

Febrile Respiratory Illness Associated with Human Adenovirus Type 55 in South Korea Military, 2014–2016¹

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An outbreak of febrile respiratory illness associated with human adenovirus (HAdV) occurred in the South Korea military during the 2014–15 influenza season and thereafter. Molecular typing and phylogenetic analysis of patient samples identified HAdV type 55 as the causative agent. Emergence of this novel HAdV necessitates continued surveillance in military and civilian populations.

Human adenovirus (HAdV) is a common cause of upper respiratory infections ranging from uncomplicated upper respiratory infections to life-threatening pneumonia. Military personnel, especially new recruits, are predisposed to respiratory infections caused by HAdV (1). The substantial effects of HAdV infection in the military have been demonstrated by the marked increase in the incidence of febrile respiratory illness (FRI) in the US military after vaccination against the virus ended in 1999; in turn, the incidence dramatically declined after the vaccine was reintroduced (2).

HAdVs are a group of nonenveloped double-stranded DNA viruses comprising 7 species (A–G) and >50 types (3). HAdV types belonging to species B (HAdV-3, -7, -11, -16, -21) and E (HAdV-4) are commonly associated with respiratory infections in adults, particularly in military personnel (4). Novel types or genomic variants, such as HAdV-14 (3) and HAdV-7 (5), have been implicated in epidemics of severe infection. HAdV-55, another emerging type reported in China, Turkey, Spain, Singapore, and Israel (5,6), has been associated with severe clinical manifestations, which often lead to respiratory failure and death (7,8).

Since fall 2014, we have observed an outbreak of FRI and pneumonia in military personnel in South Korea. In addition to the increased incidence of FRI, patients

experienced severe manifestations. We describe the epidemiologic, clinical, and molecular characteristics of FRI in the South Korea military during October 2014–May 2016.

The Study

We obtained data regarding temporal trends in FRI incidence from military sentinel surveillance, which has been monitoring weekly FRI rates since October 2011. Monthly numbers of patients with pneumonia (inpatients, outpatients, and emergency room patients) were extracted from a computerized data warehouse that stores data from all military hospitals. We identified pneumonia cases by using the International Classification of Diseases and Related Health Problems, 10th Revision, codes J12–J18. The influenza season, which starts in October and ends the following May, was used as a surrogate for the HAdV season in this study. More detailed information on FRI surveillance is available in the online Technical Appendix (<https://wwwnc.cdc.gov/EID/article/23/6/16-1848-Techapp1.pdf>).

The trends in FRI rates showed an unusual surge during the 2014–15 influenza season (Figure 1, panel A). The FRI rate increased for 15 weeks in the 2014–15 season, compared with 10 weeks in the 2012–13 season and 5 weeks in the 2013–14 season. Peak FRI rate in the 2014–15 season (10.4%) was higher than rates in the preceding 2 seasons (4.7% and 7.5%). The numbers of pneumonia cases in 2014–15 and 2015–16 seasons were 3,140 and 3,145 patients, respectively, a 191% increase from the mean number during 3 preceding seasons.

In April 2014, a multiplex real-time PCR for identifying 15 viruses from respiratory specimens was introduced at the Armed Forces Capital Hospital, the only tertiary hospital in the South Korea military health care system (detailed methods in online Technical Appendix). A total of 1,484 nonduplicate specimens were tested by the end of May 2016 (Figure 1, panel B; online Technical Appendix Table 1, Figure). HAdV was identified in 490 (33.0%) of total specimens, and it accounted for 79.7% (282/354) and 53.2% (150/282) of positive results in the 2014–15 and 2015–16 seasons, respectively. Increased HAdV activity was observed from December until the following May.

