Original Article

Epidemiologic characteristics of immunoglobulin M antibodies in lower respiratory tract infection pathogens of children: association with severe pneumonia in Chengdu city from 2019 to 2023

Jinghua Ye¹, Mei Cha¹, Haibo Yao², Qin Zhang¹, Mengjun Luo³, Yiting Du⁴, Yuanhu Lu⁵, Jinyan Chen¹, Yinghong Fan⁶, Dongmei Xiao⁷, Yixiao Peng¹, Yanyin Liu⁸, Liangyin Deng⁹

¹Department of Public Health, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ²Department of Medicine, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ³Clinical Laboratory, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ⁴Department of Emergency, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ⁵Information Center, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ⁶Department of Respiratory, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, Chengdu 611731, Sichuan, China; ⁷Department of Hospital Infection Control, The First People's Hospital of Shuangliu District (West China Airport Hospital of Sichuan University), Chengdu 610200, Sichuan, China; ⁸Department of Public Health, The Fourth People's Hospital of Sichuan Province, Chengdu 611731, Sichuan, China; ⁹Internal Medicine, Peng Zrou Harmonious Hospital, Chengdu 611731, Sichuan, China

Received May 7, 2025; Accepted June 17, 2025; Epub July 15, 2025; Published July 30, 2025

Abstract: Objective: To analyze the epidemiologic characteristics of serum immunoglobulin M (IgM) antibody aming six pathogens in children with acute lower respiratory tract infections (LRTI) and its association with severe pneumonia, thereby providing a basis for clinical diagnosis and treatment. Methods: A total of 25,693 children with lower respiratory tract infections who were hospitalized in the Department of Respiratory Medicine at Chengdu Women's and Children's Central Hospital from 2019 to 2023 were included and retrospectively analyzed in the study. The epidemiologic characteristics of serum IgM antibodies for six LRTI pathogens in the enrolled children were analyzed. Logistic regression analysis was conducted to identify risk factors for severe pneumonia. Results: The IgM positive detection rate among the six pathogens was 30.26%. In addition, the IgM positive detention rates among Mycoplasma pneumoniae (MP), Parainfluenza virus (PVI) and Influenza B virus (IFV-B) of female children were higher than those of male children (P < 0.05); Children who were younger than 1 year oldshowed the lowest IgM positive detention rates of MP and IFV-B (P < 0.05). The positivity rates of IgM antibodies for MP, IFV-B, PIV, and Respiratory syncytial virus (RSV) markedly varied across different lower respiratory tract infections (P < 0.001; P = 0.007; P = 0.004; P < 0.001). The IgM positive detection rate of two or more pathogens before COVID-19 pandemic were higher in comparison to those post COVID-19 (P < 0.05). Compared to the non-severe group, children with severe pneumonia showed a relatively lower detection rate of MP but a significantly higher detection rate of RSV. The results of multiple logistic regression analysis suggested that age and gender were independent influencing factors for severe pneumonia, with an area under the Receiver Operating Characteristics curve of 0.727. Conclusion: In Chengdu city, the positivity rates of IgM antibodies for LRTI pathogens in children exhibited seasonal, age-related, and diagnostic category characteristics-related variations. Severe pneumonia cases were characterized by RSV infection and younger age. Clinicians should take epidemiologic features into consideration to optimize their diagnostic and therapeutic strategies.

Keywords: Children, lower respiratory tract infection, pathogens, IgM antibodies, severe pneumonia

Introduction

Respiratory infections are one of the most common infectious diseases in children, with symptoms including fever, runny nose, cough, and nasal congestion. Pediatric respiratory infections are recurrent and frequent, making them one of the leading causes of death in children under the age of five. Therefore, early diagnosis of respiratory infection pathogens enables precise treatment and reduces disease burden [1-3]. Studies have shown that Respiratory Syncytial Virus (RSV), Influenza viruses (IFV), Parainfluenza viruses (PIV), and Adenoviruses (ADV) are the major viruses responsible for lower respiratory tract infections (LRTI) in children. Following the COVID-19 pandemic, increasing attention has been paid to various viral infections and Mycoplasma pneumoniae (MP), though related literature remains scarce. However, the epidemiologic characteristics of respiratory tract pathogens are associated with factors such as region, environment, and age, with reports varying especially across different regions [4, 5]. Chengdu city, as a major first-tier megacity in Southwest China, has a population exceeding 20 million and over 80% of its area is urbanized. With its large and dense population, great cross-regional population mobility, and topography-induced air pollution, the pediatric respiratory tract pathogens readily disseminate in this area. This causes a high frequency of respiratory tract infections. In addition, as a consequence of the COVID-19 pandemic, the pathogen monitoring system has undergone structural shifts, resulting in more attention to the changing epidemiologic characteristics of pathogens in clinical settings. Studies have demonstrated that the occurrence rate of pneumonia in children under 5 years old is up to 25% in developing countries, among which 7%-13% are hospitalized for severe pneumonia. This significantly complicates clinical management extra burdens the healthcare system [6]. Currently, research on the long-term changes in pathogen spectrum and predictions of severe pneumonia risk in Chengdu city following the COVID-19 pandemic remains scarce. Therefore, this study aimed to collect the IgM profiles of six pathogens in 25,693 children with acute LRTI in Chengdu city between 2019 and 2023 for the analysis of the epidemiologic characteristics of common pediatric pathogens in this region and their association with severe pneumonia. The study not only provides detailed information about LRTI pathogens to assist clinicians in implementing treatment regimens, but also offers scientific evidence to public health authorities for developing targeted prevention and control strategies, thereby effectively reducing the incidence of respiratory tact infections and their associated mortality.

