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Molecular Typing and Epidemiology Profiles of Human Adenovirus Infection among Hospitalized Patients with Severe Acute Respiratory Infection in Huzhou, China --Manuscript Draft--

Manuscript Number:	PONE-D-21-05692
Article Type:	Research Article
Full Title:	Molecular Typing and Epidemiology Profiles of Human Adenovirus Infection among Hospitalized Patients with Severe Acute Respiratory Infection in Huzhou, China
Short Title:	Human Adenovirus Infection in Hospitalized Patients in China
Corresponding Author:	Lei Ji Huzhou Center for Disease Control and Prevention Huzhou, CHINA
Keywords:	Human adenovirus; Respiratory tract infection; Epidemiology
Abstract:	<p>Background: Severe acute respiratory infections (SARI) threaten human health and cause a large number of hospitalized patients every year. However, as one of the most common pathogen that cause acute respiratory tract infection, the molecular epidemiological information relating to HAdV among patients with SARI is limited. Here, we evaluate the epidemiological and molecular characteristics of HAdV infections among hospitalized patients with SARI from January 2017 to December 2019 in Huzhou, China.</p> <p>Methods: From January 2017 to December 2019, a total of 657 nasopharyngeal swabs collected from inpatients with SARI were screened for HAdV and other common respiratory viruses by multiplex real-time PCR. All samples that tested positive for HAdV were further typed by sequencing partial sequences of hexon gene. Genotypes of HAdV were confirmed by phylogenetic analysis. Epidemiological data were analyzed using Microsoft Excel 2010 and service solutions (SPSS) 21.0 software.</p> <p>Results: 251 (38.20%) samples were positive for at least one respiratory virus. HAdV was the second common viral pathogen detected, with a detection rate of 7.08%. Infection with HAdV was found in all age groups tested (0 ~, 2 ~, 5 ~, 15 ~, 50 ~, 65 ~). Children under 15 years old accounted for 84.62% (44/52) of the infections. Higher activity of HAdV infection could be seen in spring-early autumn season. 7 different types of HAdV belonging to 4 species (HAdV-A, B, C, E) were identified in hospitalized SARI cases, with HAdV-B3 as the most prevalent HAdV types, followed by HAdV-B7 and HAdV-E4. HAdV-B3 was the most frequently detected genotype in 2017 and 2019, accounting for 75.00% (9/12) and 63.64(7/11) of typed HAdV infections in 2017 and 2019 respectively. No predominant strain was responsible for HAdV infections in 2018, although HAdV-B7 (28.57%, 2/7) and HAdV-C1 (28.57%, 2/7) were the major causative genotypes.</p> <p>Conclusions: This study revealed the prevalence and the molecular epidemiological characteristics of HAdV infections among hospitalized patients with SARI in Huzhou from January 2017 to December 2019. The HAdV prevalence is related to age and season. As the most prevalent HAdV types, HAdV-B3 was co-circulating with other types and presented an alternate prevalence pattern.</p>
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The sequences of the HAdV strains obtained in this study were deposited in the GenBank under the accession numbers MW594169-MW594198

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Title:

Molecular Typing and Epidemiology Profiles of Human Adenovirus Infection among Hospitalized Patients with Severe Acute Respiratory Infection in Huzhou, China

Short title:

Human Adenovirus Infection in Hospitalized Patients in China

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Abstract

Background: Severe acute respiratory infections (SARI) threaten human health and cause a large number of hospitalized patients every year. However, as one of the most common pathogen that cause acute respiratory tract infection, the molecular epidemiological information relating to HAdV among patients with SARI is limited. Here, we evaluate the epidemiological and molecular characteristics of HAdV infections among hospitalized patients with SARI from January 2017 to December 2019 in Huzhou, China.

Methods: From January 2017 to December 2019, a total of 657 nasopharyngeal swabs collected from inpatients with SARI were screened for HAdV and other common respiratory viruses by multiplex real-time PCR. All samples that tested positive for HAdV were further typed by sequencing partial sequences of hexon gene. Genotypes of HAdV were confirmed by phylogenetic analysis. Epidemiological data were analyzed using Microsoft Excel 2010 and service solutions (SPSS) 21.0 software.

