Changes in Critical Bronchiolitis After COVID-19 Lockdown

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Abstract

Introduction
In response to the coronavirus disease 2019 (COVID-19) pandemic, state and local governments implemented mitigation strategies, including lockdowns, thereby averting the typical fall/winter 2020 bronchiolitis season and reducing the incidence of respiratory viruses, such as respiratory syncytial virus (RSV). Florida implemented a strict lockdown from April 1, 2020, to April 30, 2020. The removal of lockdown precautions on September 25, 2020, was followed by an atypical out-of-season surge of bronchiolitis in April 2021. Anecdotally, this surge appeared to be associated with both increased poly-viral coinfections and disease severity.

Objective
To determine if the bronchiolitis out-of-season surge differed from historical seasonal case patterns.

Methods
A single-center retrospective cohort study of admissions to the pediatric intensive care unit (PICU) with International Classification of Diseases, Tenth Revision (ICD-10) codes of bronchiolitis, from December 9, 2019, to February 29, 2020 (12 weeks, pre-lockdown group or PreLD), was compared to March 29, 2021, to June 19, 2021 (12 weeks, post-lockdown group or PostLD). Variables used for comparison were gender, ethnicity, age, viral coinfections, viruses detected, PICU length of stay, hospital length of stay, mortality, maximum respiratory support needed, mechanical ventilation days, extracorporeal life support (ECLS) days, and severity of disease measured by Pediatric Logistic Organ Dysfunction-2 (PELOD-2) and Pediatric Sequential Organ Failure Assessment (pSOFA). Categorical data were analyzed using Fisher’s exact test, and a t-test was used for continuous variables. A two-sided p < 0.05 was considered significant.

Results
A total of 135 subjects were analyzed from the two cohorts. More patients were admitted during the PostLD phase (87 vs. 48). The PostLD group had a higher age at admission (11.2 ± 12.3 vs. 6.6 ± 7.5, p = 0.0075), but there were no differences in gender or race/ethnicity. The PostLD group also exhibited a higher proportion of RSV infections (73 vs. 16, p < 0.0001) and poly-viral infections (p < 0.0001). Higher coronavirus OC43 (9 vs. 0, p = 0.0265) and parainfluenza types 1-4 (human parainfluenza virus (HPIV)) (19 vs. 1, p = 0.0117) detections, yet fewer human metapneumovirus (HMPV) detections (0 vs. 4, p = 0.0147), were observed PostLD. No differences were found in hospital length of stay, PICU length of stay, mortality, mechanical ventilation days, ECLS days, or severity of illness scores based on PELOD-2 or pSOFA scores.

Conclusion
In the bronchiolitis out-of-season surge, there were an increased number of admissions to the PICU. Those patients were older, and more likely to have RSV, as well as a coinfection with coronavirus OC43 or HPIV, yet less likely to have HMPV. No difference in length of stay or disease severity was demonstrated.

Categories: Pediatrics, Infectious Disease, Other
Keywords: viral coinfections, trend, severity of disease, respiratory syncytial virus (rsv), pediatric critical care, bronchiolitis, covid-19

Introduction
In December 2019, a cluster of respiratory infections caused by a novel strain of coronavirus (SARS-CoV-2) linked to a seafood market in Wuhan, China marked the start of an unprecedented worldwide pandemic [1]. The coronavirus disease 2019 (COVID-19) pandemic was officially declared on March 11, 2020. To date, there have been over 270 million confirmed cases and more than 5 million deaths worldwide [2]. In response,
many countries implemented lockdowns (e.g. permitting only essential activities to operate, initiating in-
person school closures, and enforcing mandatory quarantine to travelers), social distancing, mandatory
mask usage, and isolation of infected patients [3]. In the United States, most states implemented lockdown
orders from late March to early April 2020, which were lifted beginning late April to early June 2020, with a
varying degree of other precautions remaining in place depending on the state [4]: In the majority of
counties in Florida, complete lockdown orders took place from April 1st through April 30, 2020, limiting
only essential businesses to operate. By May 1, 2020, non-essential businesses operated with 25% capacity;
by June 5, 2020, the allowed capacity increased to 50%, and personal care services were able to operate; by
August 31, 2020, public schools reopened in person; and by September 25, 2020, all capacity restrictions
were lifted [5].

