Research Article

Downregulation of Aquaporin 3 Mediated the Laxative Effect in the Rat Colon by a Purified Resin Glycoside Fraction from Pharbitis semen

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Background. Pharbitis Semen, the seeds of Pharbitis nil, is widely used as a traditional purgative medicine in China, Korea, and Japan. This study investigated the laxative effects of a purified resin glycoside fraction obtained in our previous study from Pharbitis semen in vivo and in vitro. Materials and Methods. After orally administering a purified resin glycoside fraction from Pharbitis Semen (RFP) to rats, the content of fecal water, AQP3, NF-κB, COX-2 expression, and the prostaglandin E2 (PGE2) concentrations in the colon were examined. Moreover, human intestinal epithelial cells (HT-29) were used to investigate the mechanism of RFP decreasing the AQP3 expression. Results. Results obtained showed that treatment with RFP increased the feces excretion and fecal water content of rats in a dose-dependent manner. More interestingly, AQP3 expression was suppressed by RFP treatment both in the rat colons and in HT-29 cells, while the NF-κB pathway-mediated PGE2 production was activated. Interestingly, pretreating rats with BAY-11-7082 (NF-κB inhibitor) or indomethacin (COX-2 inhibitor) and RFP neither induced diarrhea nor decreased the AQP3 expression in the colon. Conclusions. The purgative property of the purified resin glycoside fraction was attributed to NF-κB activation in the colon, which increased the COX-2-mediated secretion of PGE2. PGE2 decreased AQP3 expression which inhibits water absorbed from the intestine to the blood vessel side, resulting in the laxative effect of RFP.

1. Introduction

Aquaporins (AQPs) are membrane proteins that function as water/glycerol channels and play a vital role in water transport across cell membranes. To date, 13 types of AQPs channels have been identified [1]. There are 9 types of AQP in the intestine, such as AQPs: 1-4 and AQPs: 7-11 [2, 3]. Among them, AQP3 is the most important aquaporin of the colon. It is located in the colonic villus epithelial cells and contributed to the transport of water [4]. It has been reported previously that many factors could lead to laxative effect through altering AQP3 expression to prevent the water transport in colon, such as gut hormone, 5-hydroxytryptamine, bacterial pathogens, and laxative MgSO4 [5–7].

Resin glycosides, as the characteristic constituents of Convolvulaceae, are reported to be responsible for the drastic purgative behavior of all the important morning glory family species used in traditional medicines. Pharbitis semen, the seeds of Pharbitis nil (Convolvulaceae), are widely used to treat constipation [8]. Previous studies only proposed that Pharbitis Semen triggered diarrhea through accelerating peristalsis in the colon resulting in water elimination [9]. However, the material basis and molecular mechanism involved in the laxative effect of Pharbitis Semen have not yet been elucidated. Many studies have demonstrated that aquaporins (AQPs) mediated the cathartic effect of laxatives [5, 10, 11]. The nuclear factor-kappa B (NF-κB) is important in regulating cellular responses. There are many known NF-κB pathway activators such as stress, activation of oncoproteins and kinases, cytokines, and dysregulation of cell receptors [12]. It was reported that the expression of AQP3 in the rat colon was downregulated via transcription factors NF-κB activation [13]. Another research reported that PGE2, a synthetic product of COX-2, is of vital importance for AQP3 regulation to generate the laxative effect in rhubarb extract [14]. We hypothesized that the laxative effect of Pharbitis Semen may...
be related to AQP3 regulation in the colon mediated by NF-
κB activation and possibly involvement of PGE₂ production.

In this study, we investigated the material basis of Phar-
bitis Semen induction of diarrhea and explored the mech-
anism of a purified resin glycoside fraction from Pharbitis
Semen (RFP) on AQP3 protein expression in vitro and in
vivo.

