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Original Research Article

Immunological response in health care workers post COVID-19 vaccination

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ABSTRACT

Background: To curb SARS-CoV-2, various vaccines received Emergency Authorization Use by copious agencies. These vaccines are administered to health-care workers, front-line workers, elderly individuals and many others. Health care workers were the first to be administered with vaccines, thus, evaluating the antibody response helps to portray a picture of the burden of SARS-CoV-2 infection amid Health Care Workers.

Materials and Methods: The present study aimed to investigate the production of total antibodies against SARS-CoV-2 before and after vaccination (2 doses) among Health Care Workers along with associated factors. The present study was conducted on 60 healthcare workers as cases who got vaccinated for COVID-19. The semi-quantitative detection of anti-SARS-CoV-2 IgG Ab in the samples was detected using Erba Lisa COVID-19 total antibody indirect ELISA kit using standard protocol as described by the manufacturer. **Results:** In the present study, co-morbidities such as hypertension was presented in only 8 (12.3%) individuals, diabetes in 4 (6.1%) and Coronary Artery Disease in 1 (1.5%). After first dose of administration, 86.67% HCWs were found seroprositive for the anti-SARS-CoV-2 Ab while 13.33% were seronegative, however, after second dose 100% seroprevalence was observed amid HCWs. The statistical analysis depicted a significant difference (p<0.0001) in antibody titres among individuals before and after vaccination. The mean of total antibody levels detected at zero week was (4.1), 4th week, (39.09), 8th week (79.39) and 12th week (105).

Conclusion: The present study provides important information on the detectable levels of antibodies induced by COVISHIELD against the SARS-CoV-2 which was independent of gender and age.

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1. Introduction

Severe acute respiratory syndrome corona-virus 2 (SARS-CoV-2), the causative agent of COVID-19, has led to a global pandemic, affecting 290 million people and 5.4 million deaths till 7^{th} January 2022. To contain this brunt, various novel vaccines received an Emergency Use Authorization (EUA) by the U.S.A Food Drug Administration (FDA), European Medicine Agency (EMA), U.K. Medicines and Healthcare Products Regulatory

Agency (MHRA) and Indian Central Drugs Standard Control Organization (CDSCO) as well as Drugs Controller General of India (DCGI).¹ About 910 million people are administered with vaccine doses. These vaccines are administered to health-care workers, front-line workers, elderly individuals including those with comorbidities. Since health care workers are at the front line of the ongoing pandemic, they were the first to receive the vaccines, thus, evaluating the antibody response aids to render a snapshot of the burden of SARS-CoV-2 infection amid HCWs.² Various studies on immune response for SARS-

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CoV-2 showed that antibodies were present 7-14 days after the onset of symptoms.^{3–5} These antibodies were found to decline with the passage of time but remain detectable for several months following infection.^{6,7} It was observed that SARS-CoV-2 antibodies have neutralizing capacity in-vitro and shield against reinfection in study models.⁸ Till today, early data has been published on immunogenicity and safety of the vaccine, however, associated factors such as duration of immune response and rate of antibody production is yet to uncover.^{9,10} Therefore, the present study aimed to investigate the production of total antibodies against SARS-CoV-2 before and after vaccination (2 doses) among HCWs along with associated factors such as co-morbidities and adverse reaction after vaccination.

2. Materials and Methods

The present study was conducted on 60 healthcare workers (34 males and 26 females within the age group of 22-64 years) as cases who got vaccinated for COVID-19 vaccine at Government Medical College and Hospital, Amritsar. Before commencement of the study, ethical permission and approval was obtained from the Ethics Commission at Viral Research and Diagnostic Laboratory (VRDL), Government Medical College (GMC), Amritsar, India. Individuals vaccinated with COVID-19 vaccine were enrolled as cases while individuals with previous SARS-CoV-2 history and/or were symptomatic at the time of sample collection were excluded. About 5ml of blood sample was withdrawn to detect COVID-19 specific IgG Ab base line titre. The first blood samples were taken at day 0 before vaccination followed by 4^{th} week, 8^{th} week, 12^{th} week, 24th week post vaccination. The semi-quantitative detection of anti-SARS-CoV-2 IgG Ab in the samples was detected using ErbaLisa COVID-19 IgG indirect ELISA kit using standard protocol as described by manufacture. Antibody titre was evaluated on the basis of cut off values following comparison between negative control and sample. The sample was evaluated negative if its value was similar to the negative control whereas if the value obtained was atleast 2 times higher than cut off value of negative control it was considered as positive.

