Original article

COVID-19 and recurrent respiratory infections in children of Kazakhstan

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Abstract: The study goal was to investigate the COVID-19 clinical course in children with recurrent respiratory infections (RRI) in Nur-Sultan, Kazakhstan. Material and Methods — we conducted the retrospective analysis of 94 children with RRI, diagnosed with COVID-19, in Nur-Sultan, Kazakhstan. The study involved 53 males and 41 females. The inclusion criterion for the study was the frequency of RRI at least six times per year. In the course of our study, we split the patients among three groups and identified two phenotypes. These groups included children with RRI and atopic phenotype (Group 1), with D-deficiency phenotype (Group 2), and control group (Group 3) encompassing children with RRI lacking these phenotypes.

Results — The most common symptoms of 94 pediatric patients were dry cough (94.7%), fever (81.9%), along with a loss of appetite and fatigue (76.6%). Malaise was observed in 74.5% cases, rhinorrhea was noted in 71.2% of patients, sore throat was detected in 64.9 % of children, and dyspnea was established in 45.7% of cases. We observed no statistical differences in clinical manifestations of COVID-19 among three groups of children. However, duration of hospitalization period, of fever, and of the catarrhal period differed significantly among the groups (p<0.001).

Conclusion — In children with RRI and vitamin D deficiency, who were diagnosed with COVID-19, the course of the infection was unfavorable (which was confirmed by a longer hospital stay and catarrhal period), and a more severe intoxication syndrome was observed. In the group of children with atopic phenotype, a prolonged residual cough was detected.

Keywords: COVID-19, children, symptoms, recurrent respiratory infections.


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Introduction

At the end of December 2019, 41 cases of pneumonia of unknown origin had been registered in Wuhan city of Hubei province, China, which have led to the development of SARS. The investigation had uncovered a new type of coronavirus, SARS-CoV-2, which have caused the pandemic. Extremely crowded Huanan seafood market have been considered the nidus of this coronavirus infection [1, 2]. The Research Group of International Committee on Taxonomy of Viruses had specified this new coronavirus as SARS-CoV-2, and World Health Organization (WHO) has named this coronavirus infection COVID-19 [3, 4]. Due to rapid spread of SARS-CoV-2 infection worldwide, WHO has announced the pandemic of COVID-19 on March 11, 2020 [5].

The Republic of Kazakhstan is located in Central Asia and has borders with Russia, China, Kyrgyzstan, Uzbekistan and Turkmenistan. The first case of COVID-19 was registered here on March 13, 2020. On March 22, 2020, the Government adopted strict quarantine measures. Consequently, big cities of Kazakhstan have been closed. According to official data, at the time of the manuscript preparation, 187,970 confirmed COVID-19 cases, 168,768 treated cases and 2,556 deaths have been registered in Kazakhstan from the very beginning of the pandemic [6].

The first case of COVID-19 infection among children was described in January, 2020, in Shenzhen, China [7]. Epidemiological data from numerous publications have indicated that SARS-CoV-2 infection course in children was less severe than in adults, unless there were serious comorbidities [8]. Furthermore, according to the results of another Chinese study of 2,143 children, the COVID-19 infection course was mild in most patients, while severe cases were observed just in 5.9% of infected children. As for adult patients, the severe course of COVID-19 infection was registered in 18.5% [9].

Fever and cough were the most frequently observed clinical symptoms in children [4]. Another study has reported that cough was present in 48.5% of patients, pharyngeal erythema was observed in 46.2%, fever was detected in 41.5%, tachypnea was noted in 28.7%, whereas diarrhea and nasal congestion were established in just 10% of those [10]. Regarding adult patients, most common clinical signs were fever (98.6%), fatigue (69.6%), dry cough (59.4%) [11], and shortness of breath (31%) [12].

When diagnosing SARS-CoV-2 infection in children, taking a thorough medical history, conducting accurate physical
examination, PCR and other laboratory tests are recommended. In blood test, increase or decrease of leucocytes and lymphocytes may be observed, while in biochemical test, increased levels of C-reactive protein, procalcitonin, and liver enzymes may be registered. The X-ray findings may indicate the signs of unilateral or bilateral infiltrative pneumonia. On CT scan of lungs, such sign as ground-glass opacities may be detected [13, 14].

