Original Article

Prevalence of Human Coronaviruses NL63, HKU1, 229E, OC43, MERS, and SARS-CoV-2 among Hospitalized Patients with Acute Respiratory Infection in Tehran

Seyed Jalal Kiani¹, Atefeh Kachooei Mohaghegh¹, Masoud Eslami², Ahmad Tavakoli³, Mohammad Hadi Karbalaie Niya^{1,4}, Alireza Javan⁵, Sheida Alizadeh^{6,7}, Zahra Salavatiha¹, Mahdieh Hosseini¹, Zahra Safaie¹, Mohammad Reza Rezvani⁸, Seyed Hamidreza Monavari^{1*}

- 1. Department of Medical Virology, School of Medicine, Iran University of Medical Sciences, Tehran, Iran.
- 2. Maham Foundation for Respiration Science (MFRS), Neuroscience Research Centre (NRS), Iran University of Medical Sciences, Tehran, Iran.
- 3. Research Center of Pediatric Infectious Diseases, Institute of Immunology and Infectious Diseases, Iran University of Medical Sciences, Tehran, Iran.
- 4. Gastrointestinal and Liver Diseases Research Center, Iran University of Medical Sciences, Tehran, Iran.
- 5. Student Research Committee, School of Medicine, Iran University of Medical Sciences, Tehran, Iran,
- 6. Department of Bacteriology and Virology, Shiraz University of Medical Science, Shiraz, Iran
- 7. Gastrointestinal and Liver Diseases Research Center, Iran University of Medical Sciences, Tehran, Iran.
- 8. Department of Hematology and Blood Transfusion, School of Allied Medical Sciences, Iran University of Medical Sciences, Tehran, Iran.

Abstract

Background and Aims: Coronaviruses cause upper respiratory ailments and sporadically lower tract sickness in vulnerable populations. This study examined the prevalence of six human coronaviruses, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), OC43, 229E, HKU1, NL63, and Middle East respiratory syndrome coronavirus (MERS-CoV) during the 2019 pandemic caused by SARS-CoV-2 in Tehran, Iran.

Material and methods: Specimens were collected from 204 adult patients with acute respiratory illness. The specimens were examined for the presence of six human coronaviruses using consensus and subtype-specific Real-time reverse-transcription polymerase chain reaction (PCR) assays. The demographic and clinical characteristics associated with coronavirus infection were examined retrospectively.

Results: Coronaviruses were identified in 204 adult patients. The gender ratio was 104/100 male and female, respectively. HCoV-229E, HCoV-NL63, HCoV-OC43, HCoV-HKU1, MERS-CoV, and SARS-CoV-2 were detected in 0.5%, 1.0%, 4.0%, 7.4%, 0.0%, and 22.5% of patients, respectively. The acute clinical features were similar across subtypes. There was no report of fatality incidence among the population during the investigation period.

Conclusions: HCoVs could play a significant role in causing upper respiratory tract infections among adults and older children. Based on the findings of this study and those of others, more extensive studies using other diagnostic methods and higher sample sizes are suggested.

Keywords: Human coronavirus, Pandemic, Iran, Prevalence

Introduction

he seven strains of human coronaviruses

*Corresponding author:

Seyed Hamidreza Monavari, Ph.D Department of Virology, School of Medicine, Iran University of Medical Sciences Email: monavari.hr@iums.ac.ir

Tel: +98 2188602205

(HCoVs) belong to the genus Alphacoronavirus or Beta coronavirus of the Coronavirinae subfamily and include HCoV-229E, HCoV-NL63, HCoV-OC43, HCoV-HKU1, severe acute respiratory syndrome coronavirus (SARS-CoV), MERS-CoV, and SARS-CoV-2 (1-3). All human coronaviruses, except for SARS-CoV, MERS-CoV, and SARS-CoV-2, are frequently linked to minor gastrointestinal and respiratory illnesses (4,5). The highly deadly

viruses, SARS-CoV, MERS-CoV, and SARS-CoV-2, have resulted in severe illnesses or death in humans (6, 7).

Poor growth and poverty of cytopathic effects in cell cultures have been the main limits for HCoVs research. However, with the development of polymerase chain reaction (PCR) technology, there has been a broad and rapid expansion in the field of corona virology (8,9). According to early studies, HCoV-229E and HCoV-OC43 were known to account for 5 to 30% of human respiratory tract infections (10), and HCoV-NL63 was found to exist in 2 to 3.6% of respiratory samples in several recent studies (11-16) Despite various studies during the last few years on the etiology of coronaviruses (15, 17), our understanding of the status of these viruses in Iran is limited.