2. Materials and Methods

2.1. Extraction and Isolation of the Purified Resin Glycoside Fraction (RFP). The purified resin glycoside fraction from
Pharbitis Semen (RFP) was obtained from our previous work
[15]. The MeOH-insoluble fraction (Fr.C’) was derivatized by
NH₄ silica gel on-column catalyzation to obtain individual
constituents. Eleven acylated resin glycosidic acid methyl
esters were obtained in this fraction [15]. Therefore, Fr.C’
was characterized as a resin glycoside fraction from Pharbitis
Semen (RFP).

2.2. Materials. RFP was dissolved in carmellose sodium
(CMC-Na) or dimethyl sulfoxide (DMSO) before the admin-
istration of rats or cells, respectively. Indomethacin and
MgSO₄ were obtained from Aladdin (Shanghai, China). BAY II-7082 (BAY) was purchased from MCE (HY-13453, NJ,
USA). Prostaglandin E₂ (PGE₂) was purchased from Santa
Cruz Biotechnology (sc-201225A, TX, USA). PGE₂ ELISA
kit was purchased from R&D Systems, Inc. (St. Louis, MO,
USA). The following primary antibodies were used: AQP3
(abi25229, Abcam, Cambridge, UK), COX-2 and p-p65
(Cell Signaling Technology, Boston, USA), Glyceraldehyde
Phosphate Dehydrogenase (GAPDH) and Proliferating Cell
Nuclear Antigen (PCNA) (Beyotime, Nanjing, China), and
mouse IgG HRP and Alexa Fluor® 647 Goat Anti-Rabbit IgG
(Fcmrccs, Nanjing, China).

2.3. Animals. Nine-week-old male specific pathogen-free SD
rats (220-250 g) were obtained from the Experimental Ani-
mal Center of Yangzhou University (Yangzhou, China). Keep
these animals in climate-controlled facilities with automatic
light and dark cycles and allow free access to water and
standard food. It is kept and treated in strict accordance with
the obligations of the Animal Ethics Committee of China
Pharmaceutical University and the guidelines for the care and
use of laboratory animals of the National Institutes of Health.

2.4. Animals Treatment. The rats were divided separately into
two groups randomly. The first group (N1 = 30) was divided
separately into five small groups with the equal number:
the normal control group, the RFP (31.25, 62.5, and 125
mg/kg, respectively)-treated groups, and MgSO₄ (2 g/kg)-
treated group. RFP or MgSO₄ was orally administered to the
five group rats (water was provided ad libitum) after fasting
for 18 h. The second group (N2=24) was divided separately
into four small groups (vehicle, RFP-treated vehicle, BAY,
or indomethacin combined with RFP administration group).
BAY (10 mg/kg) or indomethacin (20 mg/kg) was intraper-
toneally administered to rats 1 h before RFP treatment. After
different treatment for 6 h, all these rats were sacrificed and
their colons were removed. The colon was washed with PBS
and then rapidly frozen with liquid nitrogen and stored at
-80°C. Fecal samples were disposed as described formerly
[14]. The water content of feces was calculated based on the
difference between wet and dry fecal weights.

2.5. Immunohistochemistry and Histological Examination of
Rat Intestine. Rats were sacrificed at 6 h after RFP admin-
istration. The colons were removed and fixed immediately for
3 h in 4% paraformaldehyde after washing with PBS. Three
tissue samples of every group dissected from colon were used.
Procedures of immunohistochemistry and hematoxylin and
eosin (H&E) staining were carried out as formerly described
[11, 16]. Tissue slices were visualized using NanoZoomer 2.0
RS (Beijing, China).

2.6. Cell Culture. HT-29 cells were purchased from the Cell
Bank of Shanghai Institute of Biochemistry and Cell Biology,
Chinese Academy of Sciences (Shanghai, China). This cell
line was cultured in PRMI-1640 with 10% fetal bovine serum
(FBS, GIBCO, USA), 100U/ml penicillin, and 100mg/ml streptomycin at 37°C with 5% CO2.

2.7. Cell Viability Assay. Cell viability was measured by MTT
assay which has been described previously [11]. Cells were
administered with a series of concentrations of RFP for 6
h. Cell viability was calculated using the following for-
mula: %Cell viability=AI/A0 x100. A1 and A0 represent the
absorbance values of the RFP treatment and control, respec-
tively.