Results: 251 (38.20%) samples were positive for at least one respiratory virus. HAdV was the second common viral pathogen detected, with a detection rate of 7.08%. Infection with HAdV was found in all age groups tested (0~, 2~, 5~, 15~, 50~, 65~). Children under 15 years old accounted for 84.62% (44/52) of the infections. Higher activity of HAdV infection could be seen in spring-early autumn season. 7 different types of HAdV belonging to 4 species (HAdV-A, B, C, E) were identified in hospitalized SARI cases, with HAdV-B3 as the most prevalent HAdV types, followed by HAdV-B7 and HAdV-E4. HAdV-B3 was the most frequently detected genotype in 2017 and 2019, accounting for 75.00% (9/12) and 63.64% (7/11) of typed HAdV infections in 2017 and 2019 respectively. No predominant strain was responsible for HAdV infections in 2018, although HAdV-B7 (28.57%, 2/7) and HAdV-C1 (28.57%, 2/7) were the major causative genotypes.

Conclusions: This study revealed the prevalence and the molecular epidemiological characteristics of HAdV infections among hospitalized patients with SARI in Huzhou from January 2017 to December 2019. The HAdV prevalence is related to age and season. As the most prevalent HAdV types, HAdV-B3 was co-circulating with other types and presented an alternate prevalence pattern.

Keywords: Human adenovirus ; Respiratory tract infection ; Epidemiology

Background

Human adenoviruses (HAdVs) are non-enveloped, double-stranded DNA viruses belonging to the genus Mastadenovirus of the Adenoviridae family[1]. HAdVs have been recognized as pathogens that cause a broad spectrum of diseases, including respiratory illness, keratoconjunctivitis, gastroenteritis, cystitis and meningoencephalitis[2, 3]. They are associated with sporadic infection, as well as with community and institutional outbreaks. As a significant causative agent of respiratory tract illnesses, HAdV accounts for at least 5 to 10% of pediatric and 1 to 7% of adult respiratory tract infections (RTIs)[4, 5].

There are currently seven different HAdV species (HAdV-A through HAdV-G), and to date, 51 serotypes and over 70 genotypes have been identified based on serology, phylogenetic analyses and whole genomic sequencing (<http://hadvvg.gmu.edu/>). Different types of HAdVs display

87 different tissue tropisms that correlate with clinical manifestations of infection. The HAdV types
88 most commonly associated with respiratory infection belong to HAdV species B
89 (HAdV-3, HAdV-7, HAdV-11, HAdV-14, HAdV-21), HAdV species C (HAdV-C1, -C2, -C5, and
90 -C6) and HAdV species E (HAdV-4)[3].

91 The predominant types of HAdV circulating at a given time differ among countries or regions and
92 change over time. Replacement of dominant viruses by new strains may occur because
93 transmission of novel strains between countries. During the last decade, outbreaks of respiratory
94 tract infections caused by novel HAdV strains have occurred frequently in many countries
95 including China[6, 7]. Therefore, clarifying the genotype of HAdV currently circulating is
96 essential for epidemiological surveillance and a better understanding of the epidemic pattern of
97 HAdV infection. At present, China has not yet established a national HAdV surveillance system.
98 Although data about HAdV associated with respiratory infection in China can be found in several
99 studies, most studies are performed with specific groups, especially for children[8-13]. There is a
100 lack of epidemiological analyses of HAdV associated respiratory infection among patients in all
101 age groups in China. The aim of this study was to evaluate the epidemiological and molecular
102 characteristics of HAdV infections among hospitalized patients with severe acute respiratory
103 infection (SARI) from January 2017 to December 2019 in Huzhou, a medium-sized city located in
104 eastern China.

105

106 **Materials and methods**

107 **Ethics statement**

108 This study was part of the national SARI surveillance program and was approved by the human
109 research ethics committee of Huzhou Center for Disease Control and Prevention. The only human
110 materials used were nasopharyngeal swabs collected from patients for routine detection. Data
111 records and collected clinical specimens were deidentified and anonymous. Oral informed
112 consents were obtained from each participant.

