

# Neurologic Manifestations in a Prospective Unselected Series of Hospitalized Patients With COVID-19

Juan Carlos García-Moncó, MD, PhD, Antonio Cabrera Muras, MD, Markel Erburu Iriarte, MD, Patricia Rodrigo Armenteros, MD, Alejandra Collía Fernández, MD, Javier Arranz-Martínez, MD, Solange Kapetanovic, MD, Ana Lorenzo-García, MD, Amaia Bilbao González, PhD, and Marian Gomez-Beldarrain, MD

**Correspondence**  
Dr. García-Moncó  
hospit05@sarenet.es

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## Abstract

### Background

An increasing number of neurologic problems are being described in coronavirus disease 2019 (COVID-19) disease, but their frequency and type have not been defined. In this study, we sought to determine the extent of neurologic manifestations of COVID-19 in a prospective series of unselected patients admitted to the general medicine wards of our hospitals due to COVID-19 and who were examined by a team of neurologists.

### Methods

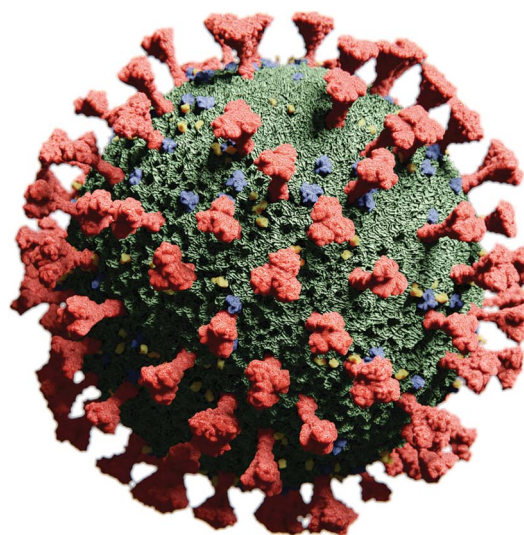
Eight neurologists provided medical attention to patients hospitalized for COVID-19 to provide medical support to other hospital units tasked with the care of an increasingly larger influx of patients with COVID-19. A series of 100 consecutive, unselected patients were evaluated systematically, including a questionnaire that collected medical information derived from the initial examination and the medical history.

### Results

Eighty-eight percent of the patients had 1 neurologic manifestation associated with COVID-19 during hospitalization. Most common were anosmia-dysgeusia and headache (44% each), myalgias (43%), and dizziness (36%). Less frequent were encephalopathy (8%), syncope (7%), seizures (2%), and ischemic stroke during the period of hospitalization (2%). Anosmia and headache associated with younger patients with less severe disease, and both were associated with each other and with serum inflammatory markers. Encephalopathy was associated with fever and syncope and with markers of inflammation.

### Conclusions

Neurologic disturbances are common in patients with COVID-19, particularly if patients are evaluated by neurologists. There is a wide variety of neurologic conditions, some of them severe, in the spectrum of COVID-19 disease that will benefit from an evaluation by practicing neurologists.



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The present pandemic of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection has a primary respiratory component, but in severe cases, the clinical manifestations have included

Osakidetza Basque Health Service (JCG-M, ACM, MEI, PRA, ACF, SK, AL-G, ABG), Basurto University Hospital, Departments of Neurology and Research Unit; and Galdakao Hospital (JA-M, MG-B), Vizcaya, Spain.

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disorders of other organ systems. A number of neurologic problems whose exact frequency and type are not well defined have been reported in the very recent literature<sup>1–7</sup> as well as anecdotally in our own pandemic environment. Clinical information derives mainly from series of respiratory patients in whom neurologic problems may have been missed or underestimated as well as from case reports that do not allow an accurate estimate of the extent and description of these neurologic disturbances.

Our hospitals serve an area that has reported nearly 2,000 cases of coronavirus disease 2019 (COVID-19) from March 20 to April 17, 2020 (our study period). This high case load in such a short period of time nearly overwhelmed the normal functioning of our hospitals making the care of COVID-19 patients a medical priority that required resources from all medical specialties. Thus, a number of neurologists were asked to participate in the care of hospitalized patients with COVID-19. This rapidly evolving pandemic also provided an opportunity to observe the clinical development of the patients from a neurologic perspective.

