
Hiren Patel a*, Hetul Patel a≡, Kartik Dholakia a≡, Takshil Shah a≡, Saloni Shah a≡ and Farhin Saiyed a≡

a Faculty of Dental Science, Dharmsinh Desai University, Nadiad, India.

Authors’ contributions
This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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Original Research Article

ABSTRACT

Background & Aim: Concerns are prevailing about the safety and side effects of the mRNA vaccine for coronavirus disease 2019 (COVID-19). Hence a study aiming the effectiveness of vaccine, post vaccination symptoms, natural progression of infection in the individuals was performed.

Methods: A randomized, cross-sectional survey was performed comprising the 757 individuals from the pool of 1000 people, to investigate the side effects of the vaccines using an independent online questionnaire gathering responses from healthcare workers (HCWs) and general population with detailed review of organ systems.

Results: A total of 757 people responded, led us to the point where 476 people (62.87 %) of people experiencing minimum one symptom following the first dose of either of the two vaccine, in comparison to 188 responses (24.8%) of people experiencing at least one symptom following second dose of either of the two vaccine.

Conclusion: Commonly reported symptoms (occurrence in descending order) were soreness,
Keywords: Covid-19; m-RNA Covid vaccine; efficacy; symptoms.

1. INTRODUCTION

The COVID-19 pandemic had a dramatic health, social, and economic impact. As of the time of this report, the level of uncertainty was extremely high and exacerbated by an excess of contradictory information [1,2]. Many lay media and websites have unreliably delivered real-time numbers on new cases and deaths, often also providing unauthorized medical advice, without waiting for confirmation. The amount and variety of news has led to a massive informative overload, generating a real infodemic.

Providing the population with evidence-based scientific data is beneficial and necessary, but does not necessarily lead to individuals correctly understanding or interpreting the information. Evidence-based data runs the risk of being self-defeating if too much information is released and the public become saturated with different facts and contradictory information, leading to emotional reactions and mistrust toward decision-makers [3]. Moreover, debates among individuals and organizations who have a strong web and media presence, often provoke conflicting opinions and negative beliefs, as in the current COVID-19 situation [4,5]. One of the most discussed topics is about prevention of SARS-CoV-2 through the development of vaccines: there is a lot of information from different sources, often conflicting, that have already caused much controversy and, in some cases, been labeled as “fake news” [6,7].

Severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) has caused a global pandemic in a short period of time, imposing challenges on medical services, researchers, epidemiologists and policy makers about the nature of the virus; and posing challenges for a successful vaccine outcome. The disease is asymptomatic or mild in most patients. However, a substantial percentage of people have more extensive pneumonia that can progress to hypoxemic respiratory failure, shock, dysfunction of organs and death. Unfortunately, no effective treatment has been demonstrated to radically change the natural history of SARS-CoV-2 infection. As immunization is one of the most successful and cost-effective health interventions to prevent infectious diseases, vaccines against COVID-19 are considered to be of great importance to prevent and control COVID-19. In the month of September 2020 two vaccines produced from Serum Institute of India and Bharat Biotech received approval from the government for vaccination in India. Along with these two, several other vaccines were ongoing trials. This unparalleled initiative in developing vaccines created many uncertainties looming around the efficacy and safety of these vaccines. This study aimed to assess the side effects and efficacy following COVID-19 vaccination [8,9].

2. METHODS

Study comprised of conducting a survey with a detailed questionnaire pertaining to effectiveness of vaccine, post vaccination symptoms, natural progression of infection and treatment modalities during hospital stay or home quarantine facility if infected after getting vaccinated [10]. Our survey was based on the vaccination drive carried out initially for healthcare workers followed by mass vaccination drive over a period eight months [11]. Our survey includes people of all age groups ranging from 18 to 91 years old. The responses were filled by the individuals during follow up interviews. The question set included responses in binary fashion along with descriptive features needed in certain categories. A total of 757 responses were collected and descriptive analysis was performed (Table 1).

3. RESULTS AND DISCUSSION

A total of 757 people responded to the questionnaire given to them, which lasted for eight months from the month of September 2020 to April 2021. Among those who responded, 59.5% were healthcare workers including doctors, nurses, technicians followed by general population (40.5%). Amongst the vaccine recipients 37.8% were male and 62.2% female. From the survey responders 698 people (92.2%) were administered Covishield (manufactured by Serum Institute, India) vaccine and 59 (7.8%) people were given Covaxin vaccine (manufactured by Bharat Biotech, India).
Table 1. Overall symptomatic response after covid-19 vaccination based on age

<table>
<thead>
<tr>
<th>Crosstab</th>
<th>Symptomatic overall</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Asymptomatic</td>
<td>Symptomatic</td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;20</td>
<td>Count 22</td>
<td>226</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 8.3%</td>
<td>46.0%</td>
</tr>
<tr>
<td>21-30</td>
<td>Count 27</td>
<td>147</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 10.2%</td>
<td>29.9%</td>
</tr>
<tr>
<td>31-40</td>
<td>Count 22</td>
<td>31</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 8.3%</td>
<td>6.3%</td>
</tr>
<tr>
<td>41-50</td>
<td>Count 37</td>
<td>26</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 13.9%</td>
<td>5.3%</td>
</tr>
<tr>
<td>51-60</td>
<td>Count 38</td>
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<td>% within symptomatic overall 14.3%</td>
<td>1.8%</td>
</tr>
<tr>
<td>61-70</td>
<td>Count 69</td>
<td>36</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 25.9%</td>
<td>7.3%</td>
</tr>
<tr>
<td>71-80</td>
<td>Count 44</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 16.5%</td>
<td>2.9%</td>
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<tr>
<td>&gt;80</td>
<td>Count 7</td>
<td>2</td>
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<tr>
<td></td>
<td>% within symptomatic overall 2.6%</td>
<td>0.4%</td>
</tr>
<tr>
<td>Total</td>
<td>Count 266</td>
<td>491</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 100.0%</td>
<td>100.0%</td>
</tr>
</tbody>
</table>

