## **Vital Surveillances**

# **Epidemiological Characteristics of Human Parainfluenza Viruses Infections — China, 2019–2023**

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## **ABSTRACT**

**Introduction**: A retrospective study based on sentinel surveillance was conducted in 10 provincial-level administrative divisions (PLADs) in China to enhance the understanding of the epidemiological characteristics of human parainfluenza viruses (HPIVs).

Methods: From January 2019 to June 2023, respiratory specimens were collected from individuals with acute respiratory infections (ARIs) and screened for four HPIVs serotypes and other common respiratory viruses using multiplex real-time polymerase chain reaction (PCR). This study analyzed the association of HPIVs infections with seasonal patterns, geographical distribution, demographic profiles, clinical features, and co-infection status.

**Results**: During the study period, a total of 12,866 ARIs were included. The overall detection rate of HPIVs was 6.15%, varying from 5.04% in 2022 to 9.70% in 2020. The median age of HPIVs-infected patients was 3 years. HPIV2 was more prevalent among individuals aged 5-17 years (42.57%), while HPIV4 was more common in those over 65 years (12.24%). HPIV3 (54.16%) and HPIV1 (27.18%) were the predominant serotypes, and their prevalence exhibited significant seasonal fluctuations postcoronavirus disease 2019 (COVID-19) pandemic. The peak of HPIV3 shifted three months later in 2020 compared to 2019 and returned to a summer peak thereafter. Two peaks of HPIV1 were observed in 2021 following the peak of HPIV3. Additionally, coinfections were frequent in HPIVs cases (overall rate: 22.12%), with human rhinovirus being the most common co-infecting virus.

**Conclusions**: The prevalence of HPIVs in China was predominantly due to HPIV3 and HPIV1, and their seasonal patterns were altered by pandemic restrictions. Hence, continuous surveillance of HPIVs is essential.

Human parainfluenza viruses (HPIVs) significant viral agents causing acute respiratory and infections (ARIs) severe acute respiratory (SARIs) in infections children globally (1-3). Currently, there are four known serotypes (HPIV1 to HPIV4) (2). HPIV1 and HPIV3 are more prevalent, with epidemics typically occurring in the fall and summer, respectively. In contrast, HPIV2 and HPIV4 are primarily active during the winter and spring months (4-5). The coronavirus disease 2019 (COVID-19) pandemic has led to observable seasonal changes in the patterns of several respiratory viruses (6-7). The objective of this study was to analyze epidemiological characteristics of HPIVs infections, aiming to offer a crucial scientific groundwork for preventing and controlling HPIVs epidemic in China.

## **METHODS**

Clinical samples were collected from 11 sentinel sites across 10 provincial-level administrative divisions (PLADs), including seven PLADs in northern China (Beijing Municipality; Inner Mongolia Autonomous Region; and Henan, Gansu, Jilin, Liaoning, and Shandong provinces) and three PLADs in southern (Chongging Municipality, Jiangxi Guangdong provinces). Following the case definitions in the guideline for hospitalized acute respiratory infections (ARIs) and severe acute respiratory infections (SARIs, 2011 version) released by the National Health Commission (8), a minimum of 20 ARIs patients per month were included in the study, with respiratory samples taken for pathogen screening. The study was approved by the Second Ethics Review Committee of the National Institute for Viral Disease Control and Prevention at China CDC, and informed consent was obtained from all patients or their guardians before sample collection.

A commercial nucleic acid detection kit (Kinghawk, Beijing, China) was used to identify twelve viral respiratory pathogens, including HPIVs (HPIV1 to HPIV4), respiratory syncytial virus (HRSV), influenza virus (IFV), human adenovirus (HAdV), human rhinovirus (HRV), human metapneumovirus (HMPV), human enterovirus (HEV), human bocavirus (HBoV), and human coronavirus (HCoV). Prior to testing, all assays underwent validation.

Categorical data were analyzed using the chi-square ( $\chi^2$ ) test and Fisher's exact test, while continuous data were assessed using median tests. Statistical analyses were conducted using R (version 4.1.3, R Core Team and the R Foundation for Statistical Computing, Vienna, Austria). Statistical significance was defined as a *P*-value <0.05.

