Introduction

β-D-glucans (hereafter referred to as “glucans”) represent part of a group of physiologically active compounds generally called “biological response modifiers.” They are highly conserved carbohydrates forming structural components of cell walls of some plants, fungi, yeast, seaweed and bacteria. Generally, glucan represents a group of chemically heterogeneous polysaccharides existing in various numbers of molecules bound together in several forms of linkage together with several forms and degrees of branching.

Glucans have a long history as natural immunomodulators. The first reports showing that some infectious diseases can have therapeutic effects on malignant processes can be found almost two hundred years ago. These studies were later followed by studies of the immunomodulating properties of lipopolysaccharide (LPS). The problems with toxicity of LPS were dismissed as a result of later investigations showing that a saccharidic moiety of LPS is non-toxic but is responsible for the immunomodulating activity. The history of glucan began app. 60 years ago with two different starting points—one originated in Europe and the United States and the second in Japan. Based on historical use, the Japanese groups investigated mushroom-derived glucans, whereas the European and American groups focused on yeast-derived glucans.

The first studies showed that glucan application significantly stimulated the phagocytic system and enhanced general defense and resistance to experimental tumors. During subsequent decades of intensive research by laboratories around the world, glucans were found to significantly stimulate defense reactions against infections and cancer. In addition, several additional effects were later shown. These included reduction of stress, hypoglycemic effects, lowering cholesterol, reduction of cytotoxic effects and improving diseases such as ulcerative colitis. Another advantage of using glucan as a stimulator of immune reactions is the fact that it has been shown to act in all species tested so far, starting with earthworms and ending with humans.

Immunity of fish

As in all gnathostomes, there are two types of immunity in fish: The innate and the adaptive. The cellular component of innate immunity represents various phagocytic cell types such as the evolutionarily ancient...
macrophages, and natural killer (NK) cells. In fish, there were two types of NK cell homologues: Non-specific cytotoxic cells and NK-like cells. Blood leukocytes primarily form further cellular component of innate immunity. They produce a row of humoral substances, mainly cationic antimicrobial peptides, complement components, lectins, cytokines, anti-inflammatory immune mediators like IL (interleukin)-10, TGF-β and many others that are able to kill immediately altered and foreign (allogeneic or xenogeneic) cells. These substances are released into body fluids and epithelial and skin mucus. These innate mechanisms are well developed in bony fish but in some cases, especially in aquacultures where it is a higher probability of spreading infectious diseases, they are not adequate without some external stimulation.

The major organs of fish adaptive immunity are the thymus, kidney, spleen and GALT (gut-associated lymphatic tissue). Conversely, to more evolutionary-advanced vertebrate taxa, they do not possess bone marrow and lymph nodes. The fish thymus is the first lymphoid organ appearing in ontogeny. Structural anlage of thymus is immediately colonized by lymphoid cells. During early ontogeny, the thymus develops in an organ with the epithelial/reticular stroma, which forms a framework for thymocytes and macrophages and other accessory cells. In comparison to tetrapod thymus, fish thymic tissue is less differentiated. In many fish species, the cortex and medulla are lacking, even if in some species the middle and inner zones of thymus tightly resemble the clearly differentiated cortex and the medulla of advanced vertebrates. It also underlies histopathological degeneration during ageing. The kidney is a paired organ consisting of pronephros and mesonephros, both hemolymphopoietic, comparable to bone marrow of tetrapods. The pronefros first become erythroid. Later, the aggregates of lymphoid cells could be found among urinary tubules. In addition to these functions, fish kidney represents an immunocompetent organ with strong phagocytic capacity where antigen processing and antibody formation take place. Similar to the situation in spleen, the plasmacytes, lymphocytes, monocytes, granulocytes and non-specific natural cytotoxic cells, can be found in kidney. Fish lack germinal centers but the architecture of kidney tissue ensures a suitable micromilieu for immune processes from phagocytosis, antigen presentation up to efficient humoral immune response. The kidney and particularly the well-developed GALT serve as analogs of bone marrow. Antigen trapping and processing also takes place within the special structures, the ellipsoid sheets, which are described in most bony fish. It is believed that they could represent analogs or evolutionary predecessors of germinal centers of mammals.