Data and methods

Patient selection

This is a retrospective study conducted at Chengdu Women's and Children's Central Hospital, a national Grade A tertiary hospital. In 2023, the hospital recorded 2.82 million patient visits and had a total of 1840 approved beds. Therefore, because it represents the region well, the data collected here should hold significant clinical value. A total of 25,693 children hospitalized in the Department of Respiratory Medicine of Chengdu Women's and Children's Central Hospital for acute LRTI between 2019 and 2023 were enrolled and divided into a severe pneumonia group (n = 6.093) and a non-severe pneumonia group (n = 19,600) based on their clinical conditions. The study was approved by the Chengdu Women's and Children's Central Hospital's Ethics Committee (Approval No.: Scientific Research Ethics Review 2025(41)). The flow chart is illustrated in Figure 1.

Inclusion and exclusion criteria: (1) Inclusion Criteria: Children were eligible for the study if they met the diagnostic criteria for LRTI as defined in the Zhu Futang's Practical Pediatrics (8th Edition) [7]; their age \geq 28 days but < 18 years; they were hospitalized in the Department of Respiratory Medicine of Chengdu Women's and Children's Central Hospital between Jan. 2019 and Dec. 2023; their fasting venous blood were collected and IgM antibody tests were completed within 24 hours after hospitalization; their demographic data, clinical records and laboratory reports were complete and obtainable. (2) Exclusion Criteria: Children were excluded from the study if their medical records were incomplete; their respiratory tract samples were not collected; they had a history of Crohn's disease, congenital heart disease, or immunodeficiency requiring long-term immunosuppressive therapy, which could interfere with IgM antibody detection results; or they were discharged or transferred to another department or hospital within 24 hours after hospitalization. It was worth noting that only the first medical record was kept for analysis if an

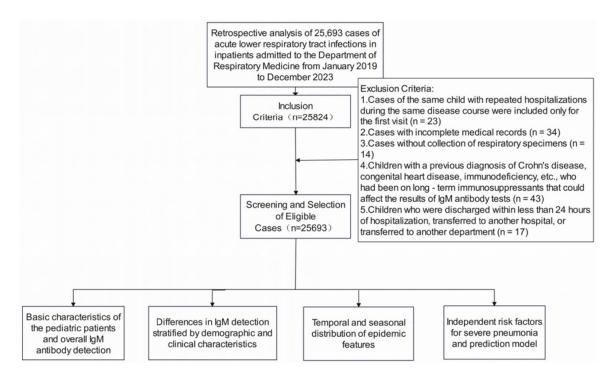


Figure 1. The flow chart of the study.

enrolled child had multiple medical records for the same disease course due to repeated hospital visits.

Data retrieval method: The Jiangsu Mandala Electronic Medical Record System (Version No.: 2023SR0407610) was employed to retrieve data of children who were hospitalized at the Department of Respiratory Medicine of Chengdu Women's and Children's Central Hospital between 2019 and 2023. Children with acute LRTI, defined by the International Classification of Diseases (10th Revision) (ICD-10) codes, were screened for the study.

Data collection

Patients' demographic data (gender, age, length of hospital stay, year of admission and season of admission), clinical diagnostic data and laboratory data [the IgM detection results for MP, influenza A virus (IFV-A), influenza B virus (IFV-B), RSV, parainfluenza virus (PIV), adenovirus (ADV)] were collected.

The IgM antibody detection kits were purchased from Zhengzhou Autobio Biotechnology Co., Ltd. (Approval No.: CFDA Imported Medical Device Registration No. 2010-3400365). The value threshold specified in instructions on the kit was used to determine IgM positive results. The data collected were checked separately by

two researchers. Patients' age, length of hospital stay, and other variables were checked to verify that they met the inclusion criteria. Patients who did not fall into the inclusion range were excluded, or for those whose relevant data were missing, their original medical records were consulted to supplement the information.

Outcome measures

Primary outcome measures: (1) The IgM antibody positive detection results in children. (2) Comparison of IgM positive detection rate of a single pathogen and co-detection rates of multiple pathogens across different clinical characteristics (e.g., gender, age, diagnostic results). (3) Comparison of IgM positive detection rate of a single pathogen and co-detection rates of multiple pathogens across different years and seasons. (4) Comparison of IgM positive detection rate of a single pathogen and co-detection rates of multiple pathogens before and after the COVID-19 pandemic. (5) Comparison of clinical characteristics and IgM positive detection rates of different pathogens between the severe pneumonia group and the non-severe pneumonia group.

Secondary outcome measures: (1) Basic demographic and clinical data of the hospitalized children. (2) Independent risk factors for severe

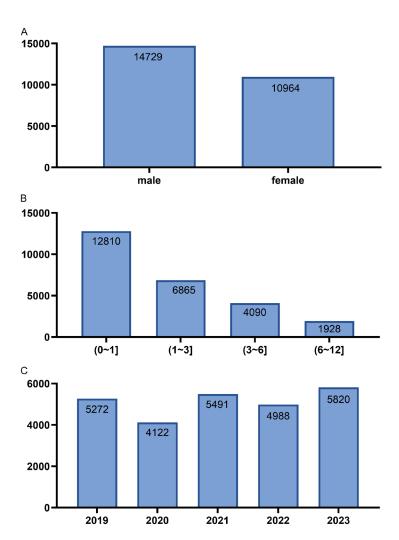


Figure 2. Basic information of hospitalized children seeking medical treatment. A: The number of hospital visiting children of different gender; B: The number of children seeking medical treatment of different ages; C: The number of children seeking medical treatment in different years.

