Effect of Quercetin on Infectious Pancreatic Necrosis Virus: in Vitro Replication Study

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Abstract

Infectious Pancreatic Necrosis Virus (IPNV) is a viral agent that causes significant losses in salmonid fish in the rainbow trout and salmon farming sectors. The present study aimed to examine the effects of quercetin on IPNV replication in Rainbow trout gonad (RTG-2) cell cultures. The half-maximal (50%) cytostatic concentration (CC₅₀) of quercetin was detected in the RTG-2 cell culture by cytotoxicity test with Cell Counting Kit-8 (CCK-8, Sigma-Aldrich) (n=3). Titration, cytopathogenic effect reduction, and qRT-PCR tests were used to determine the effect of quercetin on IPNV replication in RTG-2 cell culture. At the end of the cytotoxicity test, the non-toxic dose of quercetin was determined. Besides that, 50 μmol/L quercetin decreased the titer of IPNV in RTG-2 cells from 10⁻⁷ to 10⁻⁵, at the same time, the same dose of quercetin reduced the viral load to 40%. We concluded that quercetin is a potent antioxidant agent for IPNV infections and it can be an option in the treatment of the IPN disease.

Introduction

Aquaculture is one of the fastest growing sectors in the world, due to the increasing demand for it in food production systems (Lymbery, 2000) and rainbow trout (Oncorhynchus mykiss) is one of the most produced fish species in aquaculture both in Turkey and the world (Anonymous, 2013). IPNV is a viral agent that causes significant losses in salmonid fish in the rainbow trout and salmon farming sectors. The virus is a double-stranded RNA virus classified within the Aquabirnavirus genus of the Birnaviridae family (Anonymous, 2009). IPNV prevalence was determined many times both in Turkey and the world (Suzuki & Nojima, 1999; Candan, 2002; Nishizawa et al. 2008; Ogut & Altuntas, 2011). So, it is obvious that effective antiviral agents are necessary for successful treatment. On this matter, many investigations of antiviral activities had been done on Infectious Hematopoietic Necrosis Virus (IHNV) and Viral Hemorrhagic Septicemia Virus (VHSV) using fustin, fisetin, butine and sulfuretin flavonoids that were extracted from Rhus verniciflua (Kang et al. 2012).

Oxidative stress is the increase in amount of free radicals that called reactive oxygen species (ROS) produced by physiological events during normal metabolism in all cells and rendered harmless by antioxidants (Halliwell, 1994). Many studies have shown that eukaryotic cells are the source of these free radicals as they use oxygen to generate energy from normal metabolic activities, moreover, viruses may cause cell damage due to the oxidative stress that occurs during replication (Muller, 1992; Najafi et al. 2014; Gullberg et al. 2015) and affect pathogenicity (Stehbens & Nojima, 2004; Dikici et al. 2005; Ivanov et al. 2013). The damage of reactive oxygen products is inevitable for all fishes that breathe oxygen. Oxidative stress and antioxidant defense mechanisms in fish have been investigated in...
Antioxidants are substances that can control free radicals and minimize their effects against the damage of reactive oxygen species. Also it has been determined that flavonoids have effective antioxidant activities in this regard by preventing the formation of ROS or by removing ROS directly (Frankel et al. 1993). Quercetin is also an important aglycone flavonoid that known to be a powerful natural antioxidant both in vitro and in vivo (Nabavi, Nabavi, Esfami & Moghaddam, 2012). Quercetin has been reported to be effective for human health as antioxidant, anti-cancer, anti-diabetic, anti-inflammatory, and blood pressure regulator (Kelly, 2011; Larson et al. 2012). In recent years, the antiviral activity of quercetin on many viruses such as Influenza virus, Ebola virus, Chikungunya virus, Epstein-Barr virus, Hepatitis C virus, and Mayaro virus has also been investigated (Lani et al. 2015; Lee et al. 2015; Abdal et al. 2015; Qiu et al. 2016). In this study, the effect of quercetin on IPNV replication in the fish cell cultures, and the non-toxic dose of quercetin on Rainbow Trout Gonad-2 (RTG-2) cell were examined for the first time.