## **RESULTS**

#### **Overall Detection**

During the study period, we enrolled 12,866 cases of ARIs. Excluding the 14.18% (1,825) of cases that lacked relevant clinical information, 70.46% (9,066) were classified as SARIs, while 15.35% (1,975) were considered mild ARIs. The median age of patients with ARIs was 3 years (range: 1 day to 100 years), and 71.34% were under the age of 5 years. A majority of

the patients, 80.7% (10,378/12,866) hailed from northern PLADs (Table 1), with males accounting for 57.88%. The overall viral detection rate among the ARIs was 38.18% (4,912/12,866). HPIVs were identified in 6.15% (791/12,866) of the cases, being the third most frequently detected viruses following HRSV at 12.37% (1,592/12,866) and HRV at 9.44% (1,214/12,866). Within the HPIVs-positive ARIs, HPIV3 was the most common serotype, accounting for 54.61% (432/791), followed by HPIV1 at 27.18% (215/791), HPIV2 at 12.77% (101/791), and HPIV4 at 6.19% (49/791) (Table 1, Supplementary Table S1, available at https://weekly.chinacdc.cn/).

Out of 9,066 SARIs, the overall viral detection rate was 37.13% (3,366/9,066). The detection rates of HPIVs at 5.40% (490/9,066) in SARIs were in line with ARIs. Among HPIVs-infected individuals, HPIV3 was the most prevalent at 58.16% (285/490), followed by HPIV1 at 27.35% (134/490), HPIV2 at 8.98% (44/490), and HPIV4 at 5.92% (29/490) (Supplementary Table S2, available at https://weekly.chinacdc.cn/).

The detection rate of HPIVs among mild ARIs (6.84%, 135/1,975) was similar to that of all ARIs and SARIs. HPIV3 (40%, 54/135) was the most commonly detected type, while HPIV4 (5.93%, 8/135) was the least detected. Interestingly, in mild ARIs, HPIV2 (28.15%, 38/135) had a slightly higher prevalence than HPIV1 (25.96%, 35/135).

TABLE 1. Geographical distribution of HPIVs infections in China from 2019 to 2023.

PLADs	Period	riod No. of samples	of samples Region of China -	No	No. of HPIVs cases included in the analysis					
PLADS	ADS Period No. of samples Region of Ch		Region of China	HPIV1 (%)	HPIV2 (%)	HPIV3 (%)	HPIV4 (%)	HPIVs		
Jilin	2019–2023	2,927	Northern	60 (43.5)	1 (0.7)	78 (56.5)	0 (0)	138*		
Henan	2019–2022	2,068	Northern	24 (22.6)	22 (20.8)	45 (42.5)	15 (14.2)	106		
Shandong	2019–2023	1,509	Northern	21 (18.2)	14 (12.1)	76 (66.0)	5 (4.3)	115 <sup>†</sup>		
Gansu	2020–2023	1,446	Northern	34 (23.1)	35 (23.8)	66 (44.9)	12 (8.2)	147		
Beijing	2019–2023	1,444	Northern	36 (36.7)	14 (14.3)	40 (40.8)	9 (9.2)	$98^{\dagger}$		
Inner Mongolia	2021–2023	537	Northern	0 (0)	8 (21.6)	25 (67.6)	4 (10.8)	37		
Liaoning	2021–2022	447	Northern	0 (0)	3 (60.0)	2 (40.0)	0 (0)	5		
Jiangxi	2021–2023	1,069	Southern	4 (57.1)	0 (0)	3 (42.9)	1 (14.3)	<b>7</b> §		
Guangdong	2021–2023	841	Southern	21 (31.3)	0 (0)	46 (68.7)	0 (0)	67		
Chongqing	2020–2023	578	Southern	15 (21.1)	4 (5.6)	51 (71.8)	3 (4.2)	71* <sup>¶</sup>		
Total	2019–2023	12,866	China	215 (27.2)	101 (12.8)	432 (54.6)	49 (6.2)	791		

Note: Due to co-infection among the four HPIVs serotypes, the total number of cases counted in the table for the four serotypes was higher than the number of actual HPIVs infections.

