

Flue Vaccination does not Protect Against COVID-19 Infection; A Cross-Sectional Study.

Mohammad Hossein Abbasi

IUMS: Iran University of Medical Sciences

Shahnaz Rimaz

IUMS: Iran University of Medical Sciences

Sara Esmaeili

IUMS: Iran University of Medical Sciences

Seyed Hamid Reza Faiz

IUMS: Iran University of Medical Sciences

Taghi Riahi

IUMS: Iran University of Medical Sciences

Melika Ansarin

IUMS: Iran University of Medical Sciences

Kamran Aghakhani (✉ kamranaghakhani@gmail.com)

Iran University of Medical Sciences

Research article

Keywords: SARS-Cov-2, COVID-19, Influenza, vaccine, clinical staff

Posted Date: January 14th, 2021

DOI: <https://doi.org/10.21203/rs.3.rs-143691/v1>

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Abstract

Introduction:

Vaccination is likely to be the final solution to stop the COVID-19 pandemic which has been considered as a global public health emergency. Influenza and corona viruses have previously demonstrated antigenic cross-reactivity.

Methods:

This cross-sectional study was aimed to evaluate the transmission rate and the severity of corona virus infection among health care workers with history of previous influenza vaccination. Subjects of the study were asked about their demographics, influenza vaccination history prior to pandemic, infection with Covid-19 and the severity parameters of the disease.

Results:

Influenza vaccination has no correlation in the prevalence of Covid-19 infection rate nor in the severity of the disease process among those who received flu vaccines and those who were not vaccinated. Vaccinated and unvaccinated subjects were equal in terms of sex, age and comorbidities.

Asthma has not demonstrated to contribute to the severity of the disease.

Conclusion:

Influenza vaccination, regardless of the evidence on its antigenic cross reactivity with corona virus, is not associated with lesser involvement by or any contribution to the severity of the 2019 novel SARS-COV2 disease.

Highlights

- Influenza vaccination is not recommended for prevention from infection by COVID-19
- Influenza vaccination do not improve COVID-19 infection outcome
- Asthma is not associated with a more severe course of the disease

Introduction

SARS-COV-2 or COVID-19 virus infection was initially identified in Wuhan City, China in December 2019 and became a public health emergency of international concern (PHEIC). To date, the burden brought by this pandemic has accounted to 25 million confirmed cases and 855 thousand deaths worldwide because of rapid geographical transmission and severity of the disease ^[1]. COVID-19 is associated with 2–14 days of incubation period and a life-threatening respiratory illness especially for those elderly smoker patients with simultaneous comorbidities ^{[2][3]}.

It seems that the final solution to stop the pandemic would be a safe and efficient vaccine. Many attempts have been made to find a vaccine and one example is the non-replicating adenovirus type-5 (Ad5)-vectored COVID-19 vaccine which was introduced by Feng-cai et. al. and has shown appropriate safety and efficacy at the 2nd phase of the clinical trial [4].

Previous studies have implicated the cross-reactivities between Corona and Influenza viruses on their surface antigens [5-7] which are involved in viral invasion and spread and also the overt inflammatory response to virus [8]. There has been also evidence of cross-immunity between corona virus and BCG [9] [10]. Hemagglutinin-esterase is another similar component between corona and influenza viruses which mediates virus-cell attachment and membrane fusion [11]. So, targeting this shared viral component by anti-influenza vaccine is expected to prevent cellular invasion by the corona virus.

Influenza vaccination among clinical staff in Iran has been reported to have a coverage rate of 6% which is higher than national statistics among general population [12]. Furthermore, the higher rate of clinical staffs' exposure to confirmed cases of COVID-19 makes them an appropriate subject for this study.