In this study, we sought to determine the type and frequency of neurologic manifestations of COVID-19 by evaluating a consecutive prospective series of unselected patients admitted to the general wards of our hospitals due to COVID-19 and who were examined by a team of neurologists who assisted in the care of these patients.

## Methods

A team of 8 neurologists (4 attending neurologists and 4 residents in their last 2 years of neurology residency) provided medical attention to patients admitted to the hospital due to respiratory or systemic COVID-19. This was part of the plan established at our 2 institutions to provide medical support to other hospital units tasked with the initial care of an increasingly larger influx of patients with COVID-19. The study was conducted at the Hospital Universitario de Basurto and the Hospital de Galdakao, both in Vizcaya, Spain. Both hospitals provide health care to a pooled population of 670,000 inhabitants.

Increasing evidence for neurologic involvement in some of these patients led us to investigate this possibility with a standardized approach. To this end, we designed a questionnaire to verify the neurologic manifestations in a prospective unselected series of patients with COVID-19. The questionnaire reflected the neurologic manifestations already described in the incipient literature of the COVID-19 pandemic.<sup>1–5,8–10</sup> The questionnaire also collected medical information derived from the initial examination and the medical history of the patient, leading to a more accurate interpretation of any neurologic disturbances and also leading to a detailed neurologic examination if needed.

## Patient Selection

One hundred patients were recruited consecutively in an unselected manner by all 8 neurologists from March 20 to April 17.

## Inclusion and Exclusion Criteria

Patients were eligible if (1) they had respiratory symptoms with the need of supplementary oxygen and a positive reverse-transcriptase PCR (RT-PCR) assay for SARS-CoV-2 in a nasopharyngeal swab; (2) they had a chest X-ray or CT scan consistent with COVID-19 pneumonia, and (3) they did not require admission in the intensive care unit (ICU). The CURB-65 score for pneumonia severity was used to assess the patient's condition and help determine the need for inpatient treatment. This is a yes/no scale that evaluates confusion, blood urea nitrogen (BUN) >19 mg/dL, respiratory rate  $\geq$ 30 per min, systolic blood pressure <90 mm Hg or diastolic  $\leq$ 60 mm Hg, and age  $\geq$ 65 years. Scores of 0–1 indicate a low mortality risk, whereas 2–5 represent a moderate to high mortality risk. Exclusion criteria included an inability to communicate due to speech impairment or significant cognitive impairment.

## Variable Collection

The following variables were collected and introduced anonymously into a registry:

- Age, sex, admission date, and date of disease onset.
- Fever at onset and fever only during admission.
- History of hypertension, dyslipidemia, diabetes mellitus, tobacco use, obesity (body mass index  $>30$  kg/m<sup>2</sup>), cardiac disease (type), and respiratory disease (type).
- History of headache (type), stroke, epilepsy, dementia, and others (specify).
- Neurologic symptoms at presentation or during admission: anosmia-ageusia, headache (type), myalgias, dizziness (defined as unsteady gait with no other focal neurologic findings), vertigo, syncope (convulsive or nonconvulsive), encephalopathy (defined as impaired attention and arousal, presenting with confusion, lethargy, delirium, or coma), and seizures.
- Laboratory parameters: serum creatinine, transaminases, creatine kinase (CK), C-reactive protein (CRP), coagulation parameters (international normalized ratio [INR], activated partial thromboplastin time [aPTT]), D-dimer, ferritin, and platelet count.

## Data Analysis

A quantitative descriptive analysis based on percentages and a detailed qualitative analysis of those infrequent conditions that appeared only in a few patients were performed. We analyzed whether there was an association of the main neurologic manifestations with age, sex, laboratory parameters of systemic inflammation (serum CRP, D-dimer, fibrinogen, and ferritin), and disease severity according to the CURB-65 scoring system. We also analyzed the association between different neurologic manifestations.

## Statistical Analysis

The description of the variables was performed using frequency tables, mean and SD, or median and interquartile range (IQR). To study the association between categorical variables, the  $\chi^2$  or Fisher exact test was used. The non-parametric Wilcoxon test was used to examine the

differences in quantitative analytic variables according to dichotomous variables. The nonparametric Kruskal-Wallis test was used to examine the differences in quantitative analytic variables according to qualitative polytomous variables. Furthermore, logistic regression models were also performed to study the risk of developing neurologic manifestations according to other characteristics. The results are presented as the odds ratio (OR) and 95% confidence interval (CI).