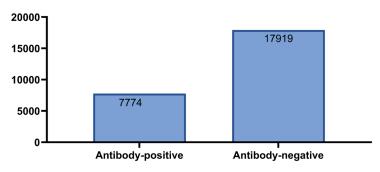


Figure 3. Overall IgM positive detection rates of pathogens.

pneumonia and the construction of a predictive model, which was evaluated using the Area Under the Receiver Operating Characteristics Curve (AUC).

Statistical analysis

Data were processed using SPSS 22.0 software. Counted data were presented as (n, %). Between-group comparisons were performed using the chi-square (χ²) test. Pairwise comparisons of the counted data among multiple groups were conducted using the Bonferroni-corrected chi-square test. Measured data were expressed as mean ± standard deviation ($\bar{x} \pm s$). Betweengroup comparisons were performed using the t-test. Analysis of influencing factors for severe pneumonia was performed using binary logistic regression method. The predictive performance of the model was evaluated using the ROC curve. A P-value < 0.05 was considered significant.

Results

Basic information of the enrolled children

A total of 25,693 children were included in this study, with males accounting for 57.3% and females accounting for 42.7%. When grouped by age, the highest proportions were observed in the \leq 1 year group (49.9%), followed by the 1 to \leq 3 years group (26.7%). When grouped by the year of admission, 20.5% of the cases were from 2019, 16.0% from 2020, 21.4% from 2021, 19.4% from 2022, and 22.7% from 2023. See **Figure 2**.

Overall IgM positive detection rates of LRTI pathogens in children

A total of 7,774 children were found positive for IgM antibodies for LRTI pathogens, with a posi-

tivity rate of 30.26%. Among them, 7,492 cases (96.37%) tested positive for IgM antibodies to a single pathogen, while 282 cases (3.63%) tested for two or more pathogens. See **Figure 3**.

Comparison of IgM positive detection rates of LRTI pathogens across different clinical characteristics

Female children exhibited higher IgM antibody positive rates for MP, PIV, IFV-B, as well as for a single pathogen and for the combination of two or more pathogens compared to male children (all P < 0.05). The IgM antibody positive rates for MP, IFV-B, PIV, and RSV exhibited significant differences across different age groups. Notably, the IgM antibody positive rate for MP increased with age (all P < 0.05). In addition, the IgM positive detection rates of MP, IFV-B, and the co-detection rate for two or more pathogens were lowest in children who were under 1 year old (all P < 0.05). Rates for MP. IFV-B, PIV, and RSV also varied markedly across different LRTI. Children diagnosed with bronchitis exhibited the highest MP positivity rate; those diagnosed with bronchiolitis exhibited the highest co-detection rate for two or more pathogens (all P < 0.05). See Table 1.

Comparison of IgM positive detection rates for a single and multiple pathogens across different years and seasons

The IgM antibody positive rates for a single pathogen, multiple pathogens as well as MP were significantly different across different years (all P < 0.05). The highest positive rates for MP and a single pathogen were observed in 2023 (all P < 0.05). Moreover, the differences in IgM positive detection rates for a single pathogen were statistically significant across different seasons, with the highest detection rates observed in both autumn and winter (all P < 0.05). See Table 2 and Figure 4.

Comparison of IgM positive detection rates of a single pathogen and multiple pathogens before and after the COVID-19 pandemic

The IgM positive rates detected for a single pathogen and for two or more pathogens were higher before the COVID-19 pandemic compared to after (P < 0.05). See **Figure 5**.

Comparison of IgM positive detection rates of LRTI pathogens before and after the COVID-19 pandemic

The IgM positive rates of MP, IFV-A, IFV-B, PIV, ADV, and RSV were all higher before the COVID-

19 pandemic compared to after, with varied reduction observed in the IgM antibody positive rates for all six pathogens after the COVID-19 pandemic (all P < 0.05). See **Table 3**.

Comparison of clinical characteristics and IgM antibody positive rates of various LRTI pathogens between the severe pneumonia group and the non-severe pneumonia group

The severe pneumonia group showed a younger mean age and longer hospitalization duration than the non-severe pneumonia group. The IgM positive detection rate of MP was lower while that of RSV was higher in the severe pneumonia group in comparison to the nonsevere pneumonia group. Moreover, the IgMpositive detection rate of a single pathogen was higher in the severe pneumonia group than that of the non-severe pneumonia group. Ratios of comorbid asthma and rhinitis cases were markedly lower when the same two groups were compared. The case distribution by year and season showed significant differences between the two groups (all P < 0.05). No statistically significant differences were observed in other pathogens or co-infection rates (all P > 0.05). See Table 4.

Independent risk factors for severe pneumonia by multiple logistic regression analysis

Logistic regression analysis was performed with the occurrence of severe pneumonia as the dependent variable and year, season, length of hospital stay, comorbid asthma, comorbid rhinitis, age, positive detection of MP and RSV, as well as any single pathogen as independent variables. The results revealed that age, gender, comorbid rhinitis and asthma, season, year, and length of hospital stay were independent influencing factors for the occurrence of severe pneumonia, with an AUC of 0.727 (P < 0.05). See Table 5 and Figure 6.