Methods And Materials

Clinical staff enrolled in this study were requested to fill-up a questionnaire. Subjects were asked regarding their demographics, influenza vaccination history (during the previous year and before testing positive for Covid-19), medical history of comorbidities (including asthma, diabetes mellitus (DM), hypertension (HTN), cardiovascular diseases (CVD) and immunodeficiencies), symptoms they experienced (respiratory and non-respiratory involvement), COVID-19 PCR or Chest CT-scan results and in case of confirmed disease, the severity parameters such as; home care, hospital admission, ICU admission, intubation, oxygen saturation, arterial blood gas results and disease duration. The subjects were segregated into three categories; mild, moderate and severe. Subjects who experienced mild clinical symptoms, those tested positive but were asymptomatic and those individuals confirmed positive through chest CT-scan but were treated at home with conservative measures were considered mild cases. Severe cases were those who were admitted to the hospital and in the intensive care unit (ICU), respiratory distress with respiratory rate above 30, Oxygenation index (OI) (calculated by partial pressure of oxygen (O_2) divided by fraction of inspired oxygen (FiO_2) less than 300 mmHg and those with oxygen saturation less than 93%. Those subjects whose clinical manifestation fall between these spectrums were considered moderate cases. This study was approved by the research committee of the Iran University of Medical Sciences (IUMS) with a code number: IR. IUMS. REC.1399.414.

Analysis:

Statistics of quantitative data were presented by means and variances while qualitative data were reported by their frequencies. Chi-square test was used to assess dependence between categorical variables. Parametric data were compared using student T-test and Mann-Whitney U Test. P-value equal

or less than 0.05 was considered statistically significant. Analyzes were performed using IBM SPSS version 22.

Results

Of the 510-healthcare staff that participated in the study, 33 were infected by SARS-COV2 accounting to a prevalence rate of 6.47 % among our hospital staff while 132 (25.9%) of the participants have history of influenza vaccination. 176 were males and 334 were females and the mean age of the participants was 28.94 with an SD deviation of 5.87 years. Furthermore, the subjects of the study were equal in terms of the basic variables such as age and sex.

Individuals in vaccinated and unvaccinated groups were not statistically significant in terms of frequency on gender categories ($P = 0.108$) and also were not significantly different in terms of age ($P = 0.441 =$). The prevalence and duration of comorbidities such as diabetes mellitus (DM), hypertension (HTN), asthma, cerebrovascular diseases (CVD) and immunodeficiencies were not statistically significant (P -value > 0.05) [Table 1, Table 2] between vaccinated and unvaccinated participants and so the samples in two groups were equal in their baseline characteristics.

Table 1

descriptive of qualitative statistics and comparison between vaccinated and unvaccinated individuals

Variable (N)		All cases	Influenza vaccination		Test	P-value
			Vaccinated	Unvaccinated		
Sex	Male	176	38	138	Chi-square	0.108
	Female	334	94	240		
Asthma	Positive	32	7	25	Chi-square	0.593
	Negative	478	125	353		
HTN	Positive	12	2	10	Fisher's Exact	0.740
	Negative	498	130	368		
CVD	Positive	11	3	8	Fisher's Exact	1.000
	Negative	499	129	370		
Immunodeficiency	Positive	11	2	9	Fisher's Exact	0.737
	Negative	499	130	369		
DM	Positive	5	1	4	Fisher's Exact	1.000
	Negative	505	131	374		

DM: Diabetes mellitus, HTN: hypertension, CVD: cardiovascular diseases

Table 2

descriptive (Mean \pm SD)/(Q1, Median, Q3) of quantitative statistics and comparison between vaccinated and unvaccinated individuals

Variable (N)	All cases	Influenza vaccination		Test	P-value
		Vaccinated	Unvaccinated		
Age	28.94 \pm 5.87	28.60 \pm 4.69	29.06 \pm 6.24	Mann-Whitney U Test	0.441
DD in Asthmatics	(0.0, 3.5, 17.0)	12.00 \pm 8.29	12.00 \pm 7.69	T-test	1.000
DD in HTN cases	(2.0, 4.5, 8.0)	6.10 \pm 6.04	(1.75, 4.5, 9.75)	T-test	0.726
DD in CVD cases	(2.0, 11.5, 27.25)	11.33 \pm 14.46	14.86 \pm 12.79	T-test	0.710
DD in Immunodeficiency cases	5.73 \pm 3.95	8.00 \pm 9.89	5.22 \pm 2.38	T-test	0.760

