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.
Air pollution might affect the clinical course of COVID-19 in pediatric patients

Piotr Rzymski a,b,*, Barbara Poniedziałek a, Joanna Rosińska a, Przemysław Ciechanowski c, Michał Peregrym c, Maria Pokorska-Śpiewak d, Ewa Talarek d, Izabela Zaleska a, Paulina Frańczak-Chmura l, Małgorzata Pilarczyk g, Magdalena Figlerowicz h, Izabela Kucharek i, Robert Flišiak j

a Department of Environmental Medicine, Poznan University of Medical Sciences, 60-806 Poznań, Poland
b Integrated Science Association (ISA), Universal Scientific Education and Research Network (USERN), 60-806 Poznań, Poland
c Department of Paediatrics and Infectious Diseases, Medical University of Warsaw, Regional Hospital of Infectious Diseases in Warsaw, 01-201 Warsaw, Poland
d Department of Paediatrics and Infectious Diseases, Wroclaw Medical University, 50-368 Wrocław, Poland
e Department of Children’s Infections Diseases, Medical University of Warsaw, Regional Hospital of Infectious Diseases in Warsaw, 01-201 Warsaw, Poland
f Department of Paediatrics and Infectious Diseases, Provincial Jan Baj Opy Hospital in Lublin, 20-089 Lublin, Poland
g Department of Infectious Diseases and Hepatology, Faculty of Medicine, Collegium Medicum, Nicolaus Copernicus University, 85-030 Bydgoszcz, Poland
h Department of Infectious Diseases and Child Neurology, Poznan University of Medical Sciences, 60-572 Poznan, Poland
i 2nd Department of Paediatrics, Centre of Postgraduate Medical Education, Department of Paediatrics and Neonatology with Allergology Center, Central Clinical Hospital of the Ministry of the Interior, 02-507 Warsaw, Poland
j Department of Infectious Diseases and Hepatology, Medical University of Białystok, 15-089 Białystok, Poland

A R T I C L E   I N F O

Keywords:
- Pandemic
- Particulate matter
- Benzo(a)pyrene
- Pediatric
- Inflammation
- Epidemiology

A B S T R A C T

Air pollution, to which children are more susceptible than adults, can promote airway inflammation, potentially exaggerating the effects of respiratory viral infection. This study examined the association between the clinical manifestation of COVID-19 in unvaccinated pediatric patients hospitalized in Poland (n = 766) and levels of particulate matter 2.5 (PM2.5) and benzo(a)pyrene (B(a)P) within a week before hospitalization. Children aged < 12 years exposed to mean and max 24 h B(a)P levels > 1 ng/m3 revealed higher odds of cough, dyspnea, fever, and increased concentrations of inflammatory markers (C-reactive protein, interleukin-6, procalcitonin, white blood cell count). In older patients (13–17 years), elevated mean 24 h B(a)P levels increased odds of dyspnea, fever, and diarrhea, and higher concentrations of C-reactive protein and procalcitonin. Exposure to max 24 h PM2.5 levels > 20 µg/m3 was associated with higher odds of cough, increased concentrations of C-reactive protein (group <12 years), and increased procalcitonin concentration (groups ≤12 years and 13–17 years). In both age groups, length of stay was extended in patients exposed to elevated levels of max 24 h PM2.5, mean and max 24 h B(a)P. This study suggests that worse air quality, particularly reflected in increased B(a)P levels, might affect the clinical course of COVID-19 in pediatric patients and adds to the disease burden during a pandemic.

1. Introduction

The COVID-19 pandemic has posed a tremendous challenge to the healthcare system, overwhelming its different sectors and branches. The hospitalized patients mainly included elderly individuals suffering from obesity and various comorbidities. Although children are susceptible to SARS-CoV-2 infection and contribute to its spread in the population, most pediatric cases are asymptomatic or mild with fever and cough but no hospitalization requirement (Jackson et al., 2022). The risk of severe disease and death in this group is low (Mania et al., 2022; Pokorska-Śpiewak et al., 2021). Despite it, some children and adolescents require hospitalization and may develop pediatric inflammatory...
multi-system syndrome due to SARS-CoV-2 infection (Ward et al., 2022).