However, during the recent pandemic caused by SARS-CoV-2, we conducted a survey study on the prevalence of six human coronaviruses, including HCoV-229E, HCoV-NL63, HCoV-OC43, HCoV-HKU1, MERS-CoV, and SARS-CoV-2 in patients clinically confirmed for severe acute respiratory syndrome presented with fever, nausea, cough, and sore throat, who were admitted to hospitals affiliated with Iran University of Medical Sciences in Tehran, Iran.

Methods and Materials

Patients and Methods

Between October 2020 and June 2021, clinical samples (i.e., throat swabs and nasopharyngeal aspirates) from 204 patients with acute respiratory infections admitted to three general hospitals, including Firroozgar, Rasool Akram, and Ali Asghar in Tehran, were collected. The epidemiology and clinical spectrum of the disease caused by HCoV-229E, HCoV-NL63, HCoV-OC43, HCoV-HKU1, MERS-CoV, and SARS-CoV-2 infection in patients hospitalized for acute respiratory illness were examined during one year.

RNA Extraction

Extraction of viral RNA from samples was carried out using an RNA extraction kit (Yekta Tajhiz Kit, Tehran, Iran). The extracted RNAs

(templates for RT-PCR) were immediately stored at -70°C until use.

RT-PCR for Coronaviruses

After RNA extraction to screen for six coronaviruses, PCR was performed using six sets of primers and probes precisely designed as described previously to amplify HCoV-229E, HCoV-OC43, HCoV-NL63, HCoV-HKU1, MERS-CoV, and SARS-CoV-2 (Table 1). Each PCR mixture contained TaqMan onestep PCR 2X Master Mix (10 ul), forward primer (1 ul), reverse primer (1 ul), probe (0.5 ul), double-distilled water (2.5 ul), sample (5 ul), in a total volume of 20 ul. The cycling program consisted of an initial denaturation at 94°C for 3 min, followed by 45 cycles of 94°C for 10s and 60°C for 30s (Table 2).

Results

A total of 204 patients with ages ranging from 1 to 90 years old (mean 42.69 years) were enrolled in this study. Specimens were collected from both female (100) and male (104) patients. HCoV RNA was detected in 72 (30.3%) of the 204 patients. Of the 72 specimens positive for HCoV RNA, 1.0%, 4.0%, 0.5%, 7.4%, and 22.5% were positive for HCoV-NL63, HCoV-OC43, HCoV-229E, HCoV-HKU1, and SARS-CoV-2, respectively, and no positive case was found for MERS-CoV (Table 3).

Additionally, we found no significant associations between the gender and age of the patients infected with these six serotypes of human coronaviruses (P>0.05). Both HCoV-NL63 positive cases had fever, vertigo, and malaise. The only patient infected by HCoV-229E presented symptoms including fever, cough, sore throat, vertigo, and malaise. Among the eight HCoV-OC43-positive cases, the most common clinical symptoms were fever (n=5) and body aches (n=4). Likewise, the most common clinical symptoms observed among the 15 patients infected with HCoV-HKU1 were body aches (n=8), fever (n=8), sore throat (n=6), vertigo (n=6), and malaise (n=6).