Prior influenza vaccination was not significantly associated with prevalence rate of infection by COVID-19 (P-value 0.067; Chi-square) [Table 3]. Furthermore, there was no significant difference in severity of the disease between vaccinated and unvaccinated patients with COVID-19 (P-value = 0.101; Chi-square) [Table 4].

Table 3

History of vaccination against influenza and involvement by COVID-19 disease

Influenza vaccination	Vaccinated	Unvaccinated	Sum
COVID-19			
Healthy	13	20	33
Diseased	119	358	477
SUM	132	378	510
P-value: 0.067 Chi-square test			

Table 4
History of vaccination against influenza and severity of COVID-19 disease

Influenza vaccination	Vaccinated	Unvaccinated	Sum
Severity			
Mild	4	12	16
Moderate to Severe	9	8	17
SUM	13	20	33
P-value: 0.101 Chi-square test			

163 of the participants have expressed that they experienced the COVID-19 symptoms during the past 6 months and 57 of these participants have subjected themselves to Covid-19 test and this accounted to a 35.0% testing rate among our hospital staff. Experiencing the Covid-19 symptoms regardless of the test result was not statistically significant between the vaccinated and unvaccinated participants (P-value = 297) [Table 5]. Results of the 114 PCRs or chest CT-scans showed that 33 of the participants were positive of Covid-19. The most prevalent manifestations of the disease were musculoskeletal pain, fever and cough. Disease duration was significantly higher on participants with history of influenza vaccination (P-value = 0.020) [Table 5]. One participant positive with Covid-19 without any history of influenza vaccine has been admitted to a general ward while 2 of the participants infected with the virus with previous flu vaccination were admitted to the ICU and were intubated. There was no significant difference noted on the oxygen saturation level (SaO₂) between the vaccinated and unvaccinated participants (P-value = 0.149) [Table 5]. Furthermore, we observed no significant difference in severity of the disease among asthmatics and non-asthmatic patients (P-value = 1.000, Fischer exact test).

Table 5
Descriptive of COVID-19 involvement statistics

Variable		All cases	Influenza vaccination		Test	P-value
			Vaccinated	Unvaccinated		
COVID-19 signs	Positive	163	47	116	Chi-square	0.297
	Negative	347	85	262		
COVID-19 test	Performed	114	N/A	N/A	N/A	N/A
	Not performed	396	N/A	N/A		
manifestations	Cough	40	11	29	N/A	
	Dyspnea	11	4	7		
	Fever	43	9	34		
	Musculoskeletal symptoms	57	16	41		
	Anosmia	28	8	20		
	GI symptoms	26	5	21		
	Asymptomatic	7	1	6		
COVID ward Admission (33)	Positive	1	0	1	Fisher's Exact	1.000
	Negative	32	13	19		
ICU admission (33)	Positive	2	2	0	Fisher's Exact	0.148
	Negative	31	11	20		
Intubation (33)	Positive	2	1	1	Fisher's Exact	1.000
	Negative	31	12	19		
Overall disease duration (33)		(9.50, 14.00, 20.00)	22.08 ± 11.68	12.53 ± 6.29	T-test	0.020
SaO2 (33)		(93.00, 94.00, 97.00)	(91.25, 94.00, 96.50)	95.00 ± 2.53	T-test	0.149

Discussion

This study found no significant correlation between influenza vaccination prior to the pandemic to the rate of infectivity and the severity of the disease caused by COVID-19.