Apart from socioeconomic and ethnicity disparities, comorbidities, such as asthma, diabetes, cardiovascular diseases, neurologic disorders, obesity, were identified as the leading risk factor of severe COVID-19 in children. In particular, medically complex individuals are more likely to be hospitalized due to SARS-CoV-2 infection (Graff et al., 2021; Woodruff et al., 2021). At the same time, it is known that air pollutants, such as particulate matter (PM), especially its PM$_{2.5}$ fraction (with aerodynamic diameter $\leq 2.5$ $\mu$m), and benzo(a)pyrene, can act as an irritant and promote inflammation in the respiratory tract (Rzymski et al., 2022). Compared to adults, children are at higher risk of the adverse effects of these pollutants due to smaller airway caliber with lower pulmonary capacity but a higher respiratory rate and more susceptible epithelium (Kurt, Zhang, and Pinkerton, 2016; Babin et al., 2007; I.-J. Wang et al., 2016).

This study aimed to analyze whether air pollution, PM$_{2.5}$ and B(a)P, is related to the severity of COVID-19 among patients aged $<18$ years ($n = 766$) who required hospitalization in selected Polish medical units over 17 months of the pandemic. Specifically, the relations between increased levels of air pollutants and the manifestation of various COVID-19 symptoms, the concentration of inflammatory markers at admission, oxygen saturation, and length of hospital stay were examined. Both pollutants considered in the present study are particularly concerning in regions such as Central Europe, where the combustion of wood and coal continues to play a role in domestic heating. This results in elevated emissions of PM and PM-bound polycyclic hydrocarbons, especially between late autumn and early spring (Aniol et al., 2021; Nazar and Niedoszytko, 2022). Moreover, contrary to many world regions, the COVID-19 lockdown in Poland was not associated with a temporary decline in PM emissions, while in selected locations, they were even increased (Rogulski and Badyda, 2021).

2. Material and methods

2.1. Patients data

All clinical data of hospitalized patients aged $<18$ years were retrieved from the SARSTer database, managed by the Polish Association of Epidemiologists and Infectiologists. A total of 766 records of children, and adolescents, hospitalized with COVID-19 in different healthcare units in Poland between March 2020 and July 2021 (17 months) were collected. All patients were unvaccinated against COVID-19. All of these individuals were diagnosed and treated according to the Polish recommendations for the management of COVID-19 (Flisiak, Horban et al., 2020a, 2021; Flisiak, Parczewski et al., 2020, 2020b; Marczyńska et al., 2020; Okarska-Napierała et al., 2021).

The clinical data collected for this study included:

(i) frequency of early COVID-19 symptoms cough, dyspnea, fatigue, fever, headache, fatigue, anosmia, nausea, diarrhea, and vomiting;
(ii) levels of inflammatory markers at admission: C-reactive protein (CRP), interleukin-6 procollagen (PCT), interleukin-6, and white blood cell (WBC) – the following levels were defined as hyperinflammation: CRP $>10$ $\mu$g/L, IL-6 $>37$ pg/mL, PCT $>0.1$ ng/mL and WBC $>11 \times 10^9$/µL (Yasuhara et al., 2020; Shafiek et al., 2021);
(iii) the parameters of the clinical course of the disease: oxygen saturation (SpO$_2$) at admission, the need for oxygen therapy and mechanical ventilation, length of hospital stay, and outcome (survival or death).

As the immune system undergoes changes in children and adolescents (Bartlett et al., 1998), two age groups were considered: $\leq 12$ years and 13–17 years. Moreover, the previous epidemiological studies have shown that individuals aged 13–17 years are far more likely to be infected with SARS-CoV-2 and experience symptomatic infection than $\leq 12$ years (Jang et al., 2022; Ryan et al., 2022). Moreover, in the present study, these two age groups had a similar sex distribution and did not differ in frequencies of comorbidities.

The study had a retrospective, non-interventional nature, and according to the Polish law (Dz.U. 2020 poz. 1291) it did not require written consent from patients and bioethical approval. All personal data were protected according to the European Union General Data Protection Regulation.