3. Results

The present study was conducted on 60 health care workers vaccinated for COVID-19 at Government Medical College, Amritsar. Co-morbidities such as hypertension was presented in only 8 (12.3%) individuals, diabetes in 4 (6.1%) and coronary artery disease in 1 (1.5%). After vaccination, 56% of individuals experienced weakness, 79% had fever while malaise, joint pains and body aches were reported in 33.33%, 3.33% and 16.67% subjects, respectively.

In this study, after first dose of administration, 86.67% HCWs were found seropositive for the anti-SARS-CoV-2

Abs while 13.33% were seronegative, however, after second dose, 100% seroprevalence was observed amid HCWs (Table 1). The statistical analysis depicted a significant difference (p<0.0001) in antibody titres among individuals before vaccination and after vaccination. The mean of total antibody levels at zero week (4.1), 4th week, (39.09), 8th week (79.39) and 12th week (105). Overall, no statistically significant difference was observed between antibody response, gender and age (Figures 1 and 2).

n-66; Day Zero:- Total Ab Index before Vaccination (p Value-); Day 28:- Total Ab index after 1st Dose 1^{st} sampling (p Value-); Day 56:- Total Ab index after 2^{nd} dose 1^{st} sampling (p Value-); Day 84:- Total Ab index after 2^{nd} Dose 2^{nd} sampling (p-value < 0.05)

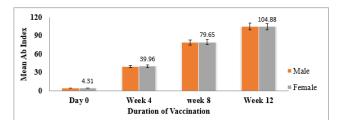


Fig. 1: Mean rise of total antibody levels in an individual with the duration of vaccination

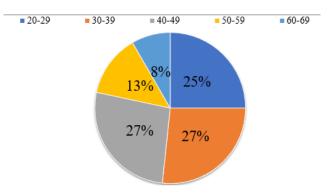


Fig. 2: Age wise distribution of healthcare workers involved in the study

4. Discussion

Vaccination plays a dynamic role in reducing the COVID-19 pandemic. The key question that needs to be addressed here is that how much protection does vaccines provide in controlling the infection by producing more antibodies. ChAdOx1 nCoV -19 (AZD1222) vaccine has been found to induce Anti-spike antibody, antibody-dependent monocyte/neutrophil phagocytosis, and cellular responses.¹¹

In this study, total antibody levels were measured by ELISA for comparing the results before and after vaccination among healthcare workers. Significantly higher

 Table 1: Total antibody index among health care workers before and after vaccination

