Retrospective Post-mortem SARS-CoV-2 RT-PCR of Autopsies with COVID-19-Suggestive Pathology Supports the Absence of Lethal Community Spread in Basel, Switzerland, before February 2020

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Abstract

Introduction: Coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has rapidly spread around the world. While the first case was recorded in Hubei in December 2019, the extent of early community spread in Central Europe before this period is unknown. A high proportion of asymptomatic cases and undocumented infections, high transmissibility, and phylogenetic genomic diversity have engendered the controversial possibility of early international community spread of SARS-CoV-2 before its emergence in China. Methods: To assess the early presence of lethal COVID-19 in Switzerland, we retrospectively performed an analysis of deaths at University Hospital Basel between October 2019 and February 2020 (n = 310), comparing the incidence of clinical causes of death with March 2020 (n = 72), the month during which the first lethal COVID-19 cases in Basel were reported. Trends of COVID-19-suggestive sequelae, such as bronchopneumonia with organization, acute respiratory distress syndrome (ARDS), or pulmonary embolisms (PE) were evaluated. In cases where autopsy was performed (n = 71), analogous analyses were conducted on the cause of death and pulmonary histological findings. Eight cases with a COVID-19-suggestive clinical history and histopathology between October 2019 and February 2020, and 3 cases before October 2019, were selected for SARS-CoV-2 RT-PCR. Results: A statistically significant rise in pulmonary causes of death was observed in March 2020 (p = 0.03), consistent with the reported emergence of lethal COVID-19 in Switzerland. A rise in lethal bronchopneumonia was observed between December 2019 and January 2020, which was likely seasonal. The incidence of lethal ARDS and PE was uniformly low between October 2019 and February 2020. All autopsy cases analyzed by means of SARS-CoV-2 RT-PCR yielded negative results. Conclusion: Our data suggest the absence of early lethal community spread of COVID-19 in Basel before its initial reported emergence in Switzerland in March 2020.

Introduction

We are currently in the midst of a coronavirus disease pandemic, COVID-19, caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). Since the WHO
declared COVID-19 a pandemic on March 11, global cases have steadily risen with a total of 10.3 million cases and 508,000 deaths as of July 1, 2020, out of which 31,000 cases and 1,600 deaths are attributed to Switzerland [1]. The first case of COVID-19 in Switzerland was reported on February 25, 2020, while the first lethal case was recorded on March 5, 2020 [2, 3].

SARS-CoV-2 is a part of the Orthocoronavirinae subfamily of Coronaviridae [4]. Currently, four subtypes of Orthocoronavirinae are described in the literature: while α- and β-coronaviruses are able to infect humans, typically causing symptoms of the common cold, γ- and δ-coronaviruses are only found in animals [5]. The zoonotic properties of coronaviruses enable host transmission from animal to human via airborne droplets [6]. In-depth studies have described a primary natural coronavirus reservoir in bats, although other secondary animal hosts include camels, mice, dogs, civet cats, ferrets, raccoons, and other rodents [7–9]. 2002 and 2012 marked key moments in our understanding of coronavirus pathogenicity, when two coronavirus outbreaks caused by severe acute respiratory syndrome coronavirus (SARS-CoV) and Middle-East respiratory syndrome coronavirus (MERS-CoV), respectively, developed into pandemics spreading to multiple countries and affecting thousands of individuals. In both cases, the novel virus originated from bats and spread to intermediate hosts (palm civets for SARS, camels for MERS) [10].

COVID-19 presents with similar clinical features as previous coronavirus-induced diseases, most commonly manifesting as a combination of cough, fever, and general malaise with a mean symptom onset of 14 days [11, 12]. A more serious disease progression with ICU admission was reported in a quarter of cases with a lethal disease outcome of about 10% amongst hospitalized patients [13–15]; interestingly, this cohort reported a higher incidence of comorbid illness such as hypertension, a history of cardiovascular disease, and chronic pulmonary conditions [16, 17]. Other suggestive predisposing features for serious disease outcome include older age [18, 19], obesity [20], a history of smoking [21], blood group A [22], and male sex [19]. In an effort to shed light on disease susceptibility, the scientific community is currently studying the pathophysiological complexities of lethal COVID-19, investigating the role of key players such as angiotensin-converting enzyme-2, a membrane protein, which has been previously shown to serve as a vital entry portal for host invasion [23–25].

