Evaluation of Ammonia Nitrogen Exposure in Immune Defenses Present on Spleen and Head-Kidney of Wuchang Bream (Megalobrama amblycephala)

Honghui Guo 1, Siqi Chen 2, Kang Ouyang 1, Yu Kuang 1, Hui Yang 1, Yingying Wang 1, Rong Tang 1,3,4,5, Xi Zhang 1,3,4,5, Dapeng Li 1,3,4,5 and Li Li 1,3,4,5,*

1 College of Fisheries, Huazhong Agricultural University, Wuhan 430070, China; honghuiGuo@webmail.hzau.edu.cn (H.G.); ouyangkang@webmail.hzau.edu.cn (K.O.); kuangyu@webmail.hzau.edu.cn (Y.K.); yanghui1325196@yeah.net (H.Y.); wangyinling@163.com (W.Y.); tangrong@mail.hzau.edu.cn (R.T.); zhangxi@mail.hzau.edu.cn (X.Z.); ldp@mail.hzau.edu.cn (D.L.)
2 Hubei Aquaculture Technology Extension Center (Hubei Aquatic Breeds Introduction and Breeding Center), Department of Agriculture and Rural Affairs of Hubei Province, Wuhan 430060, China; m1857625524163.com
3 Engineering Research Center of Green Development for Conventional Aquatic Biological Industry in the Yangtze River Economic Belt, Ministry of Education, Wuhan 430070, China
4 Hubei Provincial Engineering Laboratory for Pond Aquaculture, Wuhan 430070, China
5 Freshwater Aquaculture Collaborative Innovation Center of Hubei Province, Wuhan 430070, China
*
Correspondence: foreverlili78@mail.hzau.edu.cn

Abstract: Ammonia is one of the most important environmental factors in aquatic ecosystems. However, there are limited studies on the effects of chronic or long-term ammonia stress and its potential molecular mechanism in fish. This study aimed to investigate the immune response and molecular mechanisms in the spleen and head-kidney of fish following chronic ammonia exposure. Megalobrama amblycephala (9.98 ± 0.48 g) were exposed to different concentrations of total ammonia nitrogen (0–30 mg/L) for 30 days. Ammonia exposure caused significant increases in cortisol levels and decreases in lysozyme and complement 3/4 concentrations in the serum, indicating inhibitory effects of ammonia stress on innate immune responses. Ammonia exposure also induced concentration-dependent increases in ammonia concentrations in tissue, pathological damage and indexes of spleen and head-kidney. Additionally, the contents of immunoglobulin M (IgM), interleukin 1β (IL-1β) and tumor necrosis factor α (TNF-α) as well as mRNA levels of toll-like receptors (TLRs)/Myeloid differentiation factor 88 (MyD88)-independent signaling molecules in the spleen and head-kidney were significantly downregulated after ammonia exposure. Our findings suggested that chronic ammonia exposure caused the suppression of innate and adaptive immune responses through downregulating TLR/MyD88-independent signaling. Adverse influences of chronic ammonia stress were more severe in the spleen than in the head-kidney.

Keywords: ammonia; stress; toll-like receptors; spleen; head-kidney; immunity

1. Introduction

Ammonia is one of the most important environmental factors in aquatic ecosystems that affects the growth and health of aquatic animals [1]. Ammonia nitrogen generally comes in one of two ionized forms—NH₄⁺ and the un-ionized form, NH₃, in the aquatic environment. NH₃ is extremely toxic to fish by reason of its ability to diffuse across cell membranes easily [2–4]. Over the past decades, high levels of ammonia frequently occur in aquatic environments due to disorderly discharges of sewage effluent, agricultural run-off, and high-density aquaculture [5–7]. Although the recommended level of ammonia nitrogen in drinking water is up to 3 mg/L (NH₄/L) [8], the real ammonia level can reach much higher (>20 mg/L) at times, due to ineffective or nonexistent sewage treatment or other reasons [9,10].
In fact, the immunosuppressive effects of ammonia in fish have been proven to contribute to disease outbreaks [11,12]. In some studies on Takifugu rubripes, Scophthalmus maximus and Pelteobagrus fulvidraco, acute ammonia exposure could upregulate gene expression levels of B-cell activating factor, heat shock protein 70/90, tnf-a, and interleukin-1/1β/6/8/12 [13–15]. Wang et al. also reported that serum lysozyme activities were decreased significantly in Acanthopagrus schlegeli after 24 h of ammonia exposure [16]. Similarly, decreased mRNA levels of complement C3 and IgM were observed in the spleens of Pelteobagrus vachellii and Rhynchocypris lagowski under acute ammonia exposure [17,18]. Although much attention has been paid to the effects of acute ammonia exposure on fish immunity, there are only limited studies on the effects of chronic or long-term ammonia stress and its potential molecular mechanism. In addition, some investigators have reported adverse effects of ammonia nitrogen on mRNA expression of TLRs through transcriptional analysis [16,19–22]. Consequently, we hypothesize that persistent ammonia exposure also could affect fish immunity, in which the TLR signaling pathway might play an important role.

The spleen and head-kidney are the major immune organs in teleost fishes, which are responsible for trapping and clearing foreign particulate materials and maintaining a stable internal environment [23,24]. They are also the main sites where immune antibodies are produced [25–27]. Additionally, in fish, the innate immune system consists of fixed and mobile cells as well as a wide range of defense molecules (such as lysozyme, complement, cytokines) [28,29]. Wuchang bream (Megalobrama amblycephala) is a cyprinid fish native to the Yangtze basin of China, which was ranked 11th in the world in total annual production of this fish in 2014 [30]. As the main aquaculture species, M. amblycephala is a relatively sensitive species to ammonia nitrogen and often suffers from stress induced by ammonia nitrogen [31–33]. In light of the above, a chronic exposure experiment was conducted in Wuchang bream to elucidate the features of immune responses after chronic ammonia exposure and the mechanism behind these effects through the detection of changes in serum immune parameters, histology, and TLR pathway-related molecules in the spleen and head-kidney. Our results are also conducive to monitoring the health status and welfare of M. amblycephala in intensive aquaculture systems.

2. Results

2.1. Ammonia Content in Spleen and Head-Kidney

Figure 1 showed ammonia content in the spleen and head-kidney of M. amblycephala after 30 days of exposure to various levels of ammonia. A significant increase in ammonia level was observed in the spleen of fish exposed to 30 mg/L total ammonia nitrogen compared with the control group (p < 0.05). There was no statistical difference in ammonia levels in head kidney between all ammonia treatment groups and control group (p > 0.05).
2.2. Serum Cortisol, Lysozyme, C3 and C4 Levels

As showed in Figure 2, serum cortisol levels increased significantly in all ammonia treatment groups compared with the control group \((p < 0.05)\). Contrarily, serum lysozyme activities were significantly decreased in the fish exposed to higher concentrations of ammonia (20 and 30 mg/L total ammonia nitrogen) \((p < 0.05)\). Complement C3 and C4 levels in the serum were significantly reduced in the groups treated with 10 to 30 mg/L total ammonia nitrogen \((p < 0.01)\).

![Figure 2](image)

**Figure 2.** Effects of chronic ammonia exposure on levels of serum cortisol, lysozyme, C3 and C4 \((A-D)\) in *M. amblycephala*. Each column represents mean ± SE of six duplicates. The values of \(p < 0.05\) and 0.01 are represented as “*” and “**” above the column, indicating significant difference versus control, respectively.

2.3. Immunity Organ Indexes

Both the spleen and head-kidney indexes increased with the increase in total ammonia nitrogen concentrations after exposure (Figure 3). Compared with the control group, the spleen index was significantly elevated in the fish treated with higher total ammonia nitrogen (20 and 30 mg/L) \((p < 0.05)\). No significant difference was detected in the head-kidney index between all ammonia treatment groups and control group \((p > 0.05)\).