Sr.No.	Age	Gender	Day Zero	Day 28	Day 56	Day 84
V1	37	female	1.3 ± 0.02	70.28 ± 0.02	80.3 ± 0.04	99.2 ± 0.03
v2	44	female	2.3 ± 0.01	12.6 ± 0.03	84.7 ± 0.05	120.84 ± 0.04
v3	47	female	3.2 ± 0.01	30.74 ± 0.01	101.6 ± 0.08	109.5 ± 0.02
v4	48	male	0.6 ± 0.01	40.2 ± 0.07	85.6 ± 0.04	116.79 ± 0.05
v5	60	male	1.3 ± 0.03	80 ± 0.05	100.1 ± 0.08	121.61 ± 0.09
v6	52	female	3.5 ± 0.03	30.24 ± 0.08	84.4 ± 0.04	102.8 ± 0.04
v0 v7	53	female	6.7 ± 0.02	45.78 ± 0.03	85.6 ± 0.01	102.0 ± 0.04 100.9 ± 0.04
v8						
	63 26	male	2.1 ± 0.02	44.08 ± 0.01	83.1 ± 0.04	110.52 ± 0.09
v9	26	male	5.2 ± 0.01	24.67 ± 0.05	80 ± 0.03	108 ± 0.08
v10	64	male	0 ± 0.00	3.39 ± 0.02	82.5 ± 0.02	45.6 ± 0.05
v11	46	male	3.4 ± 0.02	41.91 ± 0.04	90.3 ± 0.04	100.1 ± 0.04
v12	58	male	7.2 ± 0.01	55.62 ± 0.06	85.6 ± 0.02	125.7 ± 0.02
v13	48	female	1.3 ± 0.02	50.32 ± 0.08	90.4 ± 0.06	124.8 ± 0.01
v16	47	male	5 ± 0.02	30.16 ± 0.05	54.1 ± 0.02	89.6 ± 0.04
v17	40	male	7 ± 0.03	40.84 ± 0.03	88.1 ± 0.01	102.2 ± 0.03
v18	57	male	4.5 ± 0.02	54.61 ± 0.06	83.1 ± 0.02	94.4 ± 0.09
v19	63	male	6 ± 0.05	38.8 ± 0.02	90.5 ± 0.09	138.8 ± 0.03
v20	28	female	2.4 ± 0.02	48.13 ± 0.01	87.2 ± 0.04	113.8 ± 0.04
v21	52	female	5.2 ± 0.01	28.02 ± 0.01	62.2 ± 0.07	83 ± 0.04
v22	32	female	6.2 ± 0.01	43.94 ± 0.07	63.1 ± 0.03	99.6 ± 0.02
v23	35	female	5.7 ± 0.02	36.14 ± 0.09	79.7 ± 0.09	102.7 ± 0.05
v23 v24	36	female	1.4 ± 0.02	9.74 ± 0.03	30.6 ± 0.02	80.9 ± 0.03
	34			9.74 ± 0.03 60.56 ± 0.04		
v25		male	4.2 ± 0.02		88.8 ± 0.03	107.2 ± 0.06
v26	45	male	4.2 ± 0.02	4.73 ± 0.01	82.8 ± 0.02	96.5 ± 0.07
v27	26	male	6.4 ± 0.01	41.28 ± 0.04	80.3 ± 0.01	137.08 ± 0.09
v29	25	male	2.4 ± 0.02	57.45 ± 0.05	84.1 ± 0.04	104.3 ± 0.03
v30	23	female	5.4 ± 0.06	29.7 ± 0.07	40.11 ± 0.01	102.3 ± 0.04
v31	24	male	7.1 ± 0.05	40.61 ± 0.07	89.4 ± 0.08	99.4 ± 0.05
v32	22	male	4.2 ± 0.03	49.85 ± 0.03	81.5 ± 0.03	103.3 ± 0.09
v33	34	female	6.4 ± 0.02	37.36 ± 0.06	87.2 ± 0.07	112 ± 0.07
v34	31	female	7.1 ± 0.01	16.61 ± 0.04	70 ± 0.06	105.7 ± 0.06
v35	31	female	3.4 ± 0.02	14.82 ± 0.05	62.5 ± 0.05	91.2 ± 0.03
v36	55	female	1.4 ± 0.02	66.2 ± 0.03	91.9 ± 0.05	109.1 ± 0.06
v37	62	male	5.2 ± 0.05	21.1 ± 0.01	82.8 ± 0.03	107.6 ± 0.04
v38	29	female	6.8 ± 0.01	40.37 ± 0.02	78.4 ± 0.09	115.23 ± 0.12
v40	32	male	1.9 ± 0.06	68.09 ± 0.07	89.4 ± 0.01	102.5 ± 0.09
v42	46	female	2.1 ± 0.02	7.19 ± 0.03	78.8 ± 0.04	45.4 ± 0.02
v43	45	female	4.5 ± 0.02	63.36 ± 0.06	82.2 ± 0.03	105 ± 0.02 105 ± 0.03
					98.4 ± 0.06	105 ± 0.05 126.52 ± 0.15
v44	50	male	4.6 ± 0.03	72.3 ± 0.06		
v45	48	female	6.4 ± 0.08	43.42 ± 0.03	83.4 ± 0.01	113.1 ± 0.12
v46	53	male	3.7 ± 0.01	61 ± 0.06	79.4 ± 0.05	111.8 ± 0.08
v48	47	male	1.7 ± 0.02	73.22 ± 0.02	81.6 ± 0.07	118.31 ± 0.10
v49	48	male	8.9 ± 0.02	43.5 ± 0.05	90.6 ± 0.04	106.6 ± 0.05
v50	35	female	2.4 ± 0.01	56.68 ± 0.01	82.5 ± 0.01	99.52 ± 0.04
v51	34	female	8.3 ± 0.01	71.43 ± 0.03	79.7 ± 0.02	89.3 ± 0.08
v52	43	female	2.8 ± 0.02	17.27 ± 0.04	83.7 ± 0.05	100.7 ± 0.08
v53	24	female	4.2 ± 0.01	41.37 ± 0.01	78.52 ± 0.04	98.52 ± 0.04
v54	34	male	2.4 ± 0.02	6.94 ± 0.04	66.9 ± 0.06	96.9 ± 0.09
v55	45	male	1.8 ± 0.02	37.52 ± 0.03	82.8 ± 0.02	116.63 ± 0.12
v56	33	male	4.8 ± 0.02	32.89 ± 0.01	80.52 ± 0.02	107.7 ± 0.03
v50 v57	48	male	7.2 ± 0.01	9.72 ± 0.02	58.5 ± 0.02	107.7 ± 0.03 102.1 ± 0.07
v58	23	male	4.2 ± 0.02	31.36 ± 0.03	60.3 ± 0.03	96.4 ± 0.10
v59	26 26	male	6.2 ± 0.03	42.45 ± 0.04	88.1 ± 0.08	132.74 ± 0.09
v60	36	male	7.3 ± 0.04	41.74 ± 0.04	77.8 ± 0.02	104.9 ± 0.04
v61	25	female	3.8 ± 0.02	57.87 ± 0.03	82.8 ± 0.03	100.6 ± 0.02
v62	27	male	4.9 ± 0.01	5.52 ± 0.01	58.6 ± 0.08	98.7 ± 0.02
v63	33	female	6.2 ± 0.02	52.17 ± 0.04	82.8 ± 0.04	121.74 ± 0.05
v64	35	male	7.3 ± 0.04	11.99 ± 0.02	70.6 ± 0.09	103.5 ± 0.08
v65	25	male	0.2 ± 0.01	9 ± 0.01	75 ± 0.04	115.25 ± 0.09
v66	27	male	3.4 ± 0.02	46.69 ± 0.03	78.1 ± 0.02	111.8 ± 0.10

