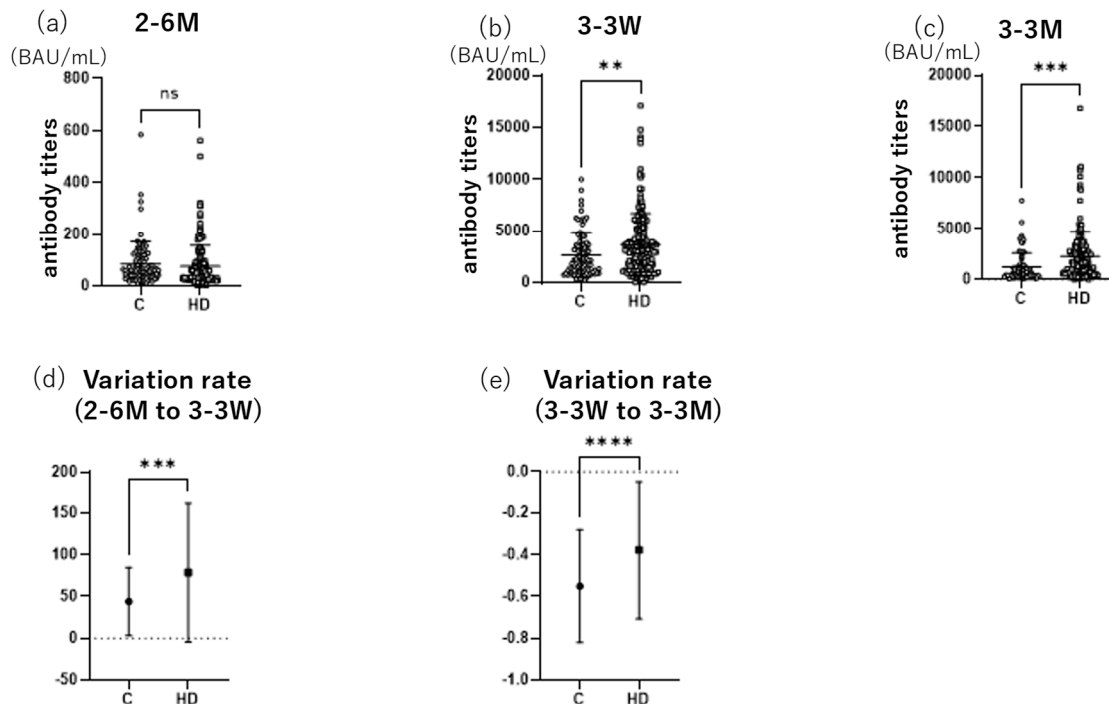
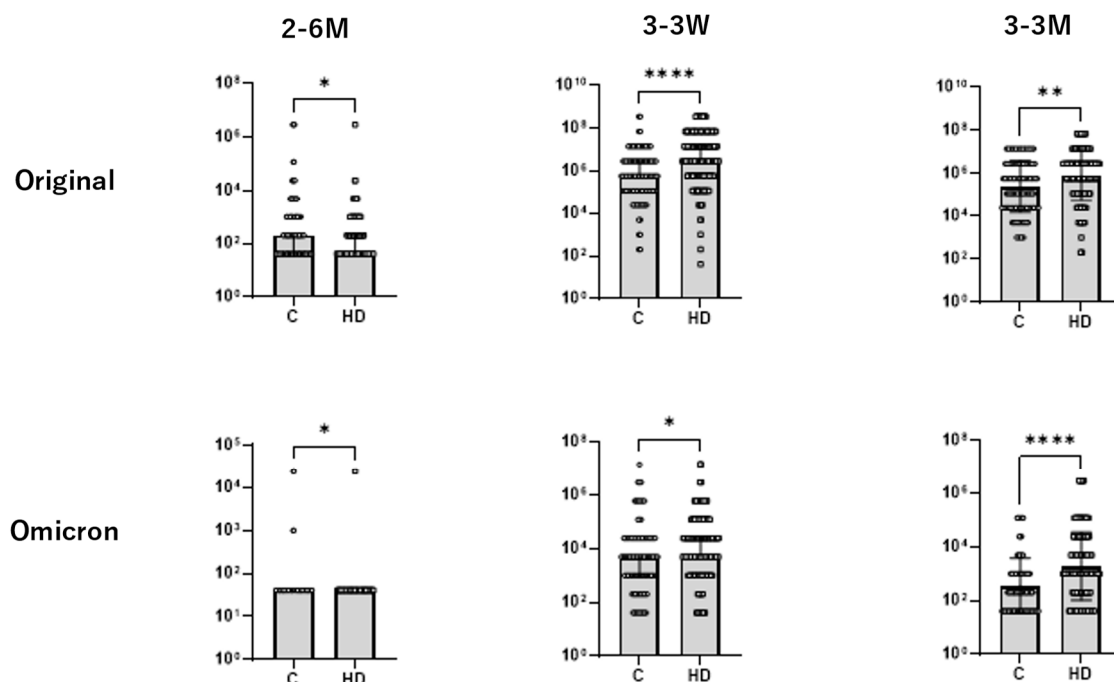




