



Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.



A Comparison Between Chinese Children Infected with Coronavirus Disease-2019 and with Severe Acute Respiratory Syndrome 2003

Xiaoli Xiong, MD^{1,*}, Gilbert T. Chua, MBBS, FHKAM(Paed)^{2,*}, Shuiqing Chi, MD³, Mike Yat Wah Kwan, MSc(Applied Epidemiology) CUHK, FHKAM(Paed)⁴, Wilfred Hing Sang Wong, PhD², Aifen Zhou, MD⁵, Chi Chiu Shek, FHKAM(Paed), FRCPC⁴, Keith T. S. Tung, MPH², Huan Qin, MMSc⁶, Rosa S. Wong, PhD², Xue Li, PhD^{2,7}, Peng Chen, MBBS⁸, Shuai Li, MD³, Celine S. Chui, PhD^{2,7}, Winnie W. Y. Tso, MBBS, BSc(Hons)², Marco H. K. Ho, MD, FRCP², Ian C. K. Wong, PhD, FRCPC(Hon)^{7,9}, Godfrey C. F. Chan, MD, FRCPC(UK)², Yu Lung Lau, MD (Hon), FRCPC², Kenneth K. Y. Wong, PhD(Lond), FRCS(Edin)¹⁰, Patrick H. Y. Chung, MS(HK), FRCSEd(Paed)¹⁰, Hui Li, MBBS¹¹, Paul K. H. Tam, FRCS(Eng, Edin, Glas, and Ire), FRCPC^{10,12}, Shao-tao Tang, MD^{3,*}, and Patrick Ip, MPH, FRCPC(UK)²

Objectives To compare the clinical and laboratory features of severe acute respiratory syndrome 2003 (SARS) and coronavirus disease 2019 (COVID-19) in 2 Chinese pediatric cohorts, given that the causative pathogens are biologically similar.

Study design This is a cross-sectional study reviewing pediatric patients with SARS (n = 43) and COVID-19 (n = 244) who were admitted to the Princess Margaret Hospital in Hong Kong and Wuhan Children's Hospital in Wuhan, respectively. Demographics, hospital length of stay, and clinical and laboratory features were compared.

Results Overall, 97.7% of patients with SARS and 85.2% of patients with COVID-19 had epidemiologic associations with known cases. Significantly more patients with SARS developed fever, chills, myalgia, malaise, coryza, sore throat, sputum production, nausea, headache, and dizziness than patients with COVID-19. No patients with SARS were asymptomatic at the time of admission, whereas 29.1% and 20.9% of patients with COVID-19 were asymptomatic on admission and throughout their hospital stay, respectively. More patients with SARS required oxygen supplementation than patients with COVID-19 (18.6 vs 4.7%; $P = .004$). Only 1.6% of patients with COVID-19 and 2.3% of patients with SARS required mechanical ventilation. Leukopenia (37.2% vs 18.6%; $P = .008$), lymphopenia (95.4% vs 32.6%; $P < .01$), and thrombocytopenia (41.9% vs 3.8%; $P < .001$) were significantly more common in patients with SARS than in patients with COVID-19. The duration between positive and negative nasopharyngeal aspirate and the length in hospital stay were similar in patients with COVID-19, regardless of whether they were asymptomatic or symptomatic, suggesting a similar duration of viral shedding.

Conclusions Children with COVID-19 were less symptomatic and had more favorable hematologic findings than children with SARS. (*J Pediatr* 2020;224:30-6).

The outbreak of coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) emerged in Wuhan, China, in December 2019. Subsequently, more patients with COVID-19 were diagnosed in other parts of Mainland China, nearby regions and countries in Asia, and beyond. It was declared a pandemic by the World Health Organization on March 11, 2020.¹ The first patient with SARS was reported in Mainland China in 2003, and subsequently other patients with COVID-19 were diagnosed in other parts of the world. Significantly fewer people were affected by SARS-CoV within the 6-month epidemic, and only 8096 cases and 774 deaths were reported worldwide.² There have been no known reported cases of SARS since 2004.

ACE2	Angiotensin-converting enzyme 2
COVID-19	Coronavirus disease 2019
CT	Computed tomography
LOS	Length of stay
NPA	Nasopharyngeal aspirate
SARS	Severe acute respiratory syndrome
SARS-CoV	Severe acute respiratory syndrome coronavirus
SARS-CoV-2	Severe acute respiratory syndrome coronavirus 2

From the ¹Department of Integrated Chinese and Western Medicine, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science and Technology, Wuhan; ²Department of Pediatrics and Adolescent Medicine, The University of Hong Kong, Hong Kong SAR; ³Department of Pediatric Surgery, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan; ⁴Department of Pediatrics and Adolescent Medicine, Princess Margaret Hospital, Hong Kong SAR; ⁵Department of Maternal Healthcare, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science and Technology, Wuhan; ⁶Institute of Maternal and Child Health, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science and Technology, Wuhan; ⁷Center for Safe Medication Practice and Research, Department of Pharmacology and Pharmacy, The University of Hong Kong, Hong Kong SAR; ⁸Department of Respiratory Medicine, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China; ⁹Research Department of Practice and Policy, UCL School of Pharmacy, University College, London, UK; ¹⁰Division of Pediatric Surgery, Department of Surgery, The University of Hong Kong, Hong Kong SAR; ¹¹Department of Hematology, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science and Technology, Wuhan; and ¹²Dr. Li Dak Sum Research Center, The University of Hong Kong-Karolinska Institute Collaboration in Regenerative Medicine, The University of Hong Kong, China.