2.2. Air pollution data

The following air pollution parameters were included in the analysis: PM$_{2.5}$ and B(a)P. The data was collected from the Polish Chief Inspectorate Of Environmental Protection database, which is legally responsible for the air pollution monitoring in Poland. The mean 24 h and max 24 h levels in the inhabited area during a week preceding the hospitalization were retrieved for each patient. If more than one air quality monitoring station was available in a particular area, the data were collected from all of them and averaged. The period of the week before hospitalization was chosen because it most likely represented a time of transition of SARS-CoV-2 infection from incubation phase to symptomatic one, a time during which the innate immune response constitutes an essential line of the antiviral defense (Rzymski et al., 2022; Kasuga et al., 2021; Diamond and Kanneganti, 2022). Its disruption can lead to cellular overactivation and hyper-inflammation, subsequently favoring a more severe clinical course of COVID-19 and worsening the prognosis (Peyneau et al., 2022; Galani and Andreakos, 2021; Blot et al., 2020; Janssen et al., 2021). In turn, air pollutants such as PM$_{2.5}$ and B(a)P have been evidenced to promote inflammation and adversely affect innate immune response (Glencross et al., 2020). As shown in experimental and epidemiological studies, the pro-inflammatory effect may also occur in the respiratory system following the short-term exposure to increased levels of these pollutants (Fan et al., 2021; Y. Li et al., 2021; Coker et al., 2021; Chen et al., 2017).

The following air quality limits were considered in this study: mean 24 h and max 24 h PM$_{2.5}$ $>20$ $\mu$g/m$^3$, and mean 24 h and max 24 h B(a)P $>1.0$ ng/m$^3$ (Directive 2004/107/EC; Directive 2008/50/EC).

2.3. Statistical analysis

The data was analyzed with Statistica v. 13.1 (StatSoft, USA) separately for patients aged $\leq 12$ and 13–17 years. For continuous variables, differences were tested with a Student’s t-test. To evaluate associations between exceedance of air pollution limits and symptomatology, biochemical parameters, and clinical course, the classical odds ratios (ORs) with a 95% confidence interval (CI) were calculated according to the formulas given by Bland and Altman using MedCalc (MedCalc, Ostend, Belgium). A p-value $<0.05$ was considered statistically significant.

3. Results

3.1. Group characteristics

The studied group constituted 766 pediatric patients aged 0–17 years hospitalized between March 2020 and July 2021 due to COVID-19. None of the individuals was vaccinated. Their demographic characteristics and general clinical data are presented in Table 1. The number of patients requiring oxygen therapy or mechanical ventilation was very low, similar to fatal cases.

3.2. Air pollution and early symptoms of COVID-19

Association between exposure to air pollution above quality limits
and the presence of selected early symptoms of COVID-19 were observed (Fig. 1). The odds of cough, fever, and dyspnea were higher in children aged ≤ 12 years exposed to high mean and max levels of 24 h B(a)P. Odds of cough were also increased in this group when exposed to elevated max 24 h B(a)P levels (Fig. 1A). Exposure to max 24 h B(a)P level exceeding the limit increased odds of fever, dyspnea, and diarrhea in patients aged 13–17 years (Fig. 1B). Exceedances of air pollution parameters did not alter the odds of other symptoms considered in the study.

### 3.3. Air pollution and inflammatory markers at admission

The levels of inflammatory markers in the studied groups are summarized in Table 2. In the group aged ≤ 12 and 13–17 years, significantly elevated IL-6 concentrations were noted in individuals exposed to max 24 h PM$_{2.5}$ (by 3.8-fold and 8.6-fold, respectively), mean 24 h B(a)P (by 5.7-fold and 8.8-fold, respectively) and max 24 h B(a)P (by 5.4-fold and 4.9-fold, respectively) at levels exceeding the limits exhibited significantly elevated IL-6 concentrations at admission. PCT concentrations were increased only in children aged ≤ 12 years exposed to high levels of mean 24 h B(a)P (by 2.0-fold) and max 24 h B(a)P (by 4.0-fold). CRP and WBC concentrations were significantly elevated in children aged ≤ 12 years exposed to increased levels of mean 24 h B(a)P (by 1.8-fold and 1.1-fold, respectively) and max 24 h B(a)P (by 2.0-fold and 1.1-fold, respectively). In addition, both parameters were higher in individuals aged 13–17 exposed to mean 24 h B(a)P exceeding the limit (by 2.0-fold and 1.1-fold, respectively).