In fish species with well-developed alimentary tract, the aggregations of lymphoid cells are often present in connective tissue of the mucosa. Lymphoid cells infiltrating gut epithelia and lamina propria form clusters but are never structurally organized like Peyer’s patches found in the mammalian GALT. On the other hand, the fish perienteral lymphoid tissue plays a similar role as an effective immunological barrier. In more advanced teleosts, aggregates of lymphocytes, plasmacytes, granulocytes, and macrophages occur in and under the intestinal epithelium. Together with the epithelial cells, these accumulations may form a microenvironment for food antigen collecting such as M cells. So the fish GALT could functionally serve as the gut barrier of mammals. Authors seeking more information on gastrointestinal microbiota in fish should see an excellent review by.

Clusters of lymphoid cells and antibody-forming cells were also identified in some other regions where the potential pathogens may invade and are phagocytized. Such predominant locations include the skin epidermis, gills and pharynx, heart, liver, and pancreas. Generally, in all above mentioned organs and tissues, the plasma cells and lymphocytes are ultrastructurally similar to those of ectotherms and endotherms. The B and T cell dichotomy in fish has been documented but in contrast to B cells, fish T cells are Ig cells. The discovery of TR genes confirmed definitely the occurrence of conventional T cells in fish. Several key T cell markers as well as the typical cytokines produced by different T subpopulations resembling the Th1, Th2 and Th17 of mammals were described. On the other hand, particular sub-populations of gut intra-epithelial lymphocytes seem to be different. New studies document that the diversity of fish naïve TCRβ expressed by CD8(+) and CD8(-) αβ T cells may regulated by different regulatory mechanisms than in mammals.

Teleostean B-cell subsets express either both IgM and IgD or only recently discovered IgT also called IgZ. In contrast to endothermic vertebrates, fish are devoid of IgA. IgM is regarded as the main molecule bearing antibody activity but it usually is a tetramer. IgM and IgT are never co-expressed by the same B cell, which identifies two distinct lineages of B cells in fish.
appears that IgT acts like an intestinal mucosal antibody against some parasites, whereas IgM antibodies are acting mainly in the serum. In addition to immunoglobulins, the non-specific factors like (hemo) lysins, (hem) agglutinins (lectins), lytic enzymes, lysozyme, C-reactive protein, antibacterial peptides, and complement are also present. In some species, immunization results in the formation of specific antibodies not only in the serum, but also in skin and gill mucus and in the gut lamina propria.

Aquacultures and protecting against infection

For nearly 4,000 years, the farming of fish in aquacultures has been a source of human food. Aquaculture production in the past decades represents the increased importance in the food supply for a continually growing world population. At present, fish farming is the fastest growing agricultural industry. Aquacultures have expanded globally with an increase not only in absolute production (kilotons/year) but also in the number of fish species being cultured in both freshwater and marine systems. The most important farmed fish species are carp, salmon, tilapia and catfish.

On the other hand, intensification and rapid increase in aquaculture activities world-wide has provided new opportunities for the emergence and transmission of aquatic pathogenic microorganisms, both bacterial and viral. The specific diseases caused by these etiological agents represent a significant limiting factor for fish aquaculture farming. Readers seeking more information on the current state of aquaculture should read the latest comprehensive.

Therefore, commercial aquacultures are not only dependent on good water quality. The achievement in preventing disease outbreak and restriction of its spreading is at least of equal importance. The use of vaccines, antibiotics, and non-specific immunostimulants are three possible methods of farmed fish protection.

Efficient vaccines were developed against only several bacterial pathogens at the end of the 20th century but no effective vaccines are available against a row of other bacterial diseases and most particularly against viral diseases.

The use of antibiotics must be considerably reduced, primarily to avoid environmental hazards and the spread of antibiotic-resistant genes. In addition, some antibiotics may suppress fish immune responses.

When fish are attacked by a pathogenic microorganism, the non-specific mechanisms of natural immunity are more important than the specific response. The non-specific defense in fish is similar to other vertebrates that are endowed by many elements such as phagocytic cells, granulocytes and humoral factors as complement, lysozyme, C-reactive protein, interferon and transferrin. A rapid defense response through the use of antimicrobial cationic peptides and other natural immune factors is much more efficient than prolonged and relatively slow production of specific antibodies that are characteristic of these cold blooded animals.

Effects of glucan on bacterial diseases

Alternative strategies to vaccination and use of antibiotics represent applications of various immunostimulatory substances as dietary supplements. Although little is known about the mechanism of their action in fish, some of them appear to enhance the non-specific killing of pathogenic microbes.

At present, non-specific immunostimulants represent the primary tools in modern fish farming. They induce and enhance resistance against bacterial and viral infectious diseases by stimulating innate humoral and cellular defense mechanisms. The stimulatory action of a row of structurally non-related substances has been studied for their suitability to prevent infections in aquacultures.