![Figure 3](image)

**Figure 3.** Effects of chronic ammonia exposure on spleen \((A)\) and head-kidney indexes \((B)\) in *M. amblycephala*. Values are expressed as Min to Max \((n = 18)\). The values of \(p < 0.05\) and 0.01 are represented as “*” and “**” above the column, indicating significant difference versus control, respectively.
2.4. Pathological Evaluation

The spleen of Wuchang bream in the control group showed a normal appearance with abundant erythrocytes and leukocytes (Figure 4A). After exposure to ammonia for 30 days, slight increases in the numbers of erythrocytes were noted along with the occurrence of melano-macrophage centers in the spleen of fish exposed to 5 mg/L total ammonia nitrogen (Figure 4B, Table 1). Similar but more serious changes were observed in fish treated with 10 to 30 mg/L total ammonia nitrogen, such as increases in the number and size of melano-macrophage centers as well as markedly increased erythrocyte numbers (Figure 4C,D). In addition, cytoplasm vacuolization was also detected in the spleen of fish exposed to 30 mg/L total ammonia nitrogen (Figure 4E, Table 1).

![Figure 4. Light microscope photographs of spleens of M. amblycephala (H&E).](image)

- **A**: Spleen tissue of control fish showing normal architecture.
- **B–D**: Spleen tissues from fish exposed to 5, 10 and 20 mg/L ammonia nitrogen showing an increase in erythrocytes and melano-macrophage centers (black arrows).
- **E**: Spleen tissue from fish exposed to 30 mg/L ammonia nitrogen showing increased melano-macrophage centers (black arrows) as well as cytoplasm vacuolization (white arrows). Scale bar = 50 µm.
Table 1. Quantitative analysis of histopathological changes in the spleen and head-kidney of *M. amblycephala* under chronic ammonia exposure.

| Tissue        | Lesions                     | 0  | 5   | 10  | 20  | 30  |
|--------------|-----------------------------|----|-----|-----|-----|-----|
| Spleen       | Melano-Macrophage centers   | 0.00 ± 0.00 | 0.00 ± 0.00 | 1.00 ± 0.58 | 1.33 ± 0.67 | 1.33 ± 0.33 |
|              | Increased erythrocytes      | 0.00 ± 0.00 | 2.67 ± 0.88 * | 5.33 ± 0.67 ** | 6.00 ± 0.00 ** | 6.00 ± 0.00 ** |
|              | Cytoplasm vacuolation       | 0.00 ± 0.00 | 0.00 ± 0.00 | 0.00 ± 0.00 | 0.00 ± 0.00 | 0.67 ± 0.33 * |
| Head-kidney  | Melano-Macrophage centers   | 0.00 ± 0.00 | 0.00 ± 0.00 | 0.67 ± 0.67 | 0.67 ± 0.33 | 2.33 ± 0.67 * |

*: Six different grades (0 unchanged–6 severe) indicated increased levels of histopathological damage. All values are expressed as mean ± SE (*n* = 3). The values of *p* < 0.05 and 0.01 are represented as “*” and “**” above the column, indicating significant difference versus control, respectively.

As for the head-kidney, Figure 5A shows a normal structure of head-kidney tissue in control fish after exposure. No obvious pathological sign was noted in the head-kidney of fish exposed to lower total ammonia nitrogen (5 mg/L) (Figure 5B, Table 1). Slight increases in the number of melano-macrophage centers were observed in fish exposed to 10 and 20 mg/L total ammonia nitrogen (Figure 5C, D, Table 1). The size and number of melano-macrophage centers were enhanced in the 30 mg/L total ammonia nitrogen group (Figure 5E, Table 1).

---

**Figure 5.** Light microscope photographs of head-kidney tissue of *M. amblycephala* (H & E). (A, B) The head-kidney of fish exposed to 0 and 5 mg/L ammonia nitrogen showing normal structure. (C–E) The head-kidney from fish exposed to 10, 20 and 30 mg/L ammonia nitrogen showing the increase in melano-macrophage centers in number and size (black arrows). Scale bar = 50 μm.
2.5. Tissue Immune Parameter Analysis

Levels of innate immune parameters (TNF-α, IL-1β) and acquired immune parameter (IgM) in the spleen and head kidney decreased along with the increase in ammonia exposure levels (Figure 6, Supplementary Table S1). In the spleen, the contents and mRNA levels of TNF-α and IgM were significantly decreased in all ammonia treatment groups compared with the controls \((p < 0.05)\). Meantime, splenic IL-1β content and mRNA expression levels were significantly down-regulated in the fish exposed to greater than 20 mg/L total ammonia nitrogen \((p < 0.05)\). By contrast, only the content and mRNA level of IL-1β as well as the expression level of \(\text{tnf-}\alpha\) in the head-kidney were markedly reduced in ammonia treatment groups relative to controls \((p < 0.05)\). There were no significant differences in the levels of IgM protein and gene expression as well as TNF-α concentration in head kidney between ammonia-treated groups and the control group \((p > 0.05)\).

Figure 6. Relative mRNA levels of TLR signaling pathway genes as well as the contents of TNF-α, IL-1β and IgM in the spleen and head-kidney of \(\text{M. amblycephala}\). Values expressed as means ± SE of six duplicates. \(\text{igm}\): immunoglobulin M, \(\text{il-1β}\): interleukin 1β, \(\text{tnf-}\alpha\): tumor necrosis factor α, \(\text{jk}\): tumor necrosis factor receptor-associated \(\text{jk}\) B, \(\text{erk}\): extracellular regulated protein kinase, \(\text{nf-}\alpha\): nuclear factor-κ gene binding, \(\text{pi}\): phosphatidylinositol 3-kinase, \(\text{akt}\): protein kinase B, \(\text{myd88}\): myeloid differentiation factor 88, \(\text{traf6}\): tumor necrosis factor receptor-associated \(\text{jk}\) 6, \(\text{tlr}\)1/2/3/4/5: toll-like receptor 1/2/3/4/5. The values of \(p < 0.05\) and 0.01 are represented as “*” and “**” above the column, indicating significant difference versus control, respectively.

To elucidate the molecular mechanism of chronic ammonia toxicity on the immune system, transcriptional changes of key genes involved with the TLR signaling pathway were determined along with the levels of TNF-α, IL-1β and IgM. In the spleen, mRNA levels of toll-like receptor genes (\(\text{tlr}\)2, \(\text{tlr}\)4), MyD88-independent pathway associated gene \(\text{traf6}\), NF-κB signaling pathway associated gene \(\text{nf-}\kappa\beta2\) and MAPK signaling pathway associated gene \(\text{erk}\)1 were significantly down-regulated in the ammonia treatment groups compared with the control \((p < 0.05)\), while there were no remarkable changes in the expression of \(\text{tlr}\)1, \(\text{tlr}\)3, \(\text{tlr}\)5, \(\text{pi}\)3, \(\text{akt}\), \(\text{myd88}\), \(\text{nf-}\kappa\beta1\), \(\text{jk}\)1 and \(\text{p}\)38 \(\alpha\) \((p > 0.05)\). Similarly, transcription levels of \(\text{tlr}\)1, \(\text{tlr}\)2, \(\text{tlr}\)5, \(\text{akt}\), \(\text{traf6}\), \(\text{nf-}\kappa\beta1\), \(\text{erk}\)1 and \(\text{jk}\)1 were significantly reduced in the head-kidney of fish following chronic ammonia exposure \((p < 0.05)\). However, there were
no significant differences in expression of \textit{tlr3}, \textit{tlr4}, \textit{myd88}, pi3 \textit{k}, \textit{nf-\textit{kb}2} and \textit{p38\alpha} in the head-kidney between ammonia treatment groups and the control group ($p > 0.05$).

