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he etiology and pathogenesis of preeclampsia (PE) are not completely understood; however, deficient trophoblast invasion seems to play a leading role.1 The resulting placental ischemia leads to an angiogenic imbalance in maternal serum, with increased antiangiogenic factors, such as soluble fms-like tyrosine kinase 1 (sFlt-1), and decreased proangiogenic factors, such as placental growth factor (PIGF).2 Some studies have shown that this angiogenic status might induce endothelial cell dysfunction, which contributes to the development of PE with its signs and symptoms (hypertension, proteinuria, thrombocytopenia, elevated liver enzymes, and neurologic disturbances).3 Several disorders have previously been demonstrated to mimic the clinical features of PE, including acute fatty liver, thrombotic thrombocytopenic purpura, hemolytic uremic syndrome, acute exacerbation of lupus erythematosus, and COVID-19.4−7. These conditions share some clinical and laboratory findings with PE, and its diagnostic criteria may overlap; therefore, to provide adequate care, a differential diagnosis, although challenging, is necessary. Some studies have shown that the sFlt-1−PIGF ratio is a highly specific marker for placental dysfunction and may help to accurately discriminate PE from some of its imitators.6,8,9 Since the outbreak of the COVID-19 pandemic, some studies have reported an increased PE incidence in pregnant women with SARS-CoV-2 infection, and several explanations for this association have been proposed.10−13 On May 2020, a small case series published by Mendoza et al1 described a PE-like syndrome induced by severe COVID-19 during pregnancy. That study showed that up to 62.5% of pregnant women with
severe pneumonia because of SARS-CoV-2 infection may go on to develop features of PE. Nevertheless, among the 5 cases with signs and symptoms of PE, only 1 case had abnormal angiogenic status, suggesting that the remaining 4 cases probably did not have PE. Of these 4 cases, only 1 patient was still pregnant after recovering from pneumonia, and all signs and symptoms of PE had disappeared. Given that PE does not resolve spontaneously, and delivery is the only definitive treatment, severe COVID-19-induced PE-like syndrome was the most plausible explanation for this case.

Therefore, only 1 confirmed case of PE-like syndrome was described in that study at the beginning of the COVID-19 pandemic; this syndrome has not been confirmed in larger studies, and its effect on perinatal outcomes has not been studied. Therefore, this study aimed to confirm the existence of a PE-like syndrome in a larger cohort of women with severe COVID-19 and to investigate its implications for pregnancy outcomes and prognosis.

Materials and Methods
This was a prospective cohort study conducted at Vall d’Hebron Barcelona Hospital Campus, Barcelona, Spain, from February 2020 to September 2021. Our site was a referral hospital for severe COVID-19 pneumonia, especially among pregnant patients. The inclusion criteria were pregnant women admitted to the intensive care unit (ICU) for severe pneumonia because of SARS-CoV-2 infection during pregnancy. SARS-CoV-2 infection was confirmed by a real-time polymerase chain reaction of nasal and pharyngeal swabs. Participants who met the PE diagnostic criteria and had normal sFlt-1−to−PlGF ratio (<38) were not delivered for PE and were classified as patients with PE-like syndrome. To establish whether the prognosis of women with PE-like syndrome was more similar to that of women with PE or to that of normotensive women, pregnancy outcomes were compared with 2 reference groups: women with true PE (met the PE criteria and had an sFlt-1−to−PlGF ratio of >38) and women who did not meet the PE criteria and did not deliver during the ICU stay (non-PE group). As comparing women with PE-like syndrome with women without PE and electively delivered because of pneumonia during the ICU stay did not provide additional information on the prognosis of patients with PE-like syndrome, these women were excluded from this study (Figure).