Discussion

The study results showed that a total of 7,774 cases tested positive for LRTI pathogen IgM antibodies among the 25,693 enrolled children, yielding an overall positivity rate of 30.26%. The IgM positive rate for MP was 28.34%, for IFV-A was 0.12%, for IFV-B was 1.33%, for ADV was 0.24%, and for RSV was

IgM in lower respiratory tract infection pathogens of children

Table 1. Comparison of IgM positive detection rates of pathogens between different clinical characteristics

	Numahau	Pathogen type					Positive cases of pathogen superimposed detection		
Classification	Number of cases	Mycoplasma pneumoniae	Influenza A virus	Influenza B virus	Parainfluenza virus	Adenovirus	Respiratory syncytial virus	Single pathogen	Superposition of two or more pathogens
Gender									
Male	10964	3537 (32.26)	15 (0.14)	187 (1.71)	103 (0.94)	29 (0.26)	68 (0.62)	3631 (33.12)	145 (1.32)
Female	14729	3745 (25.42)	16 (0.11)	155 (1.05)	95 (0.64)	33 (0.22)	105 (0.71)	3861 (26.21)	137 (0.93)
χ^2		144.209	0.214	19.933	6.752	0.284	0.675	144.692	8.562
Р		< 0.001	0.644	< 0.001	0.009	0.599	0.412	< 0.001	0.003
Age									
≤ 1 year group	12810	2167 (16.92)	13 (0.10)	99 (0.77)	82 (0.64)	25 (0.20)	161 (1.26)	2311 (18.04)	114 (0.89)
1~≤ 3 year group	6865	2463 (35.88)	8 (0.11)	130 (1.89)	76 (1.11)	24 (0.34)	7 (0.10)	2515 (36.64)	90 (1.31)
3~≤ 6 year group	4090	1675 (40.95)	9 (0.22)	82 (2.00)	29 (0.71)	10 (0.24)	3 (0.07)	1684 (41.17)	58 (1.42)
> 6 year group	1928	977 (50.67)	1 (0.05)	31 (1.61)	11 (0.57)	3 (0.16)	2 (0.10)	982 (50.93)	20(1.04)
χ^2		1809.707	4.211	62.201	14.234	5.071	130.095	1680.701	11.934
P		< 0.001	0.211	< 0.001	0.003	0.167	< 0.001	< 0.001	0.008
Classification									
Community-acquired pneumonia	23877	6761 (28.32)	30 (0.12)	303 (1.27)	174 (0.73)	59 (0.25)	153 (0.64)	6955 (29.13)	249 (1.04)
Bronchitis	1423	438 (30.78)	1 (0.07)	31 (2.18)	16 (1.12)	3 (0.21)	4 (0.28)	448 (31.48)	21 (1.48)
Bronchiolitis	393	83 (21.12)	0 (0.00)	8 (2.04)	8 (2.04)	0 (0.00)	16 (4.07)	89 (22.65)	12 (3.05)
χ^2		14.275	0.82	9.972	11.101	1.042	71.504	11.80	16.38
P		0.001	0.662	0.007	0.004	0.595	< 0.001	0.003	< 0.001
Total	25693	7282 (28.34)	31 (0.12)	342 (1.33)	198 (0.77)	62 (0.24)	173 (0.67)	7492 (29.16)	282 (1.10)

Note: The criterion for a IgM positive test of Mycoplasma pneumoniae was a serum antibody titer > 1:160; IgM positive detection rate of a single pathogen = (Number of IgM positive cases for a single pathogen/Total number of cases) × 100%; IgM positive co-detection rate of two or more pathogens = (Number of IgM antibody positive cases for two or more pathogens/Total number of cases) × 100%.

Table 2. Comparison of IgM positive detection rate of a single pathogen and of multiple pathogens across different years and seasons (n, %)

Classification Case number		IgM positive detection rates of a single pathogen	IgM positive detection rates of two or more pathogens		
Year					
2019	5272	1953 (37.04)	219 (4.15)		
2020	4122	851 (20.65)	30 (0.73)		
2021	5491	995 (18.12)	23 (0.42)		
2022	4988	1241 (24.88)	9 (0.18)		
2023	5820	2452 (42.13)	1 (0.02)		
χ^2		1145.535	583.429		
P		< 0.001	< 0.001		
Season					
Spring	6540	1692 (25.87)	87 (1.33)		
Summer	6362	1710 (26.88)	91 (1.43)		
Autumn	6426	2173 (33.82)	59 (0.92)		
Winter	6365	1917 (30.12)	45 (0.71)		
χ^2		120.994	20.60		
Р		< 0.001	< 0.001		

Note: From March to May was defined as Spring, June to August as Summer, September to November as Autumn, and December to February as Winder in accordance with meteorological changes.