113 **Patients and specimens**

114 During the influenza A H1N1 epidemic in 2009, a surveillance system for SARI was established to
115 monitor influenza infection ~~in these cases in china~~. As local SARI surveillance sentinel hospital,
116 the First People's Hospital of Huzhou was responsible for sample collection from ~~surveillance~~
117 ~~cases~~. The inclusion criteria for hospitalized SARI cases were as follows: the onset of the disease
118 has a history of fever(> 38°C), accompanied by cough, and the onset does not exceed 10 days.
119 Nasopharyngeal swabs were freshly collected and sent to Huzhou Center for Disease Control and
120 Prevention for routine detection. All the specimens were stored at - 80 °C until further processing.
121 Demographic and clinical data were obtained from the hospital's database.

122 **Detection of HAdV and other common respiratory viruses**

123 Total viral nucleic acids (DNA and RNA) were extracted from 200 µL of each specimen using
124 TIANLONG Ex Viral DNA/RNA Kit (TIANLONG Biotech, Xi'an, China) according to the
125 manufacturer's instructions. Multiplex real-time PCR kit (BioGerm, Shanghai, China) was used to
126 detect HAdV and other common respiratory virus pathogens, including Human Influenza
127 virus(HIFV), Human respiratory syncytial virus(HRSV), Human rhinovirus(HRV), Human
128 bocavirus(HBOV), Human metapneumovirus(HMPV), Human Parainfluenza Virus(HPIV) type
129 1-4 and Human coronavirus (HCoV). The qPCR cycling program was as follows: 50 °C for 10



130 min, 95 °C for 5min, followed by 40 cycles of 95 °C for 10 s, and 55 °C for 40 s. Samples with a
 131 cycle threshold (Ct) < 35 were regarded as positive.

132 **HAdV genotyping**

133 HAdV-positive samples were further molecularly typed by nested PCR amplification and
 134 sequencing of HAdV hexon gene hyper-variable regions 1–6 (HVR1–6) as described
 135 previously[14]. Primer set AdhexF1 (nt 19135–19160;5'-TICTTTGACATICGIGGIGTICTIGA-3')
 136 and AdhexR1 (nt 20009–20030;5'-CTGTCIACIGCCTGRTTCCACA-3') were used for first-round
 137 amplification; a second-round PCR was performed using primer set AdhexF2 (nt 19165–19187;
 138 5'-GGYCCYAGYTTYAARCCCTAYTC-3') and AdhexR2 (nt 19960–19985;
 139 5'-GGTTCTGTICCCAGAGARTCIAGCA-3') if insufficient DNA was amplified from the first
 140 reaction for sequencing. The PCR products were visualized by electrophoresis and sent to TaKaRa
 141 Biotechnology (Dalian, China) for further purification and sequencing.

142 **Phylogenetic analysis**

143 Partial nucleotide sequences of hexon gene obtained in this study were compared with the NCBI
 144 GenBank database (<http://www.ncbi.nlm.nih.gov>) by using online BLAST tools to preliminarily
 145 determine the genotype. Multiple sequence alignment and phylogenetic analysis were conducted
 146 using MEGA software version 6.06. The phylogenetic tree was generated using the
 147 neighbor-joining method and bootstrap analysis was performed with 1000 replications.

148 **Statistical analysis**

149 Epidemiological data were analyzed using Microsoft Excel 2010 and service solutions (SPSS)
 150 21.0 software. Statistical differences were determined using the Chi-square test and P-values
 151 <0.05 were considered to represent a statistically significant difference.

152 **Accession numbers**

153 The partial hexon gene sequences obtained in this study have been deposited in GenBank under
 154 the accession numbers MW594169-MW594198.

155


156 **Results**

157 **Characteristics of the SARI cases and the Viral infection profiles**

158 From January 2017 to December 2019, a total of 657 specimens (191 in 2017, 204 in 2018 and
 159 262 in 2019) were collected from inpatients with SARI during the study period. ~~Among those~~
 160 ~~SARI cases,~~ 361 (54.95%) were male and 296 (45.05%) were female, ~~the~~ age range was from 1
 161 month to 86 years old with 590 (89.80%) cases ~~were children younger than 15 years old.~~

162 The viral infection profiles are shown in Table 1. Overall, 251 (38.20%) samples were positive for
 163 at least one respiratory virus, the detection rate of respiratory virus was 45.54% (87/191) in 2017,
 164 36.27% (74/204) in 2018 and 34.35% (90/262) in 2019. During the study period, the most
 165 commonly detected viral pathogen in SARI cases was RSV, with a prevalence rate of 10.65%
 166 (70/657), followed by HAdV (7.91%, 52/657) and HIFV (6.09%, 40/657). HMPV was detected in
 167 30 patients (4.57%), HPIV was detected in 24 (3.65%), HBOV was detected in 21 (3.20%), HRV
 168 was detected in 11 (1.67%), and HCoV was detected in 3 patients (0.46%).