Gestational age was confirmed by fetal crown-rump length measurement at the first-trimester scan. Participants were examined at 3 different times: before severe pneumonia, during severe pneumonia (at the time of ICU stay), and after recovery from severe pneumonia. Demographic characteristics, obstetrical and maternal history, and biophysical and biochemical markers were collected. Biophysical and biochemical data included systolic blood pressure (SBP; mm Hg), diastolic blood pressure (DBP; mm Hg), hypertension (SBP of >140 mm Hg and/or DBP of >90 mm Hg), mean uterine artery pulsatility index (PI) percentile by transabdominal Doppler ultrasound, proteinuria (in 24-hour urine samples, spot urine protein-to-creatinine ratio, and/or dipstick urinalysis), platelet count (µL), aspartate aminotransferase (AST; U/L), alanine aminotransferase (ALT; U/L), lactate

FIGURE
Inclusion flowchart

| sFlt-1/PlGF ≤ 38 | sFlt-1/PlGF > 38 |
|------------------|------------------|
| Preeclampsia-like group (n=8) | Preeclampsia group (n=7) |
| No delivery during severe pneumonia (n=38) | Non-Preeclampsia group (n=53) |

PlGF, placental growth factor ratio; sFlt-1, soluble fms-like tyrosine kinase 1.

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dehydrogenase (LDH; U/L), and creatinine level (mg/dL). In women with suspected PE, maternal serum PI GF and sFlt-1 (pg/mL) were determined using the fully automated Elecsys assays for sFlt-1 and PI GF on an electrochemiluminescence immunoassay platform (cobas e analyzers; Roche Diagnostics, Rotkreuz, Switzerland). The sFlt-1-to-PI GF ratio was calculated, with ≤38 values considered normal and unlikely to be associated with an underlying placental dysfunction.16

PE was defined as new-onset high blood pressure (SBP of >140 mm Hg and/or DBP of >90 mm Hg), worsening of previous high blood pressure in addition to new-onset proteinuria (protein-to-creatinine ratio of >300), worsening of previous proteinuria, or at least 1 of the following signs and symptoms of severe PE: cerebral or visual symptoms, elevation of liver enzymes to twice the normal concentration, platelet count of <100,000/μL, serum creatinine level of >1.1 mg/dL, or pulmonary edema.17 Here, to diagnose hypertension, at least 5 of 6 consecutive measurements had to be abnormal for a 6-hour period. Hemolysis, elevated liver enzymes, and low platelet count (HELLP syndrome) is considered a variant of PE. The diagnostic criteria for the HELLP syndrome were hemolysis with increased LDH (>600 U/L), AST or ALT (≥70 U/L), and platelet counts of <100,000/μL.18 Elective delivery was recommended at ≥37 weeks of gestation in women with PE without severe features and ≥34 weeks of gestation in women with severe PE and/or HELLP syndrome. In COVID-19 cases that met the PE diagnostic criteria and had an sFlt-1-to-PI GF ratio of ≤38, elective delivery because of PE was not recommended; alternatively, these cases were managed according to the clinical evolution of COVID-19. Elective delivery because of severe pneumonia was indicated at ≥36 to 38 weeks of gestation and ≥33 to 35 weeks of gestation if invasive respiratory support was required.

This study was approved by the Vall d’Hebron Barcelona Hospital Campus Ethics Committee (PR[AMI]181/2020) on March 13, 2020. All patients included in the study gave their oral or written consent according to the criteria of our ethics committee.

**Statistical analysis**

Categorical data were reported as frequency and percentage, and comparisons among the groups were estimated by the chi-square or Fisher tests, as appropriate. Continuous variables were reported as median (interquartile range [IQR]). The Mann-Whitney U test was used to assess differences among the groups. The Wilcoxon signed-rank test was used to assess differences of repeated measurements on a single sample. The statistical significance level was set at P<.05, and the Bonferroni correction was applied to compensate for multiple comparisons. The Stata statistical software package was used for data analysis (StataCorp 2019; Stata Statistical Software: Release 16; StataCorp LLC, College Station, TX).