Abbreviation: HPIVs=human parainfluenza viruses; PLADs=provincial-level administrative divisions.

<sup>\*</sup> it included one case with HPIV1 and HPIV3 co-infection.

<sup>&</sup>lt;sup>†</sup> it included one case with HPIV2 and HPIV4 co-infection.

<sup>§</sup> it included one case with HPIV1 and HPIV4 co-infection.

<sup>&</sup>lt;sup>¶</sup> it included one case with HPIV3 and HPIV4 co-infection.

## **Season Pattern**

The detection rate of HPIVs increased from 5.07% (78/1,539) in 2019 to 9.70% (146/1,505) in 2020, decreased to 7.63% (311/4,075) in 2021, and further dropped to 5.04% (208/4,124) in 2022 (Figure 1, Supplementary Table S3, available at https://weekly.chinacdc.cn/). The epidemics of HPIV1 and HPIV3 exhibited distinct seasonal patterns.

The incidence of HPIV1 showed a peak between August and December 2019, then shifted to the winter period from December 2020 to January 2021.

Subsequently, two peaks were observed in July and October 2021. In 2021, the detection rate of HPIV1 was highest at 3.46% (141/4,075) (*P*<0.001) (Supplementary Table S3). However, from January 2022 to June 2023, the detection of HPIV1 decreased, maintaining a relatively low level at 0.51% and 0.37% without a distinct peak (Figure 1).

The incidence of HPIV3 differed from HPIV1, peaking in summer (June–August) in 2019, shifting to autumn and winter (October–November) in 2020, and returning to summer (June–August) in 2021 and 2022.

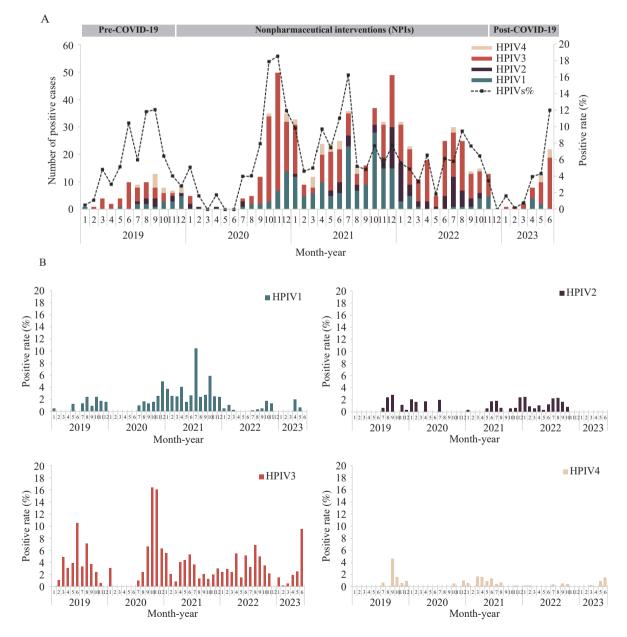


FIGURE 1. Season pattern of HPIVs in China from 2019 to 2023. (A) Detection of HPIVs by year. (B) Detection of HPIV1 to HPIV4 by year.

Abbreviation: HPIVs=human parainfluenza viruses; COVID-19=coronavirus disease 2019.

HPIV2 showed the highest detection rate in 2022 (1.24%, *P*=0.010) and was present all year except in November and December. In contrast, HPIV4 was only sporadically detected, with a higher rate in 2019 (0.78%, *P*=0.002) (Supplementary Table S3). There was no significant seasonal pattern observed for these two serotypes (Figure 1).