Study goal: to investigate the clinical course of coronavirus infection in children with recurrent respiratory infections.

Material and Methods

Study design and participants

The retrospective analysis of 94 children at the age of 3-17 years old with recurrent respiratory infections (RRI), who were diagnosed with COVID-19 and had a positive family history of SARS-CoV-2 infection, was conducted. In such children, real-time reverse transcription polymerase chain reaction (RT-PCR) was conducted for COVID-19 confirmation. All children were admitted to the Division of Pulmonary Care Medicine at the City of Nur-Sultan Children’s Hospital from June 1 to August 28, 2020. The study involved 53 males and 41 females.

Inclusion and exclusion criteria

The inclusion criteria incorporated the frequency of RRI at least six times per year; absence of comorbidities (endocrine disorders, congenital malformations, oncological diseases); under 18 years old; positive PCR test for COVID-19; parental consent, and informed consent of patients. The exclusion criteria encompassed chronic foci of ENT pathology (e.g., chronic sinusitis, pharyngitis) and primary immunodeficiencies.

Epidemiological characteristics and procedures

Children were more often infected from their family members. On the first day of hospitalization, all children underwent PCR test for COVID-19. To identify SARS-CoV-2 RNA, the sample was scraped off the posterior wall of the laryngeal cavity, using a cotton swab, which was then placed in the solution for virus preservation. The sample was transported at a temperature of 2-8 °C to the specialized facility, following the rules listed in the Sanitary and Epidemiological Requirements for Laboratories Using Potentially Hazardous Chemical and Biological Substances of November 14, 2017, No. 15990. The analysis for the viral RNA detection was carried out on an automated analyzer Tianlong (China) within 24 hours.

Examination of patients

At the hospital, a complete clinical examination, along with thorough collection of clinical and epidemiological (the quarantine regimen was introduced in Kazakhstan) anamnesis was carried out. Furthermore, first-level examination including complete blood count, urinalysis and biochemical test were conducted. The latter included identifying the content of C-reactive protein, creatinine, glucose, chloride, blood urea nitrogen, bilirubin, aspartate aminotransferase, alanine aminotransferase, total protein, electrolytes level (total calcium, potassium, sodium). Besides, blood coagulation profile and additional laboratory tests (such as lactate dehydrogenase, procalcitonin, vitamin D, and IgE) were performed, if prescribed by physicians. Some patients underwent chest X-ray, computed tomography, abdominal ultrasound, kidney ultrasound, ECHO-cardiography, and ECG. Laboratory test was performed at least once in five days; some children were tested more frequently, if necessary (based on the course of the disease). Discharge criteria for pediatric patients were improvement of clinical and laboratory data, and negative PCR test for COVID-19.

Statistical analysis

Data processing and compilation of tables was performed using Microsoft® Excel 2010 for Windows. For statistical data processing, the statistical software package Jamovi (version 1.2.27.0, www.jamovi.org) was used. The tables present clinical manifestations, gender, age, values of basic laboratory parameters and their percentages. Normally distributed variables were presented using means ± standard deviations, whereas for non-normally distributed variables, medians were used. For quantitative indicators, the one-way ANOVA test (Kruskal-Wallis test) was conducted regardless of the normal distribution. The chi-square test was employed for qualitative traits. Parameter values with p<0.05 were considered statistically significant.

Results

Demographics and study groups

Our retrospective study involved 94 patients (age range: 3-17 years old, average age and SD: 8.4±3.4). The average duration of a hospital stay was 8.4±2.8 days.

Among all observed children, 31 (33.0%) had history of allergy (total IgE >100 U/ml, a history of food allergy, hives, or seasonal hay fever). Deficiency of the vitamin D was detected in 28 (29.8%) patients (25(OH)D < 20 ng/L). In 35 (37.2%) patients (control group), those parameters were within normal ranges. That is why we split patients among three groups and compared the course of COVID-19 among the groups: children with RRI and atopic phenotype (Group 1), with D-deficiency phenotype (Group 2) and control group (Group 3) encompassing children with RRI lacking these phenotypes (Figure 1).

