Diaphragm impairment in COVID-19
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Abstract
Among patients affected by the virus COVID-19, physicians have observed ventilation disorders. It is relevant to assess neurological involvement, including the role of diaphragmatic function. Its possible impairment could be related to the systemic inflammatory response and disease progression that both typify COVID-19 infection. We distinguished two groups (severe group (SG) and mild group (MG)) according to the severity of respiratory symptomatology. We performed neurophysiological and sonography studies to evaluate the diaphragmatic function. Regarding the sonography variables, we identified statistically significant differences in the right mean diaphragmatic thickness along with the expiration, showing 1.56 mm (SEM: 0.11) in the SG vs 1.92 mm (SEM: 0.19) in the MG (p = 0.042). The contractibility of both hemidiaphragms was 15% lower in the severe group, though this difference is not statistically significant. In our examination of the neurophysiological variables, in the amplitude responses, we observed a greater difference between responses from both phrenic nerves as follows: the raw differences in amplitude were 0.40 µV (SEM: 0.14) in the SG vs 0.35 µV (SEM: 0.19) in the MG and the percentage difference was 25.92% (SEM: 7.22) in the SG vs 16.28% (SEM: 4.38%) in the MG. Although diaphragmatic dysfunction is difficult to detect, our combined functional and morphological approach with phrenic electroneurograms and chest ultrasounds could improve diagnostic sensitivity. We suggest that diaphragmatic dysfunction could play a relevant role in respiratory disturbance in hospitalised patients with severe COVID-19.

Key Words: Coronavirus; diaphragm; electromyography; phrenic nerve; ultrasound.
breathing problems, we have assessed diaphragmatic function via neurophysiological and sonography analyses.

Materials and methods

This study has been accepted by the Research Ethics Committee from La Princesa Hospital, with local reference number 4144, and it was carried out following the legislation in force regarding confidentiality. All included patients (or their representatives) were informed about the study and provided oral informed consent as proposed by the Agencia Española de Medicamentos y Productos Sanitarios because of the COVID-19 emergency.

We distinguished two groups (severe and mild) of patients affected by COVID-19 disease hospitalized at La Princesa University Hospital, according to the severity of respiratory symptoms. The severe group consisted of patients who experienced dyspnoea on mild exertion, pain of diaphragmatic origin or hiccup. The mild group consisted of patients who experienced only dyspnoea and only during severe exertion. We performed electroneurograms of both phrenic nerves and sonography of both hemidiaphragms, and we

Table 1. Demographic and clinical data comparison of mild and severe groups.

| IBM          | Mild (n = 10) | Severe (n = 9) | p-value |
|--------------|--------------|----------------|---------|
| Underweight  | 0 (0%)       | 1 (11.1%)      | 0.90    |
| Normal weight| 4 (40%)      | 1 (11.1%)      |         |
| Overweight I | 2 (20%)      | 3 (33.3%)      |         |
| Overweight II| 2 (20%)      | 3 (33.3%)      |         |
| Obesity I    | 2 (20%)      | 1 (11.1%)      |         |
| HBP Yes      | 3 (30%)      | 0 (0%)         | 0.29    |
| HBP No       | 7 (70%)      | 9 (100%)       |         |
| Diabetes Mellitus Yes | 2 (20%) | 1 (11.1%) | 0.77    |
| Diabetes Mellitus No    | 8 (80%)      | 8 (88.8%)      |         |
| Dyslipidemia Yes | 6 (60%)      | 1 (11.1%)      | 0.08    |
| Dyslipidemia No    | 4 (40%)      | 8 (88.8%)      |         |

Table shows distribution of clinical characteristics, and statistical comparison between mild and severe groups. IBM = Index Body Mass, HBP = High Blood Pressure (>120/80mmHg).

