Oxyresveratrol protective effects against deoxynivalenol-induced intestinal barrier dysfunction and bacterial translocation on porcine intestinal epithelial IPEC-J2 cells

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Abstract

Deoxynivalenol (DON) is a major mycotoxin contaminant and is known to impair intestinal barrier function. Previous experiments in our laboratory have proven that polyphenols such as resveratrol (RES) may be effective in enhancing epithelial barrier function. Due to the structural similarity of oxyresveratrol (OXY) with RES, it was hypothesized that OXY could also protect against DON-induced intestinal damage. Accordingly, this study aimed to explore potential protective effects of OXY against DON-induced epithelial barrier dysfunction and bacterial translocation on IPEC-J2 cells, in comparison to resveratrol (RES). The results showed that OXY increased transepithelial electrical resistance (TEER) and reduced FD-4 diffusion, whereas DON reduced TEER and increased FD-4 diffusion in IPEC-J2 cells. On the other hand, OXY reduced FD-4 diffusion in DON-damaged cells but showed no significant difference in terms of TEER. Such protective effects coincided with the significantly reduced \textit{E. coli} translocation in cells co-exposed to DON and OXY. Further mechanistic studies demonstrated that OXY protected against DON-induced barrier dysfunction by enhancing the expression of claudin-4 via mitogen-activated protein kinase (MAPK)-dependent pathways. Apparently, OXY worked through the same way as RES did, with results dovetailed nicely with anticipation. These results imply that OXY may share similar health benefits with RES by enhancing epithelial barrier functions and protecting against DON-induced intestinal damage.

Keywords: Bacteria translocation; Barrier function; Deoxynivalenol; Oxyresveratrol; Resveratrol; Tight junction.

1. Introduction

Mycotoxins are highly toxic low-molecular-weight secondary metabolites of some fungi. Mycotoxin contamination is a serious worldwide problem, with an estimation of up to 25% of the world’s food crop may be contaminated by mycotoxins (FAO, 2013). Mycotoxins can cause various acute and chronic effects on humans and animals, depending on the species and susceptibility of an animal within a species. In human, the toxicological syndromes caused by ingestion of mycotoxins range from death, induction of cancer, growth impairment and decreased resistance to infection (Zain, 2011).

Deoxynivalenol (DON) is a mycotoxin of the trichothecenes family that is mainly produced by \textit{Fusarium graminearum} and \textit{F. culmorum}. DON is commonly found in cereals and grains, particularly in wheat, barley, maize, oats and in their products, in
many temperate regions of Europe, North America and Asia. DON exhibits various toxic effects in humans and animals. Acute high doses of DON exposure elicit abdominal distress, increased salivation, malaise, diarrhea and emesis (vomiting) (Peska et al., 2005). Chronic low doses of DON cause anorexia, reduced weight gain, nutrients malabsorption, neuroendocrine changes and immunologic alterations (Peska et al., 2005). The intestine provides the first physical barrier to protect against ingested toxins and pathogens and is also the first line of defense against intestinal infection. The gut barrier is formed to a large extent by tight junctions that seal the paracellular space between neighboring cells and limit transport of lipids and proteins between the apical and basolateral plasma membranes (Harhaj et al., 2004). Following ingestion of mycotoxin-contaminated food, intestinal epithelial cells (IECs) could be exposed to high concentrations of toxin. Recent reviews by Pinton et al. (2014) and Maresca (2013) have summarized DON induced perturbation of the gut functions in animals and humans (Pinton et al., 2014). Previous studies have also shown that DON can alter intestinal barrier functions in vivo (intestinal epithelial cell line), in vivo (animals) and ex vivo (intestinal explants) (Diesing et al., 2012; Pinton et al., 2009; Pinton et al., 2012). A mechanistic study using IPEC-1 cells demonstrated that DON (30 μM) significantly underminded physiological barrier (a drop in TEER) by reducing expression of claudin-4 protein via activation of the p44/42 extracellular signal-regulated kinase (ERK) signalling pathway (Pinton et al., 2010). Such impairment of intestinal integrity could result in subsequent translocation of luminal antigens and bacteria, which could have major implications for human health in terms of sepsis and inflammation (Pinton et al., 2014). Polyphenols are widespread constituents of fruits, vegetables, cereals, dry legumes, chocolate, and beverages, such as tea, coffee, or wine. Many polyphenols can impact the biological system and can prevent diseases associated with oxidative stress, such as cardiovascular diseases, cancers, inflammation, neurodegenerative diseases and diabetes (Sclabbert et al., 2005). For example, resveratrol (3,4′,5-trihydroxy-trans-stilbene) (RES) is a stilbene oxide present abundantly in red wine and grape skin (Walle et al., 2004) and has been extensively studied. It has been known to provide multiple desirable bioactivities such as antioxidant, anti-inflammatory, cancer chemopreventive, and anti-aging (Pirola et al., 2008) and may be useful in treating cardiovascular diseases, cancers, pain, inflammation, tissue injury, and in reducing the risk of neurodegenerative disorders (Li et al., 2012). Studies of RES-mediated effect on intestinal barrier function remain very limited, and only few studies have shown that RES ameliorates small intestinal and colonic inflammation in mice by reducing different pro-inflammatory cytokine expression and secretion (Bereswill et al., 2010; Sanchez-Fidalgo et al., 2010). RES has a number of naturally occurring analogs (Figure 1), which are produced by diverse plant species, such as grape, mulberry and peanut. These derivatives differ mainly in the number and position of hydroxyl groups, leading to variation in solubility, stability, bioavailability and antioxidant activity (He et al., 2013). For instance, oxyresveratrol (OXY) (trans-2,3′,4,5′-tetramethoxystilbene) is a hydroxylated analog of RES and is present mainly in Morus alba bark (Shin et al., 1998). OXY is a multifunctional bioactive compound: it shows neuroprotective (Chao et al., 2008), depigmenting (skin whitening) (Kim et al., 2002), anti-inflammatory (Chen et al., 2013), as well as anti-viral effects (Galindo et al., 2011). OXY is structurally similar to RES, which has an additional hydroxyl group. Indeed, OXY shows comparable physiological properties to RES in various studies (Li et al., 2015; Xu et al., 2014), and it is suggested to be superior to RES because of its higher water-solubility and lower toxicity (Andrabi et al., 2004). Thus, OXY is a potential substance to be utilized to produce medicines, cosmetics and health supplements. Although OXY has been found to possess a wide range of pharmacological properties and of high value in health promotion, studies on OXY-mediated beneficial effects on gut health remains very limited.