*Contributed equally.

The authors declare no conflicts of interest.

0022-3476/\$ - see front matter. © 2020 Elsevier Inc. All rights reserved.
<https://doi.org/10.1016/j.jpeds.2020.06.041>

Both SARS-CoV and SARS-CoV-2 belong to the *betacoronavirus* genus and are phylogenetically related to bat SARS-like coronavirus, but relatively more distantly to Middle Eastern Respiratory Syndrome-CoV. SARS-CoV and SARS-CoV-2 share 79% genetic similarity.³ They also share similar infection pathophysiology in humans, because they both bind to the same human receptor, angiotensin-converting enzyme 2 (ACE2), for entry into host cells. However, SARS-CoV-2 has higher transmissibility, with a higher reproductive number of 2.0-2.5 compared with 1.7-1.9 for SARS-CoV.⁴ Studies from Wuhan describing the clinical characteristics of 171 children with COVID-19 showed that 15.8% of patients in their cohort were asymptomatic carriers, and only 1 patient required intensive care with ventilator support.⁵ In contrast, previous studies summarizing the clinical characteristics of pediatric patients with SARS showed they were almost all symptomatic.⁴ A serologic study showed that asymptomatic SARS cases in children was rare.⁶ Furthermore, compared with other coronavirus infections that cause milder diseases, both SARS and COVID-19 have significantly greater morbidity and mortality.⁷ Nevertheless, studies providing a direct comparison of clinical and laboratory features between children infected with SARS and COVID-19 are lacking. In this study, we investigated the clinical and laboratory features in 2 representative Chinese pediatric cohorts with SARS and COVID-19. Understanding the differences and similarities in the clinical phenotypes and laboratory measures in pediatric patients with SARS and COVID-19 will guide us in the identification, quarantine, and management of infected children in a timely manner.

Methods

This was a comparative study examining the clinical and laboratory features of Chinese children (aged ≤ 18 years at admission) with SARS in Hong Kong and COVID-19 in Wuhan, China. In Hong Kong, pediatric patients diagnosed with SARS and admitted to Princess Margaret Hospital from March 1 to April 30, 2003, were included in the study. Princess Margaret Hospital is a key public hospital that served as the major center for managing patients with SARS in Hong Kong at the beginning of the SARS-CoV outbreak in 2003. More than one-third (36%) of pediatric patients with SARS in Hong Kong were admitted to and managed in Princess Margaret Hospital. The case definitions and clinical characteristics have been previously published in a peer-reviewed journal.⁸ In Wuhan, pediatric patients who tested positive for SARS-CoV-2, as confirmed by nasopharyngeal aspirate (NPA) specimen using a reverse-transcriptase polymerase chain reaction test and who were admitted to the Wuhan Children's Hospital between January 21 and March 20, 2020, were included in this study. The Wuhan Children's Hospital is the main center assigned by the central government for treating children diagnosed with SARS-CoV-2 infection in Wuhan.⁵ Children in both cohorts were tested if symptomatic or if in contact with a confirmed case of SARS or COVID-19. Those with a positive test were hospital-

ized regardless of symptoms. Those with SARS or SARS-CoV-2 infection are referred to as SARS or COVID-19 cases, respectively, regardless of symptoms. Hospital records and laboratory results from both SARS and COVID-19 cohorts were retrieved and analyzed. Demographics, clinical symptoms, days between positive and negative NPA, length of stay (LOS) in hospital, need for oxygen and mechanical ventilation, and laboratory tests were compared between the 2 cohorts. The duration between positive and negative NPA was defined as the time between the first SARS-CoV-2 positive NPA specimen and the first of 2 consecutive NPA negative specimens. Asymptomatic patients were defined in Wuhan as no clinical symptoms and no abnormal computed tomography (CT) findings, as all patients with COVID-19 received a CT scan of the thorax before transfer or after admission to the Wuhan Children's Hospital.⁹ Both patients with COVID-19 and patients with SARS were discharged after resolution of the clinical symptoms and after receiving 2 consecutive negative NPA tests for SARS-CoV-2 and SARS-CoV, respectively. Patients with SARS were also mandated to stay in the hospital for 21 days from symptom onset.

Statistical Analyses

Continuous variables were expressed as mean \pm SD or median (IQR) and analyzed by the independent *t* test or Mann-Whitney *U* test where appropriate. Categorical variables for SARS, symptomatic COVID-19, and asymptomatic COVID-19 groups were expressed as number (%) and analyzed by a χ^2 test, Fisher exact test, or ANOVA. Age- and sex-matched references from Chinese children were used for comparing blood measures. A 2-sided α of $< .05$ was considered statistically significant. Patients with missing data will not be analyzed for the particular variables. Statistical analyses were performed using SPSS software version 19 (IBM, Armonk, New York, NY), Microsoft Excel (Microsoft, Redmond, Washington), and Statistical Analysis System v9.4 (SAS Institute Inc., Cary, North Carolina).