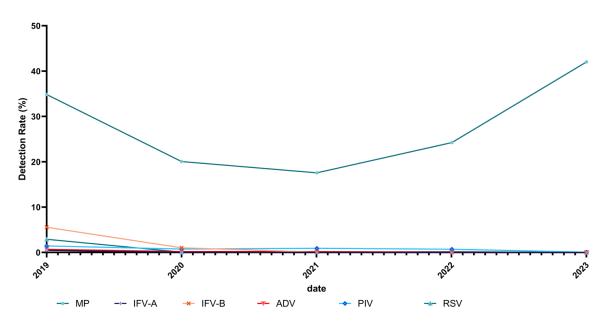


Figure 4. Changing trend of IgM positive rates of various pathogens in children in different years. Note: RSV: Respiratory Syncytial virus; ADV: Adenovirus; PIV: Parainfluenza virus; IFV-B: Influenza B virus; IFV-A: Influenza A virus; MP: Mycoplasma Pneumoniae.

0.67%. These positive detection rates were all lower than those reported by Jun-e Ma et al. and Guo et al. [8, 9]. Variations in geography, research period, subject selection, as well as the detection methods used might be the causes of this difference. Evidently, climate,

population density, and distribution of medical resources in Chengdu city differ from other regions, affecting the MP transmission and its infection rates. Additionally, variations in research period and in the subject selected for various studies may also have influenced the

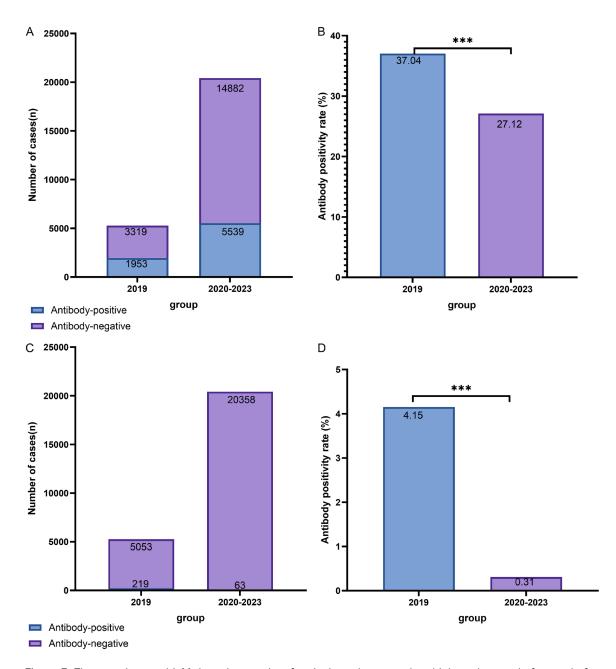


Figure 5. The superimposed IgM detection results of a single pathogen and multiple pathogens before and after COVID-19. A: IgM positive detection cases of a single pathogen before and after COVID-19; B: IgM positive rates of a single pathogen before and after COVID-19; C: IgM positive detection cases of two or more pathogens before and after COVID-19; D: IgM positive detection rate of two or more pathogens before and after COVID-19. Comparison between the two groups, ***P < 0.001.

virus detection rates. Furthermore, differences in the detection methods and standards used in studies may have led to variations in results [10, 11].

The high IgM positive detection rate for MP, IFV-B, and the high IgM positive co-detection rate of two or more pathogen in females compared

to males contradicts the findings of Peer et al., but aligns with the results of Mai and Wang [12-14]. This gender difference in IgM antibody positivity rates may be in association with strong immune responses and the distinct regulatory effects of estrogen and androgen on immune responses in females [15, 16]. Moreover, genetic susceptibility and anatomical dif-

Table 3. Comparison of IgM positive detection rates of pathogens before and after COVID-19 pandemic

Year	Case number	MP	IFV-A	IFV-B	PIV	ADV	Respiratory syncytial virus
Before the COVID-19 pandemic (2019)	5272	1836 (34.83)	24 (0.46)	293 (5.56)	76 (1.44)	37 (0.70)	154 (2.92)
After COVID-19 (2020-2023)	20421	5446 (26.67)	7 (0.03)	49 (0.24)	122 (0.60)	25 (0.12)	19 (0.09)
χ^2		137.273	61.615	902.185	39.047	58.433	501.089
Р		< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001

Note: MP: Mycoplasma pneumoniae; IFV-A: Influenza A virus; IFV-B: Influenza B virus; PIV: Parainfluenza virus; ADV: Adenovirus.

Table 4. Comparison of clinical characteristics and IgM positive detection rates of different pathogens between the severe pneumonia group and none-severe pneumonia group

Index	Severe pneumonia group (n = 6093)	Non-severe pneumonia group (n = 19600)	t/χ²	Р	
Ages (Years)	1.88 ± 2.60	2.51 ± 2.56	9.817	< 0.001	
Sex			1.333	0.248	
Male	3454	11275			
Female	2639	8325			
Years			367.327	< 0.001	
2019	1497	3775			
2020	1271	2851			
2021	1209	4282			
2022	1170	3818			
2023	946	4874			
Season			55.071	< 0.001	
Spring	1486	5054			
Summer	1541	4821			
Autumn	1370	5056			
Winter	1696	4669			
Length of hospital stay	7.06 ± 1.54	6.02 ± 1.37	46.316	< 0.001	
Pathogen detection type					
MP	1560	5722	29.508	< 0.001	
IFV-A	12	19	3.858	0.050	
IFV-B	69	273	2.400	0.123	
PIV	36	162	3.376	0.066	
ADV	13	49	0.259	0.611	
RSV	60	113	11.581	0.001	
Infection with a single pathogen			24.996	< 0.001	
Yes	1687	6087			
No	4406	13513			
Co-infection with 2 or more pathogens	58	224	1.562	0.212	
Comorbid asthma					
Yes	463	2501	121.329	< 0.001	
No	5630	17099			
Comorbid rhinitis					
Yes	107	1007	128.147	< 0.001	
No	5986	18593			

Note: MP: Mycoplasma pneumoniae; IFV-A: Influenza A virus; IFV-B: Influenza B virus; PIV: Parainfluenza virus; ADV: Adenovirus; RSV: Respiratory Syncytial Virus.