**Results**

During the study period, 106 patients with severe pneumonia because of COVID-19 were admitted to the ICU. Of those, 38 (35.8%) did not meet the inclusion criteria for needing elective delivery during the ICU stay because of clinical worsening of pneumonia and not having clinical suspicion of PE. Therefore, 68 women (64.2%) were included in the study: 7 (6.6%) with PE, 8 (7.6%) with PE-like syndrome, and 53 (50.0%) without PE features and not needing delivery during severe pneumonia (non-PE group). All 7 cases with PE required elective delivery during severe pneumonia (4 [57.1%] because of PE, 2 [28.6%] because of COVID-19 worsening, and 1 [14.3%] because of nonreassuring cardiotocography). In contrast, none of the cases in the PE-like syndrome group or the non-PE group required elective delivery during severe pneumonia. The median (IQR) gestational age at the time of admission to the ICU was significantly higher in the PE group (35.4 [30.3–36.6]) than in the PE-like syndrome group (25.6 [21.3–30]) and non-PE group (27.7 [24.5–31.3]) (P<.01). No other pregnancy baseline characteristic differed among the groups (Table 1).

**TABLE 1**

Demographic characteristics of the study population

| Characteristics         | PE like (n=8) | PE (n=7) | Non-PE (n=53) |
|-------------------------|--------------|----------|---------------|
| Age (y)                 | 31.9 (25.3–33.6) | 33.8 (30–42.4) | 33.6 (31.0–37.2) |
| BMI (kg/m²)             | 32.2 (24.8–36.1) | 28.7 (24.3–29.9) | 28.7 (26.4–32.3) |
| Chronic hypertension (%)| 0/8 (0)      | 0/7 (0)   | 1/52 (1.9)    |
| Smoking (%)             | 0/8 (0)      | 0/7 (0)   | 2/52 (3.9)    |
| Diabetes mellitus (%)   | 0/8 (0)      | 0/7 (0)   | 1/53 (1.9)    |
| Nulliparous             | 4/8 (50.0)   | 5/7 (71.4) | 11/53 (20.8)  |
| ART                     | 1/8 (12.5)   | 1/7 (14.3) | 3/45 (6.7)    |
| Race (%)                |              |          |               |
| White                   | 6/8 (75.0)   | 7/7 (100.0) | 46/53 (86.8)  |
| South-Asian             | 0/8 (0)      | 0/7 (0)   | 3/53 (5.7)    |
| Asian                   | 1/8 (12.5)   | 0/7 (0)   | 1/53 (1.9)    |
| Afro-descendant         | 1/8 (12.5)   | 0/7 (0)   | 1/53 (1.9)    |
| Mix                     | 0/8 (0)      | 0/7 (0)   | 2/53 (3.8)    |
| Gestational age at ICU admission | 25.6 (21.3–30.0) | 35.4 (30.3–36.6) | 27.7 (24.5–31.3) |

Continuous data are presented as median (interquartile range). Categorical data presented as frequency (percentage).

**ART,** assisted reproductive treatment; **BMI,** body mass index; **ICU,** intensive care unit; **PE,** preeclampsia.

*Statistically significant difference compared to the PE group.

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### TABLE 2
PE-related findings during and after severe pneumonia