Several types of immunostimulants have been used in fish cultures to induce protection against a wide range of diseases. Various compounds are being added to feed, based on their possible role as immunostimulants. During the study of their prophylactic effects, of these substances the β-glucans appeared to be the most convenient for applications in aquacultures. Therefore, in recent years, attention has focused primarily on possible immune stimulation in farmed fish by the use of β-glucans. To date, numerous studies confirming the potent immunostimulatory properties of β-glucans in many fresh and seawater fish species documenting the effects of β-glucans on the pathogen resistance, protection, survival and fish specific humoral immunity have been published. The most successful one is glucan. It is, therefore, not surprising that feed containing glucan is routinely manufactured for commercial fisheries. The most common brands are MacroGard, Vetregard and EcoActiva.

The main fish species studied were rainbow trout (Oncorhynchus mykiss), Channel catfish (Clarias Gariepinus), African catfish (Labeo rohita), turbot (Scophthalmus maximus L.), pink snapper (Pagrus auratus), sea bass (Dicentrarchus labrax), red tail black shark (Epalzeorhynchus bicolor), fathead minnows (Pimephales promelas Rafinesque 1820), atlantic cod (Gallus morhua L.), gilthead seabream (Sparus aurata), large yellow
croaker (Pseudosciaena crocea),[60] carp (Cyprinus carpio),[46,61] nile tilapia (Oreochromis niloticus),[62,63] and zebrafish (Danio rerio).[62]

Administration of glucan in carp enhanced survival, most likely via stimulation of both non-specific and specific immune reactions (superoxide anion, IL-1 secretion and antibody formation), regardless of how it was administered (intraperitoneal injection, bathing and oral administration).[64] A stimulation of complement and C reactive protein responses were found in carp.[63,64] Studies of glucan-activated macrophages in trout revealed an increased ability to kill salmonid pathogen Aeromonas salmonicida, despite its virulence.[46] A more detailed study using radioactively labeled glucan showed the transfer of this material through the intestine via the epithelial cells in the lower part of the intestine. The material was later cleared from blood.[65] The exact mechanisms of glucan action on anti-infective immunity are not fully established. Recent observations suggest the role of neutrophil extracellular traps.[66]

Improvements in the composition of vaccine used in fish are still necessary. Based on the known strong effects of glucans on fish immunity, their use as part of the vaccination process is not surprising. Early studies showed that the addition of glucan to a vaccine resulted in non-specific resistance against vibriosis and yersiniosis in salmon.[67] These early studies primarily used injected glucan, and in addition to protection against infection, glucan also increased production of cytokines, complement and lysozyme production and antibody formation.[37] The direct addition of glucan into the feed resulted in reduction of mortalities caused by infections with salmon anaemia virus and Piscirickettsia salmonis. In addition, lower attachment of sea lice to fish was also observed.[68]

Similar effects were found in infection with A. salmonicida.[69] The addition of glucan to the Aeromonas vaccine significantly increased the production of antibodies in all antigens tested.[70] However, even the elevated level of antibodies did not offer sufficient protection against Aeromonas infection.

Detailed studies of the adjuvant effects of glucan were done using vaccine against furunculosis. In all cases, vaccines enforced with glucan induced significantly stronger protection of salmon, as measured by serum antibodies levels against four different parts of the A. salmonicida.[49,71] A model using Flexibacter columnaris infection of trout also revealed a strong protection against mortality.[72]

Other authors tested the effects of vaccine VYS-2, a protein fraction of Aeromonas, with and without glucan and showed that glucan increased the protective effects in carp.[73] Vibrio damsela vaccine containing glucan was successfully used in turbot[51] and the feeding of glucan with bactericin Edwardsiella tarda showed strong protection in Japanese flounder (Paralichthys olivaceus).[74]

A different approach was used by.[80] These authors fed Indian major carp with glucan for 30 days followed by vaccination against E. tarda. The results showed that this combination strongly increased the specific immunity and reduced mortality in immunocompromised animals. Of the four tested, (glucan, levamisole, vitamin C, and vitamin E), the glucan was the most effective substance. A series of papers tested the effects of glucan in conjunction with vitamin C. Use of this combination following vaccination can serve as an example. The combination increased the activity of macrophages and specific antibody response.[75] Another study used a different feeding period to address the possibility that longer exposure to glucan might lead to exhausting the natural defense. All these studies led to the conclusion that glucan represents an ideal immunostimulant in the fish industry.[39]