Table 2. Demographic, clinical, respiratory variables at time of testing and clinical course of the mild group.

| Patient | Genre | Age | Days of symptoms | Symptoms suggesting phrenic nerve damage | Respiratory variables | Clinical course |
|---------|-------|-----|------------------|------------------------------------------|-----------------------|----------------|
|         |       |     |                  | Ventalation disturbances | Costal pain | Hiccups | Oxygen Saturation | Supplemental Oxygen |                     |
| 1       | M     | 76  | 7                | No | No | No | 95% | No | Good |
| 2       | M     | 85  | 9                | No | No | No | 97% | No | Good |
| 3       | M     | 43  | 7                | No | No | No | 98% | No | Good |
| 4       | F     | 69  | 8                | No | No | No | 96% | NC 4 L | Good |
| 5       | F     | 48  | 11               | Yes | No | No | 99% | NC 2 L | Good |
| 6       | M     | 54  | 11               | No | No | No | 96% | No | Good |
| 7       | M     | 31  | 6                | No | No | No | 96% | No | Good |
| 8       | M     | 68  | 17               | Yes | No | No | 92% | NC 4 L | ICU during 6 days -> Good at discharge |
| 9       | M     | 41  | 6                | Yes | No | No | 95% | No | Good |
| 10      | F     | 59  | 8                | No | No | No | 95% | NC 2 L | Good |

In one patient (number 9), we performed only nerve conduction of both phrenic nerves performing sonography during the hospital admission. MEP = Motor Evoked Potential, Dif = Difference, M = Male, F= Female, NC = Nasal Cannula, ICU = Intensive Care Unit.
recorded the following sociodemographic variables and primary comorbidities among patients: age, sex, body mass index (BMI), presence of high blood pressure, diabetes mellitus, previous diagnosis of polyneuropathy or myopathy (clinically or electromyographically) and history of stroke or cerebral haemorrhage prior to the infection. The inclusion criteria were:
- Age > 18 years.
- COVID-19 disease confirmed by a positive Polymerase Chain Reaction (PCR) from a nasopharyngeal swab.
- Patients (or their legal guardians) had to consent and sign the informed consent document.

The exclusion criteria were:
- History of polyneuropathy or myopathy.
- History of stroke or cerebral haemorrhage prior to the COVID-19 disease.

To determine the presence of diaphragmatic injury, we performed electromyography of both phrenic nerves and diaphragmatic sonography. The neurophysiological variables collected were as follows: the latency of motor evoked potentials of both right and left phrenic nerves, the difference in the latency in each side as the percentage of this difference, the amplitude of the motor evoked potentials of both right and left phrenic nerves, and the difference in the amplitude in each side as the percentage of this difference, as Vincent et al. previously described.\(^{14}\) The sonography variables evaluated in each group were the right and left diaphragm thickness during inspiration and expiration, hemidiaphragm displacement, and expiratory fraction from both sides.

To evaluate the diaphragmatic function, we performed both techniques either in imaging tests (e.g. ultrasounds)
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Sonography technique
We positioned the patient in the supine position to evaluate the diaphragmatic movement in the respiratory effort. We used mode B (bidimensional mode) to assess the diaphragm’s thickness and mode M (movement mode) to evaluate the diaphragmatic displacement. We measured the diaphragmatic thickness of both the right and left sides along with the maximum inspiration and at the end of the expiratory effort. Similarly, we calculated dynamic changes, the percentage of contraction of each diaphragm and each side’s expiratory fraction.

Statistical analysis
Quantitative data are shown by the mean and the standard error of the mean (SEM). The differences in latency and hemidiaphragm amplitude are shown in percentages. To compare quantitative measures related to electromyographical and sonography studies, we studied their normality via the Kolmogorov–Smirnov test. In the case of fulfilling criteria of normality, we used a t-test. In the contrary case, we used the Mann–Whitney U test and the Wilcoxon signed-rank test. The statistics were performed by the statistical software XLSTAT (Addinsoft®). We considered p ≤ 0.05 statistically significant.