# **Geographical Distribution**

The prevalence of HPIVs serotypes varied across different PLADs as shown in Table 1. HPIV1 and HPIV3 were the most common in northern and southern China, especially HPIV3 which had a higher prevalence in southern regions (4.02%, 100/2,488) compared to northern regions (3.20%, 332/10,378) (P<0.001). Conversely, the detection rates of HPIV2 (0.93%, 97/10,378) and HPIV4 (0.43%, 45/10,378) were higher in northern PLADs than in the southern PLADs (P<0.001, P=0.047) (Supplementary Table S3).

# **Demographic Profile**

Among the 791 patients infected with HPIVs, 59.29% were male, with no sex-based difference observed (Table 2). The median age of the patients was 3 years, ranging from 1 month to 90 years, with the majority being children under 5 years old (77.24%), followed by individuals aged 5–17 years (16.69%) and those aged 65 years and older (3.92%)(Table 2). Notably, a significant difference was found in the age distribution for HPIV2 and HPIV4 infections (*P*<0.001). HPIV2 was more prevalent in the 5-17 years age group (42.57%), while HPIV4 was

predominant in those over 65 years old (12.24%) (Table 2). Moreover, there were no significant variations in sex, age, or region for mono- and coinfections of HPIVs (Supplementary Table S4, available at https://weekly.chinacdc.cn/).

In 490 SARIs caused by HPIVs, the median age was 2 years (range: 1 month to 90 years). The gender and age distributions of HPIVs infections among SARIs were similar to those in ARIs (Supplementary Table S5, available at https://weekly.chinacdc.cn/).

### **Clinical Features**

The analysis included 6,442 ARIs with complete clinical information, of which 357 cases were HPIVs infections. The symptoms manifested by HPIVs patients included fever (87.39%, P=0.002), cough (83.47%, P=0.023), phlegm production (49.86%, P<0.001), sore throat (33.33%, P<0.001), headache (14.57%, P<0.001), and diarrhea (7.28%, P=0.013) (Table 3). HPIV2-infected patients were particularly more likely to experience fever (97.73%, P=0.024), produce phlegm (72.73%, P<0.001), headaches (65.91%, P<0.001), and sore throats (54.55%, P=0.008). Rhinorrhea was notably seen in HPIV4 infections (75%, P<0.001), although this observation could be influenced by the limited number of HPIV4 cases within the study (Table 3). Additionally, coinfection with HPIVs was associated with a higher incidence of headache symptoms (33.33%), which was higher than that in patients with HPIVs monoinfection (0.97%, P<0.001)(Table 3).

Among the 278 SARIs caused by HPIVs, common symptoms were "cough" (88.85%, P=0.001), "rhinorrhea" (44.24%, P=0.007), and "diarrhea"

TABLE 2. Demographic characteristics of HPIVs infections among patients with ARIs (January 2019–June 2023).

Oh a va ata viati a	HPIV1	HPIV2	HPIV3	HPIV4		HPIVs
Characteristic	<i>n</i> =215	<i>n</i> =101	n=432	<i>n</i> =49	P	<i>n</i> =791
M (Q <sub>1</sub> , Q <sub>3</sub> )	3 (1, 4)	4 (3, 6)	2 (1, 3)	3 (2, 5)	<0.001	3 (1, 4)
Age (years), N (%)					<0.001	
0–4	169 (78.60)	52 (51.49)	361 (83.56)	33 (67.35)		611 (77.24)
5–17	30 (13.95)	43 (42.57)	52 (12.04)	8 (16.33)		132 (16.69)
18–39	1 (0.47)	0 (0)	1 (0.23)	1 (2.04)		3 (0.38)
40–64	3 (1.40)	2 (1.98)	8 (1.85)	1 (2.04)		14 (1.77)
65+	12 (5.58)	4 (3.96)	10 (2.31)	6 (12.24)		31 (3.92)
Sex, N (%)					0.746	
Male	121 (56.28)	59 (58.42)	262 (60.65)	30 (61.22)		469 (59.29)
Female	94 (43.72)	42 (41.58)	170 (39.35)	19 (38.78)		322 (40.71)

Abbreviation: HPIVs=human parainfluenza viruses; ARIs=acute respiratory infections.

