Clinical features and disease course of patients with acute ischaemic stroke just before the Italian index case: Was COVID-19 already there?

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Abstract
Since the end of February 2020, Italy has suffered one of the most severe outbreaks of coronavirus disease 2019 (COVID-19). However, what happened just before the Italian index case has not yet been investigated. To answer this question, we evaluated the potential impact of COVID-19 on the clinical features of a cohort of neurological inpatients admitted right before the Italian index case, as compared to the same period of the previous year. Demographic, clinical, treatment and laboratory data were extracted from medical records. The data collected included all inpatients who had been admitted to the Neurology and Stroke Units of the Ospedale Maggiore Policlinico, Milan, Italy, from December 15, 2018 to February 20, 2019 and from December 15, 2019 to February 20, 2020. Of the 248 patients, 97 subjects (39.1%) were admitted for an acute cerebrovascular event: 46 in the 2018/2019 period (mean [SD] age, 72.3 [15.6] years; 22 men [47.8%]), and 51 in the 2019/2020 interval (mean [SD] age, 72.8 [12.4] years; 24 men [47.1%]). The number of cryptogenic strokes has increased during the 2019–2020 year, as compared to the previous year (30 [58.8%] vs. 18 [39.1%], p = 0.05). These patients had a longer hospitalization (mean [SD] day, 15.7 [10.5] days vs. mean [SD] day, 11.7 [7.2] days, p = 0.03) and more frequent cerebrovascular complications (9 [30.0%] vs. 2 [11.1%]), but presented a lower incidence of cardiocerebral risk factors (18 [60.0%] vs. 14 [77.8%]). Right before the Italian index case, an increase in cryptogenic strokes has occurred, possibly due to the concomitant COVID-19.

Keywords COVID-19 · Acute ischaemic stroke · Cryptogenic stroke · Cerebrovascular events

Introduction
Since February 20, 2020, Italy, and in particular its most populous region, Lombardy, has suffered one of the most severe outbreaks of coronavirus disease 2019 (COVID-19), caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). As of December 10, 2020, in this region, inhabited by about 10,000,000 people, the cases registered by the authorities were more than 430,000 and the total deaths were about 23,000 [1].

Early molecular epidemiological tracing suggests that COVID-19 spread in Lombardy weeks before the first reported cases of infection, confirming that SARS-CoV-2 virus entered northern Italy as early as January, 2020 [2]. A very recent paper confirmed this finding, describing the earliest evidence of SARS-CoV-2 RNA in a patient in Lombardy in early December, 2019, 3 months before Italy’s first reported COVID-19 case [3].

The clinical manifestations of COVID-19 mostly affect the respiratory system, configuring varied clinical pictures ranging from mild upper respiratory tract illness to severe pneumonia with respiratory failure and death [4, 5]. Asymptomatic infections are reported [6]. More recently,
the virus-related clinical spectrum has widened, including stroke, venous thromboembolism, pulmonary embolism and cardiac complications [7]. The main pathogenic hypothesis focuses on the severe inflammatory state secondary to the infection that leads to an alteration of hemostasis [7, 8]. Neurological manifestations appear to be quite common, including acute cerebrovascular disease, conscious disturbance, headache and olfactory dysfunctions [9, 10].

Based on the assumption that COVID-19 may have already been circulating before we knew it, we turned our attention to the period right before the first index patient, focusing on clinical and laboratory features, as well as treatment and disease course, of a cohort of neurological inpatients. To avoid comparing different epidemiological periods and different clinical protocols, we ended the patients’ enrollment one day before Italian first reported COVID-19 case.

**Methods**

**Study design and participants**

This retrospective cohort study included adult inpatients. Our study enrolled all patients who were hospitalized for neurological diseases from December 15, 2018, to February 20, 2019 (Group 1) and from December 15, 2019, to February 20, 2020 (Group 2) to the Neurology and Stroke Units of the Ospedale Maggiore Policlinico of Milan.

The study was approved by the Local Ethics Committee of the Fondazione IRCCS Ca’ Granda Ospedale Maggiore Policlinico (363_2020, approved 05.05.2020).