Results
We recruited a total of 19 patients with confirmed diagnosis of both COVID-19 and dyspnoea: 10 with mild disease (the mild group, or MG) and 9 with severe disease (symptoms suggesting diaphragmatic disturbance; the severe group, or SG). The sample consisted of 12 men (63.2%) and 7 women (36.8%),

Table 5. Demographic, clinical, respiratory variables at the time of testing and clinical course of the severe group.

| Patient | Genre | Age | Days of symptoms | Ventilation disturbances | Symptoms suggesting phrenic nerve damage | Respiratory variables | Clinical course |
|---------|-------|-----|------------------|-------------------------|------------------------------------------|-----------------------|----------------|
| 1       | M     | 65  | 93               | Yes                     | No                                       | 92%                   | ICU -> Death   |
| 2       | M     | 51  | 10               | Yes                     | No                                       | 96%                   | Chronic course: oxygen therapy and corticosteroids at discharge |
| 3       | M     | 58  | 7                | Yes                     | No                                       | 97%                   | Good          |
| 4       | F     | 33  | 7                | Yes                     | No                                       | 95%                   | Good          |
| 5       | M     | 61  | 6                | Yes                     | No                                       | 92%                   | Good          |
| 6       | M     | 39  | 9                | Yes                     | Yes                                     | 90%                   | ICU -> Death  |
| 7       | F     | 62  | 7                | Yes                     | Yes                                     | 93%                   | ICU (PTE) -> Good at discharge |
| 8       | F     | 59  | 9                | Yes                     | No                                       | 94%                   | Good          |
| 9       | M     | 42  | 6                | Yes                     | No                                       | 94%                   | Good          |

Table 6. Sonography variables of the severe group

| Patient | Right diaphragmatic thickness on inspiration | Left diaphragmatic thickness on inspiration | Right diaphragmatic thickness on expiration | Left diaphragmatic thickness on expiration | Right contraction on inspiration (%) | Left contraction on inspiration (%) | Right expiratory fraction | Left expiratory fraction |
|---------|---------------------------------------------|--------------------------------------------|--------------------------------------------|--------------------------------------------|-------------------------------------|-----------------------------------|----------------------------|----------------------------|
| 1       | 1.9                                         | 1.0                                        | 1.9                                        | 0.9                                        | 0.0                                 | 16.3                              | 0.0                        | 0.1                        |
| 2       | 2.7                                         | 2.7                                        | 1.9                                        | 1.6                                        | 42.1                                | 68.8                              | 0.4                        | 0.7                        |
| 3       | 2.2                                         | 3.5                                        | 1.2                                        | 1.3                                        | 83.3                                | 169.2                             | 0.8                        | 1.7                        |
| 4       | 2.7                                         | 3.0                                        | 1.4                                        | 2.3                                        | 96.4                                | 27.7                              | 1.0                        | 0.3                        |
| 5       | 2.6                                         | 2.5                                        | 1.8                                        | 1.6                                        | 39.9                                | 60.3                              | 0.4                        | 0.6                        |
| 6       | 1.5                                         | 1.8                                        | 1.2                                        | 1.5                                        | 28.3                                | 21.5                              | 0.3                        | 0.2                        |
| 7       | 1.8                                         | 1.4                                        | 1.3                                        | 0.8                                        | 38.3                                | 70.2                              | 0.3                        | 0.7                        |
| 8       | 2.6                                         | 2.2                                        | 1.3                                        | 1.5                                        | 100.0                               | 45.2                              | 1                          | 0.4                        |
| 9       | 2.9                                         | 3.1                                        | 2.0                                        | 2.1                                        | 49.5                                | 48.5                              | 0.5                        | 0.5                        |
with a mean age of 56.4 years and no significant differences between the groups. Regarding the clinical characteristics, we observed a predominance of normal weight in the mild group (40% vs 11.1%), but a similar number of overweighted or obese patients (6 patients in the mild group and seven patients in the severe group). Among other cardiovascular variables, Diabetes Mellitus had similar distribution between both groups; however, none of the patients from the severe group suffered from HBP, and three patients from the mild group suffered from this disease. For dyslipidemia, there are more affected patients in the severe group (8 patients) than in the mild group (4 patients). Completed raw data (sonography and electroneurography variables) and personal history of all patients are shown in the tables 1, to 8. In our assessment of the clinical symptoms in each group of patients, all 10 SG patients demonstrated dyspnoea at a low effort of exertion, 4 presented with costal pain of probable diaphragmatic origin and 1 exhibited hiccups. In the MG, only three of the nine patients presented with ventilation disorder, described as a dyspnoeic sensation at high effort, and none reported diaphragmatic pain or hiccups. Only three patients described costal pain related to cough reflex and lumbar pain because of hospital convalescence. In our examination of the neurophysiological variables, in the amplitude responses, we observed a greater difference between responses from both phrenic nerves