A result was considered statistically significant at  $p < 0.05$ . All statistical analyses were performed with SAS for Windows statistical software, version 9.2 (SAS Institute, Inc., Cary, NC).

### Standard Protocol Approvals, Registrations, and Patient Consents

Verbal informed consent was obtained, and none of the patients declined. The process of obtaining verbal informed consent required that relevant risks, benefits, and alternatives be disclosed to the patient and that there be an opportunity to have questions answered. An agreement to proceed was obtained. They were subjected to clinical history and examination as part of their routine and protocolized care. Patients with encephalopathy and seizures gave their consent once they improved enough. The study was approved by the Ethics Committee of the Basque Country.

### Data Availability

Anonymized data not published within this article will be made available by request from any qualified investigator.

## Results

### Characteristics of Patients

One hundred consecutive patients admitted for respiratory COVID-19 disease were enrolled. Seventeen patients were excluded due to the inability to obtain clinical history due to severe cognitive decline or poor general condition. Thus, 117 patients were considered to yield a series of 100 patients for this study.

The characteristics of the patients are described in table 1. Females constituted 38%, a ratio of 1.6 males to females, and median age was 61 years (range, 37–93 years). The severity of the infection according to the CURB-65 scoring system was classified into low risk (CURB-65 of 0–1) in 65 and moderate to high risk (CURB-65 of 2–5) in 35. No patient scored 5 in this study sample. Sixty-seven percent had concomitant cardiovascular risk factors with hypertension being the most frequent, followed by lipid disorders, diabetes mellitus, smoking, and obesity. Pulmonary disease was present in 24%. Twenty-three patients had prior neurologic disorders that included headache (5, 4 of them migraineurs) and moderate mild impairment that did not preclude history taking. Four patients had a prior stroke. Eight patients had other prior conditions that included Parkinson disease (2), diabetic neuropathy, essential tremor, normal pressure hydrocephalus, meningioma, peripheral neuropathy, and spondylotic myelopathy.

### Neurologic Manifestations: Frequency and Distribution

At least 1 neurologic manifestation was present in 88% of the patients. Two or more were present in 58%, and 29% of patients had 3 or more neurologic disturbances. Seventy-three percent had fever on admission and was persistent in 62% for several days after admission. The more frequent neurologic disorders during hospitalization included anosmia-dysgeusia and headache (44% each), myalgias (43%), and dizziness (36%, 2 of them had vertigo). The less frequent included encephalopathy (8%), syncope (7%), seizures (2%), and ischemic stroke during the period of hospitalization (2%) (table 2).

Smell and taste dysfunction was described in 44% of patients by regular history taking, which and probably gives a true estimate of their frequency in this context. Headache was present in 44% of the patients. Most patients reported a persistent, dull, diffuse, and oppressive pain of moderate intensity, resembling a tension-type headache. Seven patients had migraine-like headaches, but only 2 patients had a history of migraine. None of the patients reported other types of headaches as verified by the evaluating neurologists.

Nonspecific dizziness was present in one-third of the patients. Two patients had isolated vertigo of peripheral type with nystagmus and imbalance.

Myalgias were also frequent, present in 43% of patients, and were diffuse, with no specific characteristics. Many patients described back pain, possibly of muscle origin.

Encephalopathy was present in 9% of our patients and had been reported in 8%. Eight of our patients with encephalopathy had a severe disease as indicated by a CURB-65 score of 2–4. We did not find an association with inflammatory markers, although the number of patients is low and this aspect should be interpreted with caution. Encephalopathy occurred in most patients in the context of fever and hypoxemia. It coincided with a minor stroke and seizures (1 patient each). One patient developed renal failure with dehydration, metabolic acidosis, and hyperkalemia. Five of these patients had a prior diagnosis of mild cognitive impairment.

There were a number of low-frequency neurologic manifestations in our series that involved syncope, seizures, and stroke. Syncope was present in 7 patients, did not associate with myoclonic jerks, and was of the neurally mediated characteristics in all patients without evidence of cardiac origin.