| Variable | PE like (n=8) | PE (n=7) | Non-PE (n=53) | P | Value | During severe pneumonia | After severe pneumonia | P | Value | During severe pneumonia | After severe pneumonia | P | Value |
|----------|--------------|----------|---------------|---|-------|-------------------------|------------------------|---|-------|-------------------------|------------------------|---|-------|
| **SBP (mm Hg)** | 153.0 (147.3–160.8) | 115.0 (112.3–125.8) | 154.0 (145.0–156.0) | .012 | 128.0 (123.0–137.0) | .018 | 108.0 (103.5–119.0) | 104.0 (98.0–112.0) | <.001 | 69.0 (64.5–76.5) | 65.0 (60.0–71.0) | .005 |
| **DBP (mm Hg)** | 95.5 (90.8–104.5) | 76.0 (63.8–77.5) | 10.0 (97.0–109.0) | .012 | 83.0 (81.0–90.0) | .018 | 69.0 (64.5–76.5) | 65.0 (60.0–71.0) | .005 |
| **Hypertension** | 8/8 (100.0) | 0/8 (0) | 7/7 (100.0) | .005 | 2/7 (28.5) | .025 | 0/53 (0) | 1/53 (1.9) | .320 |
| **Proteinuria** | 6/6 (100.0) | 0/2 (0) | 5/7 (71.4) | .16 | 8/14 (57.1) | .160 | 0/53 (0) | 1/53 (1.9) | .320 |
| **Platelet count per μL** | 245,000 (191,500–328,300) | 273,500 (224,800–384,500) | 5/7 (71.4) | .21 | 371,000 (307,000–406,000) | .06 | 259,000 (187,000–323,000) | 290,000 (240,000–352,000) | .045 |
| **Thrombocytopenia<100,000/μL** | 0/8 (0) | 0/8 (0) | 0/7 (0) | 1.0 | 0/53 (0) | 1.0 | 2/52 (3.9) | 0/53 (0) | .160 |
| **AST (U/L)** | 99 (28.3–160.8) | 18 (16–25) | 104 (42–164) | .028 | 29 (24–31) | .022 | 74 (44–144.5) | 23 (17–43) | <.001 |
| **ALT (U/L)** | 114.5 (32.8–290.3) | 15.0 (9.0–33.0) | 85.0 (39.0–150.0) | .043 | 30.0 (20.0–45.0) | .028 | 58.8 (35.5–53.0) | 25.0 (14.0–37.0) | <.001 |
| **Transaminits more than twice the normal values** | 5/8 (62.5) | 1/7 (14.3) | 6/7 (85.7) | .083 | 0/7 (0) | .014 | 32/52 (62.5) | 6/47 (12.8) | <.001 |
| **LDH (U/L)** | 282.0 (203.8–360.3) | 223.0 (167.0–246.3) | 497.5 (347.3–538.3) | .07 | 328.5 (239.0–367.8) | .68 | 280.5 (234.0–320.5) | 203.0 (173.0–202.5) | .005 |
| **LDH>600 U/L** | 0/8 (0) | 0/8 (0) | 0/7 (0) | 1.0 | 0/7 (0) | 1.0 | 2/52 (3.9) | 0/53 (0) | .160 |
| **Creatinine level (mg/dL)** | 0.32 (0.25–0.41) | 0.41 (0.39–0.42) | 0.49 (0.47–0.64) | .036 | 0.43 (0.38–0.49) | .45 | 0.35 (0.31–0.44) | 0.4 (0.34–0.51) | .029 |
| **Creatinine level>1 mg/dL** | 0/8 (0) | 0/8 (0) | 0/7 (0) | 1.0 | 0/7 (0) | 1.0 | 0/51 (0) | 0/53 (0) | 1.0 |
| **Mean uterine artery PI >95th percentile** | 1/5 (20.0) | 1/6 (14.3) | 1/5 (20.0) | 1.0 | 0/7 (0) | 1.0 | 2/13 (15.4) | 4/29 (13.8) | .320 |
| **sFlt-1 to PlGF ratio** | 4.60 (2.56–12.80) | 11.30 (7.30–23.30) | 122.90 (49.40–225.50) | .068 | 15.20 (8.50–21.90) | .18 | 6.70 (2.30–10.90) | 5.91 (3.40–10.40) | .790 |
| **Fulfill PE criteria** | 8/8 (100.0) | 0/8 (0) | 7/7 (100.0) | .005 | 7/7 (100.0) | .008 | 0/53 (0) | 1/53 (1.9) | .310 |
| **Fulfill HELLP criteria** | 0/8 (0) | 0/8 (0) | 0/7 (0) | 1.0 | 0/7 (0) | 1.0 | 0/53 (0) | 0/53 (0) | 1.0 |

Continuous data are presented as median (interquartile range). Categorical data are presented as frequency (percentage). The Wilcoxon signed-rank test was used to assess differences of repeated measurements on a single sample.