Table 5. Influencing factors for the occurrence of severe pneumonia by logistic regression analysis

Variables	В	S.E	Wald	Р	Exp (B)	95% CI lower limit	95% CI upper limit
Sex	-1.03	0.37	7.86	0.005	0.903	0.84	0.97
Length of hospital stay	0.514	0.013	1452.102	0.000	1.673	1.629	1.717
Comorbid asthma	-0.507	0.068	55.231	< 0.001	0.603	0.527	0.689
Comorbid rhinitis	-1.093	0.13	70.97	< 0.001	0.335	0.26	0.432
Age	-0.099	0.008	138.776	< 0.001	0.905	0.891	0.921
Year_2020	-0.331	0.044	57.297	< 0.001	0.718	0.659	0.782
Season_Summer	-0.083	0.02	17.878	< 0.001	0.921	0.886	0.957
Constants	-3.721	1.03	1302.245	< 0.001	0.024		

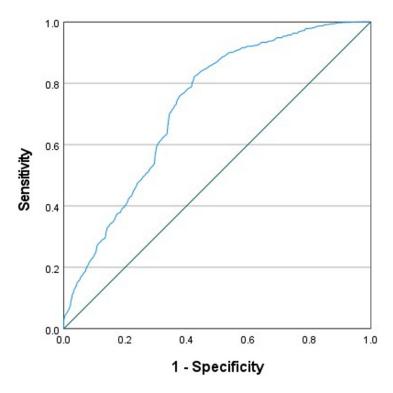


Figure 6. The ROC curve.

ferences could make female children more vulnerable to get infected by certain pathogens in comparison to male children.

The high IgM positive detection rates of MP in older groups, with the lowest detection rate in children who were no older than 1 year, was consistent with the findings of Choo et al. [17]. In contrast, the IgM positive detection rate of RSV was the highest in the same aforementioned age group, which aligned with the results of Suss et al. [18] and Bechini et al. [19]. This age-related contrast in the positivity rates might be attributed to various factors, including pathogen characteristics, immune system

development, social behavior, and environmental exposure [20, 21]. MP primarily spreads through respiratory droplets and human contact. Children over 3 years old, who undergo frequent interactions in collective settings such as a daycare center or school, are susceptible to pathogen infection, leading to a high infection rate. In addition, RSV is highly contagious. Children under 1 year old, whose immune systems are still developing, are more susceptible to infection upon first exposure. From a clinical treatment perspective, the age-related differences in pathogen distribution have important implications for the formulation of diagnostic and treatment strategies. For older children, the high IgM positive detection rate of MP suggests that clinicians should prioritize considering MP infection when evaluating chil-

dren manifesting LRTI symptoms. Early targeted tests (such as IgM antibody test or PCR) should be conducted to initiate appropriate treatment with macrolide antibiotics in a timely manner. For children under 1 year old, the high IgM positive detection rate of RSV indicates that clinicians should be vigilant for the possibility of bronchiolitis, especially during winter when RSV is prevalent. Our study also found that the IgM positive detection rate of RSV was higher in the severe pneumonia group compared to the non-severe pneumonia group, suggesting that children infected with RSV were more likely to progress to severe conditions, which aligns with the results reported by

Treston et al [22]. Therefore, clinicians should identify high-risk populations for severe conditions and rationalize medical resource allocation to reduce the incidence of severe cases and mortality [22].

Since 2019, the IgM positive detection rates of MP, RSV, ADV, IFV-A and a single pathogen infection have decreased, with MP showing the lowest IgM positive detection rate between 2020 and 2022. However, in 2023, it rebounded to the highest level, even surpassing the rate of 2019. This phenomenon may be attributed to multiple factors. Firstly, the strict public health measures implemented during the COVID-19 pandemic, such as wearing masks, social distancing, and reduced social interactions, not only suppressed the transmission of the SARS-CoV-2 virus but also significantly curtailed the spread of other respiratory tract pathogens. Moreover, while medical resources and clinical practices were primarily allocated to the detection of COVID-19 virus, less attention was paid to the detection of other respiratory tract pathogens. However, as pandemic control measures were relaxed and life returned to normal, certain populations, having been largely isolated from pathogens for an extended period, developed an "immune gap". This enabled pathogens to spread easily when they reemerged.

Among different seasons, the IgM positive detection rate of a single pathogen was higher in autumn and winter compared to spring and summer. Different pathogens exhibit varied seasonality features. In winter, the climate gets cold and dry, therefore it creates a suitable environment for viruses to grow and spread. Moreover, people tend to stay indoors due to the coldness, resulting in more close human contact and, consequently, the risk of respiratory tract pathogen infection. The impact of the flu season may also contribute to high IgM positive detection rates of other pathogens, particularly in cases of occurrence of co-infection related to influenza.