2.6. Correlation Analysis

As shown in Figure 7 and Supplementary Table S2, ammonia exposure concentration was significantly negatively correlated with the contents of TNF-\textit{\alpha}, IL-1\textit{\beta} and IgM as well as transcriptional levels of \textit{tnf-\alpha}, \textit{il-1\beta}, \textit{igm}, \textit{tlr2}, \textit{traf6}, \textit{nf-\textit{kb}2} and \textit{jnk1} in the spleen. Meanwhile, the contents and transcription levels of splenic TNF-\textit{\alpha}, IL-1\textit{\beta} and IgM exhibited significant positive correlations with each other ($p < 0.05$). Splenic IgM concentration showed a significant positive correlation with \textit{tlr2} and \textit{nf-\textit{kb}2}, and TNF-\textit{\alpha} as well as IL-1\textit{\beta} exhibited significant positive correlations with \textit{tlr2}, \textit{traf6} and \textit{nf-\textit{kb}2} ($p < 0.05$). However, in the head-kidney, ammonia exposure levels were only remarkably negatively correlated with IL-1\textit{\beta} content and the expression of genes \textit{tnf-\alpha}, \textit{il-1\beta}, \textit{tlr1/2/4/5}, \textit{traf6}, \textit{akt}, \textit{nf-\textit{kb}1}, \textit{jnk1} and \textit{erk1} ($p < 0.05$). Head-kidney IL-1\textit{\beta} was positively correlated with IgM, \textit{igm}, \textit{il-1\beta}, \textit{tlr4}, \textit{myd88}, \textit{traf6}, \textit{akt}, \textit{nf-\textit{kb}1}, \textit{erk1} and \textit{jnk1}, while IgM and TNF-\textit{\beta} exhibited a positive correlation with \textit{p38\alpha} and \textit{tlr2}, respectively ($p < 0.05$).

![Figure 7](image-url)

**Figure 7.** Spearman correlation coefficients among ammonia exposure concentration and immune parameters involved with TLR signaling pathway in the spleen and head-kidney of \textit{M. amblycephala} following chronic ammonia exposure. The values of $p < 0.05$ and 0.01 are represented as "*" and "**" above the column, indicating significant difference versus control, respectively.

2.7. IBR Indices

IBR values were calculated from the standardized data of 22 biomarkers in spleen and head-kidney in Wuchang bream under different levels of ambient ammonia after 30-day exposure (Table 2, Figure 8). As the exposure concentrations of ammonia increased, the IBR values tended to increase in both spleen and head-kidney. The values of IBR were higher in the spleen than in the head-kidney when the fish were exposed to the same treatment conditions (Table 2).
Table 2. Integrated biomarker response (IBR) indices of all measured parameters in the spleen and head-kidney of *M. amblycephala* to ammonia.

| Ammonia Nitrogen (mg/L) | Spleen | Head-Kidney |
|-------------------------|--------|-------------|
| 0                       | 0.00   | 0.00        |
| 5                       | 0.34   | 0.34        |
| 10                      | 0.48   | 0.47        |
| 20                      | 0.69   | 0.42        |
| 30                      | 0.87   | 0.95        |

Figure 8. Star plots for biomarker responses in spleen and head-kidney in *M. amblycephala* exposed to ammonia nitrogen.

3. Discussion

How the immune system of fish responds to different kinds of environmental pollutants has become a hot topic [34–37]. As a ubiquitous toxicant, ammonia may have an unexpected impact on aquatic animal health since fish often suffer from chronic ammonia stress in realistic situations. Our present study clarified the influences of long-term exposure to ammonia on fish immunity and its mechanism.

In our present experiment, serum cortisol increased significantly with the increase in ammonia exposure concentration, indicating the occurrence of stress in Wuchang bream. Similar results were observed in turbot (*Scophthalmus maximus*), common carp (*Cyprinus carpio*), and juvenile blunt snout bream (*Megalobrama amblycephala*) [32,38,39]. Chronic ammonia exposure also led to significant decreases in serum lysozyme, complement C3 and C4 in this study. Lysozyme, as an indicator of innate immune function, is the primary immune enzyme for fighting infections [40]. The complement system, as one of the bridges between innate immunity and acquired immunity, plays essential roles in clearing immune complexes, killing disease-causing bacteria and viruses [41,42]. Some previous studies have reported that acute ammonia stress can inhibit lysozyme activities [17,43] and decrease protein and mRNA levels of C3 and C4 in fish [44–46]. Thus, our present results revealed that chronic ammonia stress subverted the innate immunity of fish.

In the present study, concentration-dependent increases in ammonia concentrations were found both in the spleen and head-kidney, but a significant difference was only de-
ected in the spleen. Similar increases were found in the spleen and head-kidney indexes, indicating that ammonia stress caused severe damage to two main immune organs. Histological findings further revealed that the accumulation of ammonia might be the main cause of pathological injuries in the spleen and head-kidney, which were characterized by increased melano-macrophage centers and cytoplasm vacuolization. Melano-macrophage centers are known as macrophage aggregates, and their main functions are in the storage of cell-derived phospholipid and iron following erythrophagocytosis, deposition of resistant pathogens, and antigen processing in immune responses [47–51]. The increased size and frequency of melano-macrophage centers often indicates that fish are experiencing stress or damage or growing older [36,48,52]. The research of Kwon and Chang reported that a 5-day ammonia exposure (4.0~10.4 mg/L total ammonia nitrogen) induced severe hemosiderin deposition and increased melanin-macrophages in the spleen of black seabream [53]. Similar melano-macrophage assembly was observed in the spleen of *Pelteobagrus vachellii* exposed to 1 and 5 mg/L total ammonia nitrogen [17]. In addition, quantitative evaluation of histopathological alterations showed that the degree of pathological injury caused by persistent ammonia stress was higher in the spleen than in head-kidney.

Immune organ damage in fish is always associated with alterations of immune molecules [35]. In the innate immune system, the inflammatory cytokines IL-1β and TNF-α are the important regulators of the initiation and modulation of inflammatory response [54–57]. Decreased levels of cytokine molecules (TNF-α, IL-1β) represent a reduction of cellular immunological function, which ultimately results in a high infection rate under external stressors [58–60]. Limited studies have documented that acute exposure to 40 mg/L total ammonia nitrogen can elevate transcriptional levels of cytokines such as IL-1β and TNF-α in turbot (*Scophthalmus maximus*), whereas chronic exposure to high ammonia (50 mg/L total ammonia nitrogen) induced the opposite trend in crucian carp (*Carassius auratus*) [14,60]. Our present results showed that chronic ammonia exposure caused marked decreases in protein and transcriptional levels of splenic IL-1β and TNF-α, which were further supported by the Spearman correlation analysis between ammonia concentrations and the levels of splenic IL-1β and TNF-α. Similar results were also found in the head-kidney. Our current data indicated that chronic ammonia exposure disrupted the innate immune defense by inhibiting transcription and protein synthesis of inflammatory molecules IL-1β and TNF-α in two immune organs, the spleen and head-kidney.