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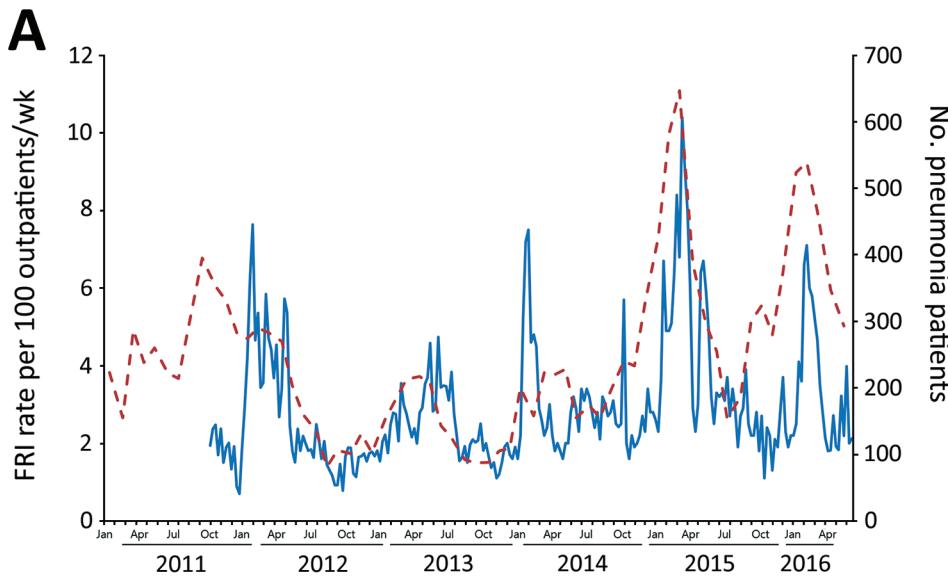
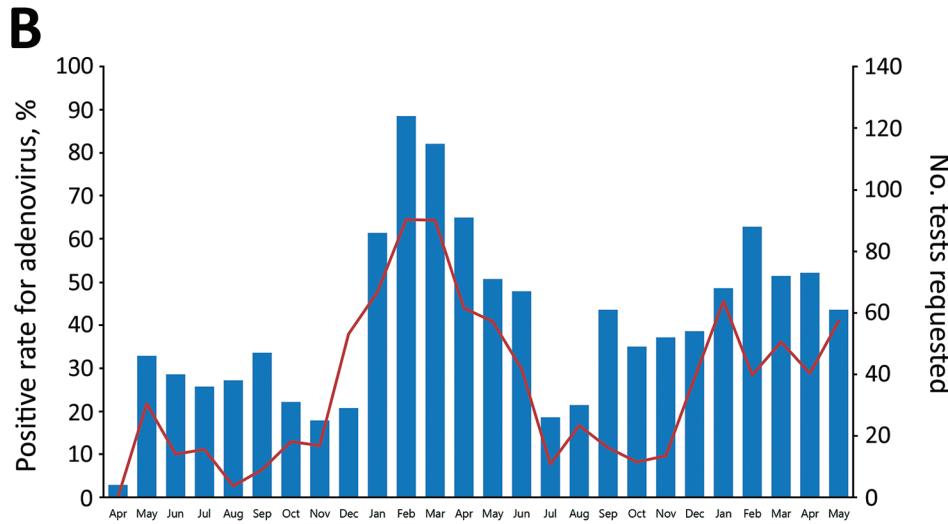


Figure 1. A) Weekly febrile respiratory illness (FRI) rate (solid line) and monthly number of pneumonia patients (dashed line) in the South Korea military, 2011–2016. B) Positive rate of human adenovirus from respiratory specimens (red line) and the number of respiratory virus PCR requested (blue bar) from a tertiary military hospital, South Korea, 2014–2016. The rate and number for each month are shown in the table at bottom.



Year	2014												2015												2016					
	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	
Positive rate, %	0.0	21.7	10.0	11.1	2.6	6.4	12.9	12.0	37.9	47.7	64.5	64.3	44.0	40.8	29.9	7.7	16.7	11.5	8.2	9.6	27.8	45.6	28.4	36.1	28.8	41.0				
No. tested	4	46	40	36	38	47	31	25	29	86	124	115	91	71	67	26	30	61	49	52	54	68	88	72	73	61				

We reviewed the demographic and clinical information of 878 military patients with FRI or pneumonia who were tested for respiratory viruses from October 2014 through May 2016 (Tables 1, 2). Soldiers of lower rank were markedly more likely to be infected with HAdV; soldiers serving in the Air Force were less likely. Patients who had been referred from other hospitals were twice as likely to be HAdV-infected than patients who visited the Armed Forces Capital Hospital directly. Rhinorrhea, sore throat, diarrhea, and nausea/vomiting were more common in patients with HAdV infection. The proportion of patients with pneumonia and the hospitalization rate did not differ between those with and without HAdV infection. However,

HAdV-infected patients had a significantly higher risk of requiring intensive care or mechanical ventilator support. In the HAdV-infected group, 8 patients required intubation and 1 died; no one in the noninfected group died or required intubation. Length of hospital stay was also significantly longer among those in the HAdV-infected group than among those in the noninfected group (12.6 vs. 9.4 days).