(9.35%, P=0.010). Conversely, non-SARIs were more likely to experience symptoms such as "fever" (96.20%, P=0.013), "phlegm" (69.62%, P=0.001), "headache" (56.96%, P=0.010), or "sore throat" (51.90%, P=0.001) (Supplementary Table S6, available at https://weekly.chinacdc.cn/).

## **Co-Infection Status**

Co-infection was frequent in HPIVs infections, with co-infection rate of 22.12% (175/791). Dual co-infections were the most common (132/175, 75.43%), followed by triple co-infections (26/175, 14.86%) and quadruple or higher co-infections (17/175, 9.71%). The top three pathogens co-infected with HPIVs were HRV (69/791, 8.72%), HRSV (43/791, 5.44%), and HAdV (40/791, 5.06%). However, the rates of co-infection between HPIVs and associated viruses varied by year (*P*=0.04). The highest co-infection rate with

HPIVs was in 2019 (25/78, 32.05%), with HCoV (9/25, 36%) being the most common co-infected virus. The lowest co-infection rate was in 2020 (26/146, 17.81%), with HAdV (10/26, 38.46%) as the predominant co-infected virus. HRV was the primary co-infected virus in 2021 (26/60, 43.33%) and 2022 (25/49, 51.02%). During January–June 2023, both HRV (5/15, 33.33%) and IFV (5/15, 33.33%) were the major co-infected viruses (Table 4).

The co-infection rate among SARIs caused by HPIVs was 26.9% (132/490), which was not different from ARIs (P=0.050). Dual co-infections were predominant in HPIVs-related infections.

The most common viruses co-infected with HPIVs in ARIs, SARIs, and mild ARIs were HRV, HRSV, and HADV. The association between combined viral infections and increased disease severity was not observed.

TABLE 3. Clinical characteristics of HPIVs infections.

	HPIVs inf	ection in 6,442	2 ARIs	4 types o	f distributio	n in 357 HPI\	/s-positive	ARIs	HPIVs i	nfections sta	tus
Symptoms	HPIVs- positive	HPIVs- negative	P	HPIV1	HPIV2	HPIV3	HPIV4	P	Mono- HPIVs	Co- pathogens	P
	( <i>n</i> =357)	( <i>n</i> =6,085)		( <i>n</i> =108)	(n=44)	( <i>n</i> =197)	( <i>n</i> =8)		( <i>n</i> =207)	( <i>n</i> =150)	
Fever	312 (87.39)	4,904 (80.59)	0.002	98 (90.74)	43 (97.73)	164 (83.25)	7 (87.50)	0.024	175 (84.54)	137 (91.33)	0.081
Cough	298 (83.47)	4,762 (78.26)	0.023	90 (83.33)	32 (72.73)	169 (85.79)	7 (87.50)	0.204	180 (86.96)	118 (78.67)	0.053
Sore throat	119 (33.33)	1,243 (20.43)	<0.001	30 (27.78)	24 (54.55)	61 (30.96)	4 (50.00)	0.008	60 (28.99)	59 (39.33)	0.053
Rhinorrhoea	114 (31.93)	2,128 (34.97)	0.045	30 (27.78)	14 (31.82)	94 (47.72)	6 (75.00)	<0.001	80 (38.65)	64 (42.67)	0.513
Phlegm	178 (49.86)	2,410 (39.61)	<0.001	64 (59.26)	32 (72.73)	79 (40.10)	3 (37.50)	<0.001	95 (45.89)	83 (55.33)	0.098
Chest pain	2 (0.56)	70 (1.15)	0.437	0 (0)	1 (2.27)	1 (0.51)	0 (0)	0.361	2 (0.97)	0 (0)	0.511
Dyspnea	19 (5.32)	416 (6.84)	0.317	5 (4.63)	3 (6.82)	10 (5.08)	1 (12.50)	0.570	12 (5.80)	7 (4.67)	0.818
Headache	52 (14.57)	437 (7.18)	<0.001	15 (13.89)	29 (65.91)	7 (3.55)	1 (12.50)	<0.001	2 (0.97)	50 (33.33)	<0.001
Abdominal pain	12 (3.36)	195 (3.20)	0.993	5 (4.63)	1 (2.27)	5 (2.54)	1 (12.50)	0.281	8 (3.86)	4 (2.67)	0.747
Diarrhea	26 (7.28)	263 (4.32)	0.013	4 (3.70)	1 (2.27)	21 (10.66)	0 (0.00)	0.074	13 (6.28)	13 (8.67)	0.516

Abbreviation: HPIVs=human parainfluenza viruses; ARIs=acute respiratory infections.