The change in the implemented precautions led to a historically unusual and unexpected change in the
incidence of respiratory infections. In the United States, respiratory viruses commonly follow a seasonal
pattern from late fall through early spring. However, observations have demonstrated an initial decrease in
respiratory syncytial virus (RSV) infections in the fall–winter months of 2020, followed by a significant
increase in April 2021 [6]. Influenza viruses and human metapneumovirus (HMPV) infections remained low
from March 2020 through May 2021, but increased infections by parainfluenza (types 1–4), adenovirus, and
non-SARS-CoV-2 coronaviruses have been noted in the spring of 2021, followed by a decline in early 2020
[7], as well as an overall decreased incidence in invasive bacterial respiratory diseases in the winter of 2020
[8].

Bronchiolitis, which is most commonly caused by RSV, is the most common lower respiratory tract infection
in children younger than two years of age. It is characterized by irritability, low-grade fever, nasal
congestion, wheezing, and increased work of breathing and can cause severe respiratory symptoms,
especially in children with underlying conditions [9]. Representing an important disease burden,
bronchiolitis is the most common cause of hospitalization in the United States in children younger than two
years of age. It comprises 18% of overall pediatric admissions and is the second most common cause of
infectious deaths in infants in developing countries. Typically, seasonal outbreaks of bronchiolitis occur in
the fall/winter months in the Northern Hemisphere including in the United States [10,11], with the exception
of Florida having the longest outbreaks compared to the rest of the nation, with expected seasons from
September through March in the northern region of the state [12].

Anecdotally, the pandemic-related out-of-season outbreak of bronchiolitis has been associated with
multiple viral coinfections, increased severity of disease, more admissions to the pediatric intensive care
unit (PICU), more interventions, and longer hospital stays. This study aimed to determine if this upsurge of
cases from March through June 2021 did, in fact, have increased the severity of illness and medical support
requirement compared with the typical seasonal spike of bronchiolitis from December 2019 through
February 2020 at our institution. While the surge in cases post-lockdown has been described, no studies have
yet been published examining the severity of illness comparisons.

Materials And Methods
We conducted, via chart review, a single-center retrospective cohort study of critically ill patients admitted
to the PICU at UF Health Shands Children’s Hospital in Gainesville, Florida with a documented bronchiolitis
diagnosis (International Classification of Diseases, Tenth Revision (ICD-10) codes: J21.9, J11.1, J21.8, J21.0,
and J21.1), from December 1, 2019, through February 29, 2020 (pre-lockdown group or PreLD), and March
29, 2021, through June 19, 2021 (post-lockdown group or PostLD).

All data were entered and retained in the REDCap® database designed for this study. Demographic and viral
indicators collected included patient sex, race/ethnicity, age at admission, RSV results, viral coinfections,
and other viruses detected. Hospitalization and outcome metrics included PICU length of stay, hospital
length of stay, and death before discharge. Illness severity was described by Pediatric Logistic Organ
Dysfunction-2 (PELOD-2) [13] score at PICU admission, in addition to the Pediatric Sequential Organ Failure
Assessment (pSOFA) [14] score for up to five sequential days following PICU admission, maximum
respiratory support required, invasive mechanical ventilation days, and extracorporeal life support (ECLS)
days.

Data management and analysis were conducted using SAS® 9.4 (SAS Institute Inc., Cary, NC). A
predetermined level of α = 0.05 was chosen to evaluate statistical significance. Descriptive statistics were
provided for all demographics and key hospitalization and illness severity metrics: frequencies and
percentages for categorical data and mean ± SD for continuous measures. Comparisons were made between
the “pre-lockdown” and “post-lockdown” time periods. Fisher’s exact tests and t-tests were used for
comparison between the two periods for categorical and continuous outcomes, respectively. This study was
approved by the Institutional Board Review of the University of Florida (RB202101398), and consent was
waived.