In addition to the innate immune defense system, adaptive immunity also plays a crucial role in fish immune defense. IgM is the first antibody secreted by the adaptive immune response to a new infection or to a foreign antigen, which are mainly expressed in the spleen and head-kidney of fish [61,62]. Lower levels of IgM may be associated with weak adaptive immunity [63]. In the present study, significant decreases in protein and transcriptional levels of IgM were detected in the spleen but not in the head-kidney, indicating that ammonia exposure has an inhibitory effect on the production of splenic IgM. Qin et al. reported that exposure to 1 mg/L total ammonia nitrogen for 48 h and 96 h significantly decreased transcriptional levels of *igm* in the spleen of *Pelteobagrus vachellii*, but not in the head-kidney [17]. Decreased IgM levels were also found in the gill, spleen and brain of *Rhyynchocypris lagowski* exposed to 0.99 mg/L un-ionized ammonia for 96 h [18]. Our Spearman correlation analysis further showed that ammonia exposure concentration was significantly negative correlated with splenic IgM, but not significantly correlated with head-kidney IgM. Thus, our study suggested that chronic ammonia stress diminished the adaptive immunity only in the spleen. That might suggest that the spleen is more vulnerable to ambient ammonia stress than the head-kidney. One reason for this hypothesis may be the discrepant impairment of the structure of the spleen and head-kidney induced by ammonia stress in our present study.

In the present study, decreased transcriptional levels of TLRs were detected in the spleen and head-kidney of Wuchang bream, implying that ammonia could cause negative effects via the TLR signaling pathway. Indeed, the inhibition of TLRs might be related to the
Int. J. Mol. Sci. 2022, 23, 3129

Increases in serum cortisol induced by ammonia exposure. Carrizo et al. proved that cortisol treatment decreased mRNA levels of TLRs (tlr1/5 m/9/22) in Oncorhynchus mykiss myotubes [64]. Susarla et al. found that cortisol could decrease transcriptional levels of cytokines (VEGF, CCL5, IFN-γ, CXCL-10, IL-8 and GCSF) after either or both TLR3 and TLR4 stimulation of primary human corneal fibroblasts [65]. Furthermore, the TLR inhibition might imply a block of TLR signaling pathways. The MyD88-dependent pathway is regulated by all TLRs, and the MyD88-independent pathway is peculiar to the TLR3 and TLR4 signaling pathway [66]. TLR signaling also leads to PI3 K-AKT pathway activation, which in turn activates B cells to produce antibody and synthesize proinflammatory molecules such as IL-1β and TNF-α through the NF-κB signaling pathway [67–69]. It is worth noting that transcriptional levels of traf6 were decreased in both spleen and head-kidney after ammonia exposure, while no significant changes were detected in the expression of myd88 and pi3 k. Thus, our results indicated that ammonia exerted interference with the MyD88-independent pathway by inhibiting TRAF6 in both spleen and head-kidney. Additionally, the TRAF6 activation can trigger the activation of downstream NF-κB signaling and MAPK signaling (p38, JNK and ERK), which ultimately induces the production of various inflammatory cytokines including TNF-α and interleukin-1β/6/12 (IL-1β/6/12) [70,71]. In addition, TRAF6 is also the crucial mediator for CD40 signaling that regulates IL-6 and Ig secretion [72]. In our study, the significant down-regulation of splenic erk1 and nf-kb2 mRNAs as well as head-kidney erk1, jnk1 and nf-kb1 mRNAs suggested that ammonia exposure suppressed the downstream NF-κB and MAPK signaling molecules in the spleen and head-kidney. Moreover, ammonia exposure down-regulated NF-κB signaling by suppressing gene nf-kb1 in the spleen but gene nf-kb2 in the head-kidney, which reflects the tissue-specific inflammatory response between spleen and head-kidney. As for the MAPK signaling pathway, ammonia exposure significantly inhibited the expression of splenic erk1 and head-kidney erk1 and jnk1. After performing Spearman correlation analysis, we found that ammonia exposure concentration was significantly negatively correlated with the levels of immune parameters (IL-1β, TNF-α and IgM) as well as the expression of TLR signaling-related genes in the spleen and head-kidney. Therefore, our results suggested that chronic ammonia exposure could impair innate and adaptive immunity via the TLRs/MyD88-independent signaling pathway.

High IBR values reflected the enhanced biological responses and poor health condition of the organisms [73]. The present results for IBR values increased with the increases in ambient ammonia concentration in both spleen and head-kidney. This result was consistent with the data on histopathology and serum-immune parameters. Moreover, under the same concentration of ammonia stress, the IBR value was higher in spleen than in head-kidney, which further proved that the influences of ammonia stress on the spleen were more severe than on the head-kidney of Wuchang bream.

4. Materials and Methods

4.1. Animal Maintenance and Experimental Protocol

Juvenile Wuchang bream with a mean weight of 9.98 ± 0.48 g from Tuanfeng Fishery (Hubei, China) were moved to 300 L fiberglass tanks, where they were supplied with running de-chlorinated and continuously aerated water. They were acclimated for 14 d by feeding commercial diets twice a day (9:00 am and 15:00 pm). After a preliminary experiment, the 96 h LC50 and safe concentrations of total ammonia nitrogen for bream juveniles were found to be 46.013 mg/L and 4.601 mg/L, respectively. Therefore, the exposure range of ammonia was set as 0 mg/L, 5 mg/L, 10 mg/L, 20 mg/L and 30 mg/L total ammonia nitrogen. Fish were randomly distributed into 5 treatment groups with triplicate tanks, and the stoking density was 20 juveniles per tank. The experimental concentrations of ammonia were obtained from a stock solution of 10 g/L made with reagent-grade ammonia chloride (BASFR, 99.5%). To maintain stable ammonia treatment concentrations, the 1/2 experiment solution was renewed every day by a new and equal concentration of ammonia nitrogen solution. Meanwhile, the control group underwent
similar manipulation with aerated tap water. The real total ammonia nitrogen levels for each treatment were measured by nesslerization every day. During the whole experimental period, the daily monitoring data by HQ40 D Water Analyzer (Hach, Loveland, CO, USA) showed that water temperature was 25.0 ± 0.60 °C, dissolved oxygen concentrations were above 5.0 mg/L, and pH was kept at 7.55 ± 0.04, with slight adjustments using 10% H₂SO₄ and 10% NaOH. Fish were fed a commercial diet twice per day as same as the acclimation.

4.2. Sample Collection and Preparation

After a 30-day experimental exposure, the fish were fasted for 48 h firstly and then were anesthetized in MS-222 solution. The body weight of each juvenile was measured before dissection, and the blood was collected immediately from the caudal veins to separate the serum for subsequent biochemical parameter determination. The tissues (spleen and head-kidney) were extracted and weighed. The spleen and head-kidney indexes were calculated using the formula [weight of tissue (g)/body weight (g) * 100%]. Three individual spleens and head-kidneys from each group were used for histopathological analysis, and the others were frozen immediately at −80 °C for the analysis of tissue ammonia concentrations, immunity parameters and gene transcriptional levels. This work conducted on M. amblycephala was approved by the Animal Care and Use Committee (IACUC) of Huazhong Agricultural University, Wuhan, China (HZAUFI-2019-018).

4.3. Ammonia Detection in Spleen and Head-Kidney

For the analysis of ammonia levels, the pretreatment of tissue samples and ammonia detection was carried out according to previous studies [74,75]. The detailed test steps can be found in Supporting Information (Supplementary Text S1).

4.4. Serum Immune Parameters Assay

Blood samples were centrifuged (845 g, 20 min, 4 °C) for preparation of serum samples. Each experimental group had six replicates and one replicate included the serum from 10 individuals of the same tank. Serum cortisol was determined by radioimmunoassay (RIA) using the kit of Beijing North Institute of Biotechnology Co., Ltd. (Beijing, China) (www.bnibt.com, accessed on 15 March 2020) according to previous study [76]. The contents of complement C3 and complement C4 and the activity of lysozyme were measured using commercial kits produced by Jiancheng Bioengineering Institute (Nanjing, China) (www.njcbio.com, accessed on 15 March 2020).