We conducted molecular typing by the sequencing of hexon and fiber genes with 74 HAdV-positive respiratory specimens collected from March through June 2016 (methods and general characteristics of the patients are available in the online Technical Appendix Table 2). Among them, 49 samples were successfully sequenced (GenBank numbers in

Table 1. Epidemiologic characteristics of patients with or without identification of HAdV from respiratory specimens by PCR, South Korea, 2014–2016*

Epidemiologic characteristic	Patients with HAdV, n = 447	Patients with other virus PCR negative for HAdV, n = 431	OR (95% CI)	p value
Year				
Apr 2014–May 2015	274 (65.4)	145 (34.6)	3.13 (2.37–4.12)	<0.001
Jun 2015–May 2016	173 (37.7)	286 (62.3)		
Rank				
Recruit or private	251 (70.5)	105 (29.5)	3.98 (2.98–5.31)	<0.001
PFC or higher	196 (37.5)	326 (62.5)		
Service				
Army	423 (52.1)	391 (47.9)	N/A	<0.001
Navy/Marine Corps	20 (50.0)	20 (50.0)		
Air Force	1 (4.8)	20 (95.2)		
Region				
Seoul/Gyeonggi-do	376 (50.0)	376 (50.0)		0.055
Gangwon-do	46 (65.7)	24 (34.3)		
Chungcheong-do	12 (38.7)	19 (61.3)		
Gyeongsang-do	10 (58.8)	7 (41.2)		
Jeolla-do	3 (37.5)	5 (62.5)		
Route of visit				
Direct	330 (47.3)	367 (52.7)	2.03 (1.45–2.85)	<0.001
Referral	117 (64.6)	64 (35.4)		
Age, y, mean (SD)	20.8 (2.0)	22.2 (5.0)		<0.001

*Values are no. (%) except as indicated. HAdV, human adenovirus; NA, not available; OR, odds ratio; PFC, private first class.

online Technical Appendix). Phylogenetic analyses showed that all 49 HAdV strains from South Korea clustered with HAdV-55 strains from China, Singapore, Taiwan, Spain, and the United States (Figure 2).

Conclusions

We describe an outbreak of FRI associated with HAdV in the South Korea military. HAdV is a well-known major cause of FRI in the military, accounting for >50% of FRI and pneumonia cases in military recruits (1). Our study also confirmed the predominance of HAdV, which was identified in 49.1% of specimens from patients with FRI or pneumonia. These findings are similar to those of previous studies from South Korea and the United States (9,10).

The most notable finding of our study is the emergence of HAdV-55 in the South Korea military. HAdV-55 is a novel type that has been associated with a severe clinical course and death in healthy young adults (7,8). We also found that HAdV infection was associated with intensive care, mechanical ventilator support, and longer hospital stay. In addition, we found that the only patient who died was HAdV infected. From a molecular perspective, HAdV-55 is a novel type with a hexon gene recombination between HAdV-11 and HAdV-14 (11). Phylogenetic analysis by using the hexon and fiber gene sequence of 49 strains collected in our study showed that they clustered with previously reported HAdV-55 strains.

Our findings have implications beyond military settings. Spread of infection of traditionally military-associated

Table 2. Clinical characteristics of patients with or without identification of HAdV from respiratory specimens PCR, South Korea, 2014–2016*

Clinical characteristic	Patients with HAdV, n = 447	Patients with other virus or PCR negative for HAdV, n = 431	OR (95% CI)	p value
Presenting symptoms				
Cough†	423 (94.6)	395 (91.9)	1.56 (0.91–2.67)	0.102
Rhinorrhea‡	229 (51.3)	192 (44.7)	1.31 (1.00–1.71)	0.047
Sore throat§	286 (64.3)	207 (48.1)	1.94 (1.48–2.54)	<0.001
Dyspnea¶	60 (13.5)	41 (9.5)	1.48 (0.97–2.25)	0.070
Diarrhea#	125 (33.8)	60 (17.2)	2.46 (1.73–3.49)	<0.001
Nausea/vomiting**	115 (31.0)	58 (16.6)	2.25 (1.58–3.22)	<0.001
Pneumonia	231 (51.7)	250 (58.0)	0.77 (0.59–1.01)	0.060
Hospitalization	277 (62.0)	270 (62.6)	0.97 (0.74–1.28)	0.836
Intensive care	70 (25.3)	30 (11.1)	2.71 (1.70–4.31)	<0.001
Mechanical respiratory support	25 (9.0)	5 (1.9)	5.26 (1.98–13.95)	<0.001
Intubation	8 (2.9)	0	NA	0.005
Death	1 (0.4)	0	NA	0.323
Length of stay, d, mean (SD)	12.6 (9.7)	9.4 (5.0)	NA	<0.001

