

PORK SAFETY

Title: Immunomodulatory and Growth Effects of the Seaweed *Ascophyllum nodosum* in Pigs – **NPB #00-063**

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Abstract:

A series of *in vitro* studies was conducted to evaluate the effect of *A. nodosum* extract (**ANE**) on activation of pig splenocytes and alveolar macrophages. In the first set of studies, splenocytes and macrophages were incubated with concentrations of **ANE** from .1-100 µg/mL. Concentrations of TNF α and PGE₂ were evaluated as measures of macrophage activation and interleukin 10 (IL-10) and interferon gamma (INF γ) were evaluated as measures of splenic lymphocyte activation. As positive controls, macrophages were stimulated with bacterial lipopolysaccharide (LPS) and splenocytes were incubated with the mitogen concanavalin A (Con A). **ANE** failed to stimulate secretion of PGE₂ or any cytokine by cultured immune cells. In a subsequent set of experiments, concentrations of **ANE** as high as 10 mg/mL were evaluated. Again, **ANE** did not alter IL-10 production by splenocytes. However, PGE₂ was increased ($P < .05$) relative to control wells in macrophages treated with 10 mg/mL **ANE** after 3 and 24 hours in culture. The results of our *in vitro* studies demonstrate conclusively that **ANE**, when presented directly to immune cells across a wide range of concentrations, generally failed to activate macrophages and lymphocytes. These immune cells are key players in both cell-mediated and antibody-mediated immune function. Very high concentrations of **ANE** enhanced production of the inflammatory mediator PGE₂, but never to the degree achieved by bacterial LPS.

An *in vivo* study also was conducted with dietary **ANE**. A total of 95 pigs (initially 15 lb and 17 d of age) was used in a 28 d growth experiment to determine the effects of **ANE** on weanling pig growth performance and immune function in response to enteric disease challenge with *Salmonella typhimurium* (**ST**). Experimental treatments were arranged in a 2 x 4 factorial with main effects of disease challenge (control vs. **ST** challenge) and dietary addition of **ANE** (0, 0.5, 1.0, and 2.0% of diet). Results suggest little beneficial effect of dietary **ANE** on growth performance or immune response in the presence or absence of **ST** challenge.

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