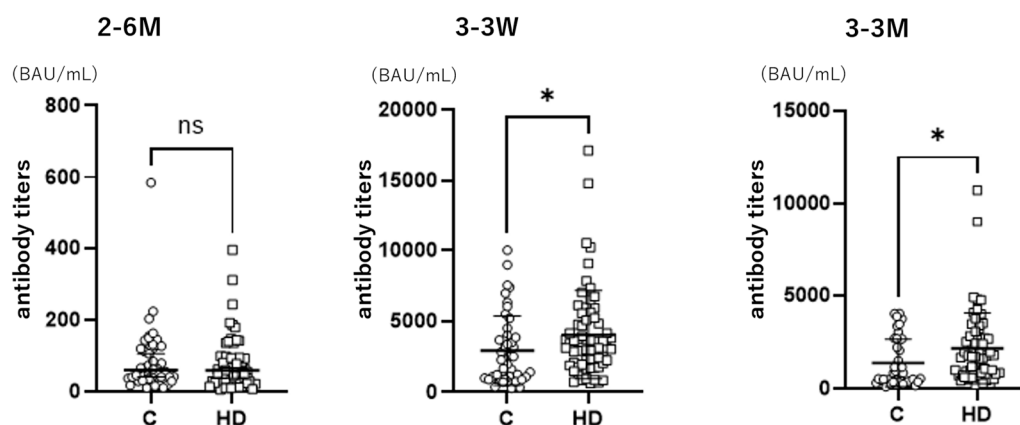
## Supplementary Materials



**Figure S1.** Comparison of antibody titers between the control and HD groups (only for participants who participated in all blood tests). At 6 months after the primary series, blood was obtained from 297 participants (control group,  $n=103$ ; HD group,  $n=194$ ); however, 29 participants withdrew from the study at 3 months after the booster dose. Blood was obtained from 258 participants (control group,  $n=82$ ; HD group,  $n=176$ ) at 3 months after the booster dose. (a) 6 months after the primary series. (b) 3 weeks after the booster dose. (c) 3 months after the booster dose. (d) Value at 3 weeks after the third vaccine - value at 6 months after the primary series/value at 6 months after the primary series. (e) Value at 3 months after the third vaccine - value at 3 weeks after the third vaccine/value at 3 weeks after the third vaccine. Bars indicate the median with a 95% confidence interval. \* $P \leq 0.05$ , \*\* $P \leq 0.01$ , \*\*\* $P \leq 0.001$ , \*\*\*\* $P \leq 0.0001$ . C: control; HD: hemodialysis. 2-6M: 6 months after the primary series; 3-3W: 3 weeks after the booster dose; 3-3M: 3 months after the booster dose.