2.8. Immunofluorescence Assay. Immunofluorescence assay
was done according to the procedures used before [11]. After
staining, the cells images were analyzed by ImageXpress®
Micro Confocal (Molecular Devices, USA).

2.9. Prostaglandin E₂ Measurement. Prostaglandin E₂ (PGE₂)
in the tissue of rats colon or cell supernatant was measured
using the PGE₂ ELISA kit. The PGE₂ extraction in the colon
or cell supernatant was performed according to the manufac-
turer's protocol included in the PGE₂ ELISA kit. The content
of PGE₂ in the samples was estimated from the standard
curve generated using known concentrations of PGE₂.

2.10. Preparation of Tissue and Cell Protein for Western Blot
Analysis. The removed colons from rats at 6 h after the
administration of RFP or MgSO₄ were used. Since AQP3 is
mainly expressed in the plasma membrane and the surface of
cells, the crude fraction mainly containing the AQP3 is
prepared as formerly reported [11].

The protein of AQP3 in HT-29 cells was also extracted.
Briefly, RFP-treated HT-29 cells were collected and sus-
pended in RAPI dissecting buffer. The fraction containing
AQP3 was obtained as procedures used before [17].

The phosphorylation of p65 is primarily translocated
into the nucleus. Therefore, the nuclear fraction was pre-
pared using a nuclear and cytoplasmic protein extraction kit
(Beyotime, Nanjing, China) according to the manufacturer's
protocol. Cells and tissues were treated with cytoplasmic
extraction buffer and placed on ice for 5 minutes. It was then homogenized and centrifuged (4000 x g, 4°C, 10 minutes). After discarding the supernatant, a nuclear extraction buffer was added to the particles. The suspension was centrifuged (16,000 x g, 4°C for 30 minutes) to obtain a supernatant as a core portion. Other proteins were extracted by lysis in RIPA buffer to obtain total protein. The sample was then kept on ice for 30 minutes and then centrifuged at 15,000 g for 10 minutes at 4°C.

All these protein concentrations extracted from tissue or cell were determined using a BCA protein assay kit. Western blot analyses were conducted according to the instruction manual included in the primary antibodies kit. The protein bands were detected using the ChemiDOC™ system (Bio-Rad, Hercules, CA).

2.11. Statistical Analysis. The numerical data are expressed as the means ± standard deviation and performed in triplicate. Data from multiple groups were analyzed by one-way ANOVA, followed by Tukey's Multiple Comparison Test. For all the tests, the level of significance was * P < 0.05, **P < 0.01, and ***P < 0.001 and “ns” represented that there is no significance.

3. Results

3.1. RFP Induced Diarrhea in Rats. As shown in Figure 1(a), macroscopic image of colon was observed. The stool in RFP-treated rats (31.25-125 mg/kg) was markedly less than the normal control rats, suggesting severe diarrhea happened after RFP treatment. In addition, the fecal water content after RFP administration increased in a dose-dependent manner (Figure 1(b)). The laxative effect of positive comparison group (MgSO₄/2 g/kg) is equal to the group of middle dose of RFP (62.5 mg/kg), suggesting that RFP had effective laxative activity.

3.2. RFP Decreased the Expression of AQP3 in the Colon of Rats. Changes in protein expression of AQP3 were observed in immunohistochemistry, which exists mainly in mucosal epithelial cells of the colon. An obvious decrease in the expression of AQP3 after RFP administration was observed (Figure 2(a)). Western blotting further confirmed this result (Figure 2(b)). Altogether, these results revealed that RFP decreased AQP3 expression in the mucosal epithelial cells of rat colon.

