Correlation between Infection with Multiple Respiratory Viruses and Length of Hospital Stay in Patients from Cheonan, Korea

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INTRODUCTION

Increased severity of viral infections has recently been reported to prolong the length of hospital stay (LOS). Further, the LOS may differ depending on the virus type [1]. However, controversy exists regarding whether these viruses increase the severity of illness and LOS [2]. It has been reported that, while co-infections do not affect LOS or mortality, influenza virus (INF) infection is associated with increased LOS and mortality [3,4]. Nonetheless, the impact of co-infections is, in general, still unclear [2,5-7]. This study examined whether the LOS for respiratory virus infections differed by sex, age, causative virus, or viral co-infection.
MATERIALS AND METHODS

1. Participants
A total of 8,860 patients who were treated at Dankook University Hospital for respiratory symptoms between December 2006 and February 2014 were retrospectively included in the study. Patients were admitted either via the emergency department or via outpatient services. Molecular tests for the presence of respiratory viruses were performed on all patients.

2. Ethical considerations
The present study was approved by the institutional review board (IRB) of Dankook University (Date of IRB approval: 2015.10.13; IRB approval No.: 2015-09-009). The work described in this study has been carried out in accordance with The Code of Ethics of the World Medical Association (Declaration of Helsinki).

3. Sample collection and extraction of viral nucleic acid
Within 24 hours of admission, nasopharyngeal secretions were collected by inserting a mucus extractor, and connected to a sterile 8-French catheter (Sewoon Medical, Cheonan, Korea) 5 ∼ 7 cm into the nostril and applying a suction pressure of 60 ∼ 80 mmHg. Samples were stored at 4°C until nucleic acid extraction was performed. Nucleic acid extraction was performed using a QIAamp DNA Mini Kit (Qiagen, Hilden, Germany) — according to the manufacturer’s protocol — using QIAcube (Qiagen, Hilden, Germany), which is automated equipment for nucleic acid extraction. The extracted nucleic acids were stored at −70°C until the tests were performed.

4. Respiratory virus detection
Respiratory viruses were detected using multiplex reverse transcriptase–polymerase chain reactions (RT-PCR), which can simultaneously detect 12 types of respiratory virus. Ribonucleic acid (RNA) was isolated from nasopharyngeal secretions and reconstructed as complementary deoxyribonucleic acid (cDNA). The cDNA was then amplified with the Seeplex RV detection kit-1 (Seegene, Seoul, Korea) to test for one DNA virus, adenovirus (ADV), and 11 RNA viruses that cause respiratory infection: respiratory syncytiatal virus (RSV) A and B; INF A and B; parainfluenza virus (PIV) types 1, 2, and 3; human rhinovirus (HRV); human coronavirus (hCoV) NL63, and OC43; and human metapneumovirus (hMPV). PCR was performed using the PTC 200 PCR system (MJ Research, Watertown, MA, USA) with a program of 40 cycles of: 30 s at 94°C, 90 s at 60°C, and 90 s at 72°C, followed by 1 cycle of 10 min at 72°C. The amplified PCR products were analyzed after 30 min of electrophoresis at 100 ∼ 150 V in 2% agarose gels stained with ethidium bromide.

5. Statistical analysis
Patient characteristics at the time of admission, including age, sex, and LOS, were analyzed retrospectively. A p-value by statistically analysis, <0.05 was considered to be significant.

RESULTS
Of 8,860 patients, 6,492 respiratory viruses were detected in 5,310 virus-positive patients. A total of 3,111 male and 2,199 female patients were positive for respiratory virus infections; their median age was 1.5 years (range, 2 days to 96.2 years, Table 1). Of the patients admitted to the hospital and referred for PCR testing for respiratory viruses, the average LOS among patients who tested positive was 7.2 days; 7.3 and 7.0 days for men and women, respectively (p =0.109) (Table 2). The average LOS for the group of patients 0 ∼ 10 years of age was 6.5 days; however, the group aged 30 ∼ 39 years was admitted for an average of 10.7 days, and the group aged 50 ∼ 59 years was admitted for an average of 13.7 days, suggesting that age is related to longer LOS (Figure 1). Regression analysis showed a significant proportional relationship between LOS and age (p<0.01).