Owing to the fact that DON and OXY are food-derived components, they may be present simultaneously in the gastrointestinal tract. There is a potential interactive effect of DON with OXY, which may result in unexpected intestinal effects in intestine. A previous study has demonstrated that co-exposure of DON and RES protected Caco-2 cells from cytotoxicity (Cano-Sanchez et al., 2015) and it has also previously been shown in our laboratory that RES may be an effective polyphenol enhancing intestinal barrier function. Due to its structural similarity with RES, it was hypothesized that OXY could also protect against DON-induced intestinal damage and bacterial translocation. Accordingly, the aim of this study was to explore the potential protective effects of OXY in comparison with RES against DON-induced intestinal barrier dysfunction and bacterial translocation of the porcine jejuna epithelial cell line, IPEC-J2. The underlying mechanism(s) was/were also elucidated.

2. Materials and methods

2.1. Chemicals and reagents

DON (3α, 7α, 15-trihydroxy-12, 13-epoxytrichothece-9-en-8-one) and RES (3,4′,5-trihydroxy-trans-stilbene) were purchased from Sigma Chemical Company (St. Louis, MO, USA). They were dissolved in dimethyl sulfoxide (DMSO) (Merck, Darmstadt, Germany), and stored at −20 °C before use. OXY (trans-2,3′,4,5′-tetramethoxystilbene) was extracted and purified from old twig of Morus alba in our group as described previously (Li et al., 2007). Phosphate buffered saline (PBS), sodium bicarbonate (NaHCO3), diethyl pyrocarbonate (DEPC), ammonium persulfate (APS), N,N,N′,N′-tetramethyl-ethane-1,2-diamine (TEMED), acrylamide, 4 kDa Fluorescein isothiocyanate-dextran (FITC-dextran), bovine...
serum albumin (BSA), protease inhibitor cocktail, phenylmethylsulfonyl fluoride (PMSF), radio-immunoprecipitation assay (RIPA) lysis buffer and Luria-Bertani (LB) agar were purchased from Sigma. Dulbecco’s modified Eagle medium (DMEM)–Ham’s F-12 (1:1) w/HEPES (DMEM/F-12), fetal bovine serum (FBS), penicillin/streptomycin, amphotericin B, 0.25% (wt/vol) trypsin and Tween 20 were provided by Gibco-Life Technology (Egggenstein, Germany). A cell counting kit-8 (CCK-8) colorimetric assay was purchased from Dojindo (Tokyo, Japan). 2X Laemmli sample buffer, Triton X-100 and 2-amino-2-(hydroxymethyl)-1,3-propanediol (Tris) were provided by BioRad (Richmond, CA, USA). The final concentration of DMSO used as solvent in the culture medium was 0.5%.