Ethics

The study protocol was approved by the University of Hong Kong/Hospital Authority Hong Kong West Cluster Institutional Review Board (Reference number: UW 20-292) and the Research Ethics Board of the Wuhan Children's Hospital (Reference number: WHCH 2020022).

Results

Patient Demographics

A total of 43 pediatric patients with SARS and 244 pediatric patients with COVID-19 were recruited into the study. Demographic and clinical characteristics are shown in **Table I**. The median age of the COVID-19 cohort was 82 months and the SARS cohort was 160.8 months (**Figure 1**). There were a total of 85 (34.8%) asymptomatic patients with COVID-19 compared with no asymptomatic patients with SARS at presentation ($P < .001$). Among asymptomatic patients with COVID-19, 94.1% (48/51) had family members with COVID-19 and were identified by screening

Table I. Characteristics and symptoms in pediatric patients with COVID-19 and SARS-CoV

Characteristics	Wuhan COVID-19 2020 (n = 244)			Hong Kong SARS-CoV 2003 (n = 43)	P value [†]
	Symptomatic (n = 193)	Asymptomatic (n = 51)	P value*		
Demographics and epidemiology					
Age, mo	67.0 (106)	116.0 (71)	<.001	160.8 (90)	<.001
Male sex	120 (62.2)	30 (58.8)	.746	20 (46.5)	.0852
Epidemiologic link identified	160 (82.9)	48 (94.1)	.047	42 (97.7)	.0083
Duration of NPA turning negative, d	8.2 ± 5.8	8.9 ± 5.1	.434	–	–
Length of hospital stay, d	13.0 ± 6.0	11.4 ± 5.2	.066	20.6 ± 3.6	<.001
Symptoms					
Fever	99 (51.3)	–	–	42 (97.7)	<.001
Chills	2 (1.0)	–	–	14 (32.6)	<.001
Myalgia	9 (4.7)	–	–	16 (37.2)	<.001
Malaise	9 (6.5)	–	–	25 (58.1)	<.001
Poor feeding	8 (4.1)	–	–	3 (7.0)	.426
Coryza	24 (12.4)	–	–	17 (39.5)	<.001
Sore throat	10 (5.2)	–	–	7 (16.3)	.019
Cough	120 (62.2)	–	–	28 (65.1)	.862
Sputum	25 (13.0)	–	–	17 (39.5)	<.001
Diarrhea	15 (7.8)	–	–	7 (16.3)	.141
Nausea	23 (11.9)	–	–	12 (27.9)	.015
Abdominal pain	4 (2.1)	–	–	3 (7.0)	.116
Headache	10 (5.2)	–	–	15 (34.9)	<.001
Dizziness	3 (1.6)	–	–	5 (11.6)	.006
Supportive care					
Oxygen supplementation	9 (4.7)	–	–	8 (18.6)	.004
Mechanical ventilation support	3 (1.6)	–	–	1 (2.3)	.559

Data are median (IQR), number (%), or mean ± SD.

*Comparison between the COVID-19 symptomatic and asymptomatic groups.

†Comparison between the COVID-19 symptomatic and SARS groups.