*Values are no. (%) except as indicated. HAdV, human adenovirus; NA, not available; OR, odds ratio. †n = 877.

‡n = 876.

§n = 875.

¶n = 876.

#n = 719.

**n = 720.

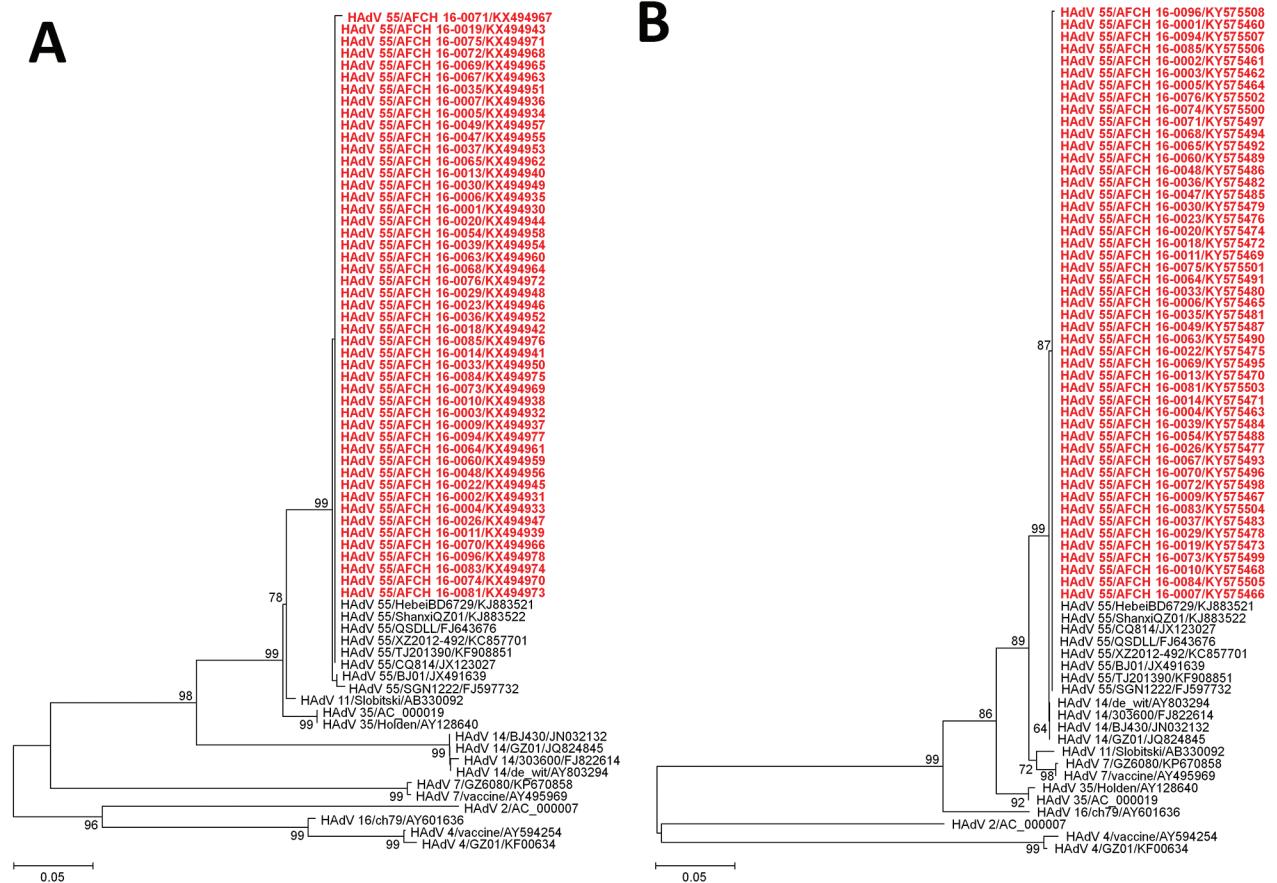


Figure 2. Phylogenetic analysis of human adenoviruses based on the partial nucleotide sequences of hexon (A) and fiber (B) genes, South Korea, 2016. Phylogenetic trees were generated by the neighbor-joining method, using the Kimura 2-parameter method. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (1,000 replicates) are shown next to the branches. Red indicates viruses identified in this study. Scale bars denote the number of base substitutions per site.