Results
A total of 135 patients were admitted to the PICU with an ICD-10 code of bronchiolitis during the two time
frames analyzed. As seen in Table 1, almost twice as many patients with bronchiolitis were admitted during the PostLD phase compared with the usual winter spike of cases (PreLD). The PostLD group had a higher age of admission (11.2 ± 12.3 vs. 6.6 ± 7.5, p = 0.0075) but no difference in gender or race/ethnicity.

|                  | Overall | Pre-lockdown | Post-lockdown | P-value |
|------------------|---------|--------------|---------------|---------|
| No. of patients  | 135     | 48           | 87            |         |
| Sex: male, n (%) | 71 (52.6)| 29 (60.4)   | 42 (48.3)     | 0.1763  |
| Race/ethnicity   |         |              |               | 0.8202  |
| Black, n (%)     | 37 (27.4)| 13 (27.1)   | 24 (27.6)     |         |
| Hispanic, n (%)  | 17 (12.6)| 5 (10.4)    | 12 (13.8)     |         |
| White, n (%)     | 73 (54.1)| 28 (58.3)   | 45 (51.7)     |         |
| Unknown, n (%)   | 8 (5.9) | 2 (4.2)     | 6 (6.9)       |         |
| Age at admission (in months, mean ± SD) | 9.6 ± 11.0 | 6.6 ± 7.5 | 11.2 ± 12.3 | 0.0075 |

TABLE 1: Demographics of the 135 patients admitted with bronchiolitis to the PICU between the two time frames.

Presented: categorical data: n (%); continuous data: mean ± SD.

† Statistical testing: Fisher’s exact tests for categorical data; t-tests for continuous data.

PICU: pediatric intensive care unit.

Microbiologically, the PostLD group had a higher proportion of RSV infections (83.9% vs. 33.3%, p < 0.0001). The distribution of RSV and the presence or absence of other coinfections also differed between PreLD and PostLD periods; a greater proportion had other infections without RSV during the PreLD period, and in contrast, during the PostLD period, a greater proportion had RSV with or without coinfection. In the PostLD period, there were a higher number of bronchiolitis cases caused by coronavirus OC43 (10.3% vs. 0.0%, p = 0.0263) and parainfluenza types 1-4 (21.8% vs. 2.1%, p = 0.0017), yet a lower proportion of HMPV (0.0% vs. 8.3%, p = 0.0147) (Table 2).
|                                | Overall | Pre-lockdown | Post-lockdown | P-value |
|--------------------------------|---------|--------------|---------------|---------|
| No. of patients                | 135     | 48           | 87            |         |
| RSV+, n (%)                   | 89 (65.9) | 16 (33.3)   | 73 (83.9)     | <0.0001 |
| RSV and other viruses         |         |              |               |         |
| RSV+/other+, n (%)            | 38 (28.2) | 5 (10.4)    | 33 (37.9)     | <0.0001 |
| RSV+/other-, n (%)            | 51 (37.8) | 11 (22.9)   | 40 (46.0)     |         |
| RSV-/other+, n (%)            | 38 (28.2) | 24 (50.0)   | 14 (16.1)     |         |
| RSV-/other-, n (%)            | 8 (5.9)  | 8 (16.7)    | -             |         |
| Average count of other viruses detected, mean ± SD | 0.7 ± 0.8 | 0.6 ± 0.5 | 0.8 ± 0.9 | 0.1963 |
| **Other viruses**             |         |              |               |         |
| SARS-CoV-2/coronavirus, n (%) | 2 (1.5)  | -            | 2 (2.3)       | 0.5383  |
| Adenovirus, n (%)             | 12 (8.9) | 4 (4.2)      | 10 (11.5)     | 0.2119  |
| Coronavirus 229E, n (%)       | -       | -            | -             |         |
| Coronavirus HKU1, n (%)       | 1 (0.7)  | 1 (2.1)      | -             | 0.3556  |
| Coronavirus NL63, n (%)       | 2 (1.5)  | -            | 2 (2.3)       | 0.5383  |
| Coronavirus OC43, n (%)       | 9 (6.7)  | -            | 9 (10.3)      | 0.0263  |
| Human metapneumovirus, n (%)  | 4 (3.0)  | 4 (8.3)      | -             | 0.0147  |
| Influenza A, *all subtypes, n (%) | 1 (0.7) | 1 (2.1)     | -             | 0.3556  |
| Influenza B, n (%)            | 1 (0.7)  | 1 (2.1)      | -             | 0.3556  |
| Parainfluenza, *all subtypes, n (%) | 20 (14.8) | 1 (2.1) | 19 (21.8) | 0.0017  |
| Human rhinovirus/enterovirus, n (%) | 46 (34.1) | 20 (41.7)  | 26 (29.9)    | 0.1874  |

**TABLE 2: Viruses detected in the 135 patients admitted with bronchiolitis to the PICU between the two time frames.**

Presented: categorical data: n (%); continuous data: mean ± SD.

† Statistical testing: Fisher’s exact tests for categorical data; t-tests for continuous data.