3.3. RFP Activated NF-κB Pathway and Their Downstream Proteins. Since NF-κB is involved in various kinds of diarrhea, NF-κB activation has been reported to be important to the downregulation of the AQP3 channel [16, 18]. The translocation of NF-κB into the nucleus and downstream protein COX-2 were analyzed. As shown in Figure 3(a), RFP induced NF-κB phosphorylation significantly, indicating the activation of NF-κB signal in rat colon. Similarly, RFP increased the expression of COX-2, which plays an important role in the regulation of intestinal ion secretion and barrier integrity of the colon through the action of its product prostaglandin E₂ (PGE₂). In addition, immunohistochemistry also showed the similar results. RFP treatment increased NF-κB nuclear translocation and the COX-2 activation (Figure 3(b)). To further confirm this result, the production of PGE₂ was detected. RFP administration actually increased the level of PGE₂ in the colon (Figure 3(c)). These results showed that RFP could activate NF-κB and induced PGE₂ production via COX-2 activation. As NF-κB and COX-2 are inflammatory mediators, the H&E staining after RFP administration was done. Results obtained showed that inflammation cell infiltration was observed, indicating that it was accompanied by tissue inflammation when RFP exerted purgative activity (Figure 3(d)). Moreover, RFP decreased the number of goblet cells (vacuoles) in enterocytes lining. This phenomenon might attribute to the production of PGE₂, which was reported to promote the secretion of mucins from goblet cells, thus leading to the decrease of number of vacuoles in the colon [19, 20].
3.4. NF-κB and COX-2 Inhibitors Suppressed RFP-Induced Laxative Effect. To determine the role of NF-κB and COX-2 in RFP-induced laxative effect, BAY 11-7082 (NF-κB inhibitor) and indomethacin (COX-2 inhibitor) were applied. The feces in the colon of rats and the fecal water content after RFP administration were almost recovered to the normal level by BAY 11-7082 or indomethacin pretreatment (Figures 4(a) and 4(b)). Moreover, RFP-induced activation of COX-2 and production of PGE₂ were effectively inhibited by BAY 11-7082 or indomethacin (Figures 4(c) and 4(d)). Interestingly, the RFP-induced activation of NF-κB was suppressed by BAY 11-7082 but not indomethacin, suggesting that NF-κB might be the upstream signal of COX-2. Additionally, the RFP-induced decrease in the expression of AQP3 was significantly inhibited by BAY 11-7082 or indomethacin. This observation may be ascribed to the inhibitory effect of NF-κB and COX-2 by BAY 11-7082 or indomethacin, respectively.

3.5. RFP Decreased the Expression of AQP3 in HT-29 Cells. To investigate the mechanisms of laxatives and diarrhea development, HT-29 cells which were derived from human colon cancer were used. The expression of AQP3 was suppressed in a concentration-dependent manner after RFP treatment for 6 h (Figure 5(a)). Similar to western blot analysis, immunofluorescence assay further confirmed that RFP decreased the expression of AQP3 (Figure 5(b)). Next, we examined whether the reduction effect of RFP is due to its cytotoxicity. There was almost no cytotoxicity on HT-29 cells after RFP (2-8 μg/ml) treatment for 6 h under our present experimental conditions (Figure 5(c)), suggesting that RFP might explicitly decrease AQP3 expression in vitro.

3.6. RFP Activated NF-κB Pathway and Their Downstream Proteins in HT-29 Cells. Since NF-κB was activated in vivo after RFP treatment, we investigated whether RFP could induce similar effects in HT-29 cells. Consistently in vivo, RFP caused a significant increase in the expression of NF-κB and COX-2 as well as PGE₂ production (Figures 6(a) and 6(b)). A further experiment was designed to test the role of NF-κB and COX-2 in RFP-induced decrease expression of AQP3. Pretreatment cells with BAY 11-7082 or indomethacin before RFP stimulation, NF-κB, and COX-2 as well as the production of PGE₂ were inhibited while AQP3 protein level was significantly increased compared with RFP alone (Figures 6(c) and 6(d)). Results obtained suggested that inhibition of RFP-induced activation of NF-κB or COX-2 could prevent the decrease of AQP3 expression in HT-29 cells.

3.7. PGE₂ Decreased the Expression of AQP3 in HT-29 Cells. It has been shown that PGE₂ could induce the decrease expression of AQP3 [14]. Based on the fact that RFP could induce the production of PGE₂, PGE₂ was added to HT-29 cells. Fluorescence intensity results showed that the cell membrane and cytoplasm expression of AQP3 decreased significantly after PGE₂ treatment for 6 h (Figure 7(a)). Western blot results further showed that PGE₂ markedly reduced the expression level of AQP3 in HT-29 cells (Figure 7(b)).