Table 1. Primers and probes used in this study

Coronavirus	Primer direction	sequence (5'-3')		
	Forward primer	5'- CAGTCAAATGGGCTGATGCA -3'		
HCoV-229E	Reverse primer	5'- AAAGGGCTATAAAGAGAATAAGGTATTCT -3'		
	Probe	5'-FAM CCCTGACGACCACGTTGTGGTTCA-BHQ1 -3'		
	Forward primer	5'-CGATGAGGCTATTCCGACTAGGT-3'		
HCoV-OC43	Reverse primer	5'-CCTTCCTGAGCCTTCAATATAGTAACC-3'		
	Probe	5'-FAM TCCGCCTGGCACGGTACTCCCT-BHQ1 -3'		
	Forward primer	5'- GACCAAAGCACTGAATAACATTTTCC -3'		
HCoV-NL63	Reverse primer	5'- ACCTAATAAGCCTCTTTCTCAACCC -3'		
	Probe	5'-FAM ATGTTATTCAGTGCTTTGGTCCTCGTGAT-BHQ1 3'		
	Forward primer	5'- CCTTGCGAATGAATGTGCTC -3		
HCoV-HKU1	Reverse primer	5'- TTGCATCACCACTGCTAGTACCAC -3		
	Probe	5'-FAM TGTGTGGCGGTTGCTATTATGTTAAGCCTG-BHQ1-3'		
	Forward primer	5'- GCAACGCGCGATTCAGTT -3		
MERS-CoV	Reverse primer	5'- GCCTCTACACGGGACCCATA -3		
	Probe	5'-FAM CTCTTCACATAATCGC CCCGAGCTCG-BHQ1 -3'		
	Forward primer	5'- GGG GAA CTT CTC CTG CTA GAA T -3		
SARS-CoV-2	Reverse primer	5'- CAG ACA TTT TGC TCT CAA GCT G -3		
	Probe	5'-FAM TTG CTG CTG CTT GAC AGA TT-BHQ1 -3'		

Table 2. The chart for PCR Real-Time

No.	Cycle	Step	Tem.	Time
1	1	Reverse	50 °C	15 min
		Transcription		
2	1	Initial Denaturation	94 °C	3 min
3	15	Denaturation	94 °C	10 sec
	45	Annealing-Extension	60 °C	30 sec
4	1	Device Cooling	25 °C	1 sec

Table 3. Prevalence of six studied coronaviruses among 204 patients

No.	HCoV	No. of Positive Case (%)				
1	NL63	2 (1.0%)				
2	229E	1 (0.5%)				
3	OC43	8 (4.0%)				
4	HKU1	15 (7.4%)				
5	MERS	0 (0.0%)				
6	SARS-CoV-2	46 (22.5%)				
7	Negative	132 (64.6%)				

Among the 46 patients infected with SARS-CoV-2, fever (n=42), chills (n=16), and malaise (n=16) were the most prevalent clinical presentations. The other clinical symptoms observed in all patients infected with the six

human coronaviruses are shown in Table 4.

Discussion

Our understanding of the epidemiology of six human coronaviruses in Iran is limited to the clinical symptoms of the respiratory system. However, the objective of this study was to survey the number of patients who might be infected with some of these six strains during the 2019 pandemic SARS-CoV-2 in the country. A total of 204 patients with ages ranging from 1 to 90 years old (mean 42.69 years) were enrolled in this study. Specimens were collected from both female (100) and male (104) patients. The genomes of human coronaviruses were detected in 72 out of 204 patients (30.3%). Among the positively confirmed samples, SARS-CoV-2 (22.5%), HCoV-HKU1 (7.4%), HCoV-OC43 (4%), HCoV-NL63 (1%), and HCoV-229E (0.5%) were responseble for the outbreak, while none of these samples showed to be positive for HCoV-MERS (Table 3). Additionally, one sample was positive for both HCoV-NL63 and HCoV-229E. In 2015, Madhi and colleagues studied 170 samples collected from hospitalized patients

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Table 4. Virus-specific general characteristics and clinical symptoms of the study population

	HCoV						
Parameters	NL63	229E	OC43	HKU1	MERS	SARS- CoV-2	Total
No. of detected	2 (1.0%)	1 (0.5%)	8 (4.0%)	15 (7.4%)	0 (0%)	46 (22.5%)	72 (30.3%)
Age range (years)	37-47	47	19-39	40-59	0	1-83	1-90
Age Mean	42	47	33.8	45.5	0	47.2	42.69
Gender (M/F)	1/1	1/0	3/5	5/10	0	23/23	104/100
Symptoms							
Loss of taste	1 (50%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1/62 (1.6%)
Loss of smell	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	2 (4.3%)	2/62 (3.2%)
Fever	2 (100%)	1 (100%)	5 (62.5%)	8 (53.3%)	0 (0%)	42 (91.3%)	58/62 (93.5%)
Cough	1 (50%)	1 (100%)	3 (37.5%)	5 (33.3%)	0 (0%)	13 (28.3%)	23/62 (37.1%)
Sore throat	0 (0%)	1 (100%)	3 (37.5%)	6 (40%)	0 (0%)	6 (13%)	16/62 (25.8%)
Dyspnea	0 (0%)	0 (0%)	2 (25%)	5 (33.3%)	0 (0%)	7 (15.2%)	14/62 (22.6%)
Chest pain	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	4 (8.7%)	4/62 (6.4%)
Chill	1 (50%)	0 (0%)	2 (25%)	5 (33.3%)	0 (0%)	16 (34.8%)	24/62 (38.7%)
Rhinorrhea	1 (50%)	0 (0%)	1 (12.5%)	2 (13.3%)	0 (0%)	3 (6.5%)	7/62 (11.3%)
Sneeze	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (2.2%)	1/62 (1.6%)
Vertigo	2 (100%)	1(100%)	3 (37.5%)	6 (40%)	0 (0%)	2 (4.3%)	14/62 (22.6%)
Headache	1 (50%)	0 (0%)	2 (25%)	5 (33.3%)	5 (33.3%)	0 (0%)	13/62 (20.9%)
Malaise	2 (100%)	1 (100%)	3 (37.5%)	6 (40%)	6 (40%)	0 (0%)	28/62 (45.1%)
Body pain	1 (50%)	0 (0%)	4 (50%)	8 (53.3%)	8 (53.3%)	0 (0%)	24/62 (38.7%)