PICU: pediatric intensive care unit; RSV: respiratory syncytial virus.

There were no differences between groups in hospital length of stay, PICU length of stay, death before discharge, mechanical ventilation days, ECLS days, or severity of illness scores based on the calculated PELOD-2 score and pSOFA (Table 3).
|                                | Overall | Pre-lockdown | Post-lockdown | P-value |
|--------------------------------|---------|--------------|---------------|---------|
| No. of patients                | 135     | 48           | 87            |         |
| Hospital length of stay (days, mean ± SD) | 5.8 ± 9.2 | 5.4 ± 5.4 | 6.1 ± 10.8 | 0.6194 |
| PICU length of stay (days, mean ± SD)     | 4.4 ± 7.8 | 3.9 ± 3.5 | 4.7 ± 9.3 | 0.4890 |
| Deaths, n (%)                  | 2 (1.5) | 1 (2.1)      | 1 (1.2)       | 1.000   |
| Maximum respiratory support (all) |         |              |               | 0.2777  |
| Low flow nasal cannula, n (%)   | 4 (3.0) | -            | 4 (4.6)       |         |
| High flow nasal cannula, n (%)  | 53 (39.3) | 17 (35.4) | 36 (41.4)     |         |
| CPAP, n (%)                    | 9 (6.7) | 5 (10.4)     | 4 (4.6)       |         |
| BiPAP, n (%)                   | 46 (34.1) | 20 (41.7) | 26 (29.9)     |         |
| Mechanical ventilation, n (%)  | 21 (15.6) | 6 (12.5)  | 15 (17.2)     |         |
| ECLS, n (%)                    | 2 (1.5) | -            | 2 (2.3)       |         |
| Intubated patients (M.V. or ECLS), n (%) | 23 (17.0) | 6 (12.5)  | 17 (19.5)     | 0.3470  |
| Mechanical ventilation only, n (%) | 21 (15.6) | 6 (12.5)  | 15 (17.2)     | 0.6211  |
| pSOFA score, PICU day of admission (mean ± SD) | 2.0 ± 2.2 | 2.0 ± 2.2 | 2.0 ± 2.2 | 0.8892  |
| PELOD-2 score, PICU day of admission (mean ± SD) | 1.3 ± 2.0 | ± 2.1     | 1.4 ± 1.9 | 0.2193  |
| Mechanical ventilation days (mean ± SD) | 6.7 ± 7.4 | 6.8 ± 5.1 | 6.7 ± 8.1 | 0.9719  |
| ECLS days (n = 2), * individually reported | 1.0 25.0 | -           | 1.0 25.0 | -       |
| Patients with 5 days of pSOFA scoring (n) | 31      | 12          | 19            |         |
| Among those with 5 days of pSOFA scoring: average scores (mean ± SD) | 3.6 ± 3.4 | 3.3 ± 3.6 | 3.8 ± 3.3 | 0.7097  |

TABLE 3: Respiratory support requirement, length of stay, and measured severity of illness of the 135 patients admitted with bronchiolitis to the PICU between the two time frames.

Presented: categorical data: n (%); continuous data: mean ± SD.
† Statistical testing: Fisher’s exact tests for categorical data; t-tests for continuous data.

PICU: pediatric intensive care unit; M.V: mechanical ventilation; ECLS: extracorporeal life support; CPAP: continuous positive airway pressure; BiPAP: bilevel positive airway pressure; pSOFA: Pediatric Sequential Organ Failure Assessment; PELOD-2: Pediatric Logistic Organ Dysfunction-2.

**Discussion**

After easing precautions intended to mitigate the COVID-19 spread, a significant decline in the expected seasonal respiratory illnesses followed by an upsurge of out-of-season bronchiolitis cases has been reported in many parts of the world [6,7,12,15]. There has been speculation that cases during this upsurge were more severe, requiring more interventions and longer hospital stays. However, to our knowledge, no studies have closely examined the data between pre-lockdown and post-lockdown bronchiolitis patterns. Our study attempts to answer this uncertainty.