ALT, alanine aminotransferase; AST, aspartate aminotransferase; DBP, diastolic blood pressure; HELLP, hemolysis, elevated liver enzymes, and low platelet count; LDH, lactate dehydrogenase; PE, preeclampsia; PI, pulsatility index; PlGF, placental growth factor; SBP, systolic blood pressure; sFlt-1, soluble fms-like tyrosine kinase 1.

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We found that some PE-related features worsened similarly during severe pneumonia across all 3 groups; however, most of these features improved significantly after recovery from severe pneumonia, without requiring elective delivery in most cases (Tables 2 and 3). Among the 68 women included in this study, there were 15 cases (22.1%) of hypertension, 19 cases (27.9%) of proteinuria, and 43 cases (63.2%) of elevated liver enzymes. There was no case of thrombocytopenia or abnormal LDH or creatinine. Moreover, 15 women (22.1%) met the PE diagnostic criteria during severe pneumonia. When comparing the incidence of PE-related features among groups, hypertension was present in the PE and PE-like syndrome groups (100%) and not in the non-PE group (0%). In addition, the sFlt-1-to-PIGF ratios were higher in the PE group than in the PE-like syndrome and non-PE groups (Table 3). No other significant difference was observed across groups regarding PE-related features. When comparing the PE-like syndrome and PE groups during pneumonia, the sFlt-1-to-PIGF ratios (4.6 [2.6–12.8] vs 122.9 [49.4–225.5]; \( P<.008 \)) and creatinine values (0.32 [0.25–0.41] mg/dL vs 0.49 [0.47–0.64] mg/dL; \( P=.015 \)) differed significantly, although creatinine values were within the normal range in both groups. In contrast, when comparing the PE-like syndrome and non-PE groups, no significant difference was found, except for hypertension (8/8 (100%) vs 0/53 (0%); \( P<.001 \)), SBP (153 [147.3–160.8] mm Hg vs 108 [103.5–119] mm Hg; \( P<.001 \)), and DBP (95.5 [90.8–104.5] mm Hg vs 69 [64.5–76.5] mm Hg; \( P<.001 \)).

When we compared the groups after the resolution of severe pneumonia because of COVID-19, only blood pressure was significantly different among groups, being higher in the PE group, although always within the normal range (Table 4).

Regarding perinatal and neonatal outcomes, the PE-like syndrome group had significantly fewer cesarean deliveries (37% vs 100%; \( P=.015 \)), higher gestational age at delivery (39.1 [38.2–40.7] weeks vs 36.3 [32–36.8] weeks; \( P=.001 \)), and higher birthweight (2737.5 [2637.3–3037.5] g vs 2412.5 [1683.8–2580] g; \( P=.004 \)) than the PE group. No difference was found between the PE-like syndrome and non-PE groups. More information can be found in Table 5.

**Discussion**

**Principal findings**

During severe pneumonia, up to 14.2% (15/106) of women met the PE diagnostic criteria; however, 8 of 15 cases (53.3%) improved after pneumonia without elective delivery. These 8 cases had normal sFlt-1-to-PIGF ratios, better outcomes than the 7 women with PE, and similar pregnancy outcomes to those of normotensive women with severe COVID-19 who did not deliver...
during severe pneumonia. This suggests that the sFlt-1—to—PlGF ratio may be a useful marker to identify a subset of patients with PE-like syndrome that otherwise may be misdiagnosed with PE and who would benefit from a more conservative management.

In Table 2, we confirm that in both the PE-like syndrome group and the PE group, most PE features observed during severe pneumonia improved after recovery, especially hypertension and transaminitis. Improvement in the PE group can be attributed to the fact that all women delivered because of COVID-19 and/or PE during ICU stay. In contrast, in the PE-like group, PE-related features improved without delivery. Given that PE does not resolve spontaneously, but only after delivery, the most plausible explanation for this outcome in the PE-like group is a PE-like syndrome induced by severe COVID-19.