**Data collection**

Demographical, clinical, laboratory and treatment data were extracted from electronic medical records using standardized anonymized data collection forms. All data were extracted by eleven physicians (A.M.P, S.L, E.A, R.B, G.C, G.L, E.M, M.P, L.S, G.V and D.V). Routine blood examinations comprised complete blood count, erythrocyte sedimentation rate, serum biochemical tests, including C-reactive protein, liver and renal function, lactate dehydrogenase, creatine kinase and coagulation profile. Instrumental examinations ranged according to clinical need. For acute cerebrovascular disease, they comprised as needed computed tomography (CT) scan of the brain, brain magnetic resonance imaging (MRI), vessel imaging and cardiovascular examinations, including extra- and intracranial ultrasound and transthoracic and transesophageal echocardiography.

Cryptogenic strokes were identified using the TOAST classification [11] by consensus of two independent observers (A.M.P; S.L).

**Statistical analysis**

The patients’ characteristics were described using percentages for discrete variables, mean and standard deviations for continuous variables. Differences between two groups were analyzed using F test from univariate ANOVA and Pearson’s chi-squared test.

Focusing on subjects with stroke, differences in prevalence of cryptogenic events between groups were evaluated using a logistic regression adjusted for age, gender, presence of cardiocerebrovascular (CCV) risk factors and related home drug therapy.

Statistical significance was set up to 0.05; as there are no preliminary data, it is not possible to predict whether the size of the collected sample reached the appropriate statistical power.

**Results**

Two hundred forty-eight patients were included. Of these, about 40% was hospitalized for acute cerebrovascular events: 46 in the 2018/2019 period (Group 1) and 51 in 2019/2020 (Group 2). The fraction of patients admitted to the Stoke Unit was not significantly different in the two period and there were no differences in access to acute phase therapies as intravenous fibrinolysis, endovascular treatment or bridging therapy nor in hemorrhagic complications in relation to acute reperfusion treatments. No significant divergences between Groups 1 and 2 regarding demographical characteristics and comorbidities, including hypertension, diabetes, hypercholesterolemia and coronary heart disease were observed (Table 1A). No significant differences were seen in the diagnostic workup of the two periods, targeted from time to time based on the clinical suspicion.

Group 2 patients had longer duration of hospitalization (mean [SD] day, 15.7 [10.5] days vs mean [SD] day, 11.7 [7.4] days, p = 0.04) and higher incidence of cerebrovascular complications (13 [25.5%] vs 4 [8.7%], p = 0.03) (Table 1A), which include new ischaemic events and/or hemorrhagic infarctions. During admission, treatment modifications, such as addition of anticoagulant thromboprophylaxis with low-molecular weight heparin and/or switching antiplatelet drugs, occurred more frequently in Group 2 (46 [90.2%] vs. 30 [69.8%], p = 0.008) (Table 1A).

Fever and laboratory reports were similar between the two groups. Fibrinogen, d-dimer and serum ferritin were not assessed, as not part of the haematochemistry routine during pre-COVID-19 era.

The frequency of cryptogenic strokes was increased during the 2019/2020 period (30 [58.8%] vs. 18 [39.1%], p = 0.05) (Fig. 1a). Group 2 patients had a longer hospitalization (mean [SD] day, 15.7 [10.5] days vs. mean [SD] day,
11.7 [7.2] days, \( p = 0.03 \), more frequent cerebrovascular complications (9 [30.0%] vs. 2 [11.1%]), although presenting a lower incidence of CCV risk factors (18 [60.0%] vs. 14 [77.8%]) (Fig. 1b–d). In these patients, a certain increase in respiratory complications, including pulmonary infection (6 [20.0%] vs. 2 [11.1%]), was observed (Fig. 1e).