Seizures were present in 2 patients, one of focal onset and the other one tonic-clonic. The former patient had a fever and mild encephalopathy and presented repeated seizures of focal (frontotemporal) onset that resolved with levetiracetam. The second one had fever and mild encephalopathy and presented a single tonic-clonic seizure that did not recur and did not need specific medication. None of them had a history of epilepsy.

**Table 1** Baseline Characteristics of Patients Hospitalized With COVID-19

Variable	All patients (N = 100) N and % <sup>a</sup>
Sex (females)	38
Age, mean (SD), median (range)	63.5 (13.9), 61 (37–93)
Fever at admission	73
Fever during admission	62
<b>CURB-65 score</b>	
0	37
1	28
2	30
3	3
4	2
<b>Cardiovascular risk factors</b>	67
Hypertension	45
Dyslipidemia	33
Diabetes mellitus	20
Smoking	16
Obesity (BMI $\geq 30$ kg/m <sup>2</sup> )	10
History of cardiovascular disease <sup>b</sup>	21
History of pulmonary disease <sup>c</sup>	24
<b>Prior neurologic disease</b>	23
No	77
<b>Moderate cognitive impairment</b>	6
<b>Headache</b>	5
Tension type	1
Migraine	4
Other	0
<b>Stroke/TIA</b>	4
<b>Others<sup>d</sup></b>	8

Abbreviations: BMI = body mass index; COPD = chronic obstructive pulmonary disease; COVID-19 = coronavirus disease 2019.

<sup>a</sup> Because there were 100 patients, the number and percentage coincide.

<sup>b</sup> History of ischemic cardiopathy or arrhythmias.

<sup>c</sup> COPD or asthma.

<sup>d</sup> Included Parkinson disease, essential tremor, normal pressure hydrocephalus, meningioma, peripheral neuropathy, and spondylitis myelopathy.

Two patients had an ischemic stroke. In one of them, the stroke occurred just before admission and in the other while under our care. Both were considered minor strokes, and while one of them showed neurologic focal findings suggestive of lacunar infarction (hemisensory syndrome with normal brain CT), the other one presented as an encephalopathy without focal findings with neuroimaging (brain CT scan) of

**Table 2** Neurologic Manifestations Associated With COVID-19 Disease

Symptom/signs	N and % (all patients = 100) <sup>a</sup>
<b>At least 1 neurologic manifestation</b>	88
<b><math>\geq 2</math> manifestations</b>	58
<b><math>\geq 3</math> manifestations</b>	29
<b>Anosmia/dysgeusia</b>	44
<b>Headache</b>	44
<b>Tension type</b>	37
<b>Migrainous</b>	7
<b>Myalgias</b>	43
<b>Dizziness (unsteady gait)</b>	36
<b>Vertigo, peripheral type</b>	2
<b>Encephalopathy</b>	9
<b>Syncope</b>	7
<b>Seizures</b>	2
<b>Focal onset</b>	1
<b>Generalized</b>	1
<b>Ischemic stroke during or before admission (1 each)</b>	2

Abbreviation: COVID-19 = coronavirus disease 2019.

<sup>a</sup> Because there were 100 patients, the number and percentage coincide.

infarction. Both patients had a favorable outcome, and neither had relevant coagulation abnormalities.

### Associations Between Clinical Manifestations and Variables of the Study

The analytical parameters revealed the presence of liver dysfunction in 38%, kidney failure in 17%, and increased CK in 25% (table 3). Coagulation parameters (aPTT and INR) were within normal limits, but D-dimer levels were notably increased in most patients. Other markers of inflammation (CRP, fibrinogen, and ferritin) were also markedly elevated.

When grouped by sex (table 4), headache and myalgia were significantly more frequent in women (OR [95% CI] = 3.60 [1.54–8.40] and 2.68 [1.17–6.17], respectively), whereas anosmia-dysgeusia and dizziness showed no sex preference. Other neurologic manifestations were not analyzed for associations with other clinical manifestations or laboratory results due to the low number of patients, unsuitable for statistical analysis.