Figure 1. The research process.
Table 1. Clinical manifestations of COVID-19 in children with RRI at the time of hospital admission in Nur-Sultan, Kazakhstan

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>COVID-19</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients</td>
<td>Group 1</td>
<td>n=31</td>
</tr>
<tr>
<td>Gender:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>18/31 (58.1%)</td>
<td>19/28 (67.9%)</td>
</tr>
<tr>
<td>Female</td>
<td>13/31 (41.9%)</td>
<td>9/28 (32.1%)</td>
</tr>
<tr>
<td>Age (years) Mean (SD)</td>
<td>6.4 ± 3.8</td>
<td>6.8 ± 3.3</td>
</tr>
<tr>
<td>Clinical symptoms:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dry cough</td>
<td>29 (93.5%)</td>
<td>27 (96.4%)</td>
</tr>
<tr>
<td>Fever</td>
<td>24 (77.4%)</td>
<td>27 (96.4%)</td>
</tr>
<tr>
<td>Loss of appetite</td>
<td>25 (80.6%)</td>
<td>24 (85.7%)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>25 (80.6%)</td>
<td>24 (85.7%)</td>
</tr>
<tr>
<td>Malaise</td>
<td>21 (67.7%)</td>
<td>24 (85.7%)</td>
</tr>
<tr>
<td>Rhinorrhea / Nasal congestion</td>
<td>21 (67.7%)</td>
<td>20 (71.4%)</td>
</tr>
<tr>
<td>Sore throat</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dyspnea</td>
<td>20 (64.5%)</td>
<td>19 (67.9%)</td>
</tr>
<tr>
<td>Vomiting / Nausea</td>
<td>14 (45.1%)</td>
<td>16 (57.1%)</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>3 (9.7%)</td>
<td>1 (4.3%)</td>
</tr>
<tr>
<td>* For these averages, medians with lower and upper quartile values were used, since the distributions of these parameters were not normal: Me (LQ, UQ).</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td></td>
<td></td>
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<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duration (days) of:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospital stay</td>
<td>7 ± 0.9</td>
<td>12.3 ± 1.5</td>
</tr>
<tr>
<td>Fever</td>
<td>4.8 ± 0.7</td>
<td>5.7 ± 0.7</td>
</tr>
<tr>
<td>Catarhal period</td>
<td>5.7 ± 0.9</td>
<td>7.3 ± 1.3</td>
</tr>
</tbody>
</table>

Table 2. Results of laboratory and instrumental tests in children with COVID-19 at the time of hospital admission

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>COVID-19</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Laboratory examination:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lymphocytes (%)</td>
<td>19 (20.2%)</td>
<td>24 (25.5%)</td>
</tr>
<tr>
<td>Decrease</td>
<td>21.1 ± 9.9</td>
<td>24.5 ± 10.9</td>
</tr>
<tr>
<td>Erythrocyte sedimentation rate (mm/h) Increase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C-reactive protein level (mg/L) Increase</td>
<td>18 (19.1%)</td>
<td>19 (20.2%)</td>
</tr>
<tr>
<td>Vitamin 25(OH)D (ng/mL) Decrease</td>
<td>16 (17.0%)</td>
<td>18 (19.1%)</td>
</tr>
<tr>
<td>Decrease</td>
<td>31.4 ± 2.7</td>
<td>20.2 ± 1.5</td>
</tr>
<tr>
<td>IgE total (kU/L) Increase</td>
<td>10 (10.6%)</td>
<td>28 (29.8%)</td>
</tr>
<tr>
<td>Increase</td>
<td>203.2 ± 88.5</td>
<td>12.9 ± 12.4</td>
</tr>
<tr>
<td>Hemoglobin (g/L) Decrease</td>
<td>31 (33.0%)</td>
<td>0 (0.0%)</td>
</tr>
<tr>
<td>Increase</td>
<td>119.5 ± 13.7</td>
<td>119.3 ± 12.6</td>
</tr>
<tr>
<td>White blood cells (&lt;10³) Decrease</td>
<td>8 (8.5%)</td>
<td>9 ± 4.2</td>
</tr>
<tr>
<td>Increase</td>
<td>26 (21.3%)</td>
<td>26 (21.3%)</td>
</tr>
<tr>
<td>Neutrophils (%)</td>
<td>55.9 ± 12.9</td>
<td>59.7 ± 14</td>
</tr>
<tr>
<td>Increase</td>
<td>4 (4.3%)</td>
<td>4 (4.3%)</td>
</tr>
<tr>
<td>Lactate dehydrogenase (U/L) Increase</td>
<td>270 (208, 352)*</td>
<td>313 (255, 553)*</td>
</tr>
<tr>
<td>Procalcitonin (ng/mL) Increase</td>
<td>1 (1.1%)</td>
<td>3 (3.2%)</td>
</tr>
<tr>
<td>Alanine aminotransferase (U/L) Increase</td>
<td>0.3 ± 0.3</td>
<td>0.3 ± 0.3</td>
</tr>
<tr>
<td>X-Ray findings</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bilateral patchy shadowing</td>
<td>11 (35.5%)</td>
<td>25 (89.3%)</td>
</tr>
<tr>
<td>Local patchy shadowing</td>
<td>9 (29.0%)</td>
<td>3 (10.7%)</td>
</tr>
<tr>
<td>Ultrasound examination</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diffuse and reactive changes in the liver</td>
<td>1 (1.1%)</td>
<td>2 (2.1%)</td>
</tr>
</tbody>
</table>