**Figure S2.** Comparison of neutralizing antibody titers between the control and HD groups. Neutralizing antibody titers in the control and HD groups were compared at three-time points: (a) 6 months after the primary series (control group,  $n=103$ ; HD group,  $n=194$ ); (b) 3 weeks after the booster dose (control group,  $n=97$ ; HD group,  $n=182$ ); and (c) 3 months after the booster dose (control group,  $n=82$ ; HD group,  $n=176$ ). Bars indicate the median with a 95% confidence interval \* $p\leq 0.05$ , \*\* $p\leq 0.01$ , \*\*\* $p\leq 0.001$ , \*\*\*\* $p\leq 0.0001$ . 2-6M: 6 months after the primary series; 3-3W: 3 weeks after the booster dose; 3-3M: 3 months after the booster dose; C: control; HD: Hemodialysis.



**Figure S3.** Comparison of antibody titers of the participants in the control and HD groups evaluated using the T-SPOT®.COVID test (T-SPOT). The severe acute respiratory syndrome coronavirus 2

(SARS-CoV-2) immunoglobulin (IgG) levels of the control and HD groups were compared. (a) 6 months after the primary series. (b) 3 weeks after the booster dose. (c) 3 months after the booster dose. Bars indicate the median with a 95% confidence interval \* $P \leq 0.05$ . C: control; HD: hemodialysis. 2-6M, 6 months after the primary series; 3-3W, 3 weeks after the booster dose; 3-3M, 3 months after the booster dose.

**Table S1.** Questionnaire about adverse reactions after the booster vaccination

Questionnaire code	
<b>Questionnaire on adverse reactions to booster vaccination.</b>	
I received the booster dose of the vaccine on date ____ month ____ 2021.	
Please answer the following questions about adverse reactions to the booster vaccination.	
1. Did you have any pain at the injection site?	Yes (lasted for ____ days) / No
2. Did you experience redness at the injection site?	Yes (lasted for ____ days) / No
3. Did you experience swelling at the injection site?	Yes (lasted for ____ days) / No
4. Did the injection site become itchy?	Yes (lasted for ____ days) / No
5. Did you feel tired all over your body?	Yes (lasted for ____ days) / No
6. Do you have headache?	Yes (lasted for ____ days) / No
7. Do you have muscle pain?	Yes (lasted for ____ days) / No
8. Do you feel chilly?	Yes (lasted for ____ days) / No
9. Did you have fever?	Yes (max. °C, lasted for ____ days) / No
10. Did you have any joint pain?	Yes (lasted for ____ days) / No
11. Did you have nausea or vomiting?	Yes (lasted for ____ days) / No
12. Do you have diarrhea?	Yes (lasted for ____ days) / No
13. Do you have stomachache?	Yes (lasted for ____ days) / No
14. Did you have allergic symptoms or anaphylaxis? Please select from the following.	
A: Allergic symptoms occurred (please specify: _____)	
B: Anaphylaxis occurred (please specify: _____)	
C: I was able to receive the inoculation without any particular problem.	
Thank you very much for your cooperation.	

**Table S2.** Characteristics of participants in the T-SPOT study.

	Control group (n=48)	HD group (n=64)	p value
Male, n (%)	31 (64.6)	35 (54.7)	0.2
Age, years ( $\pm$ SD)	62.9 $\pm$ 7.9	69.4 $\pm$ 4.8	<b>0.02</b>
BMI, kg/m <sup>2</sup> ( $\pm$ SD)	24.1 $\pm$ 4.8	22.0 $\pm$ 4.6	<b>0.02</b>
Diabetes mellitus, n (%)	9 (18.8)	21 (32.8)	0.07
Hypertension, n (%)	25 (52.1)	22 (34.4)	0.05
Malignant tumor, n (%)	6 (12.5)	13 (20.3)	0.2
Cerebrovascular disease, n (%)	7 (14.6)	14 (21.9)	0.23
Cardiovascular disease, n (%)	7 (14.6)	11 (17.2)	0.5
COPD, n (%)	3 (6.3)	4 (6.3)	0.65
Interval time* average $\pm$ SD	208.9 $\pm$ 15.1	215 $\pm$ 16.9	<b>0.03</b>

BMI: body mass index; COPD: chronic obstructive pulmonary disease; HD: hemodialysis; SD: standard deviation. \*Interval time: days from primary series to booster vaccination.