who suffered respiratory infections using the RT-PCR technique to determine the presence of four human coronaviruses: HCoV-OC43, HCoV-229E, HCoV-HKU1, and HCo V-NL-63. This study showed that the number of human coronaviruses was 15 (5.5%), and all were HCoV-E229 (18). In 2015, Sultani et al. sur-veyed samples from 172 hospitalized patients due to respiratory infections using the RT-PCR method for the presence of four human corona-viruses: HCoV-NL63, HCoV-229E, HCoV- SARS, and HCoV-OC43, and only HCoV-NL63 was detected in their study

(0.58%) (19). These findings might be considered due to two reasons: firstly, sampling in their study was conducted before the occurrence of the SARS-CoV-2 pandemic, while in our study, sampling was carried out after the appearance of the SARS-CoV-2 pandemic; secondly, during that period of time, the strains HCoV-229E and HCoV-NL63 were dominant in Iran among the other strains.

Zeng et al. conducted surveillance on respiratory infections in children using the Real-time PCR technique. Among 11,399 collected samples, 489 samples were found to be positive

(4.3%). Meanwhile, among the 506 positive cases, HCoV-OC43, HCoV-229E, HCoV-NL63, and HCoV-HKU1 serotypes were detected in 346, 65, 60, and 38 patients, respectively (20). Also, we found no significant relationships between the gender and age of the patients infected with these six serotypes of human coronaviruses (P>0.05). The clinical syndrome observations, along with the causative agents of the six human coronaviruses, are confirmed as the etiologic agents shown in Table 5. For example, in two positive cases infected with HCoV-NL63, fever and dizziness were found, and the remaining fifty percent were in the face of cough, headache, and body ache. 15% of infected patients with HKU1 showed symptoms of fever (53.3%), cough (33.3%), dizziness, and sore throat (40%). Four percent of patients confirmed the infection with OC43: fever (62.5%), cough, sore throat, and malaise (37.5%) (Table 4).

Conclusions

We found that the six human coronaviruses (HCoVs) were associated with acute respiratory tract infections in Tehran. Additionally, our data suggests that HCoVs could play a significant role in causing upper respiratory tract infections among adults and older children. Based on the findings of this study and those of others, more extensive studies using other diagnostic methods and a larger sample size are recommended.

Acknowledgment

None

Disclosure

None

Conflict of Interest

No conflict of interest is declared.

Funding

This study was financially supported by Iran University of Medical Sciences, Grants Number: 98-5-4-17227.

Ethics Approval and Consent to Participate

This study was approved by Ethics Committee of the Iran University of Medical Sciences (Approval No. IR.IUMS.REC.1399.022).

Data Availability Statement

All data generated or analyzed during this study are included in this article.