HAdV types into civilians has been recently reported in the United States and China (3,12,13). Thus, surveillance of HAdV types among both military and civilian populations is warranted; such measures are being implemented by the US Centers for Disease Control and Prevention (Atlanta, GA, USA) (12).

Our study has some limitations. First, our findings may not be generalizable due to the retrospective nature of the study. However, the military health system in South Korea provides healthcare exclusively to all military personnel; therefore, epidemiologic information gathered from our surveillance is accurate and comprehensive. Second, we conducted molecular typing with samples collected from February 2016, which was substantially later than the onset of the epidemic. However, HAdV-55 had already been identified in a case series from our center during June 2014–May 2015 (8). Because evidence shows that HAdV-55 has been already circulating since early 2014, we believe we can assume that HAdV-55 was the causative agent of the outbreak described in this study. Previously, the HAdV

typing study conducted in 2007 reported HAdV-7 as the most prevalent type (14). Lack of continuous surveillance makes it difficult to estimate exactly when this novel type was introduced into South Korea.

Further genomic analysis of the collected samples and enhanced surveillance, including of civilian populations, would provide more information on the epidemiology of HAdV infection. In addition, studies are needed on the efficacy of previous vaccines against HAdV-55.

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Dr. Yoo is a pulmonary physician and medical officer at the Armed Forces Capital Hospital of South Korea. His research interests include pulmonary infectious diseases.

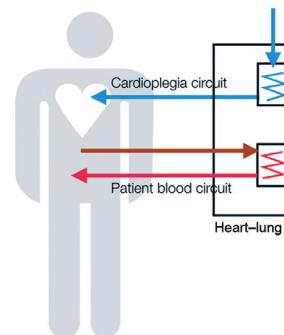
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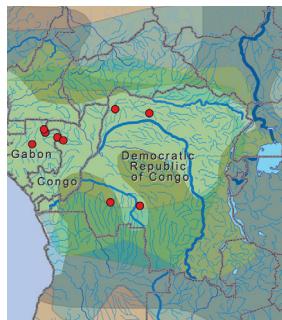
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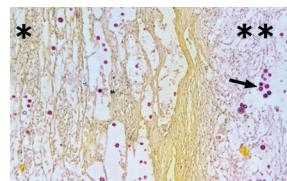


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**EMERGING
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Febrile Respiratory Illness Associated with Human Adenovirus Type 55 in South Korea Military, 2014–2016

Technical Appendix

Methods

Military Surveillance for Febrile Respiratory Illness (FRI)

Military FRI surveillance, activated in October 2011, collects data from 43 walk-in clinics that provide primary care exclusively to military personnel. The sentinel sites were selected to represent all 3 departments of the Armed Forces and various geographic areas of South Korea. FRI was defined as a body temperature $\geq 38^{\circ}\text{C}$ and the presence of a cough and/or sore throat. The FRI rate was calculated by dividing the number of FRI patients by the number of total outpatients each week. Each sentinel clinic was required to report their weekly FRI rates using a web-based report system. “Increased activity” was defined as occurring when an FRI rate of a certain week was higher than the upper limit of 95% confidence interval of the pre-season mean FRI rate.

General Characteristics of the Armed Forces Capital Hospital

The Armed Forces Capital Hospital is a 660-bed acute care hospital that is the only tertiary care hospital providing referral service to the entire military health care system in South Korea. It also provides primary and secondary care to the service members stationed in the Seoul/Gyeonggi-do metropolitan area. The military health care system in South Korea provides medical care exclusively to all military service members, except for the small number of patients who choose to use civilian clinics; in that case, they still need to visit a military walk-in clinic first for a referral.