levels of antibodies were observed after first and second dose of vaccination as compared to prior vaccination among all healthcare workers (p<0.0001). A study conducted on 515 individuals who were administered both the doses of vaccination showed that 95.0% were observed to be seropositive for the anti-SARS-CoV-2 antibodies.¹² A comparative study on HCW and non-HCW demonstrated no significant difference in seroprevalence among health care workers.¹³ Zhang et al., 2021¹⁴ observed higher Ab measurements in subjects who had SARS-CoV-2 infection followed by 2 doses of vaccination as compared to individuals who got vaccinated without prior infection. A cohort study on 2607 subjects also observed significantly higher antibody levels (p<0.001) after vaccination.¹⁵ A meta-analysis of 49 studie showed overall seroprevalence of 8.7% among HCWs.16 A previous study performed at Oxford University Hospital reported that seroprevalence of SARS-CoV-2 antibodies after first dose in participants was 94% within 15-21 days and 97% in 22-28 days.¹⁷

Our findings suggest that there is no correlation between Ab response, gender and age. A study by Uysal et al., 2021.¹⁸ and Singh et al., 2021¹² showed similar findings indicating no relationship between age and antibody levels. However, Kataria et al., 2021¹³ found that age had significant impact on Ab responses which was less in individuals with age >60 years (30.6%) as compared to age \leq 60 years (46.4%). Study by Salvong et al., 2021¹⁹ showed that age and gender were independent predictors of antibody levels. Few other studies have reported gender differences in response to COVID-19 vaccinations.^{20–23} These differences in correlation among Ab response, gender and age can be attributed to variation in population demographics in different regions or countries depicting weaker or strong immune responses in individuals of different age groups.

