中文題目: 乳癌病患接受三劑 COVID-19 疫苗後之體液及細胞免疫反應 英文題目: Humoral and cellular immune response after 3rd dose of COVID-19 vaccination in breast cancer patients 作 者: 蘇柏嘉¹, 鍾為邦², 楊舜如²

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Background

The novel coronavirus, SARS-CoV-2 are corresponding to coronavirus disease (COVID-19) since 2019. Patients with cancer have more complication and higher mortality rate in COVID-19. Vaccination provide protection from COVID-19 infection and decrease severity of disease. However, previous study showed that cancer patients had poor immune response to COVID-19 vaccine, especially in patients who receive cytotoxic agents or patients with hematological malignancy. Fortunately, humoral immunity improves after two dose of vaccination and had benefit on third dose of COVID-19 vaccination. However, T cell response to vaccination in cancer patients is still investigating.

Breast cancer is the most common malignancy in the world. There are different treatment strategies in different breast cancer subtypes. Medical treatment includes endocrine treatment, chemotherapy, target therapy, antibody drug conjugate, and immunotherapy. We would like to study immune response in breast cancer patients received 3rd dose of CIVID-19 vaccination. To investigate the influence on immune response in different treatment and disease status.

Method

We collected patients with breast cancer in single institution in south Taiwan. All patients are ≥ 20 years-old, without history of COVID-19 infection, received 3rd dose of COVID-19 vaccination at least two weeks ago. We had informed consent with all patients. We checked anti-spike-RBD Ab (anti-S Ab) to evaluate humoral immunity, anti-nucleocapsid Ab (anti-N Ab) for diagnosis of infection with SARS-CoV-2. We used cPass assay to measure neutralizing antibody for Wuhan strain, Delta variant, and Omicron BA.1 variant. To evaluate cellular immunity, we used Covi-Feron 500, to detected IFN γ responses to SARS-CoV-2 and its variants specific proteins (including Wuhan. Alpha, Beta, and Gamma strains).

Results

Twenty-eight female patients were enrolled. The characteristics and result as shown in Table 1. All of them (28/28, 100%) had anti-S antibody and no patients had positive anti-N antibody. Twenty-six (26/28, 93%) patients had neutralizing antibody for Wuhan strain; twenty-six (26/28, 93%) patients had neutralizing antibody for

Delta strain; Four (4/28, 14%) patients had neutralizing antibody for Omicron BA.1 strain.

For cellular immunity, we evaluate T cell response by IFN γ responses to SARS-CoV-2 and its variants specific proteins. The result revealed twenty-two patients had reactive T cell response. Among the six patients with non-reactive response, five patients of them had advanced breast cancer, and three patients had progressive disease. All patients with early breast cancer who received endocrine treatment had reactive T cell response.

Conclusion

Breast cancer patients received 3rd dose of COVID-19 vaccination had good response to Wuhan strain and Delta variant but poor response to Omicron BA.1 variant. Patients who had poor cellular response may related to disease status and cancer treatment. We are going to evaluate the response to omicron BA.4 and BA.5, and the response after patient who received 4th dose vaccination.

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Table 1

PR: Partial response; SD: stable disease; CR: complete response; PD: Progressive disease Anti-HER2 Tx: anti-HER2 treatment; C/T: chemotherapy; ET: endocrine treatment; CDK4/6 i: CDK4/6 inhibitor; ADC: Antibody drug conjugate.

Red color words: Non-reactive T cell response

Yellow highlight: Non-detectable neutralizing antibody (< 30% means there is no detectable neutralizing antibody)

All patients are negative for anti-nucleocapsid Ab (not shown in the table)

Table 1

Pt	Age	Disease		Current	Anti-S	Neutralizing	Neutralizing	Neutralizing	T cell
				treatment	Ab	Ab for	Ab for Delta	Ab for	response
					(U/mL)	Wuhan %)	(%)	Omicron	
								BA.1 (%)	
1	57	Advanced	PR	Anti-HER2 Tx	2500	97.6	96.6	<mark>-2.5</mark>	Reactive
2	66	Advanced	SD	C/T	2500	97.9	97.0	<mark>15.1</mark>	Reactive
3	52	Early	CR	ET	2500	97.9	97.0	52.4	Reactive
4	70	Early	CR	ET	2415	97.7	96.2	<mark>9.4</mark>	Reactive
5	34	Early	CR	ET	2500	97.5	96.2	<mark>-4.9</mark>	Reactive
6	56	Advanced	PD	CDK4/6 i, CT, ET	2500	97.7	96.5	<mark>-7.9</mark>	Non-reactive
7	46	Early	CR	Anti-HER2 Tx	171	76.8	60.6	<mark>2.4</mark>	Reactive
8	58	Advanced	SD	Anti-HER2 Tx	2500	97.7	97.0	58.3	Reactive
9	73	Advanced	SD	CDK4/6 i, ET	2500	97.6	96.0	<mark>-4.1</mark>	Reactive
10	49	Advanced	PD	Anti-HER2 Tx, C/T	224.8	74.8	70.0	<mark>11.5</mark>	Reactive
11	63	Early	CR	Anti-HER2 Tx, ET	2500	97.5	96.4	<mark>23.0</mark>	Reactive
12	45	Advanced	PR	CDK4/6 i, ET	1133	84.1	73.3	<mark>-13.2</mark>	Reactive
13	58	Advanced	PD	C/T	354	72.7	54.8	<mark>-6.0</mark>	Non-reactive
14	50	Advanced	PD	CDK4/6 i, ET	2500	97.5	96.4	<mark>13.1</mark>	Non-reactive
15	53	Advanced	PR	Anti-HER2 Tx	2500	97.9	96.9	34.0	Reactive
16	40	Early	CR	C/T	2500	97.8	96.7	<mark>-2.6</mark>	Reactive
17	52	Early	CR	Anti-HER2 Tx, C/T	2500	97.7	95.6	<mark>6.6</mark>	Reactive
18	46	Early	CR	ET	2500	97.4	92.8	<mark>23.5</mark>	Reactive
19	48	Early	CR	ET	2500	97.8	96.7	<mark>-1.5</mark>	Reactive
20	42	Advanced	PR	Anti-HER2 Tx, C/T	1222	97.5	95.7	<mark>15.7</mark>	Reactive
21	31	Advanced	CR	Anti-HER2 Tx, C/T	24.48	91.1	83.7	<mark>4.7</mark>	Non-reactive
22	52	Early	CR	Anti-HER2 Tx, C/T	1964	97.4	92.7	<mark>2.8</mark>	Non-reactive
23	83	Advanced	SD	CDK4/6 i, ET	2500	97.6	85.6	<mark>8.7</mark>	Non-reactive
24	81	Advanced	SD	ADC	20.51	<mark>10.7</mark>	<mark>0.9</mark>	<mark>7.3</mark>	Reactive
25	64	Advanced	PR	CDK4/6 i, ET	2500	97.5	96.2	<mark>29.0</mark>	Reactive
26	52	Advanced	PR	Anti-HER2 Tx, C/T	2500	97.5	95.8	<mark>0.1</mark>	Reactive
27	42	Advanced	CR	C/T	2500	<mark>17.6</mark>	<mark>10.9</mark>	<mark>8.4</mark>	Reactive
28	49	Early	CR	ET	2500	97.7	96.6	77.0	Reactive