Since its emergence in December 2019, a number of COVID-19 post-mortem studies have been performed. Major histopathological findings from whole body autopsy series include diffuse alveolar damage (DAD) with superimposed bacterial pneumonia in some cases [26–29], consistent with autopsy findings from SARS and MERS [30–32]. Most strikingly, a concurrence of COVID-19 with thromboembolic events and microvascular dysfunction, a characteristic that seems to set SARS-CoV-2 apart from its coronavirus predecessors, has been observed at autopsy [26–28, 33, 34]. In one of the largest post-mortem cohorts to date, the primary cause of death was respiratory failure histologically imposing as DAD with extensive capillary congestion and microthrombi of pulmonary capillaries in 45% and acute pulmonary embolism (PE) in 19% of cases. Superimposed bronchopneumonia was observed in 48% and, interestingly, 28% of cases were diagnosed with senile ATTR amyloidosis upon autopsy [26]. These findings are crucial to fully understand COVID-19 pathophysiology and highlight the urgent importance of managing anticoagulation treatment, especially in patient groups with thrombosis risk factors [35].

The scope of clinical and pathological studies enables an in-depth mapping of the epidemiological dynamics and origins of COVID-19. Current evidence traces SARS-CoV-2 back to the Chinese province of Hubei, where a series of pneumopathies of unknown etiology were reported in December 2019 [36]. The purported origin of the initial outbreak was a wet market in Wuhan, where close proximity between humans and a multitude of potential intermediate hosts may have facilitated host transmission [37]. However, the exact timeline of viral origins remains to be elucidated. Although Chinese authorities initially reported the emergence of the outbreak on December 31, the first case in Wuhan had shown symptoms as early as December 1 and had no direct contact to the market [12]. In view of its unspecific clinical presentation, mild disease progression, high transmissibility [38], and considerable proportion of asymptomatic cases [39, 40], unnoticed community spread before December is plausible and may have occurred; a lack of surveillance and contact tracing would have enabled the virus to spread uncontrollably to other continents. This premise is supported by a retrospective RT-PCR analysis of nasopharyngeal swabs in France, which revealed a positive case dated December 27, 1 month before the first reported case in Europe on January 24 [41, 42]. Additionally, phylogenetic analyses have revealed a high global genomic diversity of SARS-CoV-2, pointing to extensive worldwide transmission early on in the pandemic and estimating that zoonotic transfer occurred approximately between October and December 2019 [43].
Currently, evidence of early COVID-19 transmission and lethality in Central Europe is scarce. To access the presence of lethal community spread in Basel before its initial recorded emergence, we retrospectively compared the clinical cause of adult deaths at University Hospital Basel and autopsies performed at its Institute of Pathology between October 2019 and February 2020 with data from March 2020 when lethal COVID-19 was first reported in Switzerland. Autopsies with COVID-19-suggestive pathology were systematically selected for retrospective SARS-CoV-2 RT-PCR analysis.

**Materials and Methods**

**Study Cohort and Patient Selection**

A systematic retrospective analysis was performed on clinical causes of death (n = 382) at the University Hospital Basel and its subsidiaries between October 2019 and March 2020. In all cases, this was defined as the disease or condition leading directly to death according to the death certificate. Trends in COVID-19-suggestive sequelae, such as bronchopneumonia with organization, acute respiratory distress syndrome (ARDS), or PE were recorded (Fig. 1); trends in clinical cause of death for all cases are shown in online supplementary Figure 1 (for all online suppl. material, see www.karger.com/doi/10.1159/000512563). Next, all autopsies (n = 71) performed during this period at the Institute of Pathology were retrospectively analyzed. Table 1 provides an overview of the cause of death according to post-mortem report, categorized by organ system.

An in-depth study of autopsy reports and clinical history was performed on cases with a direct pulmonary cause of death between October 2019 and February 2020 (n = 13). Cases with COVID-19-suggestive sequelae were selected for retrospective RT-PCR (n = 8). Histopathological inclusion criteria were exudative and/or proliferative DAD with or without superimposed bronchopneumonia, acute PE, thrombotic microangiopathies, cases with coagulopathies of unclear etiology, and amyloidosis. Three other cases with highly suggestive histopathological findings and disease history from an earlier period (March, June, and July 2019, respectively) also underwent RT-PCR analysis. In cases with isolated tracheobronchitis and/or bronchopneumonia with a clear microbiological etiology, exudative neutrophilic inflammation without alveolar fibrin deposition and/or perceptible thromboses of pulmonary vessels, retrospective RT-PCR was not performed. The clinical and pulmonary histology findings of selected patients are presented in Tables 2 and 3. Histological findings were analyzed independently by two pathologists (J.D.H., A.T.) on HE sections; additional immunohistochemical analyses included a fibrin stain to visualize microthrombi, and a subtyping of amyloidosis according to previously described proto-
cols [44, 45]. Pulmonary findings were either graded by means of a severity scale (− no, + mild, ++ moderate, +++ extensive presence) or as a binary value (yes/no for hyaline membranes and acute PE).