Among single infections, infection with hCoV OC43 was associated with the longest LOS (10.1 days), followed by infection with INF A (8.9 days). ADV infections had the
Table 1. Analysis of cases according to the respiratory infection type, including positive ratios and length of hospital stay

| Infection type          | No. of patients | Ratio, % | Men       | Women      | Median age (average age) | Average LOS, days |
|-------------------------|-----------------|----------|-----------|------------|-------------------------|------------------|
| Submitted               | 8,860           | 100.0    | 5,292     | 3,568      | 2.2 (7.4)               |                  |
| Positive                | 5,310           | 59.9     | 3,111     | 2,199      | 1.5 (9.3)               | 7.2              |
| Single infection        | 4,250           | 48.0     | 2,496     | 1,754      | 1.5 (10.3)              | 7.3              |
| Multiple infections     | 1,060           | 12.0     | 615       | 445        | 1.4 (5.1)               | 6.7              |
| Double infections       | 943             | 10.6     | 550       | 393        | 1.4 (5.1)               | 6.7              |
| Three or more infections| 117             | 1.3      | 65        | 52         | 1.5 (5.2)               | 6.6              |

Abbreviation: LOS, length of hospital stay.

Table 2. Analysis of cases according to single respiratory virus type, including positive ratios and length of hospital stay

| Virus     | No. of patients | Length of hospital stay, days | Ratio, %*
|-----------|-----------------|------------------------------|----------|
| INF A     | 466             | 8.9                          | 11.0     |
| INF B     | 134             | 6.5                          | 3.2      |
| RSV A     | 803             | 7.3                          | 18.9     |
| RSV B     | 546             | 6.6                          | 12.8     |
| hMPV      | 305             | 7.6                          | 7.2      |
| PIV 1     | 162             | 6.9                          | 3.8      |
| PIV 2     | 49              | 6.3                          | 1.2      |
| PIV 3     | 262             | 7.7                          | 6.2      |
| HRV       | 794             | 7.1                          | 18.7     |
| hCoV NL63 | 117             | 7.2                          | 2.8      |
| hCoV OC43 | 109             | 10.1                         | 2.6      |
| ADV       | 503             | 6.2                          | 11.8     |

*Number of each virus/Total positive number.

Abbreviation: INF, influenza; RSV, respiratory syncytial virus; hMPV, human metapneumovirus; PIV, parainfluenza virus; HRV, human rhinovirus; hCoV, human coronavirus; ADV, adenovirus.

The shortest LOS, at 6.2 days (Table 2). The LOS for patients infected with hCoV OC43 was significantly different from that of patients infected with INF A virus (p<0.03). Among patients positive for respiratory viruses, those with single infections had an average LOS of 7.3 days, compared with 6.7 days in patients with double infections and 6.6 days in patients with three or more identified respiratory viruses (Table 1). Differences in LOS between patients with single and the combined group of patients with double or three or more infections were statistically significant (p<0.01). However, the difference between double infections and three or more infections was not significant (Table 1, p=0.858).

DISCUSSION

This study found no difference in LOS based on sex, but found that the LOS increased with increasing age. Previous studies have reported increasing emergency room admissions with increasing age [8]. And longer LOS among the elderly, with observations that the LOS remains constant despite the administration of antibiotics for viral infections [9]. However, longer LOS in the elderly could also be due to smoking, cardiovascular and/or respiratory diseases, and decreased immunity [10].