There was a significant association between having headaches and having anosmia-dysgeusia (OR [95% CI] = 2.56 [1.14–5.78] of having anosmia-dysgeusia with the presence of headache), whereas the other main manifestations (dizziness



**Table 3** Laboratory Analyses

Parameters	All patients = 100
<b>Blood chemistry values</b>	
Increased liver enzymes (GOT >31 U/L, GPT >31 U/L, and GGT >42 U/L), N and %	38
Acute renal failure (Cr > 1.1 mg/dL), N and %	17
Increased CK (>189 U/L), N and %	25
Median (IQR)	320.5 U/L (178–1387)
<b>Coagulation and inflammatory markers (reference values)</b>	
INR (0.9–1.2)	1 (1–1.1)
aPTT (25–40 s)	36 (32–39)
Fibrinogen (200–450 mg/dL)	682 (583–783)
D-dimer (0–500 ng/mL)	1028.5 (495–3080)
Platelets per $\mu$ L (135,000–450,000)	244,000 (194,000–332,500)
CRP (0–5 mg/L)	90 (37.9–215)
Ferritin (30–400 $\mu$ g/L)	895 (379–1713)

Abbreviations: aPTT = activated partial thromboplastin time; CK = creatine kinase; CRP = C-reactive protein; GGT = gamma-glutamyltransferase; GOT = transaminases alanine transaminase; GPT = aspartate transaminase; INR = international normalized ratio; IQR = interquartile range.

and myalgias) did not show a significant association. Among the less frequent manifestations, patients with encephalopathy had a higher risk of having syncope (OR [95% CI] = 10.88 [1.97–60.15]). The low numbers of patients with seizures and stroke precluded a statistical analysis to verify their potential association with the rest of the variables included in the study.

When the neurologic disorders were analyzed in relation to the inflammatory markers (table 5), headache was found to be associated with a higher CRP ( $p = 0.0063$ ), serum D-dimer ( $p = 0.002$ ), and fibrinogen levels ( $p = 0.0159$ ), whereas anosmia-dysgeusia, myalgia, and dizziness were not.

**Table 4** Analysis of the Main Neurologic Symptoms According to Sex<sup>a</sup>

Symptom (n)	Females, n (%)	Males, n (%)	OR (95% CI) <sup>b</sup>
Anosmia-dysgeusia (44)	20 (52.63)	24 (38.71)	1.76 (0.78–3.98)
Headache (44)	24 (63.16)	20 (32.26)	3.60 (1.54–8.40)
Myalgia (43)	22 (57.89)	21 (33.87)	2.68 (1.17–6.17)
Dizziness (36)	18 (47.37)	18 (29.03)	2.20 (0.95–5.10)

Abbreviations: OR = odds ratio; CI = confidence interval.

<sup>a</sup> Encephalopathy (n = 8), syncope (n = 7), seizures (n = 3), and vertigo (n = 2) were not included due to their low frequency.

<sup>b</sup> The OR and 95% CI of the presence of each symptom of females vs males.

Regarding patients' age, the younger the age, the higher the risk of having anosmia-dysgeusia (OR [95% CI] = 1.95 [1.02–1.09]) and headache (OR [95% CI] = 1.07 [1.03–1.11]), whereas there were no differences in myalgia and dizziness.

The variables were also analyzed in terms of disease severity as reflected by the CURB-65 scoring system. Of interest, patients with a low CURB-65 score (0–1) showed a higher risk of having anosmia-dysgeusia (OR [95% CI] = 4.03 [1.59–10.24]) or headache (OR [95% CI] = 4.79 [1.83–12.56]) than those with a more severe condition (CURB-65 2–5).

Myalgias were present in 43% of patients, although CK was increased in only 23%, with a median value of 320.5 U/L. We found no association between myalgias and CK values. Eleven of the patients (84.6%) with elevated CK levels compared with only 4 (40%) of the patients with normal CK levels had myalgia.<sup>11</sup> No case of rhabdomyolysis was observed.

## Discussion

We present the results of a prospective unselected series of neurologic manifestations in 100 consecutive patients hospitalized for respiratory COVID-19. Our hospitals serve an area of an extremely high case load, one of the largest of this current pandemic.

We found that neurologic manifestations are very frequent in hospitalized patients with COVID-19. Eighty-eight percent of patients present at least 1 neurologic manifestation. This in contrast with the 36% described in a series from Wuhan, the city origin of the initial outbreak.<sup>5</sup> In their series, data were retrospective, and their patients were not evaluated directly by neurologists, so this may account for this large difference.

