

# S35-7 Effect of supplement of marine algae, $\beta$ -glucan, yeast on immunity traits of growing pigs

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## INTRODUCTION

Addition of antibacterial substances to domestic animal feed will increase the resistance to antibiotics of bacterium. Such practices present the dangerous possibility of bacterium infection of humans. Corresponding to such a situation, the additives to the feed that activates pig's immunity and can be substituted with the antibacterial addition material were demanded.

## OBJECT

In order to evaluate the effect of the additive to the feed that activates pig's immunity, **marine algae,  $\beta$ -glucan, and yeast were added to feed from 20 to 70kg body weight of the piglet and the influence on the pig growth and immunity were examined.**

## MATERIALS AND METHOD

### 1) Animals : 16 Duroc pigs from three litters

### 2) Experimental design :

| Group           | Feed content                                    | Number              |
|-----------------|-------------------------------------------------|---------------------|
| Control         | Grower ration without antibacterial substances  | Gilts 2<br>Barrow 2 |
| Marine algae    | Grower ration + 0.8% marine algae (160g/20kg)   | Gilts 2<br>Barrow 2 |
| $\beta$ -glucan | Grower ration + 0.1% $\beta$ -glucan (20g/20kg) | Gilts 2<br>Barrow 2 |
| Yeast           | Grower ration + 0.5% yeast (100/20kg)           | Gilts 2<br>Barrow 2 |

### 3) Blood collecting and body weight measurement schedule

| 9w            | 11w           | 13w           | 15w                  | 16w                  | 17w           |
|---------------|---------------|---------------|----------------------|----------------------|---------------|
| Collect blood | Collect blood | Collect blood | 1st SRBC inoculation | 2nd SRBC inoculation | Collect blood |
| Measure BW    | Measure BW    | Measure BW    | Collect blood        | Measure BW           | Measure BW    |

4) **Measurement growth and immune traits:** To compare antibody production to a specific antigen, every group of pigs was inoculated sheep red blood cell (SRBC) at 15 and 16w of age. Phagocyte activity (PA), complement alternative pathway activity (CAPA), total leukocyte (WBC), ratio of granular leukocytes to lymph cells (RGL) and Sheep Red Blood Cell (SRBC) specific IgG (sIgG) were measured.

## RESULTS

1) **Growth traits:** There was no significant difference in body weight among groups. But, the feed intake was lower in Marine algae group than other group.

### 2) Immune traits:

(i) At 17w of age, sIgG and WBC of the marine algae group were significantly higher than the control group.

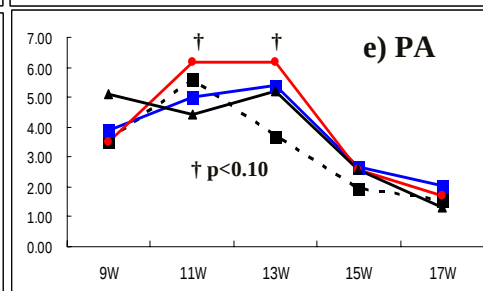
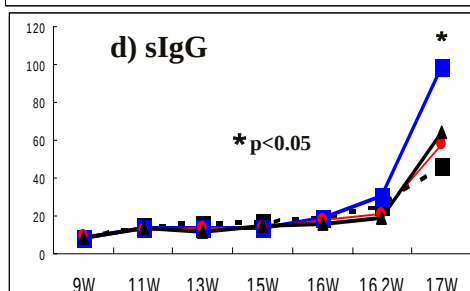
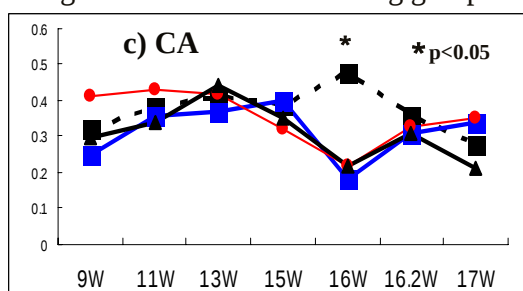
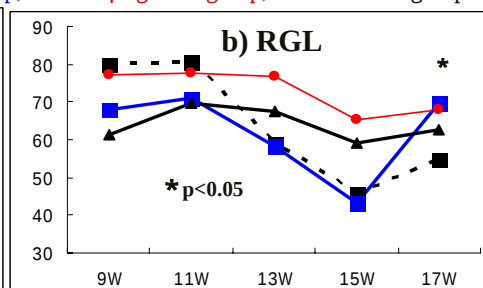
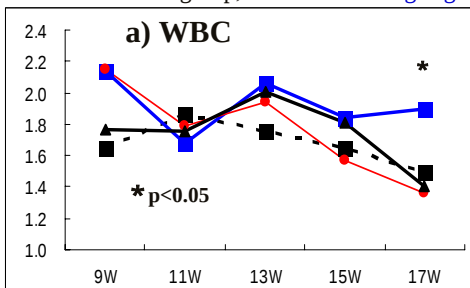
(ii) RGL of the marine algae and  $\beta$ -glucan groups was significantly higher than the control group at 17w of age.

(iii) PA of the  $\beta$ -glucan group was higher than that of the control group at 11w and 13w, however, there were no significant differences among groups.

Table 1. Body weight and feed intake of each group from 9w to 17w

| Group        | 9w   | 11w  | 13w  | 15w  | 16w  | 17w  | DG: g/d | FID:kg/pig |
|--------------|------|------|------|------|------|------|---------|------------|
| Control      | 15.4 | 19.7 | 27.4 | 38.4 | 51.0 | 63.7 | 786     | 101.9      |
| Marine algae | 15.7 | 19.6 | 28.6 | 39.1 | 51.2 | 64.5 | 803     | 99.1       |
| B glucan     | 15.3 | 19.7 | 29.6 | 39.6 | 51.9 | 64.8 | 807     | 101.3      |
| Yeast        | 15.8 | 19.8 | 29.9 | 40.2 | 52.1 | 65.5 | 815     | 103.1      |

■ -- ■ : Control group, ■ -- ■ : Marine algae group, ● -- ● :  $\beta$ -glucan group, ▲ -- ▲ : Yeast group



## CONCLUSION

These results suggest that addition of the marine algae to feed activate the pig's immune function. Usually, marine algae is not digested in pig's intestine. So, we postulated that the bacillus in small and large intestine digest the marine algae and the produced organic acids activated the mucosal immunity and the entire immunity consequently. Now, we have the plan to compare the organic acids composition and the amount of IgA in contents of intestines and the anatomical comparison of intestine tissue.