Nucleocapsid protein (N protein) and spike protein (S protein) are two major surface components of Corona virus which are involved in its pathogenesis [8]. Both proteins are also present on the surface of influenza virus ^{[13][14]}. Since N protein is associated with viral assembly and budding and S protein is involved in inflammatory reactions by inducing the host immune response, influenza vaccination was expected to decrease both proliferation and the inflammatory response caused by corona virus. Hemagglutinin-esterase (HEs) is another shared viral capsid component of influenza and corona viruses which mediated host cell membrane invasion and fusion ^[11].

Regardless of the mentioned shared antigenic components between corona virus and influenza, the novel SARS-COV2 (COVID-19) virus is not identical in its antigenic components with conventional corona virus which has made it more pathogenic and consistent with a more severe life-threatening disease. S glycoprotein on the surface of SARS-COV2 has 12.8 % antigenic variety with SARS-COV, with this in mind that the spike protein accounts for immune response against the virus ^[15]. There are also additional structural loops on receptor binding (S1) and fusion (S2) domains of the spike protein on SARS-COV2 ^[16].

Our results suggest that previous influenza vaccination has no correlation with Covid-19 infection nor the severity of novel SARS-COV2 (COVID-19) disease regardless of the previously reported antigenic similarity between influenza and corona viruses could be explained by the antigenic variety of novel 2019 corona virus from its conventional form which accounts for its higher pathogenicity and severity.

The equality of the baseline characteristics such as demographic parameters and the presence of comorbidities among the vaccinated and unvaccinated participants were ensured and the confounding biases were addressed.

Also, results of the study indicated that there is no significant difference in the severity of the disease between asthmatics and non-asthmatics which is compatible with previous studies ^[17] and this could be explained by the fact that eosinophils have a prominent role in immune response against viral illnesses such as influenza virus as a determinant of the severity ^[17].

Conclusion

Influenza vaccination is not recommended for the prevention of COVID-19. It is neither effective in reducing the rate of infection nor decreases the severity of the 2019 novel SARS-COV2 disease.

Declarations

Ethics approval and consent to participate:

This study is approved by Ethics Committee of Vice Chancellor for Research & Technology, of the Iran University of Medical Sciences (IUMS) by code number: IR.IUMS.REC.1399.414. All patients and control subjects signed the informed consent.

Consent for publication

Informed consent were obtained from all patients whom clinical data were reported in this article to participate in the study.

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Conflicts of interest/Competing interests

Authors declare no financial or non-financial conflict of interest in subject matters of this study.

Authors contributions

Mohammad Hossein Abbasi: Conceptualization, Formal analysis, Investigation, Writing - Original Draft, Writing - Review & Editing, Project administration

Shahnaz Rimaz: Methodology, Investigation, Writing - Review & Editing, Project administration.

Sara Esmaeili: Conceptualization, Investigation, Writing - Original Draft, Writing - Review & Editing,

Seyed Hamid Reza Faiz: Investigation, Writing - Original Draft, Writing - Review & Editing, Project administration

Taghi Riahi: Writing - Original Draft, Writing - Review & Editing.

Melika Ansarin: Investigation, Writing - Original Draft, Writing - Review & Editing, Project administration

Kamran Aghakhani: Conceptualization, Project administration, Writing - Original Draft, Writing - Review & Editing, Project administration

Funding

This study is supported by vice chancellor for research affairs of the Iran University of Medical Sciences. This study did not receive any specific grant from any companies, funding agencies in the public, commercial, or not-for-profit sectors.

Acknowledgement:

This study has received funding from the vice chancellor of research and technology of the Iran University of Medical Sciences (IUMS) which is greatly appreciated. We would also thank Dr. Noori for his meaningful support.