TABLE 4. Co-infection status of HPIVs infections in China from 2019 to 2023.

<b>T</b> !	Time UDD/- teftime			8 respiratory viruses co-infected with HPIVs					Co-infection status			
ı ime	Time HPIVs infections		HAdV	HRV	HMPV	HEV	HBoV	HCOV	IFV	Dual	Triple and more	Total [n, (%)]
2019	78	0	6	7	1	6	2	9*	1	19	6	25 (32.05)
2020	146	5	10*	6	1	7	6	4	1	17	9	26 (17.81)
2021	311	24	13	26*	4	13	13	0	1	43	17	60 (19.29)
2022	208	11	9	25*	3	4	3	2	8	40	9	49 (23.56)
2023 (Jan-Jun)	48	3	2	5*	0	0	2	0	5*	13	2	15 (31.25)
Total	791	43	40	69*	9	30	26	15	16	132	43	175 (22.12)

Abbreviation: HPIVs=human parainfluenza viruses; HRSV=respiratory syncytial virus; HAdV=human adenovirus; HRV=humanrhinovirus; HMPV=human metapneumovirus; HEV=human enterovirus; HBoV=humanbocavirus; HCoV=human coronavirus; IFV=influenzavirus.

<sup>\*</sup> represents the most common co-infected virus per year.

## **CONCLUSION**

The recent surge of ARIs in children within China has garnered international concern (9), epidemic trends of HPIVs were examined in this study. The current analysis found an overall HPIVs detection rate of 6.15% in ARIs and 5.40% in SARIs, aligning with the results of earlier research (10-11). HPIVs ranked third among the most commonly identified respiratory pathogens, following HRSV and HRV. Among four HPIVs serotypes, HPIV3 and HPIV1 were the most frequently observed strains, suggesting these two serotypes chiefly influence HPIVs epidemiology in China. Interestingly, the study also noted HPIV2 as prevalent in mild ARIs, a finding which warrants further verification. Additionally, nonpharmacological interventions (NPIs) might influence HPIVs epidemic patterns. Notably, fewer HPIVs infections were recorded in the first half of 2020; however, cases surged in the latter half of the year following the easing of NPIs.

This study identified notable alterations in the seasonal patterns of HPIV3 and HPIV1 following the COVID-19 pandemic. Specifically, the 2020 epidemic peak for HPIV3 shifted to the autumn and winter months, a three-month postponement from its usual summer peak observed in 2019. Post-2020, HPIV3 levels stabilized. In 2021, two successive epidemic peaks of HPIV1 were recorded in July and October, which directly succeeded the peak of HPIV3. A comparable delay in HPIVs seasonal peaks has been documented in South Africa post-pandemic (12). These timing shifts may be attributed to the enactment of NPIs, changes in societal behaviors, and the presence of immunity gaps among the population. No conspicuous seasonal trends were identified for HPIV2 and HPIV4, and their epidemiological patterns remain largely undefined within China, underscoring the imperative for ongoing monitoring.

Although co-infections with HPIVs were common, their incidence varied annually, peaking at 32.05% in 2019 and dropping to 17.81% in 2020. The decrease may be attributed to the reduced presence of common co-infecting viruses due to NPIs. HRV was frequently identified as a co-infecting agent, showing a positive correlation with HPIVs (13). The high occurrence of HPIVs and HRV co-infections could be linked to their transmission modes via direct contact, which might have been facilitated by lower adherence to preventive measures during NPIs. The co-infection rate of HPIVs rose significantly following the relaxation of NPIs.