Our results provide the spectrum of neurologic manifestations in unselected patients admitted with respiratory COVID-19 disease, which included anosmia-dysgeusia and headache in 44% of patients, myalgia in 43%, and dizziness in 36%. Less common were encephalopathy (8%), syncope (7%), seizures

**Table 5** Association of Inflammatory Parameters With the Main Neurologic Symptoms<sup>a</sup>

Symptom (n)	CRP	Serum D-dimer levels	Serum ferritin levels	Serum fibrinogen levels
	Median (IQR)	Median (IQR)	Median (IQR)	Median (IQR)
<b>Anosmia-dysgeusia</b>				
No	91.8 (47.6–215)	1235 (598–3148)	728.5 (376.5–1606.5)	690 (604.5–778)
Yes	87.5 (28.58–197.5)	950 (480–1975)	969 (382–1974)	673 (548–783)
<b>p Value</b>	0.5538	0.4204	0.3465	0.7371
<b>Headache</b>				
No	143.9 (57.6–237.5)	1515 (734.5–3740)	962 (412–1713)	709.5 (629–799.5)
Yes	61.6 (21–107.4)	700 (410–1650)	677 (291–1567)	629 (533–724)
<b>p Value</b>	0.0063	0.0020	0.3410	0.0159
<b>Myalgia</b>				
No	91.4 (37.9–215)	1330 (520–3940)	959.5 (389–1678.5)	705 (604.5–832)
Yes	80 (32.1–213.5)	820 (490–1840)	628 (360–1971)	650 (526–758)
<b>p Value</b>	0.5267	0.1440	0.4896	0.0510
<b>Dizziness</b>				
No	84 (37–180)	1080 (495–3080)	780 (358–1772)	677.5 (609–794)
Yes	91 (37.9–232)	985 (500–2815)	988.5 (447.5–1618)	688 (505–775)
<b>p Value</b>	0.8456	0.8635	0.5455	0.5120

Abbreviations: CRP = C-reactive protein; IQR = interquartile range.

For the comparison of the inflammatory parameters according to neurologic symptoms, the nonparametric Wilcoxon test was used.

<sup>a</sup> Encephalopathy (n = 8), syncope (n = 7), seizures (n = 3), and vertigo (n = 2) were not included due to their low frequency.

(2%), and stroke (2%). We believe that examination of patients with COVID-19 by neurologists resulted in the higher prevalence of neurologic manifestations in this series.

Headache and myalgia were significantly more frequent in women, whereas anosmia-dysgeusia and dizziness showed no sex preference. Anosmia-dysgeusia and headache patients were younger, whereas no differences were present in myalgia and dizziness. Headache was associated with increased inflammatory parameters (serum CRP, D-dimer, and fibrinogen), but anosmia-dysgeusia, myalgia, and dizziness were not. Anosmia-dysgeusia and headache were significantly more frequent in patients with less severe disease, and the opposite was true with headache in a retrospective Chinese series, but in neither case, the difference reached statistical significance.<sup>5</sup> Myalgias were present in 43%, whereas CK was increased in only 23%.

The frequency and variety of the neurologic problems associated with COVID-19 come from the initial series of respiratory patients of this pandemic. For the purposes of comparison, we pooled data from 7 series.<sup>1–5,8,10</sup> The aggregated results revealed a lower prevalence of myalgias (median of 27.5%) and headache in 8%. Anosmia was not mentioned. That these patients were attended by non-neurologists likely led to an underestimation of the neurologic problems of which only the most severe were

recognized. Of interest, when data from a retrospective study of 214 patients were reanalyzed by neurologists, the numbers of and types of manifestations increased.<sup>5</sup>