In our study, only 12.3% individuals had hypertension, 6.1% had diabetes and 1.5% suffered from CAD, however, none of these co-morbidities had any impact on Ab titre. Singh et al., 2021, demonstrated that people with type 2 diabetes showed less seropositivity rate in comparison to those without T2D (p<0.001) after completion of both doses. Post vaccination effects were seen in many cases and the most common being weakness followed by fever, malaise, joint pains and body aches in the present study. Other studies indicated mild to moderate side effects post vaccination.¹² Local and systemic adverse reactions, including injection site pain, chills, muscle ache, and headache, have also been reported by Folegetti et al., 2020.²⁴ According to Sasso et al., 2021,¹⁵ 1184 out of 2607 subjects displayed one or more adverse reactions after vaccination which includes muscular pain and systemic effects. Jeong et al., 2021²⁵ observed that adverse reactions were significantly associated with positivity rate and these variables may be used as parameters to estimate the formation of anti-SARS-CoV-2 antibodies.

5. Conclusion

This study renders paramount information on the detectable levels of antibodies induced by ChAdOx1 nCOV-19 against the SARS-CoV-2 after first dose and second dose in the population of Punjab. Robust immunity induced by the above said vaccine against COVID-19 measured by total IgG ELISA was also reported previously.²⁴ Since the vaccines for SARS-CoV-2 have been recently developed, questions pertaining to the durability of protection over the period after vaccination and determination of effect of booster dose to extend the duration of immunity for COVID-19, are yet to be answered. Thus, it is important to understand that serological testing is important prior to vaccination to analyse the humoral immune response of the subject, so that the vaccination protocol could be adapted. This approach could accelerate vaccination regimes, facing new variants, achieving immunity quickly, stopping the spread and hindering emergence of new variants.

6. Contribution

Manuscript writing and analysis: Amandeep Kaur, Mohan Jairath, Savjot Kaur Analysis, Writing and Review: Shailpreet Kaur, Kanwardeep Singh

7. Source of Funding

None.

8. Conflict of Interest

None.

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References

- Chou R, Dana T, Buckley DI, Selph S, Rongwei F, Totten A. Epidemiology and risk factors for coronavirus infection in health care workers: a living rapid review. *Ann Intern Med.* 2020;173(2):120–56.
- Long QX, Liu BZ, Deng HJ, Gui CW, Deng K, Chi CY. Antibody responses to SARS-CoV-2 in patients with COVID-19. *Nat Med.* 2020;26(6):845–53.
- Wang Y, Zhang L, Sang L, Feng Y, Ruan S, Zhong B, et al. Kinetics of viral load and antibody response in relation to COVID-19 severity. *J Clin Invest*. 2020;130(10):5235–79.
- Ni L, Ye F, Cheng ML, Feng Y, Qian DY, Zhoa H. Detection of SARS-CoV-2-specific humoral and cellular immunity in COVID-19 convalescent individuals. *Immunity*. 2020;52(6):971–8.
- Okba N, Muller MA, Li W, Chunyan W, Kessel G, Corman C. Severe acute respiratory syndrome coronavirus 2-specific antibody responses in coronavirus disease patients. *Emerg Infect Dis.* 2020;26(7):1478– 88.
- Ibarrondo FJ, Fulcher JA, Meza DG, Elliott J, Hofmann C, Hausner M. Rapid decay of anti-SARS-CoV-2 antibodies in persons with mild COVID-19. N Engl J Med. 2020;383(1):1085–92.