We found a larger number of admissions and higher age of admission in the PostLD group; this singularity can be explained potentially by the diminished exposure to common viruses during the winter months of 2020-2021, leading to an “immunity debt” [16] with no development of protective immunity, and in consequence, leaving a greater proportion of the population susceptible for infections like bronchiolitis at older ages [7,17]. This theory may also explain the larger proportion of patients with RSV detection and other co-viral infections, as well as the different sets of viral pathogens detected between groups, with higher detections of coronavirus OC43 and parainfluenza viruses and lower detections of HMPV in the PostLD group. It is noteworthy that viral co-infections with RSV are not known to be associated with increased illness severity, with the exception of HMPV, which is associated with increased risk of PICU admission and longer hospital stay [18], a detection that was absent during the PostLD period.
There was no significant difference in the hospital length of stay, PICU length of stay, or respiratory support requirement. However, in our cohort of patients, there was a slightly larger proportion of patients that required mechanical ventilation (17.2% vs. 12.5%) and ECLS (2.3% vs. 0%) support in the PostLD group. Though not statistically significant, further investigation with a larger sample size may be warranted.

No difference was found between groups in disease severity measured by PELOD-2 and pSOFA scores. It is worth noting that the PELOD-2 score is validated to detect multiorgan dysfunction syndrome (MODS) severity in the PICU [13]. However, in most cases, bronchiolitis presents with only respiratory symptoms and not MODS. Moreover, 87% of our subjects did not have arterial blood samples, thus potentially underestimating the respiratory dysfunction portion of the PELOD-2 score. The pSOFA score can potentially solve the problem of the lack of arterial blood samples since the oxygen saturation/fraction of inspired oxygen (FiO2) ratio can be used to measure oxygenation; however, hypoxemia is not always present in bronchiolitis. Frequently, patients with bronchiolitis require increased respiratory support based on their work of breathing and/or ventilation issues, which are parameters not captured by the pSOFA score component. Moreover, the pSOFA score is validated to discriminate pediatric in-hospital mortality [14], which is not optimal, considering the low mortality in our cohorts. Other validated scales for assessment of pediatric disease severity like the Pediatric Multiple Organ Dysfunction Score (P-MODS) [19] and the Pediatric Risk of Mortality III (PRISM III) [20] would have had the same limitations. A bedside clinical scale assessment tool such as the modified Wood’s Clinical Asthma Score (M-WCAS) [21] might have provided a better evaluation of the severity of illness in bronchiolitis; nevertheless, the retrospective design of the study made it impossible. Hence, demonstrating a difference in disease severity in bronchiolitis with a validated scale in our study was not feasible.

Our study is subject to several limitations. It was conducted at a single center with a small sample size, using a retrospective design, which limited available metrics. The disease severity might not have been well assessed by the scores used as described earlier, and the time frames analyzed in both cohorts do not include the entire bronchiolitis season. Complementary multi-centered studies are needed to further evaluate this unique out-of-season upsurge of bronchiolitis observed during the COVID-19 pandemic. A deeper understanding of the clinical and epidemiologic changes could provide useful information to guide future management of critical bronchiolitis cases, both during and after the current pandemic.

Conclusions

In this single-center retrospective cohort study of critically ill patients admitted with bronchiolitis to the PICU, we found that in the PostLD out-of-season surge, there were an increased number of admissions with bronchiolitis to the PICU. Those subjects had a higher age of admission, more frequent RSV infections, more viral co-infections, a higher number of detections of coronavirus OC43 and all subtypes of parainfluenza, and less detections of HMPV. No difference in hospital length of stay or disease severity was demonstrated.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. University of Florida issued approval RB202101398. This study was approved by the Institutional Board Review of the University of Florida and consent was waived. Animal subjects: All authors have confirmed that this study did not involve animal subjects or tissue. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Zhu N, Zhang D, Wang W, et al.: A novel coronavirus from patients with pneumonia in China, 2019. N Engl J Med. 2020, 382:727-33. 10.1056/NEJMoa2001017
2. WHO coronavirus (COVID-19) dashboard. (2021). https://covid19.who.int/
3. Chumchiam SH, Ahmed QA, Gozzer E, Schlangenhaus P, Memish ZA: COVID-19 and community mitigation strategies in a pandemic. BMJ, 2020, 368:m1066. 10.1136/bmj.m1066
4. COVID-19 restrictions. Map of COVID-19 case trends and restrictions. (2021). https://www.usatoday.com/storytelling/coronavirus-reopening-america-map/
5. Florida Department of Health. Florida COVID-19 response. Plan for Florida’s recovery. (2022). https://floridaplanrecovery.org/
6. CDC. RSV national trends. (2021). Accessed: November 15, 2021: https://www.cdc.gov/surveillance/rsvnationaltrends.html
7. Olsen SJ, Winn AK, Budd AP, et al.: Changes in influenza and other respiratory virus activity during the COVID-19 pandemic — United States, 2020-2021. MMWR Morb Mortal Wkly Rep. 2021, 70:1015-9. 10.15585/mmwr.mm7029a1
8. Brueggemann AB, Jansen van Rensburg MJ, Shaw D, et al.: Changes in the incidence of invasive disease due
to Streptococcus pneumoniae, Haemophilus influenzae, and Neisseria meningitidis during the COVID-19 pandemic in 26 countries and territories in the Invasive Respiratory Infection Surveillance Initiative: a prospective analysis of surveillance data. Lancet Digit Health. 2021, 3:e360-70. 10.1016/S2589-7500(21)00077-7