**Table 7. Neurophysiological variables of the severe group**

| Patient | Neurophysiological variables |
|---------|-----------------------------|
|         | Right MEP Lat | Left MEP Lat | Right MEP Amp | Left MEP Amp | Lat Dif (%) | Amp Dif (%) |
| 1       | 7.4 ± 0.46     | 7.2 ± 0.46   | 1.7           | 0.7          | 1.4         | 41.7        |
| 2       | 8.7 ± 0.46     | 8.3 ± 0.46   | 0.3           | 0.1          | 2.3         | 50.0        |
| 3       | 6.9 ± 0.46     | 6.5 ± 0.46   | 0.3           | 1.2          | 3.0         | 60.0        |
| 4       | 7.8 ± 0.46     | 8.3 ± 0.46   | 0.4           | 0.2          | 3.1         | 33.3        |
| 5       | 7.8 ± 0.46     | 8.1 ± 0.46   | 0.6           | 0.4          | 1.9         | 20.0        |
| 6       | 9.3 ± 0.46     | 9.2 ± 0.46   | 1.9           | 2.8          | 0.5         | 19.1        |
| 7       | 9.8 ± 0.46     | 9.9 ± 0.46   | 1.7           | 1.7          | 0.5         | 0.0         |
| 8       | 7.1 ± 0.46     | 7 ± 0.46     | 0.4           | 0.4          | 0.7         | 0.0         |
| 9       | 5.2 ± 0.46     | 5.2 ± 0.46   | 1             | 1.2          | 0.0         | 9.1         |

**Table 8. Nerve conduction study and variables of severe and mild groups**

|                  | Mild group (mean ± SEM) | Severe group (mean ± SEM) | p-value |
|------------------|-------------------------|---------------------------|---------|
| Right Latency (ms) | 7.83 ± 0.46             | 7.78 ± 0.46               | 0.759   |
| Left Latency (ms)  | 8.13 ± 0.46             | 7.74 ± 0.48               | 0.843   |
| Right Amplitude (mV) | 0.46 ± 0.08             | 0.92 ± 0.22               | 0.703   |
| Left Amplitude (mV) | 0.60 ± 0.11             | 0.97 ± 0.29               | 0.716   |
| Differences in Latency (%) | 3.09 ± 1.24           | 1.50 ± 0.38               | 0.231   |
| Differences in Amplitude (%) | 16.28 ± 4.38             | 25.92 ± 7.22               | 0.711   |
| Right thickness during inspiration (mm) | 3.11±0.40 | 2.34±0.16 | 0.945 |
| Left thickness during inspiration (mm) | 2.97±0.33 | 2.36±0.27 | 0.547 |
| Right thickness during expiration (mm) | 1.92±0.19 | 1.56±0.11 | 0.042 |
| Left thickness during expiration (mm) | 1.69±0.15 | 1.51±0.16 | 0.742 |
| Right contractibility (%) | 62.15±5.58 | 53.10±11.15 | 0.742 |
| Left contractibility (%) | 73.34±6.74 | 58.62±15.36 | 0.195 |
| Right expiratory fraction | 0.59±0.05 | 0.53±0.11 | 0.674 |
| Left expiratory fraction | 0.73±0.07 | 0.58±0.15 | 0.195 |