Molecular Analyses for Respiratory Viruses

DNA was extracted from sputum or nasopharyngeal swab specimens by using MagNA Pure LC 2.0 (Roche Diagnostics, Mannheim, Germany). Multiplex real-time PCR was conducted

for 15 respiratory viruses (adenovirus, rhinovirus, influenza viruses A and B, respiratory syncytial viruses A and B, coronaviruses 229E/OC43/NL63/HKU1, parainfluenza viruses 1/2/3, bocavirus, and metapneumovirus) by using Real-Q RV Detection Kit (BioSewoom, Seoul, South Korea) on a LightCycler 480 II (Roche Diagnostics) following the manufacturer's instructions.

Since March 2016, DNA samples from 74 consecutive patients with PCR-confirmed human adenovirus (HAdV) infection were stored at -70°C for further molecular analysis. Stored samples were transported regularly to the Agency for Defense Development for sequencing and genotyping. PCR was performed in 50 μL reactions containing dNTP mixture (2.5 mM), 5 U of LA Taq polymerase (Takara, Shiga, Japan), and 10 μM of each primer. Oligonucleotide primer sequences for 2 sequenced gene were as follows:

Hypervariable Region 7 (HVR7) of the Hexon Gene (1)

Hexon gene:

Forward: 5'-CTGATGTACTACAACAGCACTGGCAACATGGG-3'

Reverse: 5'-GCGTTGCGGTGGTGGTTAAATGGGTTTACGTTGTCCAT-3'

Fiber gene:

Forward: 5'-ATGACCAAGAGAGTCCGGCTCAG-3'

Reverse: 5'-ATGCAGTGACTAGGGCTCCAGT-3'

Initial denaturation was conducted at 94°C for 5 min, followed by 15 cycles of denaturation at 94°C for 40 s, annealing at 50°C for 40 s, and elongation at 72°C for 1 min, then followed by 25 cycles of denaturation at 94°C for 40 s, annealing at 52°C for 40 s, and elongation at 72°C for 1 min, in a ProFlex PCR system (Applied Biosystems, Foster City, CA, USA). PCR products were sequenced directly by using an Applied Biosystems 3500 series Genetic Analyzer (Applied Biosystems). GenBank accession numbers for the hexon and fiber genes sequences are KX494930–KX494978 and KY575460–KY575508, respectively. Phylogenetic trees were generated by the neighbor-joining method with 1,000 bootstrap replicates using MEGA 6 (2).

Statistical Analysis and Ethical Review

χ^2 or Fisher exact tests were used to compare categorical variables, whereas unpaired Student *t* tests were used to compare continuous variables. A p value <0.05 was considered statistically significant, and all tests were 2-tailed. Statistical analysis was performed by using SPSS Statistics 20 (IBM Corp., Armonk, NY, USA). This study was approved by the Institutional Review Board of AFMC (AFMC-16081-IRB-16-072) with a waiver of informed consent.

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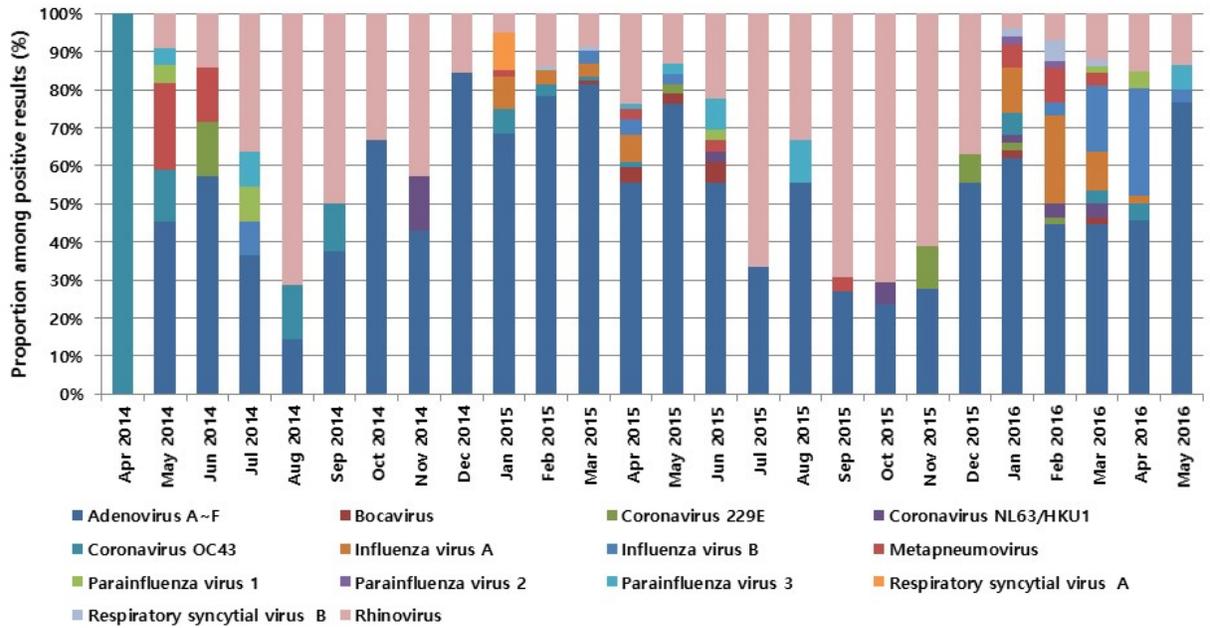
Technical Appendix Table 1. Distribution of viruses identified from respiratory specimens during flu seasons 2014–15 and 2015–16, South Korea