Given that co-infections often lead to severe conditions (14), the significance of addressing co-infections in HPIVs cases should be underscored.

HPIVs are distinguished by their infection age, clinical manifestations, and geographic prevalence (15). This study found that HPIVs predominantly infected children under five years old. However, HPIV2 was more frequently diagnosed in both children and adolescents, whereas HPIV4 was often detected in the elderly, confirming earlier reports (12). Furthermore, although HPIVs infections are known to cause a broad spectrum of clinical symptoms, certain serotypespecific manifestations such as fever, phlegm, and headache have been particularly associated with HPIV2 infections. The distribution of HPIVs across China also exhibited regional variability; for instance, HPIV3 was more commonly found in the southern PLADs, whereas HPIV2 and HPIV4 showed higher occurrences in the northern areas. Nevertheless, these observed serotype-specific epidemiological trends necessitate corroboration through extended surveillance data.

In conclusion, analysis of five years of surveillance data has unveiled the epidemic patterns of HPIVs in China. This has enhanced comprehension of the epidemiological and clinical features of HPIVs and has implications for the prevention and management of HPIVs-related respiratory diseases in China.

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# **SUPPLEMENTARY MATERIAL**

SUPPLEMENTARY TABLE S1. The population distribution by age and sex among ARIs with HPIV1–HPIV4 infections from January 2019 to June 2023.

01	HPIV1	HPIV2	HPIV3	HPIV4	HPIVs
Characteristic	n=215	<i>n</i> =101	n=432	n=49	<i>n</i> =791
Age (years), N (%)					
0–4	169 (27.66)	52 (8.51)	361 (59.08)	33 (5.40)	611
5–17	30 (22.72)	43 (32.57)	52 (39.39)	8 (6.06)	132
18–39	1 (33.33)	0 (0)	1 (33.33)	1 (33.33)	3
40–64	3 (21.42)	2 (14.29)	8 (57.14)	1 (7.14)	14
65+	12 (38.71)	4 (12.90)	10 (32.25)	6 (19.35)	31
Sex, N (%)					
Male	121 (25.80)	59 (12.58)	262 (55.86)	30 (6.39)	469
Female	94 (29.19)	42 (13.04)	170 (52.80)	19 (5.90)	322

Note: % is the ratio of the composition of the rows.

Abbreviation: HPIVs=human parainfluenza viruses; ARIs=acute respiratory infections.

SUPPLEMENTARY TABLE S2. The population distribution by age and sex among patients with SARIs caused by HPIV1 to HPIV4, from January 2019 to June 2023.

Observe at a start a	HPIV1	HPIV2	HPIV3	HPIV4	HPIVs
Characteristic	<i>n</i> =134	n=44	n=285	<i>n</i> =32	<i>n</i> =490
Age (years), N (%)					
0–4	109 (26.98)	26 (6.44)	252 (62.38)	21 (5.20)	404
5–17	13 (27.08)	13 (27.08)	17 (35.42)	5 (10.42)	48
18–39	0 (0)	0 (0)	0 (0)	0 (0)	0
40–64	1 (9.09)	2 (18.18)	7 (63.64)	1 (9.09)	11
65+	11 (40.74)	3 (11.11)	9 (33.33)	5 (18.52)	27
Sex, N (%)					
Male	74 (25.52)	28 (9.66)	171 (58.97)	19 (6.55)	290
Female	60 (0.30)	16 (0.08)	114 (0.57)	13 (0.07)	200

Note: % is the ratio of the composition of the rows.

Abbreviation: HPIVs=human parainfluenza viruses; SARIs=severe acute respiratory infections.

SUPPLEMENTARY TABLE S3. Geographic distribution and annual detection positive rate of HPIVs.