169 **Table 1 Viral infection profiles in hospitalized patients with SARI in Huzhou, 2017–2019**

Years	SARI cases	Any viral etiology	Viral infection profiles 							
			HRS V	HAd V	HIFV	HMPV	HPIV	HBOV	HRV	HCo V

2017	191	87	32	18	8	8	10	8	3	0
2018	204	74	17	8	21	14	5	7	2	0
2019	262	90	21	26	11	8	9	6	6	3
Total	657	251	70	52	40	30	24	21	11	3

170

171 **Epidemiology of HAdV**

172 ~~During our study period, HAdV was the second common viral pathogen detected in SARI cases,~~
173 ~~with a detection rate of 7.91% (52/657). As shown in Table 2,~~ among the 52 HAdV-infected
174 patients, 31 (58.33%) were male and 21 (41.67%) were female. No significant difference was
175 observed in males and females in the HAdV-infected cases ($P = 0.481$). Infection with HAdV was
176 found in all age groups tested (0~, 2~, 5~, 15~, 50~, 65~). Children under 15 years old
177 accounted for 84.62% (44/52) of the infections. There were no significant differences in HAdV
178 detection rates among different age groups ($P = 0.467$). The highest detection rate was in the 2-≤5
179 year age group (9.44%), followed by 5-≤15 years (9.13%), 15-≤50 years (7.14%), 0-<2 years
180 (5.05%), 50-<65 years (3.13%) and ≥65 years (2.86%).

181 **Table 2 HAdV-positive in hospitalized patients of different ages and gender with SARI**

Variable	Tested cases N (percentage)	SARI cases N (percentage)	HAdV-positive cases N (percentage)	HAdV-negative cases N (percentage)	Positive rate	χ^2	P
Gender						0.497	0.481
Male	361 (53.40)		31 (58.33)	330(54.55)	8.59%		
Female	296 (46.60)		21 (41.67)	275(45.45)	7.09%		
Age (years)						0.431	0.476
0~	99 (15.07)		5 (9.62)	94 (16.55)	5.05%		
2~	180 (27.40)		17 (32.69)	163 (23.51)	9.44%		
5~	241 (36.68)		22 (42.31)	219 (43.05)	9.13%		
15~	70 (10.65)		5 (9.61)	65 (6.95)	7.14%		
50~	32 (4.87)		1 (1.92)	31 (4.64)	3.13%		
65~	35 (5.33)		1 (1.92)	34 (5.30)	2.86%		
Total	657		52	605	7.91%		

182

183 **Fig. 1 Monthly distribution of HAdV infections from January 2017 to December 2019**

184

185 HAdV detection rate varied from year to year, from 9.42% (18/191) in 2017, 3.92% (8/204) in
186 2018 to 9.92% (26/262) in 2019 (Table 1). The monthly distribution of HAdV infections is shown
187 in Fig. 1. HAdV was detected in every month throughout the study period except January. Higher
188 activity of HAdV infection could be seen from spring to early autumn (April to September), and
189 the detection rate in September reached a peak of 28.17%. In contrast, lower activity of HAdV
190 infection were observed during late autumn to winter (from October to February), when the
191 average detection rate was only 2.37%.

192 Additionally, 13.46-% (n = 7) of the 52 HAdV-infected cases were co-detected with other
193 respiratory pathogens. RSV (n = 3) was the most frequently co-detected virus. HPIV (n = 2),
194 HMPV (n = 1) and HRV (n = 1) were also found to be co-infected with HAdV.