In the non-PE group, some PE-related features (such as increased liver enzymes) were observed during severe pneumonia but improved after recovery. This suggests that even in normotensive patients, severe pneumonia because of COVID-19 can induce transient systemic effects that mimic PE and that resolve completely after the resolution of pneumonia.

During severe pneumonia (Table 3), the PE-like and PE groups were identical, except for sFlt-1—to—PlGF and creatinine values that were significantly lower, although within the normal range, in all patients with PE-like syndrome. However, creatinine values were also normal in the PE group, making this finding clinically irrelevant for differential diagnosis. Therefore, the sFlt-1—to—PlGF ratio seems to be the only parameter that allows to accurately discriminate PE from PE-like syndrome.

All women with PE were electively delivered during severe pneumonia. After resolution of severe pneumonia (Table 4), most PE-related features, such as hypertension, transaminitis, and increased sFlt-1—to—PlGF ratio, improved similarly across all groups, with the remarkable difference that all cases in the PE-like syndrome and non-PE groups recovered without elective delivery. This confirms that PE-related features in these 2 groups were due to the systemic effects of COVID-19.

Regarding perinatal outcomes (Table 5), the rate of cesarean deliveries was significantly higher and the birthweight and gestational age at delivery were significantly lower in the PE group than in the PE-like and non-PE groups. No difference was found between the PE-like and non-PE groups for perinatal outcomes.

Results
Several studies have reported an association between PE and COVID-19 that may be explained by several etiopathogenic pathways.13,19—22 PE-like syndrome was described at the beginning of the pandemic in a small case series and could partly explain the higher PE

| TABLE 4 |
| --- |
| **PE-related features after severe pneumonia in the PE-like group vs the PE and non-PE groups** |

| Variables | PE-like (n=8) | PE (n=7) | Non-PE (n=53) |
| --- | --- | --- | --- |
| SBP (mm Hg) | 115 (112.3—125.8)b | 128 (123.0—137.0) | 104 (98.0—112.0) |
| DBP (mm Hg) | 76 (63.8—77.5)a | 83 (61.0—90.0) | 65 (60.0—71.0) |
| Hypertension | 0/8 (0) | 2/7 (28.5) | 1/53 (1.9) |
| Proteinuria | 0/2 (0) | — | 3/9 (33.3) |
| Platelet count per μL | 273,500 (224,800—384,500) | 371,000 (307,000—406,000) | 290,000 (240,000—352,000) |
| Thrombopenia<100,000/μL | 0/8 (0) | 0/7 (0) | 0/51 (0) |
| AST (U/L) | 15 (9—33) | 30 (20—45) | 25 (14—37) |
| LDH (U/L) | 223 (167.0—246.3) | 328.5 (239.0—367.8) | 203 (173.0—202.5) |
| Creatinine level>1 mg/dL | 0/8 (0) | 0/7 (0) | 0/53 (0) |
| Creatinine level>1 mg/dL | 0/8 (0) | 0/7 (0) | 0/53 (0) |
| Mean uterine artery PI >95th percentile | 1/7 (14.3) | 0/7 (0) | 6/47 (12.8) |
| sFlt-1—to—PlGF ratio | 11.3 (7.3—23.3) | 15.2 (8.5—21.9) | 5.91 (3.4—10.4) |

Continuous data are presented as median (interquartile range). Categorical data are presented as frequency (percentage).

ALT, alanine aminotransferase; AST, aspartate aminotransferase; DBP, diastolic blood pressure; LDH, lactate dehydrogenase; PI, pulsatility index; PlGF, placental growth factor; SBP, systolic blood pressure; sFlt-1, soluble fms-like tyrosine kinase 1.

a Statistically significant difference compared to the PE group; b Statistically significant difference compared to the non-PE group.