**RT-PCR Protocol**

In respectively selected cases with COVID-19-suggestive pathology, an RT-qPCR assay for SARS-CoV-2 was performed on archived lung samples in line with the previously performed methodology [26]. RNA was extracted from formalin-fixed paraffin-embedded (FFPE) tissue using the Maxwell RSC RNA FFPE Kit (Promega, Madison, WI, USA). The TaqMan 2019-nCoV Assay Kit v1 (Thermo Fisher Scientific), which detects three separate viral genomic regions (ORFab1, S Protein, N Protein) and the human RNase P gene (RPPH1), was then employed to detect viral genomes. The viral genome copy number was quantified by a comparative CT (ΔΔCT) method using the TaqMan 2019-nCoV Control Kit v1 (ThermoFisher Scientific), thus generating separate copy numbers for all genomic regions. CT values for any of the three viral genomic regions below 37 were considered positive. When CT values were between 37 and 40, the results were considered undetermined and the assay was repeated. Values above 40 were considered negative. Samples were always run in duplicates.

**Statistical Analysis**

Statistical analyses were performed using Microsoft Excel for Windows 2019© (Redmond, WA, USA). A Student t test (independent sample with equal variance) was utilized to compare the incidence of pulmonary causes of death and its etiologies in the period before the recorded emergence of COVID-19 (October 2019 to February 2020) and March 2020 in weekly intervals. *p* values < 0.05 were considered significant.

**Results**

**Trends in Clinical Causes of Death between October 2019 and March 2020**

University Hospital Basel recorded 382 deaths from October 2019 to March 2020 with a peak incidence between December 2019 (*n* = 71) and January 2020 (*n* = 69); during these 2 months there was an increase of cases, which enlisted bronchopneumonia as a clinical cause of death (December 11.3%, January 10.1% of deaths; Fig. 1). Twenty-seven cases (7.1%) of lethal bronchopneumonia.

### Table 1. Reported causes of death of adult autopsies performed at Basel University Hospital from October 2019 to March 2020 (*n* = 71)

| Cause of death          | October | November | December | January | February | March |
|-------------------------|---------|----------|----------|---------|----------|-------|
| Cardiovascular          | 5       | 1        | 5        | 4       | 2        | 6     |
| Heart failure           | 4       | 0        | 4        | 3       | 2        | 5     |
| Myocardial infarction   | 1       | 0        | 0        | 1       | 0        | 1     |
| Aortic dissection       | 0       | 0        | 1        | 0       | 0        | 0     |
| Pulmonary               | 3       | 1        | 5        | 3       | 1        | 11    |
| PE                      | 1       | 0        | 2        | 1       | 0        | 0     |
| Bronchopneumonia        | 1       | 1        | 3        | 0       | 1        | 0     |
| DAD                     | 0       | 0        | 2        | 0       | 9        |       |
| Other                   | 1       | 0        | 0        | 0       | 0        | 2     |
| Abdominal               | 1       | 1        | 1        | 1       | 3        | 2     |
| Liver failure           | 0       | 0        | 1        | 0       | 0        | 0     |
| Acute pancreatitis      | 0       | 0        | 0        | 1       | 0        | 0     |
| Abdominal hemorrhage    | 0       | 0        | 0        | 0       | 1        | 0     |
| Septic shock, abd. focus| 1       | 1        | 0        | 0       | 1        | 1     |
| Splenic rupture         | 0       | 0        | 0        | 0       | 1        | 0     |
| Mesenteric ischemia     | 0       | 0        | 0        | 0       | 0        | 1     |
| Cerebral                | 2       | 0        | 0        | 3       | 1        | 0     |
| PML                     | 1       | 0        | 0        | 0       | 0        | 0     |
| Traumatic brain injury  | 1       | 0        | 0        | 0       | 0        | 0     |
| Hypoxic encephalopathy  | 0       | 0        | 0        | 1       | 0        | 0     |
| Meningoencephalitis     | 0       | 0        | 0        | 1       | 0        | 0     |
| Stroke                  | 0       | 0        | 0        | 1       | 1        | 0     |
| Terminal oncological disease | 1     | 1        | 1        | 1       | 0        | 1     |
| Other                   | 0       | 0        | 2        | 0       | 1        | 1     |
| Total autopsies performed | 12   | 4        | 14       | 12      | 8        | 21    |