The table shows bilateral amplitude and latency results of both phrenic nerves and their difference and thickness of each hemidiaphragm along with inspiration and expiration, contractibility, and expiratory fraction from each hemidiaphragm.
as follows: the raw differences in amplitude were 0.40 µV (SEM: 0.14) in the SG vs 0.35 µV (SEM: 0.19) in the MG and the percentage difference was 25.92% (SEM: 7.22) in the SG vs 16.28% (SEM: 4.38%) in the MG (Table 8 and Figure 1). Sonography results of one patient from the mild group (patient number 9) were not available. Regarding the sonography variables in the remaining patients, we identified statistically significant differences in the right mean diaphragmatic thickness along with the expiration, showing 1.56 mm (SEM: 0.11) in the SG vs 1.92 mm (SEM: 0.19) in the MG ($p = 0.042$). In addition, a lower mean expiratory fraction and hemidiaphragmatic contraction were observed in the SG as follows: the right mean diaphragmatic contraction was 53.10% (SEM: 11.15) in the SG vs 62.15% (SEM: 5.58) in the MG, the left mean diaphragmatic contraction was 58.62% (SEM: 15.36) in the SG vs 73.34% (SEM: 6.74) in the MG, the right mean expiratory fraction was 0.53 (SEM: 0.11) in the SG vs 0.59 (SEM: 0.05) in the MG and the left mean expiratory fraction was 0.58 (SEM: 0.15) in the SG vs 0.73 (SEM: 0.07) in the MG (Table 8; Figures 2 and 3).

Discussion

This study has generated preliminary findings of neurophysiological and sonography data through diaphragmatic assessment. Concerning nerve conduction studies from both phrenic nerves, we observe a greater difference (25.92%) between phrenic nerves in each patient in the SG than in the MG (16.28%). However, these results are not statistically significant. Conversely, we do not observe relevant differences in the latency because of the variability in the length of the thorax among the patients included in the study. The amplitude of the motor evoked potentials evinces individual variability driven by BMI and diaphragmatic fat storage, and thus, the main parameters assessed stem from the percentage of variation between both phrenic nerves.

Sonography data demonstrate a statistically relevant outcome in comparing the right thickness during expiration between both groups, which shows a thicker diaphragm in MG. The other differences in sonography variables between both groups are not statistically significant, probably related to the small sample size. Moreover, thickness in both right and left hemidiaphragms, along with inspiration and expiration, is lesser in the SG. We should consider other sonography variables as sources of the differences in the contractibility of both hemidiaphragms because they present 10-15% lower contraction in the SG, though this difference is not statistically significant. The complexity of the symptoms and the clinical variability produced by COVID-19 require study of these symptoms' physiopathological repercussions, one of the most relevant of which is damage in respiratory function, given their importance in the deleterious progression of the disease. Previous analyses have documented cases that feature peripheral nerve damage, namely, via demyelination and axonal injury, which constitute the most frequent damage in the severest cases. These conditions also result from other viral infections, such as the varicella-zoster virus, Cytomegalovirus, human immunodeficiency virus (HIV), hepatitis E, or secondary infection with the arbovirus dengue. Based on the information described in prior studies of these viral infections, we could infer the similarity of neurological symptoms that can appear in patients infected with SARS-CoV-2. Just as generalised nerve damage may occur along the progression of COVID-19 disease, focal neuropathies are another attested consequence. Phrenic nerve damage is an example of immune-mediated focal neuropathies, which carry focal demyelination or axonal degeneration. The injury of the peripheral nerves is described in several viral infections, as the infection by
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As examined in patients and animal models infected with other coronaviruses, such as those that cause SARS and Middle East respiratory syndrome, the brain stem is one of the primarily infected nervous structures. The more typical entry of SARS-COV-2 is intranasal, hence its propagation to the central nervous system. The brain stem’s direct damage, where the pneumotaxic, apneustic and bulbar respiratory centres deteriorate, could explain the ventilatory disturbance. Diaphragm paralysis involves a loss of muscle strength of the primary muscle implied in the inspiration. We could locate the damage at the muscle fibres’ level or phrenic nerve damage. It has been described several neurological causes: polyneuropathic Guillain–Barré syndrome, inflammatory causes, neuralgic amyotrophy or Parsonage–Turner syndrome. An infection can also cause dysfunction akin to the symptoms of West Nile or dengue. We could not dismiss this possible injury in patients affected with COVID-19. The main imaging test performed in patients with suspected COVID-19 and symptoms suggesting damage in respiratory function is thorax radiography to confirm possible viral pneumonia. However, this test has a low positive predictive value in detecting the diaphragmatic damage, to evaluate its dysfunction. Given these findings, the phrenic nerve’s electroneurography helps us confirm the appropriate conduction of these nerves. Other authors have observed a high diagnostic correlation between the diaphragm’s thickness observed in sonography and the amplitude of the motor evoked potential in electroneurography. Therefore, we have performed both techniques to improve this test’s sensitivity.