- Gudbjartsson DF, Norddahl GL, Melsted P, Gunnarsdottir K, Holm H, Eythorsson E, et al. Humoral immune response to SARS- CoV-2 in Iceland. *N Engl J Med*. 2020;383(18):1724–58.
- Deng W, Bao L, Liu J, Xiao C, Liu J, Xue J, et al. Primary exposure to SARS-CoV-2 protects against reinfection in rhesus macaques. *Science*. 2020;369(6505):818–41.
- Teijaro JR, Farber DL. COVID-19 vaccines: modes of immune activation and future challenges. *Nat Rev Immunol*. 2020;21:195–7.
- Emary KRW, Golubchik T, Aley PK, Ariani CV. Oxford COVID-19 Vaccine Trial Group, et al. 2021. Efficacy of ChAdOx1 nCoV-19 (AZD1222) vaccine against SARS-CoV-2 variant of concern 202012/01 (B.1.1.7): an exploratory analysis of a randomised controlled trial. *Lancet*. 2021;397:1351–1362.
- Barrett JR, Rammerstorfer SB, Dold C. Oxford COVID Vaccine Trial Group. Phase ¹/₂ trial of SARS-CoV-2 vaccine ChAdOx1 n CoV-19 with a booster dose induces multifunctional antibody responses. *Nat Med.* 2020;27:279–88.
- Singh AK, Phatak SR, Singh R, Bhattacharjee K, Singh NK, Gupta A, et al. Antibody response after first and second-dose of ChAdOx1-nCOV (CovishieldTM!) and BBV-152 (CovaxinTM!) among health care workers in India: The final results of cross-sectional coronavirus vaccine-induced antibody titre (COVAT) study. *Vaccine*. 2021;39(44):6492–509.
- Kataria S, Phogat R, Sharma P, Deswal V, Alam S, Singh M. Is Covid-19 seroprevalence different in health care workers as per their risk of exposure? A study from a tertiary care hospital in National Capital Region of India. *BMJ*;p. 1–17. doi:10.1101/2021.02.10.21251543.
- Zhang D, Xiao S, Debes AK, Egbert ER, Caturegli E, Colantuoni E. Durability of Antibody Levels After Vaccination With mRNA SARS-CoV-2 Vaccine in Individuals with or without Prior Infection. *JAMA*. 2021;326(24):1–3.
- Sasso BL, Giglio RV, Vidali M, Scazzone C, Bivoan G, Gambino CM, et al. Evaluation of Anti-SARS-Cov-2 S-RBD IgG Antibodies after COVID-19 mRNA BNT162b2 Vaccine. *Diagnostics*. 2021;11(7):1135. doi:10.3390/diagnostics11071135.
- Galanis P, Vraka I, Fragkou D, Bilali A, Kaitelidou D. Seroprevalence of SARS-CoV-2 antibodies and associated factors in health care workers: a systematic review and meta-analysis. J Hosp Infect. 2021;108:120–54.
- Eyre DW, Lumley SF, Wei J, Cox S, James T, Justice A, et al. Quantitative SARS-CoV-2 anti- spike responses to Pfizer-BioNTech and Oxford-AstraZeneca vaccines by previous infection status. *Clin Microbio Infect*. 2021;27(10):1516–23.
- Uysal EB, Gumus S, Bektore B, Bozkut H, Gozalan A. Evaluation of antibody response after COVID-19 vaccination in health care workers. *J Med Virol.* 2021;94(3):1–7.
- Salvagno GL, Henry BM, Piazza GD, Pighi L, Nitto D, Bragantini S. Anti-SARS-CoV-2 Receptor-Binding Domain Total Antibodies

Response in Seropositive and Seronegative Healthcare Workers Undergoing COVID-19 mRNA BNT162b2 Vaccination. *Diagnostics*. 2021;11(5):832. doi:10.3390/diagnostics11050832.

- Rose D, Suther N, Makowski MS. Antibody Persistence through 6 months after the Second Dose of mRNA-1273 Vaccine for Covid-19. *N Engl J Med.* 2021;384(23):2259–61.
- Muller L, Moskorz AM, W. Age-independent immune response to the Biontech/Pfizer BNT162b2 COVID-19 vaccination. *Clin Infect Dis.* 2021;73:2065–2072.
- Shrotri M, Fragaszy E, Geismar C. Spike-antibody responses following first and second dosed of ChAdOx1 and BNT162b2 vaccines by age, gender, and clinical factors- a prospective community cohort study (Virus Watch). *Lancet*. 2021;398(10298):385–7.
- Nace NA, Kip KE, Palmer O. Antibody Responses in Elderly Residential Care Persons following COVID-19 m RNA Vaccination. J Am Med Dir Assoc. 2021;22:1593–8. doi:10.1016/j.jamda.2021.06.006.
- 24. Folegatti PM, Ewer KJ, Aley PK, Angus B, Becker S, Belij-Rammerstorfer S, et al. Safety and immunogenicity of the ChAdOx1 nCoV-19 vaccine against SARS-CoV-2: a preliminary report of a phase ¹/₂, single-blind, randomized controlled trail. *Lancet.* 2020;396:467–78.
- Jeong S, Lee N, Lee SK, Cho EJ, Hyun J. Comparing results of five SARS dose of ChAdOx1 nCoV-19 vaccine among health care workers. J Clin Microbiol. 2021;59(9):1105–26.

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