Materials and Method

RTG-2 Cell Culture Preparation

In this study, RTG-2 continuous cell culture and IPNV SDF4 strain, obtained from the stocks of Virology Department, Faculty of Veterinary Medicine, Ondokuz Mayis University, were used for the production, titration and cytotoxicity tests of quercetin. RTG-2 cells that used for virus inoculation and cell suspension were grown in Leibovitz’s L-15 Medium (1X) (Gibco Life Technologies Cat. No: 11415-056, UK) in 25 cm² cell culture flasks (Corning, NY 14831, USA) with the addition of 10% fetal calf serum (Sigma- Aldrich Cat: No: F9665, USA) with 1% penicillin-streptomycin antibiotic solution (Biological Industries, Cat, 03-33-1B, Israel). Then, the cells were removed daily from the 22°C oven to be monitored on a tissue culture microscope (Olympus, CKX41, Japan). Then the suspension of RTG-2 cells that contains 1x10⁶ cells per milliliter was prepared and used in the cytotoxicity test.

IPNV Inoculation

When the cells covered approximately 80% of the flask surface as a monolayer during the virus cultivation stage, IPNV inoculation was performed by absorbing method after pouring the cell production medium. At the end of the incubation period, L-15 medium with bovine serum was added and then was flask incubated at 15°C oven. Cells that contains the virus inoculum was checked daily for tissue cytopathic effect (CPE) by tissue culture microscope. Approximately 48-72 hours later, CPE formation was monitored, then cells that containing the virus were frozen at -80°C and thawed three times, after that centrifugation at +4°C and 3000 rpm were done, and the resulting virus supernatants were placed into vials and stored at -80°C deep freezing until used.

Calculation of Half-Maximal (50%) Cytostatic Concentration (CC 50) of Quercetin (Cytotoxicity Tests)

Quercetin, whose effect on IPNV replication, was obtained from Sigma-Aldrich (Cat. No: 849061-97-8, USA). The half-maximal (50%) cytostatic concentration (CC 50 ) of quercetin on RTG-2 cells was determined by (WST-8) viability test using Cell Counting Kit-8 (CCK-8, Sigma-Aldrich, Cat. No: 96992, USA). The test was performed on a 96-well plate (TPP zellkultur test plate 96F, Cat. No:92096, Switzerland) according to the method reported by the manufacturer. Five different concentrations of quercetin (10, 20, 50 and 100 μmol/L) were used to determine the half-maximal (50%) cytostatic concentration (CC 50) of quercetin. The plates were read at the end of the test in a microplate reader (Tecan Infinite F50) with an optical density (OD) of 450 nanometers (nm) and the OD of each well was recorded on the microplate reader. The test was repeated 3 times and CC 50 of quercetin was calculated according to the formula “[(OD test-OD blank) / (OD control-OD blank)] x 100”.

Titration Test

Microtiteration test was performed to calculate the cell infective (TCID50) of IPNV with and without quercetin at half-maximal cytostatic concentration (CC 50 ). For this purpose, a serial dilution of IPNV according to log10 was made and 100 μl was placed in 4 wells of each microtitre tablet for each dilution. The same procedure was repeated in IPNV treated with quercetin at non-toxic concentrations (10, 20 and 50 μmol/L). Then, 50 μl of the cell suspension was added to the dilutions with 300,000 live cells per mL. Microtitre plates were monitored daily under the microscope for CPE formation and when CPE was formed in all four eyes of the virus control, test results were evaluated and titers were calculated according to Spearman-Kaerber method. (Frey & Liess, 1971).

Cytotoxicopathogenic Effect (CPE) Reduction Test

The antiviral activity of IPNV in IPNV-infected cells with and without quercetin was performed by the method reported by Kujumgiev et al. (1999) using the “Cytotoxicopathogenic Effect (CPE) Reduction Test”. Test results were evaluated as score 0, 0% CPE; score 1, 0–25% CPE; score 2, 35–50% CPE; score 3, 50–75% CPE; and score 4, 75–100% CPE.
Calculation of Viral Load by Quantitative Real-Time PCR (qRT-PCR)

Viral RNA obtained from IPNV supernatants that produced in RTG-2 cells which include different doses of quercetin at 10, 20 and 50 µmol/L that determined as non-toxic dose in cytotoxicity test was extracted with GeneJET RNA Purification Kit (Thermo, K0732) according to the manufacturer’s instructions. For qRT-PCR, the number of viral particles was determined on the Biorad CFX Connect qRT-PCR using the iTaqTM Universal Probes One-Step Kit (Biorad, Cat. No: 172-5141) and IPNV primers and probes (Table 1). At the end of the test, the results were evaluated in Bio-Rad CFX Manager 3.1.