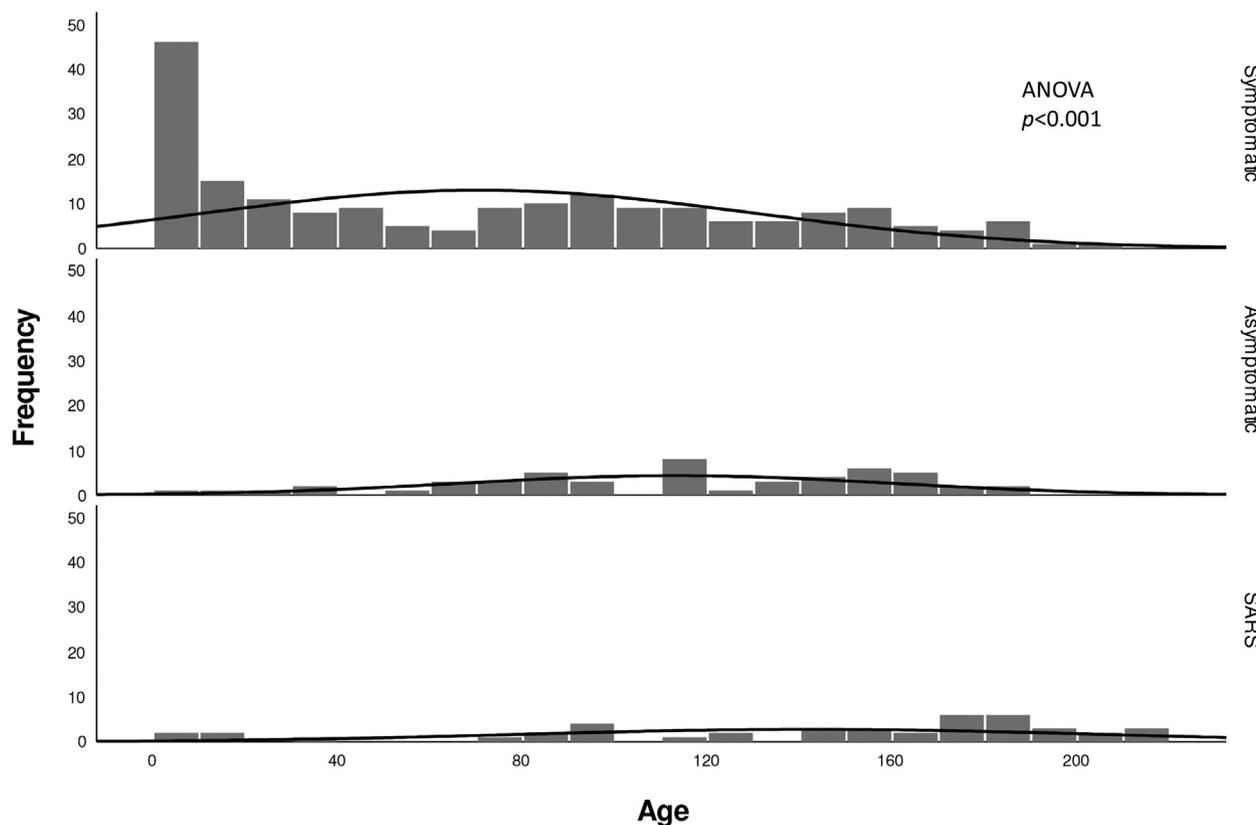


Figure 1. Age distribution of patients with COVID-19 and patients with SARS.

after family members had been confirmed positive. Among the 85 clinically asymptomatic patients with COVID-19 at presentation, 34 were considered symptomatic by definition after admission. Among them, 26.5% (9/34) developed clinical symptoms including cough, nasal congestion, vomiting, diarrhea, poor feeding. Among the symptomatic patients, 73.5% (25/34) were found to have mild abnormalities on CT imaging, including ground glass opacities and pneumonic changes; only 1 patient had both clinical symptoms and CT abnormalities.

Overall, 97.7% of patients with SARS (42/43) and 85.2% of patients with COVID-19 (208/242) had identifiable epidemiologic links in known cases. In the COVID-19 cohort, a 10-month-old boy died of intussusception and multiorgan failure, which has been reported elsewhere.⁵ No mortality was reported in the SARS cohort. There were no significant differences in the time from positive to negative NPA between the symptomatic and asymptomatic COVID-19 groups (mean of 8.2 days vs 8.9 days; $P = .434$). There were also no significant differences in the LOS in hospital between symptomatic and asymptomatic patients with COVID-19 (mean 13.0 days vs 11.4 days; $P = .066$), but the LOS in the SARS cohort (all symptomatic) was significantly longer than in the symptomatic COVID-19 cohort (mean 20.6 days vs 12.9 days; $P < .001$).

Clinical Symptoms

A comparison of symptoms between the SARS and COVID-19 cohorts are presented in [Table I](#) and [Figure 2](#). Compared with patients with COVID-19, significantly more patients with SARS developed fever ($P < .001$), chills ($P < .001$), myalgia ($P < .001$), malaise ($P < .001$), coryza ($P < .001$), sore throat

($P = .019$), sputum production ($P < .001$), nausea ($P = .015$), headache ($P < .001$), and dizziness ($P = .006$). Almost all patients with SARS developed fever, compared with only 51.3% of patients with COVID-19. Cough symptoms occurred similarly in both cohorts. More patients with SARS required oxygen supplementation than patients with COVID-19 (18.6% vs 4.7%; $P = .004$), whereas only 1.6% and 2.3% of patients with COVID-19 and patients with SARS required mechanical ventilatory support, respectively.

Laboratory Test Results

Laboratory testing results for SARS and symptomatic patients with COVID-19 are presented in [Table II](#). Patients with SARS had a significantly lower total white blood cell count ($P = .043$), lymphocyte count ($P < .001$), and platelet ($P < .001$) count, and lower albumin ($P < .001$), higher globulin ($P < .001$), higher alanine aminotransferase ($P = .01$), and lower D-dimer ($P < .001$) levels. We further explored the number of children with leukopenia, lymphopenia, and thrombocytopenia using age- and sex-matched reference ranges for Chinese and Asian children.¹⁰⁻¹² Leukopenia (37.2% vs 18.6%; $P = .008$), lymphopenia (95.4% vs 32.6%; $P < .01$), and thrombocytopenia (41.9% vs 3.8%; $P < .001$) were significantly more common in patients with SARS than in patients with COVID-19. The laboratory test results for asymptomatic patients with COVID-19 were normal.

Discussion

In our study of Chinese children infected with COVID-19 and SARS in representative pediatric cohorts from Wuhan and