Respiratory virus	2014–15			2015–16		
	No. viruses	Proportion among all samples, %	Proportion among positive PCR samples, %	No. viruses	Proportion among all samples, %	Proportion among positive PCR samples, %
Adenovirus	282	49.3	79.7	150	29.1	53.2
Bocavirus	5	0.9	1.4	2	0.4	0.7
Coronavirus 229E	1	0.2	0.3	6	1.2	2.1
Coronavirus NL63/HKU1	1	0.2	0.3	6	1.2	2.1
Coronavirus OC43	9	1.6	2.5	7	1.4	2.5
Influenza virus A	17	3.0	4.8	26	5.0	9.2
Influenza virus B	7	1.2	2.0	26	5.0	9.2
Metapneumovirus	3	0.5	0.8	10	1.9	3.5
Parainfluenza virus 1	0	0	0	3	0.6	1.1
Parainfluenza virus 2	0	0	0	2	0.4	0.7
Parainfluenza virus 3	2	0.3	0.6	2	0.4	0.7
Respiratory syncytial virus A	6	1.0	1.7	0	0	0
Respiratory syncytial virus B	2	0.3	0.6	5	1.0	1.8
Rhinovirus	54	9.4	15.3	57	11.0	20.2
No virus identified	218	38.1		234	45.3	
Total	572	100.0		516	100.0	

Technical Appendix Table 2. Epidemiologic and clinical characteristics of patients from whose respiratory samples adenovirus typing was conducted, South Korea, 2014–2016*

Characteristic	HAdV typing, n = 49 (%)	HAdV overall, n = 447 (%)	p
Epidemiologic			
Rank			
Recruits or privates	29 (59.2)	251 (56.2)	0.762
PFC or higher	20 (40.8)	196 (43.8)	
Service			
Army	45 (91.8)	423 (94.6)	0.298
Navy/Marine Corps	3 (6.1)	20 (4.5)	
Air Force	1 (2.0)	1 (0.2)	
Region			
Seoul/Gyeonggi-do	37 (75.5)	376 (84.1)	0.156
Gangwon-do	9 (18.4)	46 (10.3)	
Chungcheong-do	0	12 (2.7)	
Gyeongsang-do	3 (6.1)	10 (2.2)	
Jeolla-do	0	3 (0.7)	
Route of visit			
Direct	20 (40.8)	330 (73.8)	<0.001
Referral	29 (59.2)	117 (26.2)	
Age, y (mean, SD)	20.4 (1.9)	20.8 (2.0)	0.182
Clinical			
Pneumonia	39 (79.6)	231 (51.7)	<0.001
Hospitalization	46 (93.9)	277 (62.0)	<0.001
Intensive care	17 (37.0)	70 (25.3)	0.002
Ventilatory support	4 (8.7)	25 (9.0)	0.515
Intubation	3 (6.5)	8 (2.9)	0.085
Death	0	1 (0.4)	>0.999

*HAdV, human adenovirus; PFC, private first class.



Technical Appendix Figure. Pathogen distribution of respiratory viruses identified by multiplex real-time PCR at the Armed Forces Capital Hospital, South Korea, 2014–2016.