2.2. Cell line and culture conditions

IPEC-J2 is a nontransformed intestinal cell line originally derived from the jejunums of neonatal, unsuckled piglets (Schierack et al., 2006) and was a kind gift of Per Torp Sangild (Department of HumanNutrition/Clinical Nutrition, Faculty of Life Science, University of Copenhagen, Copenhagen, Denmark). Briefly, the cells between passages 90 and 100 were maintained in DMEM/F-12, supplemented with NaHCO3, 10% FBS, penicillin (100 UI/mL), streptomycin (100 µg/mL), and amphotericin B (0.5 µg/mL). The cells, incubated at 37 °C in humidified 5% CO2, were maintained with the media changed three times per week; and were split every two weeks using 0.25% (w/v) trypsin after washes with PBS. All cells were screened for mycoplasma contamination prior to use (Lonza, Basel, Switzerland).

2.3. In vitro bacterial translocation assay

To measure the bacterial translocation across the intestinal epithelial monolayer, a non-pathogenic clinical strain, Escherichia coli ATCC 25922 (kindly provided by W. C. Yam, Department of Microbiology, The University of Hong Kong), was used. The same strain was also used previously by other investigators for bacterial translocation study (Pinto Jr et al., 2007). Indeed, E. coli is identified to be the most prevalent species in bacterial translocation cases in humans (MacFie, 1997; Sedman et al., 1994). A non-pathogenic strain was used to ensure that any penetration across the cell monolayer depended mainly on the barrier integritystatus quo, but not due to impairment induced by bacterial endotoxins (Guo et al., 2013). The IPEC-J2 cells were seeded into a permeable Transwell membrane insert (12-mm diameter, 3.0 µm pore size, polyester, Corning, New York, NY, USA) and were grown and differentiated as described previously. After 11 days, cells were washed with PBS and incubated with media without serum and antibiotics for 12 h. The cells were then treated with OXY/RES, administrated into the apical side of the Transwell inserts for 12 h. For assay involving DON-induced damage, the procedures were the same except that 4 µM of DON was added 1 h post OXY/RES administration. After 12 h, E. coli ATCC 25922 was inoculated at 105 colony forming unit/well into the apical side of the cell monolayer. The Transwell plate was then incubated, with mild shaking (50 rpm), at 37 °C in 5% CO2. After 2 h, 100 µL media in the Transwell basolateral compartment were collected and plated on LB agar. The agar plates were incubated at 37 °C for 18 h to determine the number of E. coli present.

2.4. Measurement of trans-epithelial electrical resistance (TEER)

The IPEC-J2 cells, at a density of 5 × 104 cells/well, were seeded into a permeable Transwell membrane insert (12-mm diameter, 0.4 µm pore size, polyester, Corning). The cells were grown and differentiated as described previously. After 11 days, the cells were treated in serum-free media for 12 h, then OXY/RES (25 µM) was added into the apical Transwell compartment. At various time points, TEER was measured using an ohm-meter (model EVOM, WPI Inc., Sarasota, FL, USA). Resistance values were recorded in Ω and calculated in Ω·cm2 by multiplying the area of the membrane filter. Results were expressed as percentage of the control: % TEER = (TEERtreatment/TEERcontrol) × 100%. For DON-induced damage, the procedures were the same except DON (4 µM) was added 1 h post RES administration.

2.5. Paracellular tracer flux assay

IPEC-J2 were grown and differentiated in 0.4 µm pore inserts and treated with RES for various time points as described previously. FITC-dextran (4 kDa) was dissolved in cell culture medium and used at a final concentration of 0.5 mg/mL in the apical cell compartment. After 24 h of incubation the amount of fluorescence was measured in the basal compartment with a fluorimeter (Model VICTOR X3, Perkin Elmer, Courtaboeuf, France), with excitation and emission wavelengths of 490 and 520 nm, respectively. Levels of fluorescence were recorded in relative fluorescence units (RFU). Results were expressed as percentage of the control value: % cumulative flux of 4 kDa FITC (FD-4) = (RFUtreatment/RFUcontrol) × 100%. For DON-induced damage, the procedures were the same except DON was added 1 h post OXY/RES administration.