The table includes solely changed parameters as identified by laboratory analyses. The normal range and standard values sensu Pediatric Laboratory Values [15].
Laboratory analyses

According to the laboratory results summarized in Table 2, no statistically significant differences were found between the study groups on any laboratory parameters, except for lymphopenia (Group 1 = 20.2%, Group 2 = 25.5%, and Group 3 = 19.1%; p<0.001). Increased IgE was found only in Group 1, and in other groups its content was within the normal reference range (p<0.001) (Table 2).

Diagnostic radiology

On the chest X-ray, community-acquired bilateral pneumonia was detected more often in Group 2 (35.5%), compared with Group 1 (20.2%) and Group 3 (19.1%) (p<0.001). Ultrasound examination of the abdominal cavity and kidneys revealed no differences. On the echocardiogram and ECG, the contractile function of heart ventricles was satisfactory; no changes were detected (Table 2).

Discussion

In Kazakhstan, from March 16 until May 11, 2020, a state of emergency was introduced because of the threat of coronavirus infection spread, with a gradual weakening of quarantine measures, depending on epidemiological situation in the country. Various measures were taken to reduce the spread of coronavirus infection, such as social distancing, school and kindergarten closures, closures of potentially crowded businesses, teleworking, and restrictions on entering and exiting the country [16].

Children were more frequently infected from their family members with confirmed COVID-19 infection. SARS-CoV-2 has been causing high prevalence of pneumonia in infected individuals. According to the systematic review of 46 clinical cases of COVID-19 in children, the main manifestations of this infection was fever (64%), cough (35%), and rhinorrhea (16%). An asymptomatic course (15%) was characteristic for 15% of children. On X-ray, ground-glass opacities were detected (54%). Laboratory analyses implied lymphopenia (33%), augmented D-dimer (52%) and C-reactive protein (40%) [17].

In most cases, children had asymptomatic or mild course of COVID-19 infection. This may be explained by the lower number of angiotensin-converting enzyme type 2 receptors in children (ACE2 is abundantly expressed in the cells of the respiratory tract), compared with adults. SARS-CoV enters the host cells through ACE2 receptors [18, 19].

Clinical manifestations

The main clinical signs of COVID-19 in children with RRI in three groups were dry cough (n=89;94.7%), fever (n=77;81.9%), loss of appetite (n=72;76.6%), fatigue (n=72;76.6%), malaise (n=70;74.5%), rhinorrhea and/or nasal congestion (n=67;71.2%), and sore throat (n=61;64.9%). No statistical significance was found for clinical manifestations among the groups. (Table 1).