DAD, diffuse alveolar damage; PE, pulmonary embolism; PML, progressive multifocal leukoencephalopathy.
### Table 2. Clinical characteristics of selected adult autopsies with suggestive COVID-19 disease history and histology

| No. | Date of death | Gender | Age, years | BMI | Symptoms at presentation | Comorbidities | Presumptive COVID-19 predisposing factors | Cause of death at autopsy | Likely etiology of pneumopathy |
|-----|---------------|--------|------------|-----|--------------------------|---------------|------------------------------------------|---------------------------|--------------------------------|
| 1   | 13.03.2019    | M      | 85         | 28.4| Dyspnea, fever           | Diabetes, dyslipidemia, gout, hyperthyroidism, chronic kidney failure | Atherosclerosis, hypertension, coronary heart disease, smoker, peripheral arterial disease | DAD                       | Infectious (E. faecalis, C. albicans in sputum) |
| 2   | 19.06.2019    | F      | 48         | 18.3| Dyspnea                  | Collagenosis overlap syndrome (scleroderma and Sjögren), PAH, bronchiolitis, pulmonary hypertension, focal pulmonary AL amyloidosis | Atherosclerosis           | Right heart failure due to cor pulmonale | Rheumatological             |
| 3   | 02.07.2019    | F      | 71         | 28.8| Dyspnea, acute kidney failure | Metastatic liposarcoma, atherosclerosis, atrial fibrillation | Coronary artery disease, peripheral arterial disease, COPD, hypertension | Bronchopneumonia | Infectious (Klebsiella spp. and S. aureus in BAL) |
| 4   | 20.10.2019    | M      | 81         | 41.2| Hemoptysis               | Colon carcinoma, senile amyloidosis | Atherosclerosis, ATTR amyloidosis, obesity, COPD | Bronchopneumonia | Diffuse alveolar hemorrhage | Unclear (likely infective or due to decompensated PAH) |
| 5   | 03.12.2019    | M      | 70         | 26.0| Chronic non-productive cough, flu-like symptoms | Interstitial pneumonitis (measles virus/leukemia), sepsis | Atherosclerosis, hypertension, smoker | Bronchopneumonia | Infectious | |
| 6   | 08.12.2019    | M      | 78         | 41.5| Dyspnea, dyspnea, tachypnea | Diabetes, colitis ulcera, dyslipidemia | Atherosclerosis, coronary artery disease, obesity, COPD, hypertension, smoker | Bronchopneumonia | Unclear (likely multifactorial (obesity, immobility)) | |
| 7   | 17.12.2019    | M      | 87         | 26.0| Hyperactive delirium, tracheobronchitis | Dementia, prostate carcinoma, fracture of cervical spine, dyslipidemia | Atherosclerosis, coronary artery disease, hypertenion | Bronchopneumonia | Infectious | |
| 8   | 19.01.2020    | M      | 61         | 27.2| Respiratory failure | Lung adenocarcinoma, diabetes mellitus II | Atherosclerosis | DAD | | |
| 9   | 19.01.2020    | M      | 50         | 22.3| Post-op dizziness and dyspnea, acute disseminated hypercoagulability | Severe anaemia, insulin-dependent diabetes mellitus | Atherosclerosis | Pulmonary embolism due to unclear coagulopathy | Unclear (post-op, AML, drug induced or infectious) | |
| 10  | 20.01.2020    | M      | 64         | 22.5| Dyspnea, dry cough       | S.p. allogeneic stem cell transplantation, MDS, secondary AML, atrial fibrillation, S.p. lobectomy due to aspergillosis | Eccentric myocaridal hypertrophy | DAD | Drug induced (cytarabine) |
| 11  | 17.02.2020    | F      | 70         | 18.8| Tiredness and dyspnea    | Abdominal aortic aneurysm | Atherosclerosis, coronary artery disease, COPD | Necrotizing pneumonia | Infectious (Pseudomonas spp and Consulata spp. in sputum) | |