2.6. Protein extraction, SDS-PAGE and immunoblotting

The cells, undergoing the same treatment as described for TEER measurement, were washed with PBS and extracted for detergent-insoluble protein fractions as reported previously (Noda et al., 2012). The cells were first treated with 200 µL gentle lysis buffer [1% Triton X-100 and 5 mM EGTA in 50 mM Tris solution at pH 7.4], supplemented with protease inhibitor cocktail and PMSF, to yield cell lysates. The cell lysates were then centrifuged at 16,000 g for 10 min at 4 °C to sediment a high-density insoluble fraction that is rich in actin-associated proteins. The supernatant was discarded and the pellet was re-suspended and extracted for proteins using RIPA lysis buffer with protease inhibitor cocktail and PMSF added. Protein concentration was determined by the Bradford protein assay (BioRad, Hercules, CA, USA).

Proteins (10 µg) were loaded onto 12% sodium dodecyl sulfate polyacrylamide gel (BioRad), separated by electrophoresis (SDS-PAGE), and then blotted onto a polyvinylidene difluoride (PVDF) membrane (Millipore, Darmstadt, Germany). The membrane was blocked with 5% BSA in PBS containing 0.05% (v/v) Tween 20 (PBST) buffer. Proteins were probed by immunoblotting with diluted primary antibodies (1:1000, v/v) for claudin-4 (#32-9400, Gibco-Invitrogen, Grand Island, NY, USA), and GAPDH (#sc-166646, Santa Cruz Biotechnology, Santa Cruz, CA, USA), in combination with horseradish peroxidase (HRP)-conjugated antimouse IgG (#170-6516, BioRad) secondary antibodies. The blots were developed using Clarity Western ECL blotting kit (BioRad) and chemiluminescence was detected with a digital imaging system (Chemidoc XRS+ system with image lab software, BioRad). Quantification was performed by ImageJ software (Ver. 1.48, National Institutes of Health, Bethesda, MD, USA) by densitometric

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2.7. Kinase phosphorylation assay

To study the effects of OXY/RES on DON-induced phosphorylation of mitogen-activated protein kinases (MAPKs), quantification of total phosphorylated p38 MAPK, extracellular signal-regulated kinase (ERK), and c-Jun N-terminal kinase (JNK) was performed using a commercial enzyme-linked immunosorbent assay (ELISA) kit (ab119674, PhosphoTracer, Abcam) according to the manufacturer’s instructions. Cells were seeded in 6-well culture plates (Corning) at 2 × 10⁵ cells/well, and allowed to grow and differentiate for 11 days. After 11 days, the cells were treated with serum-free medium for 12 h. After 12 h, cells were washed and treated firstly with or without 25 µM OXY/RES for 1 h, and then with 1 or 4 µM DON for 30 min. To determine the optimal time point for the kinase phosphorylation assay, time effects of DON-induced MAPKs phosphorylation were characterized previously (data not shown). Whole cell lysates were collected, normalized for total protein and analyzed for the levels of phosphorylated p38 MAPK, ERK and JNK proteins. Treatments with OXY/RES alone or DMEM were used as controls. All samples were run in triplicate.

2.8. Statistical analysis

Data were analyzed using Minitab statistical software (ver. 16, Minitab Inc., State College, PA, USA). Values represent mean ± standard deviation (S.D.) of at least three independent experiments. Statistical analyses were performed by Student’s t-test, Chi-squared test, or 1-way ANOVA followed by Tukey’s multiple comparison test. P values less than 0.05 were considered significant.

3. Results

3.1. Choice of concentrations of DON, RES and OXY

The concentrations of DON and RES were chosen based on previous experiments using 4 µM DON and 25 µM RES as the non-cytotoxic doses (Ling et al., 2016). OXY was also assessed for cytotoxicity analysis (Schneider et al., 2012).

3.2. OXY fortified intestinal barrier and protected against DON induced damage

To determine if OXY protected a cell monolayer from DON-induced intestinal barrier damage, the protective effect of OXY on TEER and paracellular permeability was first assessed and compared with that of RES. A 25 µM RES/OXY was administrated to the cells 1 h prior to challenge with 4 µM of DON, TEER and 4-kDa FITC-dextran (FD-4) diffusion were measured after 12 h. The results showed that both RES and OXY increased TEER and reduced FD-4 diffusion, whereas DON reduced TEER and increased FD-4 diffusion in IPEC-J2 cells. On the other hand, both RES and OXY reduced FD-4 diffusion in DON-damaged cells (Figure 3) but showed no significant difference in terms of TEER (Figure 4).