Overall, respiratory viruses were detected in 59.9% (5,310/8,860) of patients collected in this study; higher than in other studies conducted at approximately the same time period [11-13]. Of the viruses identified in patients positive for respiratory viruses, HRV was the most prevalent (21.4%), followed by RSV A (15.8%) and ADV (15.2%). Similarly, Seo, et al. (2014) reported that the prevalence of HRV, ADV, and RSV A infections were 31.8%,
19.2%, and 17.4%, respectively, in Chung-nam, Korea. A study by Wishaupt, et al. (2011) in the Netherlands reported the following prevalence rates: RSV A, 23.6%; HRV, 19.4%; and RSV B infection, 11.5%. Although the virus detection rates in the present study differed from these other studies, the major viruses identified were the same, namely RSV, HRV, and ADV. In relation to virus types, the LOS of 10.1 days for hCoV OC43 was longer than those of other virus types (range, 8.9–6.2 days); this was significant (p=0.030) as was that for INF A (8.9 days), compared with the other virus types (p=0.027). However, previous studies have reported different relationships between LOS and virus types. In recently published data, the LOS for patients infected with INF A was reported to be 6.9 days, whereas LOS for patients infected with ADV or hCoV OC43 was 8.2 days and 8.1 days, respectively [14]. In another study, RSV and hMPV infections reportedly resulted in LOS of 4 days and 3 days, respectively [15]. Although RSV B and RSV A infections did not differ in severity, RSV B infection was reported to have a significantly longer recovery time as well as a longer LOS [16]. However, a study that used oxygen therapy, admission to the intensive care unit, and LOS greater than 5 days as indices for severity reported that infections with RSV A were more severe than infections with RSV B [17]. In addition, the INF A virus subtypes H7N9, H5N1, and H1N1 have been reported to have different LOS [18]. These varying LOS by study might be explained by different standards for admission and discharge at individual hospitals. In addition, differences in LOS might be influenced not only by virus type but also by underlying diseases and their severity.

Regarding the relationship between LOS and the number of infections, the LOS with single infections was longer than that with multiple infections (7.3, 6.7 and 6.6 days for double and three or more infections, respectively) by approximately 0.6 days (p=0.006). However, the LOS was similar between infections associated with double and three or more respiratory viruses. Papadopoulos, et al (2002) reported a LOS difference between RSV and HRV as single (2.8 days) and multiple infections (4.0 days).

Similarly, there have been reports of longer LOS among pediatric patients with multiple RV infections compared with single infections [19,20]. Moreover, multiple infections have been reported to induce a less favorable prognosis [8]. In contrast, Esposito, et al (2013) reported that the difference in LOS between single (6.9 days) and multiple infections (7.1 days) was not significant. Other studies have reported no difference in clinical symptoms, prognosis, or intensive care unit admission rates between single and multiple infections [21–23], with some reporting less severe disease in patients with multiple respiratory virus infections [24,25]. These results might be affected by various conditions, such as underlying diseases, type of virus detected in different geographical regions, and the combination of viruses that caused the multiple infections.

However, further research is required. LOS related to respiratory virus infection was significantly longer for older patients and those infected with hCoV OC43 or INF A. These specific viral infections can cause many complications, particularly cytopenias. Anemia, neutropenia, and thrombocytopenia are common and are reportedly caused by autoimmune mechanisms, including inhibition of progenitor cells in the bone marrow, movement of neutrophils from blood to tissue, and reduction of platelet production or increased destruction of platelets in the spleen or reticuloendothelial system [26,27]. These complications mainly develop in cases of acute respiratory infection and may affect clinical progress [28,29] because they are related both to patient recovery and disease progression. Studies on the severity of complications are important for determining effective treatment.

The present study has some limitations. The study subjects were selected only from the Cheonan area, and the analyses were based on results from a single institution, thus limiting the generalizability of the findings. In addition, as the subject selection did not involve a planned design owing to the retrospective nature of the study, additional examinations could not be performed. Further multicenter studies with a larger number of subjects are needed to clarify our findings. The strengths of
this study are the long study period of 5 years and the fact that recent data is presented.

In conclusion, the LOS for patients with multiple infections was not longer than that for patients with single infections. These results are anticipated to be useful for developing effective treatment guidelines for respiratory virus infections.

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