References

- 1. Onyekere PF, Nwankwo UV. Coronavirus pandemic: History, overview of different strains of coronaviruses and what went wrong. Coronavirus Drug Discovery: Elsevier; 2022. p. 3-16.
- 2. Keyvani H, Moghoofei M, Bokharaei-Salim F, Mostafaei S, Javad Mousavi SA, Monavari SH, et al. Prevalence of respiratory viruses in Iranian patients with idiopathic pulmonary fibrosis. J Med Microbiol. 2017;66 (11):1602-6.
- 3. Woo PCY, Lau SKP, Yip CC, Huang Y, Yuen KY. More and more coronaviruses: human coronavirus HKU1. Viruses. 2009;1(1):57-71.
- 4. Peiris J, Poon LL. Detection of SARS coronavirus. Diagnostic Virology Protocols: Springer; 2010. p. 369-82.
- 5. Kiani SJ, Ramshini M, Bokharaei-Salim F, Donyavi T, Eshrati B, Khoshmirsafa M, et al. High resolution melting curve analysis for rapid detection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) variants. Acta Virol. 2023;67(1):91-8.
- 6. Liu P, Shi L, Zhang W, He J, Liu C, Zhao C, et al. Prevalence and genetic diversity analysis of human coronaviruses among cross-border children. Virol J. 2017;14(1):230.
- 7. Dijkman R, Jebbink MF, El Idrissi NB, Pyrc K, Müller MA, Kuijpers TW, et al. Human coronavirus NL63 and 229E seroconversion in children. J Clin Microbiol. 2008;46(7):2368-73.
- 8. McIntosh K, Perlman S. Coronaviruses, including severe acute respiratory syndrome (SARS) and Middle East respiratory syndrome (MERS). Mandell, douglas, and bennett's principles and practice of infectious diseases. 2015:1928.
- 9. Myint SH. Human coronavirus infections. The Coronaviridae. 1995:389-401.
- 10. McIntosh K, Kapikian AZ, Turner HC, Hartley JW, Parrott RH, Chanock RM. Seroepidemiologic studies of

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- coronavirus infection in adults and children. Am J Epidemiol. 1970;91(6):585-92.
- 11. Arden KE, Nissen MD, Sloots TP, Mackay IM. New human coronavirus, HCoV-NL63, associated with severe lower respiratory tract disease in Australia. J Med Virol. 2005;75(3):455-62.
- 12. Bastien N, Anderson K, Hart L, Caeseele PV, Brandt K, Milley D, et al. Human coronavirus NL63 infection in Canada. J Infect Dis. 2005;191(4):503-6.
- 13. Chiu SS, Chan KH, Chu KW, Kwan SW, Guan Y, Man Poon LL, et al. Human coronavirus NL63 infection and other coronavirus infections in children hospitalized with acute respiratory disease in Hong Kong, China. Clin Infect Dis. 2005;40(12):1721-9.
- 14. Ebihara T, Endo R, Ma X, Ishiguro N, Kikuta H. Detection of human coronavirus NL63 in young children with bronchiolitis. J Med Virol. 2005;75(3):463-5.
- 15. Moës E, Vijgen L, Keyaerts E, Zlateva K, Li S, Maes P, et al. A novel pancoronavirus RT-PCR assay: frequent detection of human coronavirus NL63 in children hospitalized with respiratory tract infections in Belgium. BMC Infect Dis. 2005;5(1):1-10.

- 16. Mohammadi M, Arabzadeh SAM, Mollaei HR, Monavari SH, Nikpour N. Frequency of coronavirus NL63 infection in children with upper respiratory infection by real-time PCR. Iran J Pediatr. 2020;30(3).
- 17. Woo PCY, Lau SKP, Tsoi HW, Chan KH, Wong BHL, Che XY, et al. Relative rates of non-pneumonic SARS coronavirus infection and SARS coronavirus pneumonia. Lancet. 2004;363(9412):841-5.
- 18. Madhi A, Ghalyanchilangeroudi A, Soleimani M. Evidence of human coroanvirus (229E), in patients with respiratory infection, Iran, 2015: the first report. Iran J Microbiol. 2016;8(5):316-20.
- 19. Sultani M, Mokhrari Azad T, Eshragian M, Shadab A, Naseri M, Eilami O, et al. Multiplex SYBR green real-time PCR assay for detection of respiratory viruses. Jundishapur J Microbiol. 2015;8(8):e19041.
- 20. Zeng ZQ, Chen DH, Tan WP, Qiu SY, Xu D, Liang HX, et al. Epidemiology and clinical characteristics of human coronaviruses OC43, 229E, NL63, and HKU1: a study of hospitalized children with acute respiratory tract infection in Guangzhou, China. Eur J Clin Microbiol Infect Dis. 2018;37(2):363-9.