**Abbreviations:** ALI, acute lung injury; AML, acute myeloid leukemia; BAI, bronchoalveolar lavage; COPD, chronic obstructive pulmonary disease; DAD, diffuse alveolar damage; MDS, myelodysplastic syndrome; PAH, pulmonary arterial hypertension; post-op, postoperative; S.p., status post.
were recorded, out of which 17 cases (63%) were > 75 years of age upon death. The proportion of deaths caused by bronchopneumonia eventually decreased in February and March 2020. The incidence of lethal PE remained comparatively low during the examined period, while there was a sharp increase in cases with ARDS and/or respiratory failure in March 2020 (p < 0.05). When comparing the overall incidence of all deaths directly caused by a pulmonary etiology in weekly intervals, there was a statistically significant increase in pulmonary deaths in March 2020 (calendar weeks 10–14; p = 0.02). A total of 13 cases in March 2020 were tested positive for SARS-CoV-2 per ante-mortem nasopharyngeal swab assay. There was no overall increase in the number of deaths in March 2020 (p = 0.45).

Autopsies Performed between October 2019 and March 2020

Seventy-one autopsies were performed at the Institute of Pathology in the period investigated, with a mean autopsy rate of 15.7% between October 2019 and February 2020, which rose to 29.2% in March 2020 (annual adult autopsy rate at University Hospital Basel in 2018: 14.5%). Between October and February, 64% of autopsy cases were male (n = 35), the overall mean age was 71 years, and the mean BMI was 27; in March 2020, 57% of cases were male (n = 12), and the mean age and BMI were significantly higher (age 77 years, p = 0.015; BMI 30, p = 0.03).

Table 1 represents an overview of causes of death according to post-mortem report, sorted by organ system and month. Between October and February, the most common cause of death was cardiovascular (n = 18, 35%), predominantly heart failure (n = 13, 26%). Twenty-five percent (n = 13) of deaths had a direct pulmonary cause, most commonly bronchopneumonia (n = 6, 12%) and PE (n = 4, 8%). Lethal DAD was noted in 2 cases (4%). An unprecedented change of distribution of reported causes of death was observed in March 2020, when a sharp increase of pulmonary mortality (n = 11, 52%) predominantly caused by COVID-19-induced DAD was recorded (n = 9, 42%).

Clinical Characteristics, Histopathological Findings, and RT-PCR Results of Selected Patients with COVID-19-Suggestive Pathology

Out of the 13 cases with a direct pulmonary cause of death between October 2019 and February 2020, 8 with suggestive sequelae or microscopic findings compatible with COVID-19 were selected for retrospective SARS-CoV-2 RT-PCR. Additional three autopsies with highly suggestive COVID-19 were included. Of the 13 cases with a direct pulmonary cause of death in March 2020, two cases with suggestive COVID-19 were included. All 11 cases with COVID-19 were selected for retrospective SARS-CoV-2 RT-PCR. Additional three autopsies with highly suggestive COVID-19 were included. Overall, there was a sharp increase in cases with ARDS and/or respiratory failure in March 2020 (p < 0.05). When comparing the overall incidence of all deaths directly caused by a pulmonary etiology in weekly intervals, there was a statistically significant increase in pulmonary deaths in March 2020 (calendar weeks 10–14; p = 0.02). A total of 13 cases in March 2020 were tested positive for SARS-CoV-2 per ante-mortem nasopharyngeal swab assay. There was no overall increase in the number of deaths in March 2020 (p = 0.45).
suggestive sequelae or microscopic findings performed before October 2019 were also retrospectively analyzed. The average post-mortem interval was 30 h (range 18–69). Clinical characteristics and causes of death are listed in Table 2. This cohort was predominantly male (n = 8, 72.8%) with a mean age of 70 years. The most commonly represented cardiopulmonary comorbidities (presumably predisposing for serious COVID-19) included atherosclerosis (n = 8, 72.7%) and coronary artery disease (45.5%). Five patients (45.5%) were overweight and 2 were morbidly obese (18.2%); the overall mean BMI was 27.4. The cause of death at autopsy was most commonly bronchopneumonia (n = 4, 36.4%), followed by DAD (n = 3, 27.3%) and acute PE (n = 2, 18.2%). Other causes of death included diffuse alveolar hemorrhage and decompensated cor pulmonale. The respective etiology of pneumopathy is presented in Table 2; in some cases the etiology was unclear, with several differential diagnoses.