Anosmia or some degree of smell dysfunction has been reported in a wide range of 5.6%–85.6% of patients, but the method of detection is critical, ranging from self-reports to the administration of a standardized specific questionnaire.<sup>5,12–14</sup> Forty-four of our patients complained of anosmia-dysgeusia by clinical interview, which likely reflects the figures expected in clinical practice. It should be mentioned that anosmia is not specific for this infection. In a series of 143 consecutive patients with smell dysfunction, 87 of them have had a prior influenza-like illness, a clinical entity known as post-influenza-like hypogeusia and hyposmia, with an estimated incidence of 1 per 400 inhabitants in the United States.<sup>15</sup> The olfactory sensory neurons do not express 2 key genes coding for angiotensin-converting enzyme 2 (ACE-2) and transmembrane serine protease 2 (TMPRSS 2), two proteins involved in SARS-CoV-2 entry. In contrast, olfactory epithelial support cells and nasal respiratory epithelium express both of them, suggesting a non-neural mechanism for anosmia.<sup>16</sup>

Headache was present in 44% of the patients, and this number contrasts with previous series, where headache was

present in an average of 12% (pooled from references 1–4, 9). Headache was present in 13.1% in a retrospective series of hospitalized patients,<sup>5</sup> aimed at studying the neurologic manifestations, and was more frequent in severe patients (17% vs 10.3 in nonsevere patients). Headache characteristics have not been defined so far, and therefore, the questioning by a neurologist is of interest. More often, patients complained of a moderate-intensity headache, persistent, dull, and oppressive, resembling a tension-type headache. Most of the patients in our series did not have a history of headache; thus, the headaches may have an origin in the infectious process and did not reflect the worsening of a preexisting primary headache disorder. We found an interesting association of headache with most inflammatory markers, which could suggest a role of inflammation and/or cytokine release in its pathogenesis. Patients who complained of headache had mild-moderate disease severity; this could reflect the fact that in most severe patients, the respiratory problems overshadowed any other symptoms and would be consistent with a lower rate of headaches reported in the initial series of respiratory patients, which tended to have severe disease.

Nonspecific dizziness was present in one-third of the patients, defined as unsteady gait without other focal neurologic findings; only 2 of them had characteristics of vertigo, thus suggesting that dizziness in this population represents a nonspecific symptom in the context of a systemic infection rather than a proper vestibular dysfunction. Our frequency is higher than reported from Wuhan (9%).<sup>9</sup>

Myalgia was present in 43% of patients, was more frequent in women ( $p = 0.0185$ ), and was not associated with inflammatory markers. An average of 27% with myalgia was reported in pooled data from prior series of patients<sup>1–3,10,17,18</sup> and in 10.7% of patients in a retrospective Chinese series.<sup>5</sup> The lack of association between myalgias and CK values suggests that muscle pain was not due to muscle necrosis but responded to a different origin, perhaps to the effects of systemic inflammation and cytokine release on neuromuscular structures. This aspect will require further clarification.

Less common in this series were encephalopathy, seizures, syncope, and stroke. Although these manifestations were of low frequency, they present a challenge for their severity and life-threatening potential. These problems may be more common in particularly severe patients, often in the ICU setting,<sup>19</sup> which were not included in this study.

Encephalopathy was encountered in 9 patients, 8 of them with severe disease, and was associated with the presence of fever. Two mechanisms are plausible in this scenario, the indirect effect of the inflammatory cytokine storm of these patients, and the direct effect of the SARS-CoV-2 virus. Although direct viral CNS damage could also contribute to encephalopathy and other neurologic disturbances, RT-PCR in the CSF has been negative in most reported cases so far.

Furthermore, ACE2, the receptor for the virus has only been demonstrated in the endothelium of the nervous system, but not in parenchymal cells.<sup>20</sup> The only evidence of direct CNS involvement comes from SARS 2002–2003 where viral presence in the brain was detected with real-time RT-PCR, in situ hybridization, and electron microscopy.<sup>21,22</sup> It is possible that there was greater severity in the neurologic complications of some patients not included in our series, including encephalopathy, as we excluded 17 patients with severe cognitive issues, and we did not approach patients with COVID-19 in the ICU setting.

Seizures were present in 2 patients in our series and have been reported in a few patients to date<sup>5,8,23–25</sup>; we suspect that may be underreported. These numbers may be higher in a series of critically ill patients. No evidence of structural lesions or encephalitis has been found in these patients except for 1 patient, a situation that also occurs in influenza infections.<sup>26</sup>

Loss of consciousness with characteristics of syncope was present in 7% of the patients, but this condition has not been observed in published series so far. That the information in this study was directly obtained by neurologists in contrast to other specialties less familiar with this condition may reflect this disparity. Impaired consciousness was reported in 7.5% in a retrospective Chinese series, but it included somnolence, stupor, coma, and confusion.<sup>5</sup> Syncope did not associate with myoclonic jerks and was of the neurally mediated characteristics in all patients without clear evidence of cardiac origin. However, this needs further clarification because myocardial inflammation and arrhythmias may appear in these patients, could be potentially lethal, and could present as syncope.