The IgM positive detection rates of MP, IFV-B, PIV and RSV differed markedly among various LRTI, indicating that different pathogens may be associated with distinct types of LRTI. RSV is the primary pathogen in bronchiolitis, while MP is commonly found in bronchitis. This finding has significant implications for clinical diag-

nosis and treatment. From a clinical perspective, understanding the association between pathogens and infection type can help optimize treatment strategies. For children with bronchiolitis caused by RSV, clinicians should focus on supportive therapies, such as oxygen therapy, nebulized bronchodilators, and, when necessary, corticosteroids. For critically ill children, antiviral medications may be considered, especially in immunocompromised patients. In the case of bronchitis caused by MP infection, early diagnosis and prompt initiation of macrolide antibiotics are crucial to reduce complications and shorten disease course. Further analysis revealed that gender, age, time period before or after COVID-19, and the LRTI types were all independent factors influencing the IgM positive detection rate of a single pathogen.

The study further analyzed the clinical characteristics and IgM positivity rates of different pathogens in children with severe pneumonia. In this study, 25.6% of patients with severe pneumonia were MP positive, significantly lower than the 29.2% in the non-severe group, suggesting that while MP infection is highly contagious, it is less likely to progress to a severe condition. The RSV positive rate was markedly higher in the severe group than that of the non-severe group, reflecting the propensity of RSV infection for severe outcomes and heightened risk for younger children. This heightened severe RSV pneumonia susceptibility for younger patients can be attributed to their immature immune systems. As children grow older, their exposure to various pathogens increases, leading to the formation of immune response memory, and as a result, improved tolerance to treatment and reduced occurrence of severe conditions. This result underscored the need for heightened clinical vigilance for LRTI children under 1 year old, particularly during RSV transmission seasons. From 2019 to 2023, the severe pneumonia group showed a consistent declining trend, while the non-severe pneumonia group rebounded significantly after 2020 (the peak pandemic year) and reached its zenith in 2023. This phenomenon may stem from the effects of COVID-19 public health interventions (e.g., wearing masks and social distancing) on the transmission of respiratory tract pathogens. The results of logistic regression analysis revealed that children who had asthma presented a reduced risk of severe

pneumonia. This may be attributed to a protective mechanism in specific asthma phenotypes in response to pulmonary infections or possibly related to better home management and increased attention given to asthmatic children, leading them to receive prompt medical care when they develop LRTI [23].

In summary, the IgM positive detection rates of LRTI pathogens in children in Chengdu city exhibited seasonal, age and infection type-related differences. The characteristics of certain pathogen IgM antibodies may be associated with specific infection types. Clinicians should combine epidemiologic data with patient symptoms to select appropriate diagnostic tests, thereby optimizing treatment plans and improving therapeutic outcomes.

Acknowledgements

This study was supported by the Chengdu Science and Technology Program (Project Number: 2024-YF05-00747-SN), the 2022 Chengdu Women and Children's "Outstanding Talent Program" Project (Project Number: YC2022019) and the Chengdu Medical Research Project (Project Number: 2024136).

Disclosure of conflict of interest

None.

Address correspondence to: Qin Zhang, Department of Public Health, Chengdu Women's and Children's Central Hospital, School of Medicine, University of Electronic Science and Technology of China, No. 1617, Riyue Avenue, Qingyang District, Chengdu 611731, Sichuan, China. Tel: +86-153-90407972; E-mail: zq_2911@126.com; Dongmei Xiao, Department of Hospital Infection Control, The First People's Hospital of Shuangliu District (West China Airport Hospital of Sichuan University), No. 120, Chengbei Shangjie, Dongsheng Sub-District, Shuangliu District, Chengdu 610200, Sichuan, China. Tel: +86-13699066991; E-mail: xiaodongmei2025@163.com

References

- [1] Lv G, Shi L, Liu Y, Sun X and Mu K. Epidemiological characteristics of common respiratory pathogens in children. Sci Rep 2024; 14: 16299.
- [2] Li Y, Wang X, Blau DM, Caballero MT, Feikin DR, Gill CJ, Madhi SA, Omer SB, Simões EAF,

Campbell H, Pariente AB, Bardach D, Bassat Q, Casalegno JS, Chakhunashvili G, Crawford N, Danilenko D, Do LAH, Echavarria M, Gentile A, Gordon A, Heikkinen T, Huang OS, Jullien S, Krishnan A, Lopez EL, Markić J, Mira-Iglesias A, Moore HC, Moyes J, Mwananyanda L, Nokes DJ, Noordeen F, Obodai E, Palani N, Romero C, Salimi V, Satav A, Seo E, Shchomak Z, Singleton R, Stolyarov K, Stoszek SK, von Gottberg A, Wurzel D. Yoshida LM, Yung CF and Zar HJ; Respiratory Virus Global Epidemiology Network, Nair H; RESCEU investigators. Global, regional, and national disease burden estimates of acute lower respiratory infections due to respiratory syncytial virus in children younger than 5 years in 2019: a systematic analysis. Lancet 2022; 399: 2047-2064.