4.5. Tissue Immune Parameters Assay

Tissue samples from five fish of the same group were pooled as one duplicate and homogenized with 0.85% sodium chloride, then centrifuged at 845 × g for 15 min (4 °C) for collection of supernatant. Each experimental group had six replicates. As described in previous studies [77,78], supernatant was used to test the concentrations of IgM, IL-1β, TNF-α and total protein using commercial kits. All kits were purchased from Jiancheng Bioengineering Institute (Nanjing, China) (www.njcbio.com, accessed on the 15 March 2020).

4.6. Gene Expression Analysis

The Mrna levels of immune-associated genes in the spleen and head-kidney were tested according to the method in a previous study [79]. Detailed information on the testing procedure is provided in Supporting Information (Supplementary Text S2). The primer sequences are listed in Table 3. Melt curve analysis was performed for each primer at the end of the reaction to demonstrate the reaction specificity. After verifying that the amplification efficiencies of all selected genes ranged from 90–110%, the relative Mrna levels were calculated by the method of 2⁻ΔΔCt with β-actin as the internal control [80].
Table 3. Sequences of primers used for Qpcr amplification.

| Target Gene | Primer Sequences (from 5′ to 3′) | Accession Number and/or References | Amplification Efficiency |
|-------------|-----------------------------------|------------------------------------|-------------------------|
| igm         | F: TGGAGCAACGCCACAGCTATT R: CTCTTGGGACTCGCACCATT | KC894945 | 99.03% |
| il-1β       | F: ACGATAAGACGCCAAGGACAC | KF515511 | 104.87% |
| tnf-a       | F: TCCAAGGCCAATCATCCTTATT | KF515512 | 103.97% |
| jnk1        | F: AGCAACCCCTACTAACAACATG | MK315047 | 109.47% |
| erk1        | F: TCCCTGAAGGCGCTGAATATAC | MK315044 | 101.48% |
| p38a        | F: TGGGACCGGAATATCAACAAAC | MK315052 | 91.86% |
| nf-κβ       | F: TGATGGGACGCCAGCTATT | MK315050 | 108.68% |
| nf-κβ2      | F: AACTACCAGTGAGCCGGTGG | MK315051 | 99.24% |
| pi3 k       | F: GGCGTAAACATCCAGCTTTGC | Liang et al. [81] | 93.04% |
| akt         | F: GCCTGGTAAAGGCAAGCTTTTG | Liang et al. [81] | 101.56% |
| myd88       | F: TGGAAACAGAGCTAATCAACR | KP192128 | 109.31% |
| tlr1        | F: TCCCTGCTTACGATCTCTGTG | KP192129 | 109.86% |
| tlr2        | F: TTACTCGGCGCTTGGTGGTGG | KX196269 | 109.45% |
| tlr3        | F: TTGTGGAGAAGACGCAACACCAC | DQ986365 | 108.82% |
| tlr4        | F: TGCTGCGCGAAGCAACGTTAAG | KR092315 | 91.99% |
| tlr5        | F: GGAGGACCATCTCCTACCA | KX196271 | 97.26% |
| β-actin     | F: ACCCACACCGTGCCCATCTA | AY170122 | 108.48% |

4.7. Histopathological Evaluation

Spleen and head-kidney samples were fixed in 10% neutral-buffered formalin, then dehydrated in ethanol and embedded in paraffin. Finally, tissue slices were stained with hematoxylin and eosin (H&E). Further quantitative analysis of pathological changes was performed according to previous studies [82,83]. The detailed analysis protocol is supplied in Supporting Information (Supplementary Text S3).

4.8. Integrated Biomarker Response Analysis

IBR analysis is a method for integrating all of the measured biomarker responses into an integrative index to assess stress levels [84–86]. In the current study, it was applied to evaluate the toxic effects of different ammonia concentrations on the spleen and head-kidney of Wuchang bream after a 30-day exposure. Detailed information on the calculation procedure is provided in Supporting Information (Supplementary Text S4).

4.9. Statistical Analyses

All values were subjected to one-way analysis of variance followed by Dunnett’s post hoc test to evaluate differences between means (SPSS 22.0, Chicago, IL, USA). Spearman correlation analysis was chosen to determine the relationship among total ammonia concentrations, immune parameters and gene expression levels. Normality and variance homogeneity were previously verified. Differences were measured and considered to be significant at the $p$ value < 0.05.

5. Conclusions

Our study provides evidence that chronic ammonia stress elevated serum COR, reduced serum lysozyme and C3/C4, and decreased the protein and transcriptional levels of IL-1β, TNF-α and IgM as well as the expression levels of genes involved with TLRs/MyD88-independent signaling pathway in the spleen and head-kidney of *M. amblycephala*. In addition, the indexes of and pathological damage to the two immune organs increased with tissue ammonia accumulation. These findings indicated that ammonia accumulation in the spleen and head-kidney caused histopathological damage and induced immune suppression through inhibition of TLRs/MyD88-independent signaling. In addition, the adverse influences of chronic ammonia stress on the spleen were shown to be more severe than on the head-kidney of Wuchang bream.
Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/ijms23063129/s1.

Author Contributions: All authors contributed to the article and approved the submitted version. L.L. conceptualization, funding acquisition, methodology, project administration, resources, writing—review and editing; H.G. conceptualization, data curation, formal analysis, investigation, visualization, software, writing—original draft; S.C. data curation, formal analysis, investigation, visualization, software; Y.K., K.O., H.Y. and Y.W. investigation; R.T., X.Z. and D.L. supervision, resources. All authors have read and agreed to the published version of the manuscript.

Funding: This work was jointly supported by the National Key R&D Program of China (2019 YFD0900303), the China Agriculture Research System of MOF and MARA (CAS-45-24) and the National Natural Science Foundation of China (No. 32071621).

Institutional Review Board Statement: We adhered to all local, national and international regulations and conventions, and we respected normal scientific ethical practices. The specimens used in this study came from a population mainly of commercially fished individuals intended for human consumption. The animal protocol was approved by the Institutional Animal Care and Use Ethics Committee of Huazhong Agricultural University (Wuhan, China) (HZAUIF2019-018).

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: All datasets generated for this study are included in the article/Supplementary Materials.

Conflicts of Interest: The authors declare on conflict of interest.