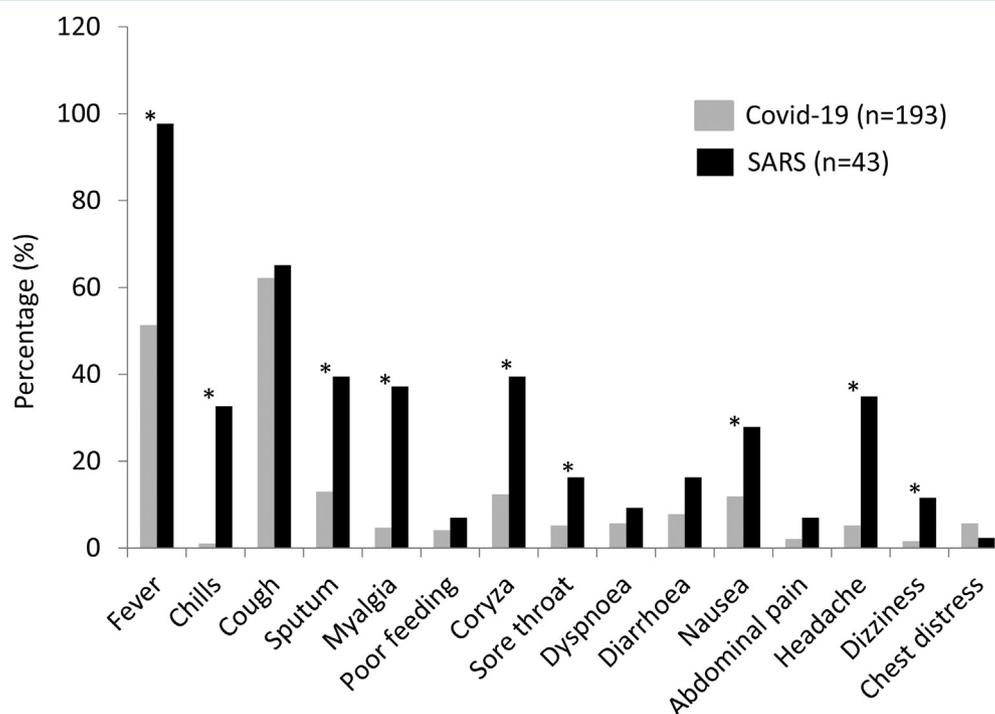


Figure 2. COVID-19 and SARS symptoms comparison. * $P < .05$.

Table II. Laboratory test results in pediatric patients with COVID-19 (symptomatic cases) and SARS

Laboratory tests	Wuhan, COVID-19 2020 (n = 193)				Hong Kong, SARS-CoV 2003 (n = 43)				
	Mean/Median	SD/IQR	Range	No.	Mean/Median	SD/IQR	Range	No.	P value
White cell count, $\times 10^9/L$	7.1	2.7	0.75-13.8	183	6.0	4.4	2.2-23.2	43	.043
Hemoglobin, g/dL	12.4	1.5	6.7-18.3	183	16.1	18.0	7.6-131.0	43	.193
Platelet, $\times 10^9/L$	291.5	128.8	14.0-751.0	183	212.0	76.5	168.0-580.0	43	<.001*
Lymphocyte, $\times 10^9/L$	3.4	1.9	0.2-11.7	184	2.1	2.8	0.6-9.4	43	<.001
Albumin, g/L	45.0	3.5	34.7-55.9	192	43.4	6.5	29.0-52.0	40	<.001
Globulin, g/L	22.8	5.0	11.1-32.2	187	34.0	5.5	23.0-44.0	40	<.001
ALT, U/L	16.0	17.0	4.0-596.0	193	17.0	20.0	12.0-80.0	41	.010*
LDH, U/L	239.0	94.2	142.0-656.0	189	262.0	168.5	147.0-1143.0	36	.285*
DD, mg/L FEU	1.2	8.9	0.01-100.8	140	-0.11	0.18	-0.2 to 0.2	11	<.001*
APTT, sec	34.3	28.6	23.4-353.0	169	32.0	4.2	25.9-37.3	29	.827

ALT, alanine aminotransferase; APTT, activated partial thromboplastin time; DD, D-dimer; FEU, fibrinogen equivalent units; LDH, lactate dehydrogenase.
*Mann-Whitney U test, results presented as median (IQR).

Hong Kong, patients with COVID-19 experienced fewer symptoms than patients with SARS. Almost all patients with SARS presented with fever compared with only one-half of patients with COVID-19, which suggests screening by body temperature will potentially miss one-half of the children infected with COVID-19. Although a similar percentage of patients in both cohorts had a cough, significantly more patients with SARS had other symptoms, including fever and respiratory difficulties.^{8,13} A study comparing the replication, cell tropism, and immune activation profile of SARS-CoV-2 and SARS-CoV infections in human lung tissue showed that SARS-CoV-2 infection generated 3.2 times more virus particles, yet induced significantly less type I, II, and III interferons and other proinflammatory cytokines, which might explain the relatively milder phenotype in COVID-19.¹⁴

Only a few children in both cohorts required oxygen therapy and mechanical ventilation. In contrast with adult patients, our results echo that children infected by either SARS-CoV-2 or SARS-CoV had a good prognosis.^{15,16} Indeed, studies have shown that a higher percentage of older adults with COVID-19 develop complications, including pneumonia, acute myocardial injury, and acute respiratory distress syndrome, with an overall mortality of $\leq 11\%$ in some populations.¹⁷ Similar observations were reported during the SARS-CoV epidemic in 2003, which had 7% and 17% mortality in Mainland China and Hong Kong, respectively.² The majority of the patients who died from SARS and COVID-19 were older patients with underlying chronic illnesses or who were immunocompromised.¹⁸ A study comparing published data in the literature also concurred with our findings that children with SARS and COVID-19 infection had favorable outcomes.¹⁹ One possible explanation why children are less affected by both SARS and COVID-19 is the different expressions of ACE2 in children and adults. The ACE2 receptor is crucial for both SARS-CoV and SARS-CoV-2 to enter into host cells.²⁰ It has been demonstrated that ACE2 expression can influence the infectivity of SARS-CoV in vitro.²¹ Nevertheless, studies demonstrating age variation of ACE2 expression currently are lacking.