As further demonstrated, patients aged ≤ 12 years exposed to max 24 h PM$_{2.5}$ levels exceeding the quality limit revealed increased odds of CRP > 10 mg/L at admission. In turn, exposure to elevated mean and max B(a)P concentrations increased odds for CRP > 10 mg/L, IL-6 > 37 pg/mL, PCT > 0.1 ng/mL and WBC > 11 × 10$^3$/μL (Fig. 2A). In case of patients aged 13–17 years, exceedances of max 24 PM$_{2.5}$ mean and max B(a)P levels were associated with higher odds of PCT > 0.1 ng/mL. Increased odds of CRP > 10 mg/L were also found for patients exposed to max 24 h level above the quality limit (Fig. 2B).

### 3.4. Air pollution and the clinical course of COVID-19

Odds for SpO$_2$ < 95% at admission were increased only in patients aged ≤ 12 years exposed to elevated mean 24 h B(a)P level (OR: 3.2, 95% CI: 1.2–10.6, p < 0.05). Children aged ≤ 12 years exposed to max 24 h PM$_{2.5}$, mean 24 h B(a)P and max 24 h B(a)P levels exceeding the limits required an extended hospital stay, on average by 40.5%, 118.5% and 120.0%, respectively (Fig. 3A). Similarly, in patients aged 13–17 years, the length of hospital stay was longer by 92.5%, 92.5% and 100.0%, respectively (Fig. 3B). Association between air pollution and death or the need for oxygen therapy and mechanical ventilation was not evaluated due to a low number of pediatric patients with these outcomes (Table 1).

### 4. Discussion

This study demonstrates the direct links between outdoor air pollution and clinical manifestation of COVID-19 in pediatric patients, indicating that exposure to pollutants such as PM$_{2.5}$ and B(a)P can increase the odds of selected symptoms manifestation, increased inflammatory response, and extended hospital stay. Although the links between the severity of COVID-19 and air pollution have already been explored, contrary to our study, previous research focused on adult or general populations, not specifically on pediatric groups (Rzymski et al., 2022; Martinez-Boubeta and Simeonidis, 2022; Bowe et al., 2021; Kogevinas et al., 2014). The present research highlights that this can...
The inflammatory markers (mean±SD) in patients aged < 18 years (n = 766) exposed to air pollutants exceeding/not exceeding the limits during a week before hospitalization.

| Parameter | Age group | mean 24 h PM$_{2.5}$ | max 24 h PM$_{2.5}$ | mean 24 h B(a)P | max 24 h B(a)P |
|-----------|-----------|----------------------|---------------------|----------------|----------------|
|           | ≤ 12      | > 20 µg/m$^3$ | ≤ 12      | > 20 µg/m$^3$ | > 1 ng/m$^3$ | ≤ 1 ng/m$^3$ | > 1 ng/m$^3$ | p/t |
| CRP [mg/L]|           | ns                  | 12.5       | 14.5       | ns 9.6       | 17.6       | ** 8.6       | 17.1 |
|           | 13–17     | ns                  | 12.5       | 14.5       | ns 9.6       | 17.6       | ** 8.6       | 17.1 |
| IL-6 [pg/mL]| ≤ 12      | ns                  | 27.2       | 32.7       | ns 20.5      | 36.6       | ± 19.1       | ± 34.9 |
|           | 13–17     | ns                  | 27.2       | 32.7       | ns 20.5      | 36.6       | ± 19.1       | ± 34.9 |
| PCT [ng/mL]| ≤ 12      | ns                  | 0.2        | 0.4        | ns 0.2       | 0.4        | ± 0.1        | ± 0.1 |
|           | 13–17     | ns                  | 0.2        | 0.4        | ns 0.2       | 0.4        | ± 0.1        | ± 0.1 |
| WBC [× 10$^3$/µL]| ≤ 12   | ns                  | 0.1        | 0.3        | ns 0.1       | 0.3        | ± 0.1        | ± 0.1 |
|           | 13–17     | ns                  | 0.1        | 0.3        | ns 0.1       | 0.3        | ± 0.1        | ± 0.1 |

ns – not significant; * - p < 0.05; ** - p < 0.01; *** - p < 0.001

Fig. 2. The odds ratio (95% confidence interval) of significantly increased inflammatory markers at admission in pediatric patients aged ≤ 12 years (A) and 13–17 years (B) in relation to the exposure to air pollution parameters exceeding limits during a week before hospitalization.