References

1. Randall, D.J.; Tsui, T. Ammonia toxicity in fish. Mar. Pollut. Bull. 2002, 45, 17–23. [CrossRef]
2. David, A.A.; Debbie, C.; Allen, W.K.; John, E.C. Interaction of Ionized and Un-Ionized Ammonia on Short-Term Survival and Growth of Prawn Larvae, Macrobrachium rosenbergii. Bioi. Bull. 1978, 154, 15–31. [CrossRef]
3. Thurston, R.V.; Russo, R.C.; Vinogradov, G.A. Ammonia toxicity to fishes. Effect of pH on the toxicity of the unionized ammonia species. Environ. Sci. Technol. 1981, 15, 837–840. [CrossRef]
4. Armstrong, B.M.; Lazorchak, J.M.; Murphy, C.A.; Haring, H.J.; Jensen, K.M.; Smith, M.E. Determining the effects of ammonia on fathead minnow (Pimephales promelas) reproduction. Sci. Total Environ. 2012, 420, 127–133. [CrossRef] [PubMed]
5. Eddy, F.B. Ammonia in estuaries and effects on fish. J. Fish Biol. 2005, 67, 1495–1513. [CrossRef]
6. Miron, D.S.; Moraes, B.; Becker, A.G.; Crestani, M.; Spanevello, R.; Loro, V.L.; Baldisserotto, B. Ammonia and pH effects on some metabolic parameters and gill histology of silver catfish, Rhamdia quelen (Heptapteridae). Aquaculture 2008, 277, 192–196. [CrossRef]
7. Yu, C.; Huang, X.; Chen, H.; Godfray, H.C.J.; Wright, J.S.; Hall, J.W.; Gong, P.; Ni, S.; Qiao, S.; Huang, G.; et al. Managing nitrogen to restore water quality in China. Nature 2019, 567, 516. [CrossRef]
8. World Health Organization (WHO). A Global Overview of National Regulations and Standards for Drinking-Water Quality, 2nd ed.; World Health Organization: Geneva, Switzerland, 2021; 66p.
9. Passell, H.D.; Dahm, C.N.; Bedrick, E.J. Ammonia modeling for assessing potential toxicity to fish species in the RIO grande, 1989–2002. Ecol. Appl. 2007, 17, 2087–2099. [CrossRef]
10. Fan, B.; Li, J.; Wang, X.; Chen, J.; Gao, X.; Li, W.; Ai, S.; Gui, L.; Gao, S.; Liu, Z. Ammonia spatiotemporal distribution and risk assessment for freshwater species in aquatic ecosystem in China. Ecotox. Environ. Saf. 2021, 207, 111541. [CrossRef]
11. Yue, F.; Pan, L.Q.; Xie, P.; Zheng, D.; Li, J. Immune responses and expression of immune-related genes in swimming crab Portunus trituberculatus exposed to elevated ambient ammonia-N stress. Comp. Biochem. Physiol. A Mol. Integr. Physiol. 2010, 157, 246–251. [CrossRef]
12. Qin, C.J.; Shao, T.; Zhao, D.X.; Duan, H.G.; Wen, Z.Y.; Yuan, D.; Li, H.T.; Qi, Z.M. Effect of ammonia-N and pathogen challenge on complement component 8α and 8β expression in the darkbarbel catfish Pelteobagrus vachellii. Fish Shellfish Immunol. 2017, 62, 107–115. [CrossRef] [PubMed]
13. Cheng, C.H.; Yang, F.F.; Ling, R.Z.; Liao, S.A.; Miao, Y.T.; Ye, C.X.; Wang, A.L. Effects of ammonia exposure on apoptosis, oxidative stress and immune response in pufferfish Takifugu obscurus. Aquat. Toxicol. 2015, 164, 61–71. [CrossRef]
14. Liu, B.L.; Jia, R.; Huang, B.; Lei, J.L. Interactive effect of ammonia and crowding stress on ion-regulation and expression of immune-related genes in juvenile turbot (Scophthalmus maximus). Mar. Freshw. Behav. Physiol. 2015, 70, 179–194. [CrossRef]
15. Li, M.; Zhang, M.; Qian, Y.; Shi, G.; Wang, R. Ammonia toxicity in the yellow catfish (Pelteobagrus fulvidraco): The mechanistic insight from physiological detoxification to poisoning. Fish Shellfish Immunol. 2020, 102, 195–202. [CrossRef] [PubMed]
16. Wang, L.; Sun, Y.; Xu, B.; Sagada, G.; Chen, K.; Xiao, J.; Zhang, J.; Shao, Q. Effects of berberine supplementation in high starch diet on growth performance, antioxidative status, immune parameters and ammonia stress response of fingerling black sea bream (Acanthopagrus schlegelii). Aquaculture 2020, 527, 735473. [CrossRef]

17. Qin, C.J.; Shao, T.; Wang, Y.M.; Yang, Q.; Bu, F. Effect of ammonia-N on histology and expression of immunoglobulin M and component C3 in the spleen and head kidney of Psettobagrus vachelli. Aquac. Rep. 2017, 8, 16–20. [CrossRef]

18. Yu, Z.; Wu, X.Q.; Zheng, L.J.; Dai, Z.Y.; Wu, L.F. Effect of acute exposure to ammonia and BFT alterations on Rhynchocypris lactoski: Digestive enzyme, inflammation response, oxidative stress and immunological parameters. Environ. Toxicol. Pharmacol. 2020, 78, 103380. [CrossRef]

19. Zhang, T.; Yan, Z.; Zheng, X.; Fan, J.; Wang, S.; Wei, Y.; Yang, L.; Wang, P.; Guo, S. Transcriptome analysis of response mechanism to ammonia stress in Asian clam (Corbicula fluminea). Aquat. Toxicol. 2019, 214, 105235. [CrossRef]

20. Cheng, C.; Ma, H.; Su, Y.; Deng, Y.; Feng, J.; Xie, J.; Chen, X.; Guo, Z. Ammonia toxicity in the mud crab (Scylla paramamosain): The mechanistic insight from physiology to transcriptome analysis. Ecotox. Environ. Safe. 2019, 179, 9–16. [CrossRef]

21. Banerjee, B.; Koner, D.; Hasan, R.; Bhattacharya, S.; Saha, N. Transcriptome analysis reveals novel insights in air-breathing magur (Clarias magur) in response to high environmental ammonia. Gene 2019, 703, 35–49. [CrossRef]

22. Yu, J.; Sun, J.; Zhao, S.; Wang, H.; Zeng, Q. Transcriptome analysis of oriental river Prawn (Macrobrachium nipponense) Hepatopancreas in response to ammonia exposure. Fish Shellfish Immunol. 2019, 93, 223–231. [CrossRef] [PubMed]

23. Renneseth, A.; Wergeland, H.I.; Pettersen, E.F. Neutrophils and B-cells in Atlantic cod (Gadus morhua L.). Fish Shellfish Immunol. 2007, 23, 493–503. [CrossRef] [PubMed]

24. Wang, X.; Xing, H.; Jiang, Y.; Wu, H.; Sun, G.; Xu, Q.; Xu, S. Accumulation, histopathological effects and response of biochemical markers in the spleens and head kidneys of common carp exposed to atrazine and chlorpyrifos. Food Chem. Toxicol. 2013, 62, 148–158. [CrossRef] [PubMed]

25. Bromage, E.S.; Kaattari, I.M.; Zwollo, P.; Kaattari, S.L. Plasmablast and plasma cell production and distribution in trout immune tissues. J. Immunol. 2004, 173, 7317–7323. [CrossRef] [PubMed]

26. Zwollo, P.; Haines, A.; Rosato, P.; Gumulak-Smith, J. Molecular and cellular analysis of B-cell populations in the rainbow trout (Oncorhyncus mykiss). Fish Shellfish Immunol. 2008, 24, 1482–1496. [CrossRef]

27. Tort, L. Stress and immune modulation in fish. Dev. Comp. Immunol. 2011, 35, 1366–1375. [CrossRef]

28. Magnadottir, B. Innate immunity of fish (overview). Fish Shellfish Immunol. 2006, 20, 137–151. [CrossRef]

29. Whyte, S.K. The innate immune response of finfish—A review of current knowledge. Fish Shellfish Immunol. 2007, 23, 1127–1151. [CrossRef]

30. Food and Agriculture Organization (FAO). World Aquaculture Production of Fish, Crustaceans, Molluscs, etc., by Principal Species [Internet]. Rome: FAO, 2020. Available online: http://www.fao.org/tempref/FI/STAT/summary/a-6.pdf (accessed on 23 February 2022).

