)                  | 5 (15.6%)  | 0.44 | 0 (    )   | 1 (3.1%) | 0.24  |
| Rhinorrhea  | 9 (20.5%)                   | 6 (18.8%)  | 0.85 | 5 (11.4%)  | 5 (15.6%)| 0.59  |
| Hypoacusis  | 9 (20.5%)                   | 3 (9.4%)   | 0.19 | 4 (9.1%)   | 1 (3.1%) | 0.30  |
| Alopecia    | 8 (18.2%)                   | 6 (18.8%)  | 0.95 | 1 (2.3%)   | 2 (6.3%) | 0.38  |
| Dysphagia   | 5 (11.4%)                   | 0 (     )  | 0.049| 2 (4.5%)   | 0 (     )| 0.22  |
| Chest pain  | 5 (11.4%)                   | 0 (     )  | 0.049| 2 (4.5%)   | 0 (     )| 0.22  |
| Anosmia     | 5 (11.4%)                   | 6 (18.8%)  | 0.37 | 0 (    )   | 1 (3.6%) | 0.24  |
| Ageusia     | 5 (11.4%)                   | 4 (12.5%)  | 0.88 | 1 (2.3%)   | 0 (     )| 0.39  |
| Odynophagia | 1 (2.3%)                    | 1 (3.1%)   | 0.82 | 0 (    )   | 0 (     )| 0.39  |
| Diarrhea    | 1 (2.3%)                    | 3 (9.4%)   | 0.17 | 0 (    )   | 0 (     )| 0.39  |
| Fever       | 0 (    )                    | 0 (     )  |     | 0 (    )   | 0 (     )| 0.39  |
| N symptoms, median (IQR) | 3 (1–6)       | 2 (1–4)   | 0.25 |

Abbreviation: IQR, interquartile range.
Overall, our results suggest that treatment with corticosteroids in patients hospitalized for COVID-19 not only improves the short-term prognosis; it also attenuates the presence of symptoms and improves the quality of life in the long term. More studies are necessary to confirm or reject this hypothesis.

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CONFLICT OF INTERESTS
The authors declare that they have no conflict of interest.

AUTHOR CONTRIBUTIONS
Ignacio Pérez Catalán: conception and design of the study, writing of the manuscript, bibliographic search, data collection and analysis, and interpretation of data. Celia Roig Martí: conception and design of the study, writing of the manuscript, bibliographic search, data collection and analysis and interpretation of data. Daniela Palomo de la Sota: data collection and bibliographic search. Alejandro Cardenal Álvarez: data collection and bibliographic search. Maria José Esteve Gimeno: data collection and bibliographic search. Sergio Fabra Juana: data collection and bibliographic search. Germán Herrero Rodríguez: data collection and bibliographic search. Elena Domínguez Bajo: data collection and bibliographic search. Nuria Tornador Gaya: conception and design of the study. Jorge Usó Blasco: conception and design of the study. Jose Manuel Ramos Rincón: conception and design of the study, writing of the manuscript, bibliographic search and analysis and interpretation of data.

DATA AVAILABILITY STATEMENT
Author elects to not share data.

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SUPPORTING INFORMATION

Additional Supporting Information may be found online in the supporting information tab for this article.

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