Duration of intoxication period and hospitalization

The duration of a hospital stay for children in Group 2 was longer (12.3±1.5) versus Group 1 (7.0±0.9) and Group 3 (6.3±0.7). The number of fever days for children varied among groups (Group 2 = 5.7±0.7, Group 1 = 4.8±0.7, and Group 3 = 4.2±0.7 (p<0.001). Children with vitamin D deficiency had prolonged catarrhal period (7.3±1.3) days. In children with allergy, the catarrhal period lasted 5.7 ± 0.9 days, while in control group, it was 5.1±0.8 days (Figure 2). In the group of children with atopic phenotype, a prolonged residual cough was observed.

The prolonged duration of clinical manifestations and hospital stay was most likely due to the fact that, in children with vitamin D deficiency, bilateral pneumonia was detected more often via X-ray examination (in 89.3% of cases) than in children with atopy (in 35.5%), or control group (in 14.3%). Obstructive bronchitis was identified in 11 patients (11.7%), and pneumonia was observed in 64.5% patients with atopic phenotype and RRI: right-sided focal lower lobe pneumonia (in 9 children; 29.0%), and bilateral lower lobe pneumonia (n=11; 35.5%). As seen in Figure 3, 2 patients had asymptomatic course of COVID-19 in the control group (5.7%); COVID-19 infection proceeded in the form of acute respiratory infection in 16 patients (45.7%); and in 48.6% patients, the infection progressed in the form of community-acquired pneumonia: right-sided focal lower lobe pneumonia (n=12; 34.3%), and bilateral lower lobe pneumonia (n=5;14.3%).

Laboratory analyses

According to the laboratory results summarized in Table 2, no statistically significant differences were found between the study groups on any laboratory parameters, except for lymphopenia (Group 1 = 20.2%, Group 2 = 25.5%, and Group 3 = 19.1%; p<0.001). Increased IgE was found only in Group 1, and in other groups its content was within the normal reference range (p<0.001) (Table 2).

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identified: atopic and D
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compared with other groups
longer recovery stage and prolonged presence of pneumonia,
during the follow
experience regarding COVID
symptoms of coronavirus
The combination of mixed infections may lead to more severe
virus-induced immune dysfunction, which could trigger further
recurrences of respiratory infections [20].

Currently, there are some published studies regarding the
severity of COVID-19 in children with vitamin D deficiency [21, 22],
as there is a relationship between vitamin D and the immune
response in conditions of various infections [23, 24]. Furthermore,
the specific data exist regarding the severity of COVID-19 course in
patients with allergy. It has been noted that such patients had
milder course of coronavirus infection, and therefore, their lungs
were affected to a lesser extent. One of the possible causes of this
finding is related to less pronounced damage of T cells in such
patients [25]. Also, it is thought that allergy is not a risk factor for
severity of COVID-19 in children [26]. Another explanation of this
pattern is that children with atopic phenotype (for example, in the
form of bronchial asthma) are less likely to suffer from severe
course of COVID-19, since inhaled steroids are used in its therapy
[27].

Our monocenter retrospective study investigated the clinical
and laboratory manifestations of COVID-19 in frequently ill
children. We discovered no significant differences among the
symptoms of coronavirus infection in our study versus global
experience regarding COVID-19 in children. It should be noted that
during the follow-up, children with vitamin D deficiency had a
longer recovery stage and prolonged presence of pneumonia,
compared with other groups, while there was a longer residual
cough after recovery in the group of children with atopy.

Conclusion

During the observation period for patients with RRI, there
were no deaths in any group, and the following phenotypes were
identified: atopic and D-deficient. An unfavorable course of COVID-
19 in the form of longer hospital stay and catarrhal period, as well
as higher severity of intoxication, was revealed in a D-deficient
group of patients infected with coronavirus. However, in the group
of children with atopic phenotype, a prolonged period of residual
cough was detected. It is possible that these etiological factors
(vitamin D deficiency and allergy symptoms) could affect the
severity and duration of coronavirus infection in children with RRI.
Further, more thorough, study is needed to confirm this
hypothesis.

Conflict of interest
The authors declare no conflicts of interest.

Ethical approval
The study was approved by the Ethics Committee: Protocol No.
2020.3.7, which was signed by the Local Bioethics Committee at Astana
Medical University. The authors complied with the ethical principles of the
Declaration of Helsinki, World Medical Association. The informed consent
of the patients was taken from all study participants.

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