- [3] Wei M, Li S, Lu X, Hu K, Li Z and Li M. Changing respiratory pathogens infection patterns after COVID-19 pandemic in Shanghai, China. J Med Virol 2024; 96: e29616.
- [4] Song LX and Zeng HW. Analysis of common viral infections in patients with respiratory tract infections in zigong fourth people's hospital from 2016 to 2019. Pract J Clin Med 2021; 18: 126-129.
- [5] Wang ZJ, Fang J, Hu XL, Hou XH, Xie YF, Zhu J, Ma YC and Zhuang Q. Pathogen and epidemiological characteristics of atypical pneumonia in children. Pract J Clin Med 2020; 17: 156-158.
- [6] Turyasiima M, Kiconco G, Egesa WI, Twesigemukama S and Nduwimana M. Prevalence and outpatient clinical diagnostic approaches for common acute respiratory tract infections in children under five years of age: a crosssectional study. Pediatric Health Med Ther 2024; 15: 49-57.
- [7] Jiang ZF, Shen KL and Shen Y. Zhu Futang's practical pediatrics (8th edition). Beijing. People's Medical Publishing House, 2015.
- [8] Ma JE, Ma QF, Wang W, Zeng Y, Cui XH, Wang GL, Liu SY, Wang Z, Xiao X, Xia ZN, Zhu HC and Zhu CL. Analysis of common respiratory infected pathogens in 3100 children after the coronavirus disease 2019 pandemic. Curr Med Sci 2022; 42: 1094-1098.
- [9] Guo Y, Yang D, Cao Y, Ding X, Chen L, Huo B and Li Y. Influence of COVID-19 public health restrictions on community-acquired pneumonia pathogens in children in Henan, China: a multicenter retrospective study. BMC Infect Dis 2024: 24: 1381.
- [10] Ma PC and Chen Y. Research progress in detection methods of respiratory pathogens. Chin J Infect Chemother 2024; 24: 118-122.
- [11] Chen X, Wang F, Fu Y, Huang L, Li F, Zhao H, Guan X, Li Q, Li Q, Wang Y, Guo Y and Xie Z. Development and evaluation of a multiplex

- digital PCR method for sensitive and accurate detection of respiratory pathogens in children. Virology 2024; 590: 109948.
- [12] Peer V, Mandelboim M, Jurkowicz M and Green MS. Sex differences in acute respiratory tract infections-multi-year analysis based on data from a large tertiary care medical center in Israel. Front Public Health 2025; 13: 1502036.
- [13] Mai W, Ren Y, Tian X, Al-Mahdi AY, Peng R, An J, Lin Q, Hu X, Wang G, Sun C, Lu Z, Du J, Xiao M and Yin F. Comparison of common human respiratory pathogens among hospitalized children aged ≤ 6 years in Hainan Island, China, during spring and early summer in 2019-2021. J Med Virol 2023; 95: e28692.
- [14] Wang C, Zhao Y, Mu D, Zhang X, Li T, Kong H and Pu D. Epidemiological characteristics of respiratory pathogens in Changchun, Jilin Province, China, following the pandemic. Int J Environ Health Res 2025; [Epub ahead of print].
- [15] Lingasamy P, Modhukur V, Mändar R and Salumets A. Exploring immunome and microbiome interplay in reproductive health: current knowledge, challenges, and novel diagnostic tools. Semin Reprod Med 2023; 41: 172-189.
- [16] Horn C, Sprute R, Kretschmer AC, Do C, Cornely OA, Jung N, Lehmann C and Fischer J. Sex in infectious diseases-How sex differences influence the immune response to infections. Inn Med (Heidelb) 2023; 64: 752-757.
- [17] Choo S, Kim SH and Lee E. Clinical significance of Mycoplasma pneumoniae specific IgM titer in children hospitalized with Mycoplasma pneumoniae pneumonia. BMC Infect Dis 2022; 22: 470.

- [18] Suss RJ and Simões EAF. Respiratory syncytial virus hospital-based burden of disease in children younger than 5 years, 2015-2022. JAMA Netw Open 2024; 7: e247125.
- [19] Bechini A, Salvati C, Bonito B, Del Riccio M, Stancanelli E, Bruschi M, Ionita G, Iamarino J, Bentivegna D, Buscemi P, Ciardi G, Cosma C, Stacchini L, Paoli S, Conticello C, Bega M, Schirripa A, Bertizzolo L, Muzii B, Azzi MV, Parisi S, Trippi F, Bonanni P and Boccalini S. Respiratory Syncytial Virus associated hospitalisations in children up to 6 years of age in Italy: a systematic review. Ann Ig 2025; 37: 241-254.
- [20] Cao J, Pan J, Yang X, Liu J, Zhu M, Zhao Z, Chen L, Chen T and Ye H. Trends in respiratory infectious pathogens in children under the age of 14 Xiamen City, Fujian Province, China, 2017-2023. China CDC Wkly 2024; 6: 143-147.
- [21] Han T, Wang Y, Zhang D, Li Y, Zhang L, Yan J, Li C, Yang S, Guo L and Yan H. Changes in infant respiratory pathogens pre-, during, and post-COVID-19 non-pharmacological interventions in Beijing. Ital J Pediatr 2025; 51: 8.
- [22] Treston B and Geoghegan S. Exploring parental perspectives: maternal RSV vaccination versus infant RSV monoclonal antibody. Hum Vaccin Immunother 2024; 20: 2341505.
- [23] Espejo D, Pilia F, Romero-Mesones C, Ojanguren I, Cruz MJ and Muñoz X. Respiratory sequelae in patients with bronchial asthma after SARS-CoV-2 pneumonia: a retrospective study in a large academic centre in 2020. ERJ Open Res 2025; 11: 00510-2024.