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

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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 evidences 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 from 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 immunodefiencies 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.

| 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 ± 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          | Mann-Whitney U Test   |         |
| Age                           | 28.94 ± 5.87       | 28.60 ± 4.69          | 29.06 ± 6.24          | 0.441   |
|                               |                    |                       | Mann-Whitney U Test   |         |
| DD in Asthmatics (0.0, 3.5, 17.0) | 12.00 ± 8.29       | 12.00 ± 7.69          | T-test               | 1.000   |
| DD in HTN cases (2.0, 4.5, 8.0) | 6.10 ± 6.04        | (1.75, 4.5, 9.75)     | T-test               | 0.726   |
| DD in CVD cases (2.0, 11.5, 27.25) | 11.33 ± 14.46      | 14.86 ± 12.79         | T-test               | 0.710   |
| DD in Immunodeficiency cases  | 5.73 ± 3.95        | 8.00 ± 9.89           | 5.22 ± 2.38          | 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 = 0.297) [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 (SaO$_2$) 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             |           |                       |            | N/A     |
| Performed                 | 114       | N/A                   | N/A        | N/A     |
| Not performed             | 396       | N/A                   | N/A        |         |
| manifestations            |           |                       |            | N/A     |
| Cough                     | 40        | 11                    | 29         |         |
| 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 a 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

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