Fig 2. Ultrasound images of a patient from the mild group. Right hemidiaphragm displacement measured during deep inspiration, using M-mode (A) and B-Mode (B) ultrasonography. Diaphragm thickness measured at end-expiration using B-mode (C).

Fig 3. Ultrasound images of a patient from the severe group. Right hemidiaphragm displacement measured during deep inspiration, using M-mode (A) and B-Mode (B) ultrasonography. There is a shorter displacement compared to the patient from Figure 2, suggesting phrenic dysfunction. Diaphragm thickness, measured at end-inspiration using M-mode (C) and B-mode ultrasound (D).
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diagnostic sensibility in detecting the diaphragmatic dysfunction and the phrenic nerve damage. The patients with more severe respiratory symptoms have dyspnoea, cough, thoracic pain and hiccups.37,38 We observe a particular trend in the presentation of dyspnoea and thoracic pain among patients with diaphragmatic dysfunction, all of whom have dyspnoea at low effort (100%) and of whom have thoracic pain in this group (44.4%). In the MG, only three patients exhibit dyspnoea, always at high effort (30%), and without symptoms suggesting diaphragmatic-origin pain in any case. As described in other reports, we note the possibility of disturbance of the nervous system in the disease’s progression. In our data series, we observe an amplitude of differences between both conduction studies from phrenic nerves, which could be explained by axonal degeneration in the acute phase of this viral infection, as happened with other viruses.27–29 Moreover, we identify a greater difference in the contraction of both hemidiaphragms in patients affected with a moderate degree of the disease and a lesser thickness in the SG, as described in neuromuscular damage from other viral infections.4,13
Because this study features a small sample size, data observed have insufficient power to detect a statistical significance between mild and severe groups. Nevertheless, these results allow us to point out the importance of the diaphragm’s role in patients with more respiratory dysfunction, which should engender stronger rehabilitation measures to improve the progression of patients infected with COVID-19. This report aims to raise awareness about the role of diaphragmatic damage in patients affected by COVID-19. During treatment of these patients, it is important to bear in mind the possible diaphragmatic injury that may necessitate rehabilitation therapies. We should assess this disturbance’s progression and each patient’s evolution in the long term as a respiratory sequel. In conclusion, diaphragmatic dysfunction could play a relevant role in respiratory disturbance in hospitalised patients with severe COVID-19. Although diaphragmatic dysfunction is difficult to detect, the combined functional and morphological approach with phrenic electroneurograms and chest ultrasounds could improve diagnostic sensitivity. If the implication of diaphragmatic dysfunction in COVID-19 disease is confirmed, the ventilatory rehabilitation approach will allow an additional therapeutic option in treatment of this serious disease.

List of acronyms
BMI - Body Mass Index
COVID-19 - coronavirus disease 2019
HIV - Human Immunodeficiency Virus
MG - Mild Group
SARS - Severe Acute Respiratory Syndrome
SEM - Standard Error of the Mean
SG - Severe Group

Contributions of Authors
LL-V: Design and conceptualized study, analyzed the data, interpreted the data and drafted the manuscript for intellectual content; JV-V, ER-M, PG-G, RW-R: Acquisition of data and revised the manuscript for intellectual content; EdlRS: Acquisition of data; JMG-R: Design and conceptualized study and revised the manuscript for intellectual content.

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Ethical Publication Statements
We confirm that we have read the journal’s position on ethical issues involved in publication and affirm that this report is consistent with those guidelines.

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