Fig. 3. The length of hospital stay (mean ± SD) in patients aged ≤ 12 (n = 557) (A) and 13–17 years (n = 209) (B) exposed to air pollutants exceeding/not exceeding the limits during a week before hospitalization. The red bar – pollution level exceeding the quality limit; the green bar – pollution level within the quality limit.

increase the overall COVID-19 burden related to pediatric patients.

Instead of using generalized population-based data as in numerous previous analyses of the relationship between air pollution and COVID-19, the levels of PM$_{2.5}$ and B(a)P before hospitalization were matched for each patient allowing to postulate the existence of a causative relationship between exposure to air pollutants and clinical course of the disease. A week preceding hospitalization represents a period of transition of SARS-CoV-2 infection from incubation phase to symptomatic one. During this time, the antiviral defense in unvaccinated individuals is solely based on the innate immune response. It includes a repertoire of dendritic cells, recognizing pathogen-associated molecular patterns and macrophages, natural killer cells, monocytes, neutrophils, and dendritic cells, recognizing pathogen-associated molecular patterns and
infectious compared to adult subjects (Loske et al., 2021). As postulated, this is, at least partially, responsible for the higher rate of asymptomatic and mild cases of COVID-19, faster recovery, and lower incidence of long-term consequences of disease in children (Ding, Yan, and Guo, 2020; Falah, Abdoli, and Kenarkoohi, 2021; Molteni, Sudre, Canas, Bhopal, Hughes, Antonelli et al., 2021). Significantly in this context, PM$_{2.5}$ and B(a)P are well documented to affect innate immune responses adversely and trigger hyper-inflammation in the respiratory tract, effectively jeopardizing its function (Xing et al., 2016; Lewis et al., 2005). Furthermore, some studies demonstrated that children’s nasal epithelium and bronchial tissue reveal significantly lower expression of receptor angiotensin-converting enzyme 2 (ACE2), acting as a cellular receptor for SARS-CoV-2, and lower expression of TMPRSS2, which along with furin is essential for its proteolytic activation (Bunyavanich, Do, and Vicencio, 2020; Saheb Sharif-Akari et al., 2020). In turn, exposures to PM$_{2.5}$ and B(a)P have been shown to up-regulate ACE2 and TMPRSS2 (Borro et al., 2020; H.-H. Li et al., 2021; G. Wang et al., 2021). Therefore, exposure to elevated levels of air pollutants overlapping the SARS-CoV-2 infection can aggravate the clinical severity of COVID-19 in children. This may also explain the previously observed increase in the number of children with symptomatic SARS-CoV-2 infections and pediatric patients requiring hospitalization due to COVID-19 during the second pandemic wave (September-December 2020) when compared to the first wave (March-August 2020) (Pokerska-Spievak et al., 2021).

Although air pollution affected symptomatology, inflammatory markers, and length of hospitalization in both age groups of pediatric patients considered in the study, broader effects were observed for children ≤12 years. This can be to several interlinked factors characterizing this group compared to older children, such as increased minute ventilation, immature immune response, dynamically developing respiratory system, and tendency to spend more time outdoors, ultimately resulting in greater exposure and its health consequences (Buka, Koranteng, and Osorno-Vargas, 2006; Dixon, 2002; Gilliland et al., 1999; Pinkerton and Joad, 2000). In addition, during the pandemic lockdowns and periods of remote learning, children may be increasingly exposed to household air pollution. During periods of high outdoor concentrations, air pollutants can infiltrate indoors and be trapped to varying degrees depending on buildings’ topography, configuration, and ventilation (Nandasena, Wickremasinghe, and Sathikumar, 2013). Importantly, the present study shows that children’s exposure to elevated levels of B(a)P may have more profound effects on the COVID-19 clinical course than in the case of PM$_{2.5}$. However, the majority of previous research on air pollution and COVID-19 and other respiratory diseases focuses not on polycyclic aromatic hydrocarbons such as B(a)P, but on particulates, ozone, sulfur dioxide, and nitrous oxides infections (Rodrigues et al., 2019; Wrotek et al., 2021; Nenna et al., 2017; Mele et al., 2021; Mendy et al., 2021; Veronesi et al., 2022). Among multifaceted harmful effects of B(a)P on human health, including promotion of cancerogenesis, its short-term exposure was shown to induce airway epithelial injury and enhance the release of epithelial-derived pro-inflammatory cytokines, such as thymic stromal lymphopoietin (a distant paralog of IL-7) and IL-6 (Fan et al., 2021). In turn, IL-7 was linked to more severe COVID-19 as it facilitates inflammation in the respiratory tract by inducing the release of innate pro-inflammatory cytokines, i.e., IL-1β, IL-6, TNF-α, IL-12, and IL-23 (Markovic et al., 2021). Moreover, thymic stromal lymphopoietin, which, along with its receptor, mediates the release of pro-inflammatory cytokines (including IL-6, and chemokines, ultimately contributing to airway inflammation, has also been shown to correlate with more severe COVID-19 (Caterino et al., 2021; Shan et al., 2010). Notably, in the present study, pediatric patients (particularly ≤12 years) exposed to elevated levels of B(a)P before hospitalization exhibited higher concentrations of different inflammatory markers at admission, including IL-6. All in all, further studies on air pollution and the severity of COVID-19 and other respiratory diseases in pediatric and adult patients should include B(a)P as a separate risk factor.