4. Discussion

Recently, herbal plants have received increasing attention as new therapeutic drugs for the treatment of constipation and related diseases [21, 22]. In order to develop drugs for the treatment of constipation, we studied the therapeutic effect of Pharbitis Semen (Convolvulaceae), which is used as a traditional stimulant laxative herb in Korea, China, and Japan [23]. It is reported that resin glycosides are the material basis that account for the purgative action of convolvulaceous species as traditional purgative medicine throughout the world [8]. However, mechanism investigations on the laxative effect of resin glycosides from Pharbitis Semen have yet not yet been documented. In present study, we clarified the mechanism of the laxative effect of RFP. Here we have
Figure 3: RFP activated NF-κB pathway and their downstream proteins. (a) The expressions of phosphorylated NF-κB (p-p65) in nuclear fractions and cytosolic COX-2 were subjected to western blot analysis. PCNA and GAPDH were used as loading controls of nuclear fraction and cytosolic fraction respectively. *P < 0.05, **P < 0.01 vs control group. (b) Immunohistochemical technique was used to analyze the expression of p-p65 and COX-2 in the colon. The p-p65 and COX-2 positive result presents brown, low resolution (20 x 2.5, bars: 100 μm) and amplifying image (20 x 10, bars: 25 μm). (c) The PGE₂ content was detected using the ELISA method. The control group was indicated as 100%. The data represented the means ± SDs for six rats. *P < 0.05, **P < 0.01, ***P < 0.001 vs control group. (d) The rats colons after RFP treatment were stained with H&E, low resolution (20 x 2.5, bars: 100 μm) and amplifying image (20 x 10, bars: 25 μm).
discovered and demonstrated the effects of RFP on AQP3 in vivo and in vitro. The main findings of our research are as follows: (1) RFP had the laxative effect via increasing water elimination in the colon. (2) RFP decreased the AQP3 protein expression in the colon of rats and HT-29 cells. (3) RFP activated the NF-κB and COX-2 expression in vivo and in vitro; PGE₂ acted as a performer to decrease the expression of AQP3.

Since tight junctions in colonic epithelial cells are rigid, AQPś play important roles in water transfer from the colon to the body [24]. AQPś 1-4 and AQP8 are found to be expressed in colon of animals [25, 26]. Among them, AQP3 has been extensively studied and is considered to function as a channel protein in dehydrating fecal contents [27, 28]. In this study, we found that RFP obviously decreased the AQP3 expression both in vivo and in vitro, which was confirmed by western blot analysis and immunofluorescence. In addition, a correlation was observed between a decrease in the expression level of AQP3 and an increase in the fecal water content. These results indicated that RFP administration may result in
Figure 5: RFP decreased the expression of AQP3 in HT-29 cells. (a) Cells were lysed after treatment with RFP for 6 h. Western blotting was done to measure the protein expression level of AQP3. ∗ P < 0.05, ∗∗∗ P < 0.001 vs control group. (b) HT-29 cells were incubated with RFP (0, 2, 4, 8 μg/ml) and then immunofluorescence assay was applied to observe the AQP3 expression (red) in HT-29 cells, bars: 10 μm. (c) Cells were treated with RFP at indicated concentrations for 6 h. The cellular viabilities were assessed by MTT. The data represented the means ± SDs for three experiments, ns vs control group.

A decrease in AQP3 levels in colonic mucosal epithelial cells, which prevents water from reabsorption from the luminal side and eventually leads to diarrhea.