195 **HAdV genotyping and phylogenetic analysis**

196 Of the 52 HAdV--positive samples confirmed by real-time RT-PCR, 30 samples were successfully
197 sequenced and genotyped by nested-PCR. Phylogenetic analysis based on partial hexon sequences
198 indicated that 4 species (A, B, C, E) of HAdV, including 7 different types were identified
199 throughout the study period, ~~see Fig. 2~~. HAdV-B3 (n = 17, 56.67 %) was the most prevalent HAdV
200 types, followed by HAdV-B7 (n = 5, ~~16.367~~%) and HAdV-E4 (n = 3, 10.00 %). HAdV-C1 (n = 2,
201 6.67 %), HAdV-C2(n =1,3.33 %), HAdV-B21(n =1,3.33 %) and HAdV-B55(n =1,3.33 %)were
202 also detected. The genotype distribution of HAdV infections in each month is shown in Fig. 3. The
203 predominant genotypes of HAdV during our study period varied according to surveillance year.
204 Overall, HAdV-B3 was the most frequently detected genotype in 2017and 2019, accounting for
205 75.00% (9/12) and 63.64(7/11) of typed HAdV infections ~~in 2017 and 2019~~, respectively. ~~5~~
206 different types were detected in 2018, including HAdV-B7 (n = 2), HAdV-C1 (n = 2), HAdV-B3
207 (n = 1), HAdV-B55 (n = 1) and HAdV-C2 (n = 1).No predominant strain was responsible for
208 HAdV infections in 2018, although HAdV-B7 (28.57%, 2/7) and HAdV-C1 (28.57%, 2/7) were
209 the major causative genotypes.

210

211 **Fig. 2 Phylogenetic analyses based on partial hexon sequences of HAdV strains.** The trees
212 were generated using the neighbor-joining method, validated by 1000 bootstrap replicates.
213 Bootstrap values $\geq 70\%$ are shown on the branch. HAdV sequences identified in this study are
214 indicated by closed circles.

215 **Fig. 3 Distribution of HAdV genotypes detected according to month.**

216

217 **Discussion**

218 SARI is one of the most common diseases in human and the leading cause of hospitalization in
219 children worldwide[15, 16]. Because the early clinical symptoms of respiratory infections caused
220 by viruses are similar, and the imaging findings lack specificity, pathogen detection is very
221 important in clinical diagnosis and epidemiological monitoring. The present study was carried out
222 from January 2017 to December 2019 among hospitalized patients with SARI in Huzhou, China.
223 During the study period, a total of 657 hospitalized SARI cases were enrolled, of which 80.57%
224 were children under 15 years of age. ~~It suggests~~ that SARI is still an important factor affecting the
225 health of local children. ~~38.20%~~ of hospitalized SARI cases in our study exhibited at least one
226 respiratory virus, which was consistent with previous reports from China (33.44%-41.50%)[17,
227 18]and other countries (37.57%-41.8%)[19, 20].

228 HAdV was the second common viral pathogen detected, with a detection rate of 7.08%, which is
229 lower than the finding in SARI cases of hospitalized children in Beijing (11.90%) and Shanghai
230 (14.70%)[21]. Previous studies have indicated that HAdV is the major pathogen that causes
231 respiratory tract infections in children, especially for children younger than 5 years[8, 10]. As
232 expected, we found that HAdV infection mainly occurred in children under 15 years of age
233 (84.62%), and the detection rate reached a peak (9.44%) in children aged 2 to ≤ 5 years.

234 Previous studies have shown that the epidemic peak seasons of HAdV-associated respiratory
235 infections varies in different parts of China, and even in different monitoring years in the same
236 region. Our study revealed that HAdV showed higher activity in the relatively high temperature
237 seasons (spring to early autumn), which is similar to what has been found in Beijing (Northern

238 China)[8]and Guangzhou(Southern China)[13], where HAdV infections occurred throughout the
239 year with the highest prevalence in the summer. However, this finding is discordant with other
240 studies conducted in Northern China that have reported seasonal peaks for HAdV infections in
241 winter and spring[9, 12]. It is worth mentioning that the surveillance period of the
242 above-mentioned studies conducted in different regions of China varies, and the predominant
243 HAdV types circulated are also different. A recent study from Hunan indicates that different
244 HAdV types showed a different seasonal distribution patterns: HAdV-3 was the predominant type
245 of HAdV infection during summer, while HAdV-7 had the highest detection rate during spring[11].
246 Based on the above research, we speculate that the discrepant seasonal peak for HAdV infections
247 are not only related to regional differences, but also related to the major types of HAdV circulating
248 locally.