Although the present findings provide important data for public health policy, study limitations must be stressed. The research was focused on two major air pollutants, PM$_{2.5}$ and B(a)P, in regions such as Poland, where coal and wood combustion plays a significant role in domestic heating (Aniol et al., 2021; Nazar and Niedoszytko, 2020). However, studies show that other air pollutants, i.e., nitrogen oxides and ozone, may also affect the COVID-19 severity and mortality (Mele et al., 2021; Khorsandi et al., 2021; Achebak et al., 2021). One should also note that the present investigation did not include the potential differences in socioeconomic factors, health behaviors (e.g., physical activity) and presence of comorbidities (e.g., chronic lung disease, diabetes, heart disease, immune deficiencies, seizure disorders, obesity,) that may affect the susceptibility of children to COVID-19 (Choi, Choi, and Yun, 2022; Sondal et al., 2022). Moreover, the patient data were collected over 17 months, during which various SARS-CoV-2 variants emerged and dominated (Hryhorowicz et al., 2021; Jablotska et al., 2021). The shifts in viral variants were not included in the study because the efficient nationwide genomic surveillance was not available in Poland, while the number of SARS-CoV-2 sequences deposited by Polish institutions in the GISAID database was very low (Furuse, 2021). However, one should note that the epidemiological study conducted in Poland during the same period did not show significant increases in the admission of individuals <18 that would otherwise indicate changes in virus pathogenicity (Flištak, Rzyszmski et al., 2021). Similarly, a study comparing the illness profile in children infected with B.1.1.7 (alpha) variant, dominant at the beginning of 2021, and B.1.617.2 (delta) variant, predominately circulating since May 2021, did not show any relevant differences in symptoms prevalence, disease duration, and burden (Molteni, Sudre, Canas, Bhopal, Hughes, Chen et al., 2021). Children diagnosed with COVID-19 caused by delta variant did not show significant differences in inflammatory markers when compared to those infected from February to March 2020 (Sheng, Shao, and Wang, 2021). Therefore, the SARS-CoV-2 evolution appears insufficient in explaining the differences in the clinical parameters found in the present study between children exposed to and not exposed to air pollutants exceeding quality standards. One should also note that the clinical course of COVID-19 can be, to some extent, influenced by meteorological factors such as temperature and humidity (Bochenek et al., 2022), which were not included in the present study. Moreover, the data on the daily duration of outdoor activity and indoor pollution levels, which may affect the exposure to B(a)P and PM$_{2.5}$ (Christian et al., 2022; Rivas et al., 2014), were not available for the studied group. Last but not least, the pediatric group in this study was constituted only of unvaccinated individuals. Whether COVID-19 vaccination may suppress the effect of air pollution on the inflammation levels during SARS-CoV-2 infection and the clinical severity of the disease require further investigations.