Furthermore, we explored the mechanism involved in RFP-induced decrease of AQP3 expression. In previous studies, NF-κB activation is of great importance for the downregulation of AQP2 channel [26]. Another research reported that the COX-2 activation and PGE2 production are involved in the decrease expression of AQP3 [13]. Our investigations discovered that when NF-κB or COX-2 activation was inhibited by pretreating rats with BAY 11-7082 or indomethacin, the laxative effect of RFP was alleviated and AQP3 protein expression almost recovered to the normal level in the colon (Figure 4). These results indicated NF-κB or COX-2 activation might be involved in RFP-induced diarrhea. Additionally, we found that PGE2, which is the synthetic product of COX-2, increased significantly in a dose-dependent manner after RFP administration, revealing that PGE2 production might play an important role in RFP-induced diarrhea. This result was further confirmed by the inhibiting effect of BAY 11-7082 or indomethacin on PGE2 production when they relieved RFP-induced diarrhea.

In in vitro experiments, we examined the effect of RFP on HT-29 cells. Although they are derived from human colon cancer, they have been widely used to study the mechanism of diarrhea and laxatives due to the normal physiological state of the colon they represent [29]. Our examination showed that RFP administration to HT-29 cells significantly reduced the protein expression level of AQP3 and activated the NF-κB pathway. Consistent to experiment in vivo, we found that
Figure 6: RFP activated NF-κB pathway and their downstream proteins in HT-29 cells. (a, b) Indicated concentrations of RFP were applied to treat HT-29 cells. (a) Western blot analysed the transposition of p-p65 in the nuclear and cytoplasm COX-2. PCNA and GAPDH were used as an equal loading control. ** *P < 0.001, ns vs control group. (b) PGE₂ concentration was determined with ELISA kit, and the control cells presented as 100%. The data represented the means ± SDs for three experiments, * * *P < 0.001, ns vs control group. (c, d) Cells were pretreated with NF-κB inhibitor BAY 11-7082 (BAY) (10 μM) or COX-2 inhibitor indomethacin (20 μM) for 1 h and further treated with RFP (4 μg/ml). (c) The expressions of p-p65, COX-2 and AQP3 were detected by western blotting. ** *P < 0.01, * * *P < 0.001, ns vs control group. (d) PGE₂ concentration was measured with the same method above. The data represented the means ± SDs for three experiments, * * *P < 0.001, ns vs control group.
RFP-induced decreased expression of AQP3 and increased production of PGE₂ recovered to the normal level when pretreating cells with BAY 11-7082 or indomethacin. PGE₂, as a downstream signal of NF-κB, might directly regulate the expression of AQP3. Additionally, PGE₂ was used to stimulate HT-29 cells; as a consequence, the expression of AQP3 was significantly downregulated. These results suggested the RFP-induced diarrhea was mediated by PGE₂ secretion. Although the mechanism by which PGE₂ reduces AQP3 expression remains unclear, it may increase endocytosis and degradation of AQP3.

NF-κB is a quick responder of cellular responses because it is a primary transcription factor that can be activated by various stimuli without need for new protein synthesis [12]. In the present study, although RFP acting on NF-κB directly or indirectly still needs to be confirmed, we found that RFP could activate NF-κB, which was observed in vitro and in vivo. NF-κB triggered the expression of COX-2 to accelerate the secretion of PGE₂, resulting in reduced water transport from intestine to blood vessel, leading to a laxative effect of RFP.

Pharbitis Semen was wildly prescribed for intractable constipation in many countries [9]. In this study, we found the presence of the inflammation response after the RFP-treated rats. Therefore, the dosage of administration and repeated administration of this purgative medicine should be noted.

5. Conclusions

In conclusion, the present study demonstrated that RFP induced its laxative effect by decreasing AQP3 expression, which was involved in water reabsorption from the intestinal tract to the vascular side in the colon. The RFP-induced decrease in the levels of AQP3 was confirmed in vivo and in vitro. Furthermore, we found that the decreased expression of AQP3 was caused by NF-κB activation. NF-κB triggered the expression of COX-2 to accelerate the secretion of PGE₂, resulting in decrease in the expression of AQP3. These findings may explain the underlying diarrhea mechanisms of RFP and provide a sound basis for developing new therapy for constipation.

Data Availability

The datasets analyzed during the current study are available from the corresponding author on reasonable request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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References

Evidence-Based Complementary and Alternative Medicine