| Variables, \( N(\%) \) | Years 2019–2020 | Years 2018–2019 | \( p \) value |
|-------------------------|-----------------|-----------------|------------|
| (A)                     |                 |                 |            |
| Gender (female)         | 27 (52.9)       | 24 (52.2)       | 0.93       |
| Age, mean (SD)          | 72.8 (12.4)     | 72.3 (15.7)     | 0.86       |
| Age > 65 years          | 38 (74.5)       | 34 (74.0)       | 0.95       |
| Cardiocerebrovascular risk factors | 35 (68.6) | 35 (76.1) | 0.41       |
| Comorbidities (at least 1) | 43 (84.3) | 40 (87.0) | 0.71       |
| Type of stroke (TOAST Classification) |     |                 | 0.33       |
| 1. Large-artery atherosclerosis | 5 (9.8)   | 4 (9.7)         |            |
| 2. Cardioembolism        | 8 (15.7)       | 13 (28.3)       |            |
| 3. Small-vessel occlusion| 3 (5.9)        | 5 (10.9)        |            |
| 4. Stroke of other determined etiology | 5 (9.8) | 6 (13.0) |            |
| 5. Stroke of undetermined etiology | 30 (58.8) | 18 (39.1) |            |
| TOAST classification     |                | 0.05            |            |
| 1–4                     | 21 (41.2)       | 28 (60.9)       |            |
| 5                       | 30 (58.8)       | 18 (39.1)       |            |
| Stroke complications     | 20 (39.2)       | 10 (21.7)       | 0.06       |
| Cerebrovascular complications | 13 (25.5) | 4 (8.7)        | 0.03       |
| Other complications     | 20 (39.2)       | 19 (41.3)       | 0.83       |
| Cardiac complications   | 3 (5.8)         | 7 (15.2)        | 0.13       |
| Infective complications | 14 (27.5)       | 11 (23.9)       | 0.69       |
| Pulmonary infections    | 11 (21.6)       | 5 (10.87)       | 0.16       |
| Respiratory complications| 11 (21.6) | 6 (13.0) | 0.27       |
| Drug therapies at home, mean (SD) |     |                 | 0.15       |
| Antiplatelet agents     | 21 (41.2)       | 22 (48.9)       |            |
| Anticoagulants          | 2 (3.9)         | 7 (15.5)        |            |
| Dual antiplatelet therapy| 1 (2.0)   | 0               |            |
| Antplatelets and anticoagulants | 1 (2.0) | 0           |            |
| Changes in therapy during hospitalization |     |                 | 0.008      |
| Nothing                 | 5 (9.8)         | 15 (33.3)       |            |
| Horizontal shift        | 23 (45.1)       | 17 (37.8)       |            |
| Anticoagulant shift     | 9 (17.7)        | 8 (17.8)        |            |
| Dual antiplatelets shift| 4 (7.9)        | 5 (11.1)        |            |
| Heparin prophylaxis     | 3 (5.9)         | 0               |            |
| Heparin + antiplatelets | 7 (13.7)        | 0               |            |
| Days of hospitalization, mean (SD) | 15.7 (10.5) | 11.7 (7.4) | 0.04       |
| Deaths                  | 1 (2.0)         | 2 (4.4)         | 0.50       |

| Variables                 | Odds ratios (confidence interval) | \( p \) value |
|---------------------------|----------------------------------|--------------|
| (B)                       |                                  |              |
| Group (years 2019–2020 vs. years 2018–2019) | 2.33 (0.98–5.23) | 0.052        |
| Age                       | 1.01 (0.97–1.04)                | 0.74         |
| Gender (female vs. male)  | 1.12 (0.47–2.67)                | 0.79         |
| CCV risk factors          | 0.45 (0.17–1.26)                | 0.13         |
| Home drugs therapy (yes vs. no) | 2.18 (0.84–5.64) | 0.11         |

Bold value indicates statistically significant (\( p < 0.05 \)
A

Years 2019-2020

5. stroke of undetermined etiology
4. stroke of other determined etiology
3. small-vessel occlusion
2. cardioembolism
1. large-artery atherosclerosis

Years 2018-2019

5. stroke of undetermined etiology
4. stroke of other determined etiology
3. small-vessel occlusion
2. cardioembolism
1. large-artery atherosclerosis

B

Hospitalization

Days

C

Cerebrovascular complications

% 35

D

CCV risk factors

% 90

E

Respiratory complications

% 25
negative. All cryptogenic strokes tested resulted positive (6
izations were considered. Seven were positive and four were
in May, 2020. Only 11 patients had undergone swab for
val were then contacted on the phone. The calls occurred
as compared to those of Group 1 (Table 1B).

Forty-three out of 51 total strokes in the 2019/2020 inter-
interval were then contacted on the phone. The calls occurred
May, 2020. Only 11 patients had undergone swab for
for COVID-19 or COVID-19 antibody test. To avoid later infec-
tions, only tests carried out within one month of hospitali-
were considered. Seven were positive and four were
negative. All cryptogenic strokes tested resulted positive (6
of 6), while none of the negatives was a cryptogenic stroke.