References

1. <https://www.worldometers.info/coronavirus/>.
2. Li H, Liu SM, Yu XH, Tang SL, Tang CK. *Coronavirus disease 2019 (COVID-19): current status and future perspective. International Journal of Antimicrobial Agents. 2020 Mar 29:105951..*
3. Giovannoni G. *Anti-CD20 immunosuppressive disease-modifying therapies and COVID-19. Multiple Sclerosis and Related Disorders. 2020 Apr 18..*
4. Zhu FC, Guan XH, Li YH, Huang JY, Jiang T, Hou LH, Li JX, Yang BF, Wang L, Wang WJ, Wu SP. *Immunogenicity and safety of a recombinant adenovirus type-5-vectored COVID-19 vaccine in healthy adults aged 18 years or older: a randomised, double-blind, placebo-controlled, phase 2 trial. The Lancet. 2020 Jul 20.*
5. Zheng J, Perlman S. *Immune responses in influenza A virus and human coronavirus infections: an ongoing battle between the virus and host. Curr Opin Virol 2018;28:42–52..*
6. Abdella R, Aggarwal M, Okura T, Lamb RA, He Y. *Structure of a paramyxovirus polymerase complex reveals a unique methyltransferase-CTD conformation. Proc Natl Acad Sci 2020;117(9):4931–41..*
7. Zeng Q, Langereis MA, van Vliet AL, Huizinga EG, de Groot RJ. *Structure of coronavirus hemagglutinin-esterase offers insight into corona and influenza virus evolution. Proceedings of the National Academy of Sciences. 2008 Jul 1;105(26):9065-9..*
8. Amawi H, Abu Deiab GA, Aljabali AA, Dua K, Tambuwala MM. *COVID-19 pandemic: an overview of epidemiology, pathogenesis, diagnostics and potential vaccines and therapeutics. Therapeutic delivery. 2020 Apr;11(4):245 – 68..*
9. Ozdemir C, Kucuksezer UC, Tamay ZU. *Is BCG vaccination affecting the spread and severity of COVID-19?. Allergy. 2020 Apr 24..*
10. Kumar J, Meena J. *Demystifying BCG vaccine and COVID-19 relationship. Indian Pediatr. 2020 Apr 30.*
11. Zeng Q, Langereis MA, Van Vliet AL, Huizinga EG, De Groot RJ. *Structure of coronavirus hemagglutinin-esterase offers insight into corona and influenza virus evolution. Proceedings of the National Academy of Sciences. 2008 Jul 1;105(26):9065-9..*
12. Askarian M, Khazaeipour Z, McLaws ML. *Influenza vaccination uptake among students and clinical staff of a university in Iran. International journal of infectious diseases. 2009 Jul 1;13(4):476 – 82..*
13. Earnest JT, Hantak MP, Park JE, Gallagher T. *Coronavirus and influenza virus proteolytic priming takes place in tetraspanin-enriched membrane microdomains. Journal of virology. 2015 Jun 1;89(11):6093 – 104..*
14. Newcomb LL, Kuo RL, Ye Q, Jiang Y, Tao YJ, Krug RM. *Interaction of the influenza A virus nucleocapsid protein with the viral RNA polymerase potentiates unprimed viral RNA replication. Journal of virology. 2009 Jan 1;83(1):29–36..*
15. Kumar S, Maurya VK, Prasad AK, Bhatt ML, Saxena SK. *Structural, glycosylation and antigenic variation between 2019 novel coronavirus (2019-nCoV) and SARS coronavirus (SARS-CoV).*

Virusdisease. 2020 Mar 5:1–9..

16. Jaimes JA, André NM, Millet JK, Whittaker GR. *Structural modeling of 2019-novel coronavirus (nCoV) spike protein reveals a proteolytically-sensitive activation loop as a distinguishing feature compared to SARS-CoV and related SARS-like coronaviruses*. *arXiv preprint arXiv:2002.06196*. 2020 Feb 14..
17. Lindsley AW, Schwartz JT, Rothenberg ME. *Eosinophil responses during COVID-19 infections and coronavirus vaccination*. *Journal of Allergy and Clinical Immunology*. 2020 Apr 25..