5. Conclusion

The results of this study suggest that exposure to air pollutants, particularly B(a)P, at a time of SARS-CoV-2 infection might affect the clinical manifestation of COVID-19 in pediatric patients, with a more significant broad effect seen in those aged ≤12 years. Overall, the findings call for more epidemiological studies on the effect of B(a)P on the severity of COVID-19 and other respiratory diseases and highlight that successful mitigation of the COVID-19 burden should also include measures to decrease the exposure to air pollution.
Funding

This research was supported by the Medical Research Agency in Poland, grant number 2020/ABM/COVID19/PTELICHZ, the Polish Association of Epidemiologists and Infectiologists, and the Department of Environmental Medicine (Poznan University of Medical Sciences, Poland).

CRediT authorship contribution statement

Piotr Rzymski, Barbara Poniedziałał, Robert Flisiak: Conceptualization.
Piotr Rzymski, Barbara Poniedziałał, Robert Flisiak: Data curation.
Piotr Rzymski, Barbara Poniedziałał, Joanna R Rosinska: Formal analysis.
Barbara Poniedziałał, Robert Flisiak: Funding acquisition.
Piotr Rzymski, Barbara Poniedziałał, Joanna R Rosinska, Przemysław Ciechanowski, Michał Peregrym, Maria Pokorska-Śpiękaw, Ewa Talarek, Izabela Zaleska, Paulina Franczak-Chmura, Małgorzata Pilarczyk, Magdalena Figlerowicz, Izabela Kucharek, Robert Flisiak: Investigation.
Piotr Rzymski, Barbara Poniedziałał, Robert Flisiak: Methodology.
Robert Flisiak: Project administration.
Barbara Poniedziałał, Robert Flisiak: Resources.
Piotr Rzymski, Robert Flisiak: Supervision.
Piotr Rzymski: Roles/Writing – original draft.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