Discussion

In December 2019, January and February, 2020 a significa-
cient increase in the number of cryptogenic strokes for no
apparent reason could be observed. When compared with the
data available in the literature, in which cryptogenic strokes
account for 30–40% of ischaemic strokes [12], it is noteworthy
that, in the current year, cryptogenic subtypes were about
60% of all strokes. COVID-19 had not revealed itself yet, at
least expressly, changing the clinical routine and the diag-
nostic protocols of our wards, which did not differ from that
of the 2018/2019 period. In our opinion, the strength of this
study lies precisely in comparing two periods in which diag-
nostic path were similar and not yet derailed due to the advent
of COVID-19. Since our wards continued with the diagnostic
routine only until the first half of March 2020, we decided not
to include in our study the patients hospitalized from Febru-
ary 21, 2020, exactly for not to compare a pre-COVID with a
COVID period. That was especially the case with our Stroke
Unit, which continued working as a “Level Stroke Unit” on
the Stroke Network until mid-March, 2020 only.

Comparing Groups 1 and 2, patients admitted this year
were neither older nor younger and did not have a higher
number of comorbidities. Instead, they presented lower
incidence of CCV risk factors, but unexpectedly had more
complicated hospitalizations. No laboratory tests seem to
explain these data. The presence, although still unknown, of
COVID-19 in the territory of Milan could hold the answer,
even though the true relationship between COVID-19 and
cryptogenic stroke incidence remains to be determined.

By now, the current evidence suggests that COVID-19 may
predispose patients to thrombotic disease due to excessive
inflammation, platelet activation, endothelial dysfunction and
stasis [7]. Recent reports described microvascular endothelial
injury in postmortem evaluation in multiple organs, consistent
with a thrombotic microangiopathy [13, 14]. This evidence
may explain the frequent reports of cerebrovascular complica-
tions, also in patients with lower CCV risk factors.

To better understand the link between cerebrovascular
accidents and COVID-19, the crucial point seems to focus
on the asymptomatic patients. Asymptomatic infections refer
to the positive detection of nucleic acid of SARS-CoV-2
by reverse transcriptase polymerase chain reaction in sam-
ple of patients who have no typical respiratory symptoms
or signs [15]. Therefore, the possible explanations of the
increased frequency of cryptogenic strokes, we have experi-
enced seem to be: first, endothelial dysfunction, involving
the thrombi formation in the cerebral vessels, may develop
even in asymptomatic patients; second, our patients were
in the early stages of their systemic viral illness that would
soon become clinically evident.

To confirm this last hypothesis, we contacted all
2019/2020 stroke cases. Some patients experienced a cough,
fever and difficulty breathing shortly after being discharged,
but unfortunately, only 11 had undergone swab for COVID-
19 or COVID-19 antibody test. Interestingly, all cryptogenic
strokes tested resulted positive, while none of the negatives
was a cryptogenic stroke. This low number of tests should
not be surprising, because most of the patients were first hos-
pitalized and then discharged before we started to consider
COVID-19 infections. Recalling all patients to test them for
antibody titre was an option. Currently, our understanding of
antibody responses following infection with SARS-CoV-2
is limited and more importantly the duration of responses
remains unclear. In the light of this, we decided not to test
our patients as a year later seemed too distant a time. More-
ever, we were worried about not introducing confounding
variables, such as post-observation infections, taking into
account, the second wave of COVID-19 which has struck
Lombardy since early October, 2020.

Limitations of the current study include the retrospective,
single centre cohort design, and the small sample size, which
could be the reason why the statistical threshold for differ-
ences could be not reached. Moreover, the data related to
swab for COVID-19 or COVID-19 antibody test was avail-
able only for a few patients.

Conclusions

Right before the Italian index case, an increase in crypto-
genic strokes has occurred. Growing evidence suggests that
the higher incidence of cryptogenic stroke in asymptomatic
COVID-19 patients or in the early stages of their systemic illness could be associated with a transient hypercoagulable state, leading to thrombi formation in the cerebral vessels. Replication in a larger cohort of patients is needed to confirm these data.

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Data availability and code availability Not applicable.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethics approval The study was approved by the local Ethics Committee of the Fondazione IRCCS Ca’ Granda Ospedale Maggiore Policlinico (363_2020, approved 05.05.2020).

Statement of human rights All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Statement of the welfare of animals This article does not contain any studies with animals performed by any of the authors.

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