249 Globally the types most commonly associated with respiratory syndromes belong to HAdV
250 species B, C or E. Many studies have reported that HAdV-B3, HAdV-B7 and HAdV-C2 are the
251 most prevalent types in China, but the predominant type distribution vary among different regions
252 and change over time. For example, most of HAdV-positive cases were caused by HAdV-B3 from
253 2012 to 2013 in Southern China[13], while HAdV-B7 dominated in Northern China during the
254 same study period[10]. However, recent reports indicated that the most predominant types have
255 changed into HAdV-B3 and HAdV-C2 in some Northern cities of China in 2017-2018[8, 9].
256 Throughout the present study period, 7 different types of HAdV belonging to 4 species (HAdV-A,
257 B, C, E) were identified in hospitalized SARI cases, with HAdV-B3 as the most prevalent HAdV
258 types, followed by HAdV-B7 and HAdV-E4. Our monitoring data showed that no type of HAdV
259 presented absolutely predominant during HAdV epidemic seasons, HAdV-B3 was co-circulating
260 with other types and presented an alternate prevalence pattern. Overall, HAdV-B3 was the most
261 frequently detected genotype in 2017. No predominant strain was responsible for HAdV infections
262 in 2018, with HAdV-B7 and HAdV-C1 as the major causative genotypes. HAdV-B3 re-emerged as
263 the predominant genotype in 2019. Similar epidemic pattern were observed in a prolonged
264 surveillance study conducted in southeastern China, where HAdV-7 and HAdV-3 alternate as the
265 predominant genotypes causing pediatric pneumonia[22]. It is worth noting that in 2017 and 2019,
266 when HAdV-3 presented as the predominant type detected, the detected rate of HAdV was
267 significantly higher than that in 2017(9.42% in 2017, 3.92% in 2018 and 9.92%in 2019). The
268 reasons need to be further explored. During HAdV infection, neutralizing antibodies are formed
269 against the epitopes located in the hyper variable regions (HVRs) of the hexon protein. Just
270 recently, Haque E et al. explore the variation in HVRs of hexon among globally distributed strains
271 of HAdV-3[23]. They found that the HVRs of HAdV-3 strains circulating worldwide were highly
272 heterogeneous and have been mutating continuously since their original isolation and suggested
273 that, this heterogeneity may explain the worldwide increased prevalence of HAdV-3 respiratory
274 infections.

275 Recent HAdV epidemiology studies showed that there was very high co-infection rate between
276 HAdV and other pathogens in respiratory tract infection cases (37.50%-74.85%)[8, 9, 11]. In our
277 study, coinfection of HAdVs and other respiratory viruses was only detected in 13.46 % of the
278 SARI cases. Such discrepant co-infection rate may be caused by the different selection criteria of
279 the research objects and methodological differences.

280 Our study is limited by a single-site setting, small sample size, and especially the partial
281 genotyping of detected HAdVs. Genotyping was only successful for 57.69% (30/52) of HAdV

282 infection cases. Besides, typing of HAdV was merely performed by sequencing of partial hexon
283 gene in the present study, which is hard to find any potential recombination between different
284 types of HAdV strains.

285 **Conclusions**

286 In conclusion, this study revealed the prevalence and molecular epidemiological characteristics of
287 HAdV infections among hospitalized patients with SARI in Huzhou from January 2017 to
288 December 2019. HAdV was the second common viral pathogen detected in SARI cases, with most
289 (84.62%) HAdV-positives cases detected among children < 15 years of age. Higher activity of
290 HAdV infection could be seen in spring -early autumn season. No type of HAdV presented
291 absolutely predominant during HAdV epidemic seasons, HAdV-B3 was co-circulating with other
292 types and presented an alternate prevalence pattern. Our results provide a reliable scientific basis
293 to better understand the role played by HAdVs in SARI cases, and for the prevention and control
294 of HAdV infection.