Our study also showed there were significantly more asymptomatic carriers in the pediatric group with COVID-19; there were none in the pediatric SARS group. This finding was consistent with studies in adults with SARS, which found only a few

asymptomatic adults or healthcare workers who tested antibody positive after the epidemic or had only mild and self-limiting symptoms.^{22,23} To the contrary, asymptomatic COVID-19 infections were reported to be common among adults and younger adults without comorbidities.²⁴ Asymptomatic transmission of SARS-CoV-2 may be one of the key factors leading to the pandemic.²⁵ In our study, the average time to achieve a negative NPA and hospital LOS were similar for symptomatic and asymptomatic patients with COVID-19, which indicates that clearance of COVID-19 virus may be similar for symptomatic and asymptomatic patients. A difficulty in concluding this is that dating of the first specimen at the onset in symptomatic patients cannot be replicated in asymptomatic patients. Although we found the LOS in hospital for the SARS cohort was significantly longer than that for the symptomatic COVID-19 cohort (20.6 days vs 11.4 days; $P < .001$), a difference in the isolation policy between the 2 outbreaks likely confounded this observation. In the SARS outbreak in 2003, the mandatory hospitalization for patients was 21 days from symptoms onset in addition to a negative NPA and resolution of symptoms. More important, our study also revealed that 35.6% of children infected with SARS-CoV-2 were asymptomatic on admission and 14.9% subsequently became symptomatic. These asymptomatic patients are still carriers of the virus and have the potential to spread the disease to others, leading to further burdens on public health systems. Public health measures such as the universal use of facemasks, social distancing, early quarantine, and identification and tracing of asymptomatic carriers of COVID-19 have already been widely adopted in places with less severe outbreak such as Hong Kong, Macau, and Taiwan. Studies have shown that frequent and proper use of facemasks in public areas was associated with a 60% lower risk of contracting SARS-CoV compared with infrequent use during the SARS outbreak in 2003.²⁶ Repeated reverse-transcriptase polymerase chain reaction testing for SARS-CoV-2 should also be performed in those highly suspected of infection where there is an initial possible false-negative result. Given the evidence of a significant proportion of asymptomatic COVID-19 carriers and experience from Asia, strict preventive measures should be enacted to control the spread of COVID-19.

We found that the COVID-19 cohort had more favorable hematologic and biochemical findings, with fewer patients

with COVID-19 having abnormal test results compared with patients with SARS. The mean total white blood cell, lymphocyte, and median platelet counts in patients with SARS were all significantly lower than in patients with COVID-19. Potential mechanisms include direct infection of hematopoietic progenitor cells via cell surface CD13 or CD66a that possibly induces growth inhibition and apoptosis.²⁷ Although the COVID-19 cohort was significantly younger than the SARS cohort, significantly fewer children in the COVID-19 cohort had hematologic abnormalities when using age- and sex-matched references. Lymphopenia was observed during the early phase of the SARS epidemic, predominantly CD4⁺ and CD8⁺ lymphopenia, and adult patients with SARS-CoV who were clinically more severe or died had significantly more severe CD4⁺ and CD8⁺ lymphopenia.^{28,29} In COVID-19 infection, the degree of lymphopenia was reported to be a prognostic indicator for older patients.³⁰ The significantly lower proportion of pediatric patients with COVID-19 with lymphopenia may explain the milder phenotype compared with the SARS group.

This study had several limitations. First, this was retrospective observational study design, which may have potential recall or sampling bias. Nevertheless, the majority of children with COVID-19 under the age of 18 years in Wuhan were admitted to the Wuhan Children's Hospital, and the majority of pediatric patients with SARS in Hong Kong were admitted to Princess Margaret Hospital. Wuhan and Hong Kong were among the most seriously affected cities in the COVID-19 and SARS outbreaks, respectively. Therefore, they are representative pediatric cohorts for these 2 infections. Second, not all family contacts of SARS cases were tested in 2003. Nevertheless, in the study conducted by Lee et al, only 0.57% of asymptomatic children from the Amoy Garden were positive for SARS-CoV antibody.⁶ None of the 14 asymptomatic children who had contacts with patients with SARS were seropositive.⁶ The Amoy Garden was one of the housing estates in Hong Kong with major community outbreak. It was also where the majority of the SARS-infected children in this cohort were recruited. Therefore, despite the lack of close contact screening in 2003, SARS asymptomatic carriage was considered to be rare. Third, younger patients with COVID-19 may not be able to describe their subjective symptoms, such as anosmia, which is considered a novel symptom of COVID-19, which may lead to under-reporting of these symptoms.³¹ Fourth, CT and other radiologic findings were not reported in details as the majority of the patients with SARS did not have a CT scan performed and therefore was considered out of the scope of this manuscript. Radiologic findings of some of the patients from Wuhan's Children Hospital have been reported elsewhere.⁹ Fifth, a recent phylogenetic network study showed that the predominant SARS-CoV-2 variant (type B) in East Asia is genetically different from the variants in Europe and America (types A and C). Whether such differences lead to different clinical characteristics between Asia and the West remains uncertain, and one must be cautious in extrapolating clinical characteristics of COVID-19 in Asia to other geographic areas.³² Finally, we did not observe any children developing Kawasaki-like hyperinflammatory shock that

was reported in European children with COVID-19 in our Chinese cohort.^{33,34} Future studies combining and comparing data from international clinical centers will be meaningful to examine the demographics and clinical spectrum of pediatric patients with COVID-19 on a global perspective, and to further identify the risk factors for these severe diseases. ■

We thank all the staff members of the Department of Pediatrics and Adolescent Medicine, the Princess Margaret Hospital, and Wuhan Children's Hospital for their dedication in caring for sick children, which can be a very difficult time for both patients and their families.