Characteristic	HPIV1	HPIV2	HPIV3	HPIV4	HPIVs	ARIs
Region, N (%)						
Northern	175 (1.69)	97 (0.93)	332 (3.20)	45 (0.43)	646 (6.22)	10,378
Southern	40 (1.61)	4 (0.16)	100 (4.02)	4 (0.16)	145 (5.83)	2,488
Р	0.784	<0.001	0.041	0.047	0.470	
Time, N (%)						
2019	18 (1.17)	9 (0.58)	39 (2.53)	12 (0.78)	78 (5.07)	1,539
2020	29 (1.93)	6 (0.40)	109 (7.24)	4 (0.27)	146 (9.70)	1,505
2021	141 (3.46)	35 (0.86)	116 (2.85)	20 (0.49)	311 (7.63)	4,075
2022	21 (0.51)	51 (1.24)	133 (3.23)	6 (0.15)	208 (5.04)	4,124
P	<0.001	0.010	<0.001	0.002	<0.001	

Note: % represents the positive detection rate (N/ARIs).

Abbreviation: HPIVs=human parainfluenza viruses; ARIs=acute respiratory infections.

SUPPLEMENTARY TABLE S4. Differences in the population distribution between HPIVs mono- and co-infections.

01	Mono-infections	Co-infections	
Characteristic	n=616	n=175	P
M (Q <sub>1</sub> , Q <sub>3</sub> )	3 (1, 4)	2 (1, 4)	0.163
Age (years), N (%)			0.286
0–4	472 (76.62)	139 (79.43)	
5–17	102 (16.56)	30 (17.14)	
18–39	2 (0.32)	1 (0.57)	
40–64	14 (2.27)	0 (0)	
65+	26 (4.22)	5 (2.86)	
Sex, N (%)			0.515
Male	361 (58.60)	108 (61.71)	
Female	255 (41.40)	67 (38.29)	

Abbreviation: HPIVs=human parainfluenza viruses.

SUPPLEMENTARY TABLE S5. Demographics of HPIVs infections among SARIs (January 2019 to June 2023).

	HPIV1	HPIV2	HPIV3	HPIV4	_	HPIVs
Characteristic	n=134	n=44	<i>n</i> =285	n=32	P	n=490
M (Q <sub>1</sub> , Q <sub>3</sub> )	2 (1, 4)	4 (2.75, 7.25)	2 (0, 3)	3 (2, 5.25)	<0.001	2 (1, 3)
Age (years), N (%)					<0.001	
0–4	109 (81.34)	26 (59.09)	252 (88.42)	21 (65.63)		404 (82.45)
5–17	13 (9.70)	13 (29.55)	17 (5.96)	5 (15.63)		48 (9.80)
18–39	0 (0)	0 (0)	0 (0)	0 (0)		0 (0)
40–64	1 (0.75)	2 (4.55)	7 (2.46)	1 (3.13)		11 (2.24)
65+	11 (8.21)	3 (6.82)	9 (3.16)	5 (15.63)		27 (5.51)
Sex, N (%)					0.607	
Male	74 (55.22)	28 (63.64)	171 (60.00)	19 (59.38)		290 (59.18)
Female	60 (44.78)	16 (36.36)	114 (40.00)	13 (40.63)		200 (40.82)

Abbreviation: HPIVs=human parainfluenza viruses; SARIs=severe acute respiratory infections.

SUPPLEMENTARY TABLE S6. Clinical characteristics of HPIVs infections among patients with SARIs and non-SARIs.

Summana	357 HPI	/s infections	- P
Symptoms	SARIs (n=278)	Non-SARIs (n=79)	- γ
Fever	236 (84.89)	76 (96.20)	0.013
Cough	247 (88.85)	51 (64.56)	0.001
Sore throat	78 (28.06)	41 (51.90)	0.001
Rhinorrhea	123 (44.24)	21 (26.58)	0.007
Phlegm	123 (44.24)	55 (69.62)	0.001
Chest pain	2 (0.72)	0 (0)	0.053
Dyspnea	18 (6.47)	1 (1.27)	0.087
Headache	7 (2.52)	45 (56.96)	0.001
Abdominal pain	11 (3.96)	1 (1.27)	0.477
Diarrhea	26 (9.35)	0 (0)	0.010

 $Abbreviation: HPIVs = human\ parainfluenza\ viruses;\ SARIs = severe\ acute\ respiratory\ infections.$