31. Sun, S.M.; Ge, X.P.; Zhu, J.; Zhao, S.; Wang, H.; Zeng, Q. Transcriptome analysis of shrimp for immune function of different diets. J. Fish Biol. 2018, 80, 540–545. [CrossRef] [PubMed]

32. Zhang, W.; Sun, S.; Ge, X.; Xia, S.; Zhu, J.; Miao, L.; Ren, M.; Lin, Y.; Ge, X.; Sun, S. Growth performance, physiological response and histology changes of juvenile blunt snout bream, Megalobrama amblycephala exposed to chronic ammonia. Aquaculture 2019, 506, 424–436. [CrossRef]

33. Zhang, W.; Sun, S.; Ge, X.; Xia, S.; Zhu, J.; Miao, L.; Yu, H. Acute effects of ammonia exposure on the plasma and haematological parameters and histological structure of the juvenile blunt snout bream, Megalobrama amblycephala, and post exposure recovery. Aquac. Res. 2018, 49, 1008–1019. [CrossRef] [PubMed]

34. Aksakal, F.I.; Ciltas, A. Impact of copper oxide nanoparticles (CuO NPs) exposure on embryo development and expression of genes related to the innate immune system of zebrafish (Danio rerio). Comp. Biochem. Physiol. Part C Toxicol. Pharmacol. 2019, 223, 78–87. [CrossRef] [PubMed]

35. Lin, W.; Li, L.; Chen, J.; Li, D.; Hou, J.; Guo, H.; Shen, J. Long-term crowding stress causes compromised nonspecific immunity and increases apoptosis of spleen in grass carp (Ctenopharyngodon idella). Fish Shellfish Immunol. 2018, 80, 540–545. [CrossRef] [PubMed]

36. Lin, W.; Guo, H.; Wang, L.; Zhang, D.; Wu, X.; Li, L.; Qiu, Y.; Yang, L.; Li, D.; Tang, R. Waterborne microcystin-LR exposure induced chronic inflammatory response via MyD88-dependent toll-like receptor signaling pathway in male zebrafish. Sci. Total Environ. 2020, 702, 134969. [CrossRef]

37. Armando, E.V.; Jordan, C.; Zhao, X.; Micah, J.G.; Jaclyn, E.C. In vivo effects on the immune function of fathead minnow (Pimephales promelas) following ingestion and intraperitoneal injection of polystyrene nanoparticles. Sci. Total Environ. 2020, 735, 139461. [CrossRef]

38. Ruyet, J.P.L.; Lamers, A.; Roux, A.L.; Severe, A.; Boeuf, G.; Mayer-Gostan, N. Long-term ammonia exposure of turbot: Effects on plasma parameters. J. Fish Biol. 2003, 62, 879–894. [CrossRef]

39. Mirghaeder, A.T.; Fayaz, S.; Hoseini, S.M. Effects of dietary 1, 8-cineole supplementation on serum stress and antioxidant markers of common carp (Cyprinus carpio) acutely exposed to ambient ammonia. Aquaculture 2019, 509, 8–15. [CrossRef]

40. Demers, N.E.; Bayne, C.J. The immediate effects of stress on hormones and plasma lysozyme in rainbow trout. Dev. Comp. Immunol. 1997, 21, 363. [CrossRef]

41. Holland, M.C.H.; Lambris, J.D. The complement system in teleosts. Fish Shellfish Immunol. 2002, 12, 399–420. [CrossRef]
42. Hawlisch, H.; Köhl, J. Complement and Toll-like receptors: Key regulators of adaptive immune responses. Mol. Immunol. 2006, 43, 13–21. [CrossRef]

43. Kim, S.H.; Kim, J.H.; Park, M.A.; Hwang, S.D.; Kang, J.C. The toxic effects of ammonia exposure on antioxidant and immune responses in Rockfish, Sebastes schlegelii during thermal stress. Environ. Toxicol. Phur. 2015, 40, 954–959. [CrossRef] [PubMed]

44. Ren, Q.; Li, M.; Yuan, L.; Song, M.; Xing, X.; Shi, C.; Meng, F.; Wang, R. Acute ammonia toxicity in crucian carp Carassius auratus and effects of taurine on hyperammonemia. Comp. Biochem. Physiol. Part C Toxicol. Pharmacol. 2016, 190, 9–14. [CrossRef]

45. Xing, X.; Li, M.; Yuan, L.; Song, M.; Ren, Q.; Shi, C.; Meng, F.; Wang, R. The protective effects of taurine on acute ammonia toxicity in grass carp Ctenopharyngodon idella. Fish Shellfish Immunol. 2016, 56, 517–522. [CrossRef] [PubMed]

46. Jia, R.; Liu, B.; Han, C.; Huang, B.; Lei. J. Effects of ammonia exposure on stress and immune response in juvenile turbot Scophthalmus maximus. Aquac Res. 2017, 48, 3149–3162. [CrossRef]

47. Agius, C. The role of melanophore-macrophage centres in iron storage in normal and diseased fish. J. Fish Dis. 1979, 2, 337–343. [CrossRef]

48. Agius, C. Preliminary studies on the ontogeny of the melano-macrophages of teleost haemopoietic tissue and age-related changes. Dev. Comp. Immunol. 1981, 5, 597–606. [CrossRef]

49. Agius, C. The melanophore-macrophage centres of fish: A review. Fish Immunol. 1985, 85–105. [CrossRef]

50. Agius, C.; Agbede, S.A. Electron microscopical studies on the genesis of lipofuscin, melanin and haemosiderin in the haemopoietic tissues of fish. J. Fish Biol. 1984, 24, 471–488. [CrossRef]

51. Roberts, R.J. Cloning and expression of TNF-α in Salmo salar L. during the smolting period. Fish Shellfish Immunol. 2006, 20, 450–461. [CrossRef] [PubMed]

52. Brown, C.L.; George, C.J. Age-dependent accumulation of macrophage aggregation in the yellow perch, Perca flavescens (Mitchill). J. Fish Dis. 1985, 8, 135–138. [CrossRef]

53. Kwon, J.Y.; Chang, Y.J. Effects of ammonia concentration on histological and physiological status in black seabream (Acanthopagrus schlegeli). Korean J. Fish. Aquat. Sci. 1996, 29, 828–836.

54. Baud, V.; Karin, M. Signal transduction by tumor necrosis factor and its relatives. Trends Cell Biol. 2001, 11, 372–377. [CrossRef]

55. Ingerslev, H.C.; Cunningham, C.; Wergeland, H.I. Cloning and expression of TNF-α, IL-1β and COX-2 in an anadromous and landlocked strain of Atlantic salmon (Salmo salar L.) during the smolting period. Dev. Comp. Immunol. 2019, 56, 377–389. [CrossRef] [PubMed]

56. Zhang, A.; Chen, D.; Wei, H.; Du, L.; Zhao, T.; Wang, X.; Zhou, H. Functional characterization of TNF-α in grass carp head kidney leukocytes: Induction and involvement in the regulation of NF-κB signaling. Fish Shellfish Immunol. 2012, 33, 1123–1132. [CrossRef]

57. Yang, X.; Wei, H.; Qin, L.; Zhang, S.; Wang, X.; Zhang, A.; Du, L.; Zhou, H. Reciprocal interaction between fish TGF-β1 and IL-1β is responsible for restraining IL-1β signaling activity in grass carp head kidney leukocytes. Dev. Comp. Immunol. 2014, 47, 197–204. [CrossRef]

58. Lin, W.; Hou, J.; Gao, H.; Qiu, Y.; Li, L.; Li, D.; Tang, R. Dualistic immunomodulation of sub-chronic microcystin-LR exposure on the innate-immune defense system in male zebrafish. Chemosphere 2017, 183, 315–322. [CrossRef]

59. Qi, X.Z.; Xue, M.Y.; Yang, S.B.; Zha, J.W.; Wang, G.X.; Ling, F. Ammonia exposure alters the expression of immune-related and antioxidant enzymes-related genes and the gut microbial community of crucian carp (Carassius auratus). Fish Shellfish Immunol. 2017, 60, 485–492. [CrossRef]