295 **Authors' contributions**

296 LJ wrote the first draft and did the phylogenetic analysis. XFW and DSX participated in the
297 HAdVs detection. LPC participated in the Genomic amplification for genotyping. GTL did the
298 epidemiological investigation and performed the statistical analysis. All authors read and approved
299 the final manuscript.

300 **Acknowledgements**

301 We thank the staff of the First People's Hospital in Huzhou for collecting the samples.

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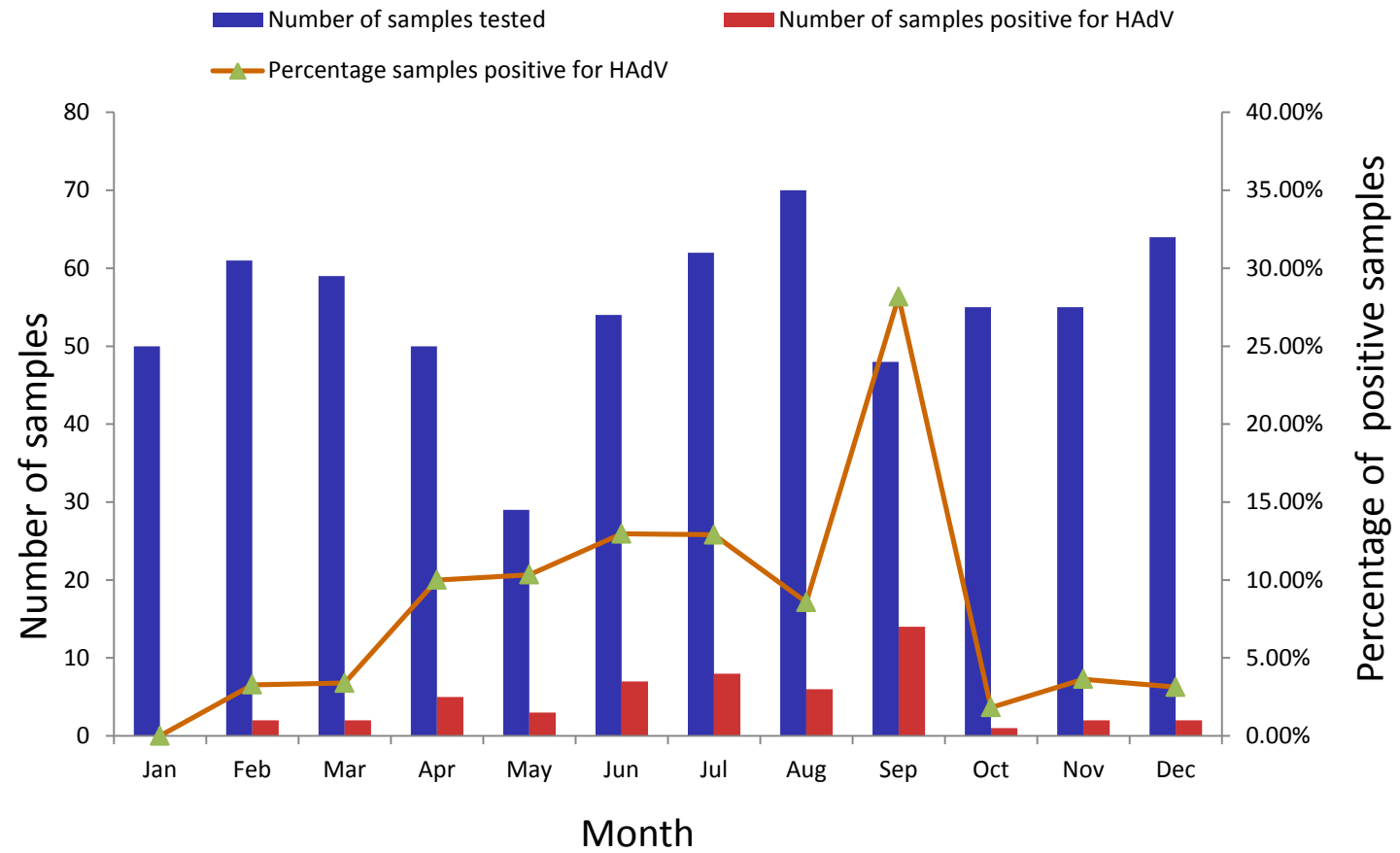


Figure2