However, not all studies were successful. Glucan with vaccine against Streptococcus bactericin was not effective in turbot[79] and glucan with vaccine against S. iniae had no effect on the Oreochromis niloticus model.[77] Trials with commercial glucan vaccine VitaStim Taito showed no effects.[78] Readers seeking more information on the role of vaccines in fish immunity should refer to[79] a comparative study of 8 different glucans found the only 1-3,1-6 β-glucans caused significant protection against A. hydrophila infection.[80] On the other hand, a comparative study of several different immunostimulators found that only glucan offered protection against white spot disease.[81]

The effects of β-glucans on bacterial infections are summarized in Table 1. It is clear that the use of glucan as part of vaccines in fish is despite decades of research, still far from conclusive. This might be explained as being the result of confusion in both delivery (oral or injected administration) and dosing. The current massive use of glucan in commercial farming therefore focuses more on general stimulation of immune response than on possible adjuvant effects.

**Effects of glucan on viral and parasitic diseases**

Main viral diseases affecting bony fish in aquacultures can occur immediately in the overwhelming majority of farmed fish.[82] Moreover, the more virulent viruses resulting in hemorrhages, ascites, and death are prone to spread globally between countries by wild fish and transmitted into new fish species [Table 1].
Additional studies focused on resistance of spotted rose snapper *Lutjanus guttatus* against dactylogyrids. Five weeks of feeding with feed including 0.05% glucan significantly reduced the number of dactylogyrids. Several parameters such as white blood counts, percentage of neutrophils, eosinophils and thrombocytes were also observed, but the connection is unclear.

The last important study focused on rainbow trout and skin-parasitic ciliate *Ichthyophthirius multifiliis*. Effects of glucan isolated from *Euglena gracilis* were dose-dependent, but clearly lowered the number of trophonts. At the same time, lysozyme activity was elevated. The trend for upregulation of some important genes was not significant.

**Mechanisms of glucan action**

In addition to direct stimulation of both specific and non-specific immunity, glucan can also influence...
expression of immune-related genes and proteins. Macrophages from Atlantic salmon and rainbow trout showed elevated levels of cytokines, but not C3. Similar data were found in plasma of tilapia.

The effects of glucan on gene expression are rapid and do not need long exposure. Four 45 min submersions in glucan each week caused enhanced gene expression of IL-1β, TNF-α, IL-6, IL-10 and TGF-β, sometimes even after first submersion. Carp treated with glucan for 15 days, followed by injection with hemorrhage virus, showed elevated MX gene expression during early stages of infection. A similar experimental model using Atlantic cod and Vibrio anguillarum challenge demonstrated that five week-long glucan immersion resulted in elevated expression of IL-1β gene in the anterior intestine and rectum, whereas the expression of IL-10 was measurable only in rectum. In the same experimental design, mannan-based oligosaccharide, additional upregulation of IL-8 and IFN-γ was observed.

Glucan exposure not only helped to show the upregulation of some genes, but even to their description. Two β-defensive genes, β-defensin 1 and β-defensin 2, were described in common carp. In addition, the expression of these genes was upregulated by glucan exposure, similar to the expression of two mucin genes. A study of inflammatory cytokines as a response to A. salmonicida infection showed that feeding with glucan resulted in reduction of gene expression of inflammation-related cytokines such as IL-1β, IL-6, IL-10, and TNF-α.

A detailed study of the effects of dietary glucan on gene expression was done in common carp. While the 7 day incubation showed very small changes, 25 day incubation increased iNOS and Bcl-2 expression in liver and head kidney. In other organs, the effects were more pronounced with a strong increase of iNOS, Bcl-2 and Nemo in gut and iNOS, Caspase 9, Bcl-2, p38 and Nemo in spleen. Despite the progress, the evaluation of the effects of glucan on genomic level are limited not only with respect of species, but also of individual genes. To better understand how glucan exposure affects individual genes, the full genomic studies need to be performed.

Conclusions

Administration of glucans through various routes including immersion, feed or injection have been found to enhance many types of immune responses, resistance to bacterial and viral infections and resistance to environmental stress. Although the efficacy of the glucan to some extent varies with type and administration, glucan used as an immunomodulatory additive has been found to be active in eliciting immunity in commercial aquaculture and is currently routinely used in commercial farming. Development of more efficient administration methods will facilitate the routine and prophylactic use of glucans as natural immunostimulants of fish. Lately, interest focused on mechanisms of action. However, nothing conclusive can be reached as a result of these studies and the effects of glucan on modulation of gene expression leading to stimulation of fish immunity still require elucidation. Therefore, the limited knowledge of mechanisms of glucan action on fish immunity does not currently allow better and more specific use of glucan in aquaculture.

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