60. Stenvik, J.; Schroder, M.B.; Olsen, K.; Zapata, A.; Jorgensen, T.O. Expression of immunoglobulin heavy chain transcripts (vh-families, igm, and igd) in head kidney and spleen of the atlantic cod (Gadus morhua L.). Dev. Comp. Immunol. 2001, 25, 291–302. [CrossRef]

61. Xia, H.; Wu, K.; Liu, W.; Gu, Y.; Wang, W.; Zhang, X. Molecular cloning and expression analysis of immunoglobulin M heavy chain gene of blunt snout bream (Megalobrama amblycephala). Fish Shellfish Immunol. 2014, 40, 129–135. [CrossRef]

62. Zhang, L.; Bassig, B.A.; Mora, J.L.; Vermeulen, R.; Ge, Y.; Curry, J.D.; Hu, W.; Shen, M.; Qiu, C.; Ji, Z.; et al. Alterations in serum immunoglobulin levels in workers occupationally exposed to trichloroethylene. Carcinogenesis 2013, 34, 799–802. [CrossRef]

63. Carrizo, V.; Valenzuela, C.A.; Zuloaga, R.; Aros, C.; Molina, A. Effect of cortisol on the immune-like response of rainbow trout Oncorhynchus mykiss) myotubes challenged with Piscirickettsia salmonis. Vet. Immunol. Immunopathol. 2021, 237, 110240. [CrossRef] [PubMed]

64. Susarla, R.; Liu, L.; Walker, E.A.; Bujalska, I.J.; Alsalem, J.; Williams, G.P.; Sreekantam, S.; Taylor, A.E.; Tallouzi, M.; Southworth, H.S.; et al. Cortisol Biosynthesis in the Human Ocular Surface Innate Immune Response. PLoS ONE 2014, 9, e94913. [CrossRef] [PubMed]

65. Akira, S.; Takeda, K.; Kaisho, T. Toll-like receptors: Critical proteins linking innate and acquired immunity. Nat. Immunol. 2001, 2, 679–680. [CrossRef] [PubMed]

66. Omori, S.A.; Rickert, R.C. Phosphatidylinositol 3-kinase (PI3K) signaling and regulation of the antibody response. Cell Cycle 2007, 6, 397–402. [CrossRef]
69. Troutman, T.D.; Hu, W.; Fulenchek, S.; Yamazaki, T.; Kurosaki, T.; Bazan, J.F.; Pasare, C. Role for B-cell adapter for PI3K (BCAP) as a signaling adapter linking Toll-like receptors (TLRs) to serine/threonine kinases PI3K/Akt. *Proc. Natl. Acad. Sci. USA* 2012, 109, 273–278. [CrossRef]

70. Kobayashi, T.; Walsh, M.C.; Choi, Y. The role of traf6 in signal transduction and the immune response. *Microbes Infect.* 2004, 6, 1333–1338. [CrossRef]

71. Akira, S.; Uematsu, S.; Takeuchi, O. Pathogen recognition and innate immunity. *Cell* 2006, 124, 783–801. [CrossRef]

72. Jalukar, S.V.; Hostager, B.S.; Bishop, G.A. Characterization of the roles of tnf receptor-associated factor 6 in cd40-mediated b lymphocyte effector functions. *J. Immunol.* 2000, 164, 623–630. [CrossRef]

73. Brooks, S.J.; Harman, C.; Hultman, M.T.; Berge, J.A. Integrated biomarker assessment of the effects of tailing discharges from an iron ore mine using blue mussels (*Mytilus* spp.). *Sci. Total Environ.* 2015, 524–525, 104–114. [CrossRef]

74. Zhang, Y.L.; Zhang, H.L.; Wang, L.Y.; Gu, B.Y.; Fan, Q.X. Changes of ammonia, urea contents and transaminase activity in the body during aerial exposure and ammonia loading in Chinese loach *Paramisgurnus dabryanus*. *Fish Physiol. Biochem.* 2017, 43, 631–640. [CrossRef] [PubMed]

75. Brown, R.H.; Duda, G.D.; Handler, P. A colorimetric micromethod for determination of ammonia; the ammonia content of rat tissues and human plasma. *Arch. Biochem. Biophys.* 1957, 66, 301–309. [CrossRef]

76. Rance, T.A.; Baker, B.I. The in vitro response of the trout interrenal to various fragments of ACTH. *Gen. Comp. Endocrinol.* 1981, 45, 497–503. [CrossRef]

77. Liang, H.; Ji, K.; Ge, X.; Ren, M.; Liu, B.; Xi, B.; Pan, L. Effects of dietary arginine on antioxidant status and immunity involved in AMPK-no signaling pathway in juvenile blunt snout bream. *Fish Shellfish Immunol.* 2018, 78, 69–78. [CrossRef] [PubMed]

78. Xia, S.L.; Li, X.F.; Abasubong, K.P.; Xu, C.; Shi, H.J.; Liu, W.B.; Zhang, D. Effects of dietary glucose and starch levels on the growth, apparent digestibility, and skin-associated mucosal non-specific immune parameters in juvenile blunt snout bream (*Megalobrama amblycephala*). *Fish Shellfish Immunol.* 2018, 79, 193–201. [CrossRef] [PubMed]

79. Guo, H.; Lin, W.; Hou, J.; Wang, L.; Zhang, D.; Wu, X.; Li, L.; Li, D. The Protective Roles of Dietary Selenium Yeast and Tea Polyphenols on Growth Performance and Ammonia Tolerance of Juvenile Wuchang Bream (*Megalobrama amblycephala*). *Front. Aquat. Physiol.* 2018, 9, 1371. [CrossRef]

80. Livak, K.J.; Schmittgen, T.D. Analysis of relative gene expression data using real-time quantitative PCR and the 2(-Delta Delta C(T)) Method. *Methods 2001, 25, 402–408. [CrossRef] [PubMed]

81. Liang, H.; Mokrani, A.; Chisomo-Kasiya, H.; Ji, K.; Ge, X.; Ren, M.; Liu, B.; Xi, B.; Sun, A. Dietary leucine affects glucose metabolism and lipogenesis involved in TOR/PI3K/Akt signaling pathway for juvenile blunt snout bream *Megalobrama amblycephala*. *Fish Physiol. Biochem.* 2019, 45, 719–732. [CrossRef]

82. Bernet, D.; Schmidt, H.; Meier, W.; Burkhardt-Holm, P.; Wahl, T. Histopathology in fish: Proposal for a protocol to assess aquatic pollution. *J. Fish Dis.* 1999, 22, 25–34. [CrossRef]

83. Corbett, P.A.; King, C.K.; Mondon, J.A. Application of a quantitative histological health index forantarctic rock cod (*Trematomus bernacchii*) from davis station, east Antarctica. *Mar. Environ. Res.* 2015, 109, 28–40. [CrossRef]

84. Beliaeff, B.; Burgeot, T. Integrated biomarker response: A useful tool for ecological risk assessment. *Environ. Toxicol. Chem.* 2002, 21, 1316–1322. [CrossRef] [PubMed]

85. Kim, W.K.; Lee, S.K.; Jung, J. Integrated assessment of biomarker responses in common carp (*Cyprinus carpio*) exposed to perfluorinated organic compounds. *J. Hazard. Mater.* 2010, 180, 395–400. [CrossRef] [PubMed]

86. Huang, X.; Lan, Y.; Liu, Z.; Huang, W.; Guo, Q.; Liu, L.; Hu, M.; Sui, Y.; Wu, F.; Lu, W.; et al. Salinity mediates the toxic effect of nano-TiO2 on the juvenile olive flounder *Paralichthys olivaceus*. *Sci. Total Environ.* 2018, 640–641, 726–735. [CrossRef] [PubMed]