The mean value of the data obtained from three replicated cytotoxicity experiments was calculated. The differences in viral load between the three replicates, controls and the different doses of quercetin were expressed as %.

Results

Production of IPNV in RTG-2 Cell Culture

RTG-2 cells used in virus generation, titration, cytotoxicity, and CPE reduction tests were photographed after tissue culture microscopy (Figure 1A). Then IPNV was inoculated on the RTG-2 cells prepared with the aim of producing the virus and the images were taken at 24, 48 and 72 hours while checking by tissue culture microscope (Figure 1 / B-C-D).

Cytotoxicity Test Results of Quercetin

The CC 50 of quercetin on RTG-2 cells was determined using (WST-8) viability test Cell Counting Kit-8 (CCK-8, Sigma-Aldrich). The mean of each concentration and control was plotted at different times for three trials. According to this; the Viability of cells treated with 50 µmol quercetin calculated 52% after 24 hours and 50% at the end of 48 hours. Whereas, the viability of the cells treated with 100 µmol quercetin was 47% after 24 hours and 49% after 48 hours (Figure 2).

Also, four flask cell cultures were grown (25 cm²) to calculate the effect of non-cytotoxic concentrations (10, 20 and 50 µmol/L) of quercetin on IPNV replication. When RTG-2 cells were occupied 80% of base flasks the cell growth medium was spilled and non-toxic doses (10, 20 and 50 µmol/L) of quercetin were added at three flasks. To prepare the virus control, L15 medium was added in one flask and all of the four flasks were incubated for 2 hours at 22˚C. At the end of 2 hours, all contents were spilled and IPNV was inoculated in the four flasks then virus growth medium was added into all flasks. After that, the CPE formations were followed and photographed for 1 week (Figure 3).

Microtitration Test Results

At the end of the microtitration test, IPNV titers were calculated as $10^{-7}/0.1$ ml for the untreated cells with quercetin, $10^{-6}/0.1$ ml for the treated cells with 10µmol/L quercetin and $10^{-5}/0.1$ ml for cells that treated with both 20 and 50 µmo/L quercetin (Table 2).

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Table 1 Primers and probe used for RT-PCR analysis of IPNV

<table>
<thead>
<tr>
<th>Name</th>
<th>Primers and Probe</th>
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<tbody>
<tr>
<td>VP3-F</td>
<td>F: 5’-TCTCCGGGCCATCAAGT-3’</td>
</tr>
<tr>
<td>VP3-R</td>
<td>R: 5’-CGGTCCGATGGTTT-3’</td>
</tr>
<tr>
<td>VP3-P</td>
<td>P:5’-FAM-CCAGAACCAGAGTGATGAGTACAT-3’-TAMRA</td>
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Figure 1. A: RTG-2 cell Culture (72h), B: 24th hour CPE image of IPNV C: 48th hour CPE image of IPNV, D: 72th hour CPE image of IPNV (x10).
Cytopathogenic Effect (CPE) Reduction Test Results

The CC50 quercetin concentration (50 /mol / L) was detected by CKK-8 assay in the RTG-2 cells and CPE of IPNV were scored by CPE reduction test under invert microscope to determine the effect of quercetine concentration on IPNV replication. At the end of the test, CPE formation was measured as score 1 (0-25%) at flask 1 and virus control flask but no change as (score 0) was observed in flasks 2 and 3 after 24 hours. In addition, 35-50% CPE formation was scored as 2 in flask 1 and IPNV virus control flask but 0-25% CPE formation was observed in the 2nd and 3rd flask and they were scored as 1 at the end of 72 hours (Figure 3).

Quantitative Real-Time PCR (qRT-PCR) Test Results

The viral load of IPNV virus control and IPNV that was treated with different quercetin concentrations were calculated using the Biorad CFXManager 3.1 program. At the end of the qRT-PCR test, it was determined that viral loads were decreased 5% after 10µmol/L quercetin application, 35% after 20µmol / L quercetin application and 40% after 50µmol/L quercetin application compared to IPNV virus control (Figure 4).
Natural compounds are used as an important source for the discovery of new antiviral drugs because of their low side effects. Flavonoids have been known to have antiviral activity since the 1940s, and there are many studies on antiviral activity of various flavonoids (Özcelik et al. 2011; Ganesan et al. 2012; Yu et al. 2014; Cheng et al. 2015; Wu et al. 2016; Rojas et al. 2016). It has been reported that quercetin which is an important flavonoid and commonly found in foods in the form of aglycone or glycoside can be used as a protecting agent against oxidative stress Kumar, Sharma, Khanna & Raj (2003) and reduces susceptibility to viral infection. (Davis, Murphy, McClellan, Carmichael & Gangemi 2008). In this study, in vitro effect of quercetin in IPNV replication was investigated in term of its antiviral effect in viral infections.