Submitted for publication May 19, 2020; last revision received Jun 9, 2020; accepted Jun 10, 2020.

Reprint requests: Patrick Ip, MPH, FRCPCH(UK), Clinical Associate Professor, Department of Pediatrics and Adolescent Medicine, The University of Hong Kong, Hong Kong SAR. E-mail: patricip@hku.hk

References

1. Coronavirus disease (COVID-19) Pandemic: World Health Organization; 2020. <https://www.who.int/emergencies/diseases/novel-coronavirus-2019>. Accessed May 10, 2020.
2. Summary table of SARS cases by country, 1 November 2002 - 7 August 2003: World Health Organization; 2003. https://www.who.int/csr/sars/country/country2003_08_15.pdf?ua=1. Accessed April 18, 2020.
3. Lu R, Zhao X, Li J, Niu P, Yang B, Wu H, et al. Genomic characterisation and epidemiology of 2019 novel coronavirus: implications for virus origins and receptor binding. *Lancet* 2020;395:565-74.
4. Petrosillo N, Viceconte G, Ergonul O, Ippolito G, Petersen E. COVID-19, SARS and MERS: are they closely related? *Clin Microbiol Infect* 2020;26:729-34.
5. Lu X, Zhang L, Du H, Zhang J, Li YY, Qu J, et al. SARS-CoV-2 Infection in Children. *N Engl J Med* 2020;382:1663-5.
6. Lee PP, Wong WH, Leung GM, Chiu SS, Chan KH, Peiris JS, et al. Risk-stratified seroprevalence of severe acute respiratory syndrome coronavirus among children in Hong Kong. *Pediatrics* 2006;117:e1156-62.
7. Lau SKP, Woo PCY, Yip CCY, Tse H, Tsoi HW, Cheng VC, et al. Coronavirus HKU1 and other coronavirus infections in Hong Kong. *J Clin Microbiol* 2006;44:2063-71.
8. Leung C-w, Kwan Y-w, Ko P-w, Chiu SS, Loung PY, Fong NC, et al. Severe acute respiratory syndrome among children. *Pediatrics* 2004;113:e535.
9. Ma H, Hu J, Tian J, Zhou X, Li H, Laws MT, et al. A single-center, retrospective study of COVID-19 features in children: a descriptive investigation. *BMC Med* 2020;18:123.
10. Ding Y, Zhou L, Xia Y, Wang W, Wang Y, Li L, et al. Reference values for peripheral blood lymphocyte subsets of healthy children in China. *J Allergy Clin Immunol* 2018;142:970-3.e8.
11. Zhang X, Ding Y, Zhang Y, Xing J, Dai Y, Yuan E. Age- and sex-specific reference intervals for hematologic analytes in Chinese children. *Int J Lab Hematol* 2019;41:331-7.
12. Nah EH, Kim S, Cho S, Cho HI. Complete blood count reference intervals and patterns of changes across pediatric, adult, and geriatric ages in Korea. *Ann Lab Med* 2018;38:503-11.
13. Stockman LJ, Massoudi MS, Helfand R, Erdman D, Siwek AM, Anderson LJ, et al. Severe acute respiratory syndrome in children. *Pediatr Infect Dis J* 2007;26:68-74.
14. Chu H, Chan JF-W, Wang Y, Yuen TT, Chai Y, Hou Y, et al. Comparative replication and immune activation profiles of SARS-CoV-2 and SARS-CoV in human lungs: an ex vivo study with implications for the pathogenesis of COVID-19. *Clin Infect Dis* 2020 Apr 9 [Epub ahead of print].
15. Ludvigsson JF. Systematic review of COVID-19 in children shows milder cases and a better prognosis than adults. *Acta Paediatr* 2020;109:1088-95.
16. Chiu WK, Cheung PC, Ng KL, Ip PL, Sugunan VK, Luk DC, et al. Severe acute respiratory syndrome in children: experience in a regional hospital in Hong Kong. *Pediatr Crit Care Med* 2003;4:279-83.