Stroke (minor) was present in 2 patients of our series. The true relationship between COVID-19 and stroke incidence remains to be determined.<sup>27</sup> Despite the fall in stroke admissions, it has been suggested that COVID-19 infection itself may cause stroke. Stroke was reported in 6 cases (2.8%) in a Wuhan series,<sup>5</sup> a frequency similar to ours; all but 1 case occurred in the severe infection group; 5 were ischemic and 1 hemorrhagic. In a New York study, 0.9% of patients hospitalized for COVID-19 infection had a stroke.<sup>28</sup>

The frequent coagulation abnormalities of patients with COVID-19 together with an exaggerated systemic inflammation and virus-related cardiac injury may all contribute to stroke.<sup>27,29</sup> Accordingly, stroke severity seems increased in patients with COVID-19 and with poorer outcomes,<sup>30,31</sup> which contrasts with our 2 patients. However, this needs further studies.

Our study had some limitations: the collection of data had to be balanced with the exigencies of patient care. Moreover, as these patients were in isolation, we were limited as to our ability to perform nonessential examinations or blood collections. As there was no follow-up, our data reflect the observations of a single point in the evolution of the hospitalized patient. We did not include patients in the ICU, so it is

## TAKE-HOME POINTS

- Neurologic manifestations are very frequent (88%) in nonselected patients hospitalized in a non-ICU setting for COVID-19 disease.
- The most frequent neurologic disturbances in patients with COVID-19 included anosmia, headache (44% each), myalgias (43%), dizziness or vertigo (36%). Less common are encephalopathy (9%), syncope (7%), and seizures and stroke (2% each).
- They represent a diagnostic challenge for the attending neurologist, particularly considering the difficulties in performing routine ancillary examinations in these patients.
- Neurologists should be alert to these neurologic complications because they may represent the debut of an undiagnosed SARS-CoV-2 infection. They can also usually represent a severe condition with evidence of systemic inflammation and coagulation abnormalities that require prompt and specific management measures.

possible that the more severe neurologic manifestations of COVID-19 disease are underrepresented.

In summary, our series disclosed a significant rate of neurologic involvement in patients with COVID-19, and when added to the increasing reports of serious cases of Guillain-Barré syndrome,<sup>32,33</sup> Miller-Fisher syndrome,<sup>34</sup> acute necrotizing encephalopathy,<sup>35</sup> myelitis,<sup>36</sup> and encephalitis (1 patient with positive RT-PCR in the CSF),<sup>25</sup> make a strong case for the neurologic consult in the management of patients with COVID-19.

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### Appendix Authors

Name	Location	Contribution
Juan Carlos García-Moncó, MD, PhD, FAAN	Basurto University Hospital	Designed and conceptualized the study; analyzed the data; and drafted the manuscript for intellectual content

### Appendix (continued)

Name	Location	Contribution
Antonio Cabrera Muras, MD	Basurto University Hospital	Major role in the acquisition of data
Markel Erburu Iriarte, MD	Basurto University Hospital	Major role in the acquisition of data
Patricia Rodrigo Armenteros, MD	Basurto University Hospital	Major role in the acquisition of data
Alejandra Collía Fernández, MD	Basurto University Hospital	Major role in the acquisition of data
Javier Arranz-Martínez, MD	Hospital de Galdakao-Usansolo	Major role in the acquisition of data
Solange Kapetanovic, MD	Basurto University Hospital	Major role in the acquisition of data
Ana Lorenzo-García, MD	Basurto University Hospital	Major role in the acquisition of data
Amaia Bilbao González, PhD	Basurto University Hospital	Performed the statistical analysis and interpreted the data
Marian Gomez-Beldarrain, MD	Hospital de Galdakao-Usansolo	Analyzed the data and drafted the manuscript for intellectual content

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