Table 3 provides an overview of histological findings of the upper respiratory tract and lungs. All 3 cases with DAD presented with both exudative and proliferative
components; in 1 case (No. 1) superimposed pneumonia was detected. All cases presented with capillary stasis and interstitial edema with varying grades of severity. One case with bronchopneumonia also presented with focal exudative DAD. Eight cases (72.7%) presented with hyaline membrane formation and 5 cases (45.5%) presented with microthrombi of pulmonary capillaries. RT-PCR assays performed on all cases yielded negative results.

Discussion

In this retrospective analysis of adult deaths, we aimed to investigate the presence of lethal community spread of SARS-CoV-2 in the region of Basel before its initial emergence in Switzerland in late February 2020. Systematic analysis of the clinical causes of death revealed an increase of deaths directly caused by bronchopneumonia in December 2019 and January 2020, followed by a regression of cases in February (Fig. 1). In line with previous investigations analyzing the incidence and mortality of community-acquired pneumonia and influenza-related illness during the winter months [46–49], these observations suggest a solely seasonal etiology. Seasonality typically affects older age groups [48]; 63% of lethal bronchopneumonia in this cohort affected individuals >75 years of age; data which are also consistent with recent health statistics reports published by the Swiss Federal Office of Statistics [50]. Between October 2019 and February 2020, the incidence of lethal ARDS, respiratory failure and PE stayed consistently low and did not show statistically significant changes over time. The incidence of lethal PE amongst autopsies (n = 6, 8%, Table 1) was lower than that of a previously retrospectively investigated post-mortem cohort [51]. As ARDS and PE constitute some of the major sequelae of severe COVID-19 symptomology [35, 52], their underrepresentation between October and February support the absence of lethal COVID-19 during this period. Furthermore, deaths caused by other etiologies did not show any significant changes in incidence in the period investigated (online suppl. Fig. 1).

A sharp, statistically significant rise in pulmonary causes of death in March 2020 (p = 0.03, <0.05), predominantly caused by COVID-19-associated DAD (Fig. 1), was paralleled amongst autopsies conducted; 52% of autopsies conducted in March had a direct pulmonary cause of death versus 25% in the months before. This abrupt rise was accompanied by a higher autopsy rate in March (29%) as compared to previous months. An increase of pulmonary mortality is uncharacteristic for this time of year and corresponds to the reported emergence of COVID-19 in February 2020. Interestingly, a statistically significant rise of mean age and BMI was recorded during March 2020, supporting a correlation between these factors and lethal COVID-19 as described in previously conducted post-mortem series [26, 27].

Retrospective RT-PCR case selection was performed according to suggestive clinical history as well as histological findings. The current literature does not identify defining histological characteristics of COVID-19 pathology. Rather, patterns of commonly described post-mortem pulmonary findings, which are unspecific but, in combination, considered as highly suggestive for COVID-19 pneumopathy were identified and used as selection criteria for RT-PCR in this study: presence and/or a combination of DAD with severe capillary stasis with or without microthrombi, superimposed bronchopneumonia and acute PE [26–28]. The etiology of DAD is markedly heterogeneous: numerous infectious agents such as viruses and atypical pathogens such as Legionella and Mycoplasma spp. have been known to produce DAD, as well as drug-induced reactions, autoimmune pneumopathies, hypoperfusion, and sepsis [53]. However, SARS-CoV-2-associated DAD differs from its other etiologies in at least one aspect: while microthrombi and capillary stasis can be a feature of DAD, they are likely a histological correlate of COVID-19-associated microangiopathic disease and coagulopathy [26, 35]. This key histological feature of COVID-19-associated DAD in comparison to its other etiologies is illustrated in Figure 2. In our cohort of retrospectively analyzed PCR cases, we have histologically observed capillary stasis in 100% and microthrombosis in 5 out of 11 cases, in line with this specific histology (Table 3). Furthermore, post-mortem analyses have revealed an unusually high representation of vascular and interstitial amyloidosis of the heart or of the lungs in patients with lethal COVID-19 [26]. The pathophysiological aspects of this observation are currently unknown, although microangiopathic vascular amyloid deposits have previously been described in some lungs with lethal ARDS, implying an association with susceptibility to acute lung injury [54].