17. Rajgor DD, Lee MH, Archuleta S, Bagdasarian N, Quek SC. The many estimates of the COVID-19 case fatality rate. *Lancet Infect Dis* 2020;20:776-7.
18. Jordan RE, Adab P, Cheng KK. Covid-19: risk factors for severe disease and death. *BMJ* 2020;368:m1198.
19. Gupta S, Malhotra N, Gupta N, Agrawal S, Ish P. The curious case of coronavirus disease 2019 (COVID-19) in children. *J Pediatr* 2020;222:258-9.
20. Guo J, Huang Z, Lin L, Lv J. Coronavirus disease 2019 (COVID-19) and Cardiovascular disease: a viewpoint on the potential influence of angiotensin-converting enzyme inhibitors/angiotensin receptor blockers on onset and severity of severe acute respiratory syndrome coronavirus 2 infection. *J Am Heart Assoc* 2020;9:e016219.
21. Jia HP, Look DC, Shi L, Hickey M, Pewe L, Netland J, et al. ACE2 receptor expression and severe acute respiratory syndrome coronavirus infection depend on differentiation of human airway epithelia. *J Virol* 2005;79:14614-21.
22. Che X-y, Di B, Zhao G-p, Alizadeh BZ, Wijmenga C, Witt M. A patient with asymptomatic severe acute respiratory syndrome (SARS) and antigenemia from the 2003–2004 community outbreak of SARS in Guangzhou, China. *Clin Infect Dis* 2006;43:e1-5.
23. Kwan MY, Chan WM, Ko PW, Leung CW, Chiu MC. Severe acute respiratory syndrome can be mild in children. *Pediatr Infect Dis J* 2004;23:1172-4.
24. He G, Sun W, Fang P, Huang J, Gamber M, Cai J, et al. The clinical feature of silent infections of novel coronavirus infection (COVID-19) in Wenzhou. *J Med Virol* 2020 Apr 10 [Epub ahead of print].
25. Bai Y, Yao L, Wei T, Tian F, Jin DY, Chen L, et al. Presumed asymptomatic carrier transmission of COVID-19. *JAMA* 2020;323:1406-7.
26. Chan KH, Yuen K-Y. COVID-19 epidemic: disentangling the re-emerging controversy about medical facemasks from an epidemiological perspective. *Int J Epidemiol* 2020 Mar 31 [Epub ahead of print].
27. Yang M, Li CK, Li K, Hon KL, Ng MH, Chan PK, et al. Hematological findings in SARS patients and possible mechanisms (review). *Int J Mol Med* 2004;14:311-5.
28. Panesar NS. Lymphopenia in SARS. *Lancet* 2003;361:1985.
29. Cui W, Fan Y, Wu W, Zhang F, Wang JY, Ni AP. Expression of lymphocytes and lymphocyte subsets in patients with severe acute respiratory syndrome. *Clin Infect Dis* 2003;37:857-9.
30. Tan L, Wang Q, Zhang D, Ding J, Huang Q, Tang YQ, et al. Lymphopenia predicts disease severity of COVID-19: a descriptive and predictive study. *Signal Transduct Target Ther* 2020;5:33.
31. Gane SB, Kelly C, Hopkins C. Isolated sudden onset anosmia in COVID-19 infection. A novel syndrome? *Rhinology* 2020;58:229-301.
32. Forster P, Forster L, Renfrew C, Forster M. Phylogenetic network analysis of SARS-CoV-2 genomes. *Proc Natl Acad Sci U S A* 2020;117:9241-3.
33. Riphagen S, Gomez X, Gonzalez-Martinez C, Wilkinson N, Theocharis P. Hyperinflammatory shock in children during COVID-19 pandemic. *Lancet* 2020;395:1607-8.
34. Jones VG, Mills M, Suarez D, Hogan CA, Yeh D, Segal JB, et al. COVID-19 and Kawasaki disease: novel virus and novel case. *Hosp Pediatr* 2020;10:537-40.

50 Years Ago in *THE JOURNAL OF PEDIATRICS*

Neonatal Mortality: Making the Nonpreventable Preventable

Valdes-Dapena MA, Arey JB. The causes of neonatal mortality: an analysis of 501 autopsies on newborn infants. *J Pediatr* 1970; 77:366-75.

Valdes-Dapena and Arey showed us that the main causes of neonatal mortality are diseases secondary to prematurity and those associated with inflammatory diseases. But as medicine continues to advance, the frequency and epidemiology of diseases tend to change as well. Such is the case with neonatal mortality, which according to the World Health Organization is showing a downward trend in recent years, from 0.43% in 2008 to 0.36% in 2015, with indications that this trend will continue in the coming years.¹

Fifty years ago, the trends toward an increase in conditions associated with prematurity such as pulmonary hyaline membrane disease or intraventricular hemorrhage and a decrease in cases of fetal hypoxia and fetal trauma were already evident. However, there was no decrease in mortality secondary to inflammatory lesions compared with previous years. Although it is true that most preventable diseases, such as inflammatory lesions, are considered problems of the past, the rates of death and complications associated with prematurity remain high.

As the years pass and neonatal critical care and technologies improve, there will be an increasing number of newborns of lower gestational age and of lower birth weight. We are better prepared to deal with premature (and extremely premature) newborns, which translates into decreased mortality. Thanks to the progress and innovations in medicine, the preventable causes of mortality seem to have remained in the past. Preventing premature births seems to be more complex, but 50 years from now we might view it as something easily accomplished.

Ramón Alanís Álvarez, MD

Department of Pediatrics

Programa Multicéntrico de Especialidades Médicas ITESM–SSNL

Monterrey, México

Reference

1. World Health Organization. Newborns: reducing mortality. 2020. www.who.int/news-room/fact-sheets/detail/newborns-reducing-mortality. Accessed February 25, 2020.