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incidence in COVID-19 pregnancies. Unfortunately, none of the studies investigating the association between both conditions have examined the sFlt-1−to−PlGF ratio or excluded women with PE-like syndrome from the analyses, making their results difficult to interpret. After that small case series, PE-like syndrome was no further investigated in larger cohorts. For this reason, the exact relationship between PE and COVID-19 remains unknown, as some studies might have misdiagnosed some cases of PE. Here, we provided further evidence that PE-like syndrome is an actual condition that may affect 7.5% (8/106) of women with severe COVID-19 and more than 50% of women meeting the diagnostic criteria for PE. According to the first case series describing the PE-like syndrome, the incidence of this syndrome in women with severe COVID-19 was 50.0%, whereas the incidence was only 7.5% in this study. This difference may be explained by the larger sample size in this study. Another plausible explanation may be the longer study period (19 months) in this study compared with the case series (31 days). Both situations may offer a wider scope of the clinical characteristics of PE-like syndrome through different management protocols and different clinical manifestations because of the increasing vaccine administration rates and the evolving SARS-CoV-2 variants.

Clinical implications
This study has important clinical implications, as it shows that PE-like syndrome is a transient condition that spontaneously resolves after recovery from severe pneumonia and has similar pregnancy outcomes to those of normotensive women with severe COVID-19. Therefore, patients with PE-like syndrome may benefit from a more conservative management. Most women with PE-like syndrome may be erroneously diagnosed with PE with severity features because of the systemic effects of COVID-19 and, therefore, may be electively delivered at ≥34 weeks of gestation. In this context, the sFlt-1—to−PlGF ratio may help to discriminate true PE from PE-like syndrome, thus improving clinical management and pregnancy outcomes for cases with PE-like syndrome by reducing the risk of iatrogenic complications and prematurity.

Research implications
In addition, our results have important research implications, as we show that
some cases classified as true PE in previous studies might instead be cases with PE-like syndrome. Therefore, to better assess the association between true PE and COVID-19, in future studies, PE-like syndrome should be excluded using the sFlt-1-to−PlGF ratio.

Strengths and limitations

One of the main strengths of this study was the large number of pregnant women with severe pneumonia because of COVID-19. In addition, this study confirmed the existence of a PE-like syndrome induced by severe COVID-19 by assessing the sFlt-1−to−PlGF ratio in a relatively large cohort of pregnant women with suspicion of PE and severe COVID-19. This study provided novel evidence for the evolution and outcomes of pregnancies with PE-like syndrome induced by severe COVID-19. Another strength of this study was that it further supports the clinical usefulness of the sFlt-1−to−PlGF ratio for discriminating between placental dysfunction and other disorders.4

The main limitation of this study was the reduced number of cases with PE and PE-like syndrome, which makes it harder to identify differences across groups for pregnancy outcomes. For instance, the median birthweight in the PE-like syndrome group tended to be lower than in the non-PE group (2373.5 g vs 3130 g, respectively), despite gestational age at delivery being the same. However, this difference was not statistically significant (P=.09), probably because of the reduced sample size in the PE-like syndrome group. Therefore, it is plausible that patients who are hypertensive with severe COVID-19 and normal sFlt-1-to−PlGF ratio (PE-like group) may have poorer perinatal outcomes than normotensive patients with severe COVID-19 (non-PE group). However, this hypothesis could not be confirmed in this study. Another limitation was that mean uterine artery PI was not recorded in any women in the PE group; for this reason, we could not investigate its role in distinguishing PE from PE-like syndrome.

Conclusions

More than 50% of women with severe COVID-19 that meet the diagnostic criteria for PE have PE-like syndrome. Therefore, future research on COVID-19 and PE should identify the cases with PE-like syndrome to better assess the association of both conditions. PE-like syndrome is a transient condition with similar pregnancy outcomes to those of normotensive women with severe COVID-19. Therefore, correct identification of cases with PE-like syndrome may improve pregnancy outcomes by reducing iatrogenic complications.

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