Chi-Square Tests

Value df P value (<0.05 is significant)
Pearson Chi-Square 256.345 7 <0.001

Table 2. Overall symptomatic response after covid-19 vaccination based on gender

<table>
<thead>
<tr>
<th>Crosstab</th>
<th>Symptomatic overall</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Asymptomatic</td>
<td>Symptomatic</td>
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<td>Gender</td>
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<td>F</td>
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<td>340</td>
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<tr>
<td></td>
<td>% within symptomatic overall 49.2%</td>
<td>69.2%</td>
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<tr>
<td>M</td>
<td>Count 135</td>
<td>151</td>
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<td></td>
<td>% within symptomatic overall 50.8%</td>
<td>30.8%</td>
</tr>
<tr>
<td>Total</td>
<td>Count 266</td>
<td>491</td>
</tr>
<tr>
<td></td>
<td>% within symptomatic overall 100.0%</td>
<td>100.0%</td>
</tr>
</tbody>
</table>

Chi-Square Tests

Value df P value (<0.05 is significant)
Pearson Chi-Square 29.353 1 <0.001
Table 3. Symptoms after first dose of covid-19 vaccine

<table>
<thead>
<tr>
<th></th>
<th>Fever After First Dose</th>
<th>Weakness after first dose</th>
<th>Headache and Body Ache after first dose</th>
<th>Malaise after first dose</th>
<th>Tachycardia after first dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>421</td>
<td>449</td>
<td>456</td>
<td>698</td>
<td>732</td>
</tr>
<tr>
<td>Yes</td>
<td>336</td>
<td>308</td>
<td>301</td>
<td>59</td>
<td>25</td>
</tr>
<tr>
<td>Total</td>
<td>757</td>
<td>757</td>
<td>757</td>
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</tr>
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</table>

Table 4. Symptoms after second dose of covid-19 vaccine

<table>
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<tr>
<th></th>
<th>Fever after Second Dose</th>
<th>Weakness after Second dose</th>
<th>Headache and Body Ache after Second dose</th>
<th>Malaise after Second dose</th>
<th>Tachycardia after Second dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>722</td>
<td>663</td>
<td>692</td>
<td>745</td>
<td>740</td>
</tr>
<tr>
<td>Yes</td>
<td>35</td>
<td>94</td>
<td>65</td>
<td>12</td>
<td>17</td>
</tr>
<tr>
<td>Total</td>
<td>757</td>
<td>757</td>
<td>757</td>
<td>757</td>
<td>757</td>
</tr>
</tbody>
</table>

Charter 1. Treatment modalities after covid-19 infection

Out of total 757 responses 92.2% received both the doses of vaccines. This includes 651 people administered Covishield and 47 administered Covaxin. The remaining 7.8% of people did not receive the second dose. 11.3% of people experienced decrease in their oxygen level and 94.3% experienced the need of oxygen during the disease course.

Post vaccination a majority of the healthcare worker falling under the age of 30 were found to more symptomatic compared to the rest of the age groups (Table 1). Females were found to be predominantly more symptomatic compared to the males irrespective of their age (Table 2).

The need of administering steroids was found in 13.6% people and other drugs 12.6%. 20.5% people were prescribed anticoagulants (Charter 1).

All over the level of acceptance of the vaccine was high and majority of the people did acknowledge the vaccine benefits.
Our healthcare workers who responded to the survey stated few to none symptoms after vaccination where fever headache body ache being the most common symptoms. None of the people were found to develop any serious post vaccination symptom or complication (Table 3 and Table 4).

Majority of vaccine recipients did not contract the disease and they found the vaccine to be effective and beneficial in order to cease the spread of the disease. The people who were infected by the virus even after vaccination showed a mild disease progression leading to no serious aftermath effect. Our findings can be co-related to the results of the various published studies and trials.

According to those receiving two standard doses of Astra oxford ChAdOx1 nCoV-19, after the prime vaccination local reactions were reported in 43 (88%) of 49 participants in the 18-55 years group, 22 (73%) of 30 in the 56-69 years group, and 30 (61%) of 49 in the 70 years and older group, and systemic reactions in 42 (86%) participants in the 18-55 years group, 23 (77%) in the 56-69 years group, and 32 (65%) in the 70 years and older group. As of now various other vaccines have been approved for by the government, one often being Sputnik V, Moderna, Zycov-D, J&J. In terms of efficacy, both Covishield and Covaxin were highly effective in the overall population [10,12].

There was a small pool of subjects due to which mass response to the vaccine cannot be formulated. The symptoms being highly subjective can also lead to overestimation of the symptoms. The follow up and responses were taken into account within two months of vaccination, leading to little or no knowledge regarding post two months.

4. CONCLUSION

Two third of the population did experience at least one symptoms post vaccination. Fever, headache and body ache being the most common symptom. The symptoms were more common with the females as compared to the males. No serious events were noted. Only 11.6% of the vaccinated people contracted the disease leading to positive outcome from the vaccine making us believe in the vaccine efficacy. The severity of the infection was mild in people getting infected even after vaccination which again leads us to the positive side of vaccination.

DISCLAIMER

The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

CONSENT

As per international standard or university standard, patients’ written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES


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