It had been reported by Cheng et al. (2015) that semi-maximal cytostatic concentration (CC50) of quercetin as 50 μmol/L in HepG2.2.15 (hepatocellular carcinoma) and HuH-7 cells was significantly inhibited Hepatitis B Virus (HBV) replication in both cells at different time intervals (2,4, and 6th day). In another study, it had been reported that HepG2 2.2.15 cells can afford the semi-maximal cytostatic concentration (CC50) of quercetin to more than 100 μmol / L (Romero et al. 2005). In this study, the semi-maximal (50%) cytostatic concentration of quercetin in RTG-2 cell at 48 hours was found to be 50 μmol/L (CC50) similar to the results of Cheng et al. (2015) who investigated the effect of quercetin on HBV replication, and detected that the inhibition of virus replication was started at 10 μmol/L quercetin concentrations, also HBV intracellular viral DNA levels in HepG2.2.15 and HuH7 cells were calculated as 68.5% and 52.9%, respectively. In addition, the level of HBV transcripts at 50 μmol L quercetin concentration was analyzed and it was recorded that it inhibited viral transcription by (51.8%) significantly (Cheng et al. 2015). Wu et al. (2016) had been stated that the inhibition effect of quercetin was 49.21% on influenza virus that was incubated with quercetin of 50 μmol/mL concentration at 4°C for 30 minutes and then cultured at 37 °C for 48 hours. Also, a study of Huh-7.5 cells that were infected with HCV JFH1 strain, revealed that viral load of the cells that were treated with 50 μM quercetin decreased as 22.6% compared to the control cells (Rojas et al. 2016).

In this study, we compared the viral load of IPNV inoculated cells with viral load of cells that were treated with different concentrations of quercetin before IPNV inoculation. Our results for the percentage of viral loads reduction of IPNV were 5% at 10 μmol / L quercetin, 35% at 20 μmol / L quercetin and 40% at 50 μmol / L quercetin. These results were found similar to other studies on different viruses that were detected antiviral

**Table 2 Infectious power of IPNV after non-quercetin and different doses of quercetin.**

<table>
<thead>
<tr>
<th>APPLICATIONS</th>
<th>DKID50 VALUES</th>
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<tbody>
<tr>
<td>IPNV</td>
<td>10⁻⁷/0.1 ml</td>
</tr>
<tr>
<td>IPNV+10μmol/L Quercetin</td>
<td>10⁻⁸/0.1 ml</td>
</tr>
<tr>
<td>IPNV+20μmol/L Quercetin</td>
<td>10⁻⁹/0.1 ml</td>
</tr>
<tr>
<td>IPNV+50μmol/L Quercetin</td>
<td>10⁻¹⁰/0.1 ml</td>
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**Figure 4. IPNV viral load after quercetin at different concentrations.**
activity of quercetin. Wu et al. (2016) had been reported that quercetin dose has reduced the HA mRNA transcription of influenza virus and 24 hours after the infection, virial NP protein synthesis in MDCK cells was inhibited. Also, another study examined the effect of quercetin on HCV JFH1 strain and proved it caused decrease in infectious titer of virus (Rojas et al. 2016). In this research, the effect of quercetin on virus replication was determined by comparing quercetin-treated and non-treated IPNV titers. Accordingly, we revealed that the infectivity of non-quercetin-treated IPNV was calculated as$10^{7.91}$ ml but infectivity for the treated IPNV with 10, 20 and 50 μmol / l quercetin were decreased depends on the doses of quercetinise as 10-6, 10-5 and 10-5, respectively. Our data was showed that the inhibitory effect of quercetin on IPNV was similar to HCV and influenza viruses studies.

As a result, this study was shown that quercetin has in vitro inhibitory effect on IPNV virus. and it may be considered as an option to treat IPN disease, but comprehensive animal experiments are needed for this matter.

References
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