References

Achebak, Hicham, Hervé Petetin, Marcos Quijal-Zamorano, Dene Bowdalo, Carlos Paredes García-Pando, and Joan Ballester. 2021. Trade-Offs between Short-Term Mortality Attributable to NO2 and O3 Changes during the COVID-19 Lockdown across Major Spanish Cities. Environmental Pollution (Barking, Essex: 1987) 286 (117220): 117220.
Anioli, Ewa, Suád, Jacek, Bihalovicz, Jan Stefan, Majewski, Grzegorz. 2021. The quality of air in polish health resorts with an emphasis on health on the effects of benzo(a)pyrene in 2015–2019. Climate 9 (5), 74.
Babin, Steven M., Howard, S., Burkom, Holtry, Rekha S., Tabernero, Nathaniel R.,–Piotr Rzymski, Barbara Poniedziałał, Robert Flisiak, Joanna R Rosińska, Katarzyna Aball, Maciej, Owczuk, Radosław, Fakah, S., Abdoli, A., Kenarkoohi, A., 2021. Claims and Reasons about Mild COVID-19 in Children. N. Microbes N. Infect. 41 (100864), 100864.
Fan, Lieyang, Li, Wei, Ma, Jixuan, Cheng, Man, Xie, Li, Ye, Zi, Xie, Yujia, et al., 2021. Benzo(a)pyrene induces airway epithelial injury through wnt5a-mediated non-canonical Wnt VAP/TAZ signaling. Sci. Total Environ. 815 (151695), 151695.
Flisiak, Robert, Horban, Andrzej, Jarosziewicz, Jerzy, Kozielewicz, Dorota, Mastalerz-Migas, Agnieszka, Owczuk, Radosław, Parczewski, Miłoś, et al., 2021. Management of SARS-CoV-2 Infection: Recommendations of the Polish Association of Epidemiologists and Infectiologists as of April 26, 2021. Pol. Arch. Intern. Med. 131 (5), 487–496.
Flisiak, Robert, Horban, Andrzej, Jarosziewicz, Jerzy, Kozielewicz, Dorota, Parczewski, Miłoś, Piekar ska, Anna, Simon, Krzysztof, Tomasiwicz, Krzysztof, Zarzeka-Michalak, Dorota, 2020a. ‘Management of SARS-CoV-2 Infection: Recommendations of the Polish Association of Epidemiologists and Infectiologists as of March 31, 2020. Pol. Arch. Intern. Med. 130 (4), 352–357.
Flisiak, Robert, Horban, Andrzej, Jarosziewicz, Jerzy, Kozielewicz, Dorota, Parczewski, Miłoś, Piekar ska, Anna, Simon, Krzysztof, Tomasiwicz, Krzysztof, Zarzeka-Michalak, Dorota, 2020b. ‘Management of SARS-CoV-2 Infection: Recommendations of the Polish Association of Epidemiologists and Infectiologists. Annex No. 1 as of June 8, 2020. Pol. Arch. Intern. Med. 130 (6), 557–558.
Flisiak, Robert, Parczewski, Miłoś, Horban, Andrzej, Jarosziewicz, Jerzy, Kozielewicz, Dorota, Piekar ska, Anna, Simon, Krzysztof, Tomasiwicz, Krzysztof, Zarzeka-Michalak, Dorota, 2021. Management of SARS-CoV-2 Infection: Recommendations of the Polish Association of Epidemiologists and Infectiologists. Annex No. 2 as of October 3, 2020. Pol. Arch. Intern. Med. 130 (10), 915–918.
Flisiak, Piotr, Rzymski, Piotr, Zarzeka-Michalak, Dorota, Rogalska, Magdalena, Rorat, Marta, Czapunya, Piotr, Lorenc, Beata, et al., 2021. Demographic and Clinical Overview of Hospitalized COVID-19 Patients during the First 17 Months of the Pandemic in Poland. J. Clin. Med. 11 (1), 117.
Furuse, Yuki, 2021. Genomic sequencing effort for SARS-CoV-2 by country during the pandemic. Int. J. Infect. Dis.: IJID: Off. Publ. Int. Soc. Infect. Dis. 103 (February), 305–307.
Galano, Joanna-Evdokia, Andreados, Evangelos, 2021. Impaired innate antiviral defences in COVID-19: causes, consequences and therapeutic opportunities. Semin. Immunol. 55 (101522), 101522.
Gavić, Tomáš, Monrad, Joshua Teperowski, Leech, Gavin, Sharma, Mrinank, Mindermann, Sören, Brauner, Jan Markus, Bhatt, Samir, Kulveit, Jan, 2021. Potential effect of SARS-CoV-2 transmission in temperate climates. BioRxiv. medRxiv. https://doi.org/10.1101/2021.06.10.21258647.
Gilliland, F.D., McConnell, R., Peters, J., Gong Jr., H., 1999. A theoretical basis for investigating ambient air pollution and children’s respiratory health. Environ. Health Perspect. 107 (Suppl 3), 403–407.
Glencross, Drew A., Tzer-reh Ho, Nuria Cami, Catherine M. Hawrylowicz, and Paul E. Pfeffer, 2020, Air Pollution and Its Effects on the Immune System. Free Radical Biology & Medicine 151 (May): 56–68.
Graf, Joseph, Smith, Christopher, Dorsa, Lori, Jung, Sarah, Curran-Hays, Shane, Jarjour, Jane, Carpenter, Lauren, et al., 2021. Risk factors for severe COVID-19 in children. Pediatr. Infect. Dis. J. 40 (4), e137–e145.
Hryhorowicz, Szymon, Ustaszewski, Adam, Kaczmarek-Ry, Marta, Lis, Emilia, Witt, Michaela, Piawski, Andrzej, Ziędkowicz, Ewa, 2021. European context of the diversity and phylogenetic position of SARS-CoV-2 sequences from polish COVID-19 patients. J. Gen. J. Med. Environ. 26 (2), 327–337.
Jabłońska, Katarzyna, Abellina, Samuel, Auguer, Pascal, Tourm, Morfali, 2021. On the Association between SARS-CoV-2 Variants and COVID-19 Mortality during the Second Wave of the Pandemic in Europe. J. Med. Access Health Policy 9 (1), 2002088.
Jackson, William M., Price, Jerri C., Eisler, Lisa, Sun, Lena S., Lee, Jennifer J., 2022. COVID-19 in pediatric patients: a systematic review. J. Neurosurg. Anesthesiol. 34 (1), 141–147.
Jang, Jinhwha, Hwang, Myung-Jae, Kim, Yoo-yeon, Park, Shin Young, You, Myeonggu, Kim, Seong-Sun, Lee, Sangwon, Chung, Dongpyok, 2022. Epidemiological Characteristics and Transmission Patterns of COVID-19 Cases among Children and