9. Meissner HC: Viral bronchiolitis in children. N Engl J Med. 2016, 374:62-72. 10.1056/NEJMra1415346

10. Staat MA, Henriksen K, Elhefni H, Groothuis J, Makari D: Prevalence of respiratory syncytial virus-associated lower respiratory infection and apnea in infants presenting to the emergency department. Pediatr Infect Dis J. 2015, 32:911-4. 10.1097/INF.0b013e31828dd6f5e3

11. Graciano S, Erviti A, Caballero MT, et al.: Mortality due to respiratory syncytial virus. Burden and risk factors. Am J Respir Crit Care Med. 2017, 195:96-103. 10.1164/rccm.201603-0658oc

12. Florida Department of Health. Respiratory syncytial virus (RSV) surveillance activity summary. Week 48 (November 28-December 4, 2021), (2021). https://www.floridahealth.gov/diseases-and-conditions/respiratory-syncytial-virus/Documents/2021-w48-rsv-summary.pdf.

13. Leteurtre S, Duhamel A, Salleron J, Grandbastien B, Lacroix J, Leclerc F: PELOD-2: an update of the Pediatric Logistic Organ Dysfunction score. Crit Care Med. 2015, 41:1761-73. 10.1097/CCM.0b013e31828a2bbd

14. Matics TJ, Sanchez-Pinto LN: Adaptation and validation of a pediatric sequential organ failure assessment score and evaluation of the Sepsis-3 definitions in critically ill children. JAMA Pediatr. 2017, 171:e172352. 10.1001/jamapediatrics.2017.2352

15. Foley DA, Yeoh DK, Minney-Smith CA, et al.: The interseasonal resurgence of respiratory syncytial virus in Australian children following the reduction of coronavirus disease 2019-related public health measures. Clin Infect Dis. 2021, 73:e2829-30. 10.1093/cid/ciaa1906

16. Hatter L, Eathorne A, Hills T, Bruce P, Beasley R: Respiratory syncytial virus: paying the immunity debt with interest. Lancet Child Adolesc Health. 2021, 5:e44-5. 10.1016/S2352-4642(21)00333-5

17. Cohen R, Ashman M, Taha MK, et al.: Pediatric Infectious Disease Group (GPIP) position paper on the immune debt of the COVID-19 pandemic in childhood, how can we fill the immunity gap. Infect Dis Now. 2021, 51:418-23. 10.1016/j.idnow.2021.05.004

18. Li Y, Pillai P, Miyake F, Nair H: The role of viral co-infections in the severity of acute respiratory infections among children infected with respiratory syncytial virus (RSV): a systematic review and meta-analysis. J Glob Health. 2020, 10:010426. 10.7189/jogh.10.010426

19. Graciano AL, Balco JA, Rahm DS, Ahmad N, Giroir BP: The Pediatric Multiple Organ Dysfunction Score (P-MODS): development and validation of an objective scale to measure the severity of multiple organ dysfunction in critically ill children. Crit Care Med. 2005, 33:1484-91. 10.1097/01.ccm.0000170943.23633.47

20. Pollack MM, Patel KM, Ruttimann UE: PRISM III: an updated Pediatric Risk of Mortality score. Crit Care Med. 1996, 24:745-52. 10.1097/00003246-199605000-00004

21. Duarte-Dorado DM, Madero-Orostegui DS, Rodriguez-Martinez CE, Nino G: Validation of a scale to assess the severity of bronchiolitis in a population of hospitalized infants. J Asthma. 2013, 50:1056-61. 10.3109/02770903.2013.854304