In our series, 3 out of 5 cases with an infectious etiology (No. 1, 3, and 11) produced a positive microbiological result in sputum or bronchoalveolar lavage (Table 2). Despite these findings, SARS-CoV-2 positivity was still plausible due to the high incidence of secondary co-infections as a result of virally induced pulmonary injury, as described clinically and in previous post-mortem series.
Despite these argumentations, RT-PCR of all retrospectively analyzed samples yielded negative results for SARS-CoV-2. This implicates other likely etiologies as predominantly causative: drug-induced or postoperative for DAD, and obesity/immobility or paraneoplastic for PE. The unclear etiology of the lethal coagulopathy in case 9 may have been due to a genetic predisposition – serology revealed a pathologically high concentration of factor VIII, which increases the likelihood of thromboembolic events [56]. For a detailed discussion of all selected cases, refer to Table 2.

Taking the latest literature into consideration, early community spread of COVID-19 is plausible due to several reasons: compared to SARS and MERS, its evolutionary dynamics suggest high human adaptability [57] in combination with a large number of asymptomatic and/or paucisymptomatic cases [40, 58]. SARS-CoV-2 is also more contagious than other coronaviruses; current estimates of the reproductive number (R0) for SARS-CoV-2 lie around 1–1.5 in Switzerland [59], which is considerably higher than SARS (0.18–1.08) [60] and MERS (<1) [61]. This may be in part due to pre-symptomatic transmission and viral shedding after recovery, although their role in contagion is currently unclear [62, 63]. Moreover, while phylogenetic analyses suggest early widespread transmission involving many independent introductions of the virus [43, 64, 65], they merely provide a temporal snapshot of a rapidly developing pandemic, warranting ongoing in-depth analyses and comparison of global genomic databases.

Most of the current evidence therefore suggests that early international emergence of SARS-CoV-2 in late 2019 is highly unlikely. The earliest case in the USA was reported on January 15, with evidence of limited early community spread in late January or early February [66]. A retrospective analysis of pediatric patients with influenza-related illness did not reveal the presence of COVID-19 in Italy from October 2019 to March 2020 [67]. Finally, lack of early community spread in Central Europe is also supported by the data of this study. Earlier European cases such as the one in France in December [41] are likely isolated events; in these cases, contact tracing is critical to analyze patterns of early temporal spread.

A few limitations apply to our study. Autopsy was not performed in the vast majority of deaths at University Hospital Basel (81.4%), which limits the reliability of this analysis. Furthermore, there are several different ways to interpret a cause of death; to achieve consistent results in this study, it was defined as the most immediate condition leading to the lethal outcome, although this does not take into consideration comorbid conditions, which may have contributed to death. In addition, there is considerable discrepancy between clinical and post-mortem diagnosis, which must be kept in mind when interpreting the results of this investigation [68]. This bias was circumvented by performing a separate statistical analysis for clinical and autopsy cause of death. Moreover, the sole evaluation of deaths is not sufficient to confirm the lack of community spread – as most cases of COVID-19 are not lethal, further analyses of nasopharyngeal throat samples or serology in a study with a similar retrospective design as this one may be worth conducting. Lastly, RT-PCR was performed according to a previously established protocol at our institute, which demonstrated consistency between results in FFPE tissue and ante-mortem nasopharyngeal swabs in a cohort of 21 patients [26]. There is currently no evidence of varying sensitivity of SARS-CoV-2 RT-PCR in FFPE tissue and its false-negative rate compared to detection in fresh samples; recent analyses were able to detect viral load in FFPE, even in subclinical cases [69]. Other factors, such as the post-mortem interval [70], hospitalization time, and type of treatment administered may impact RNA viability. In this respect, future studies on assay standardization are vital to increase the analytical sensitivity of RT-PCR testing.

In conclusion, the data acquired in this retrospective study support the absence of lethal community spread of SARS-CoV-2 between October 2019 and February 2020, indicating that its initial emergence in Switzerland occurred in March 2020. Further retrospective analyses are required to accurately map the timeline of COVID-19 epidemiology in Central Europe.

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Statement of Ethics

This study received approval from the Ethics Committee of Northwestern and Central Switzerland (ID 2020-00629).

Conflict of Interest Statement

The authors have nothing to disclose.
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