3.3. OXY reduced DON-induced bacterial translocation

The effect of OXY on DON-induced bacterial translocation was subsequently studied. After 12 h, E. coli was added. Medium (100 µL) in the basolateral compartment was collected after 2 h, and plated to determine if bacteria were present. The presence of E. coli in the basolateral compartment was deemed positive for bacterial translocation. The 2-h incubation time was determined previously in our laboratory, which demonstrated that it took at least 2 h for the E. coli to penetrate about 25% untreated cell monolayers (data not shown). In the present study, the results showed that similar to RES, OXY significantly reduced (by 36%) DON-induced bacterial translocation (Figure 5).

3.4. OXY enhanced claudin-4 expression and prevented DON-induced decreased expression of claudin-4

Immunoblot analyses revealed that both OXY and RES enhanced the protein expression of claudin-4 in untreated IPEC-J2 cells (Figures 6A and 6B); and reduced DON-induced decreased expression of claudin-4 (Figures 7A and 7B). Apparently, OXY worked through the
same pathway as RES did, with results nicely fit with anticipation.

3.5. OXY protected against DON-induced barrier dysfunction via MAPK-dependent mechanisms

We recently demonstrated that RES protected against DON-induced decreased claudin-4 expression through mitogen-activated protein kinase (MAPK)-dependent pathways. In the present study, we further investigated if OXY also protected against DON-induced decreased claudin-4 expression via MAPK pathways. Cell monolayers were pre-treated with 25 µM OXY for 1 h, then co-treated with DON for 30 min. Proteins were then immediately extracted for ELISA analysis. The results showed that OXY significantly reduced DON-activated phosphorylation of p38 (for 1 µM DON) and JNK (for both 1 and 4 µM DON) (Figures 8A & 8B), but not ERK (Figure 8C). Again, OXY showed inhibition pattern alike to that of RES found previously in our laboratory (data not yet published), in which p-JNK was the one most reduced.

4. Discussion

Polyphenols constitute a large chemical class of phytochemicals among which the stilbenoid sub-class has received growing inter-
The protective effects of oxyresveratrol due to its elaborate structural diversity and biological activities. The members of stibenoid family differ from each other mainly in the number and position of hydroxyl groups, leading to variation in solubility, stability, bioavailability and antioxidant activity (He et al., 2013; Wang et al., 1999). Human tissues are only exposed to dietary polyphenols at very low level via the blood, except for gut cells, which are constantly and directly exposed to a relatively high dose of dietary polyphenols (Kroon et al., 2004). Therefore, it is anticipated that dietary polyphenols can exert an immense effect, locally and systemically, upon the gut. However, the mechanisms of their protective effects on intestinal barrier functions are not fully understood. Previous work in our laboratory has found that RES might be effective in enhancing the intestinal barrier function and might be used as a novel mean to counteract the intestinal dysfunction and bacterial translocation following mycotoxin exposure (Ling et al., 2016). The results in the present study demonstrate that OXY, a less investigated stilbene, is a highly interesting candidate for the use of promoting intestinal health, which shows similar bioactivity but higher water solubility than RES. We evaluated its effectiveness in enhancing the intestinal barrier and protecting against intestinal damage induced by DON, one of the most commonly occurring mycotoxins in foods and feeds.

In our model system, the evaluation of possible cytotoxic effects of RES and OXY was performed. It was interesting to note that OXY showed comparable cytotoxic effects as RES, which was contradictory to few previous studies which showed lower cytotoxicity of OXY compared to RES in different model systems (Chao et al., 2008; Galindo et al., 2011). Such discrepancy may be due to differences in the concentrations and the cell types used.

Although not yet fully understood, intestinal dysfunction and the increase in bacterial translocation following DON treatment can exert adverse health effects in human and animals by increasing the susceptibility to infection as well as inflammation and sepsis (Schneider et al., 2012). In this study, we have revealed that OXY elicited potent protective effects against DON-induced intestinal barrier function and bacterial translocation in the IPEC-J2 cells. OXY displayed a comparable effect on enhancing intestinal physical barrier function as RES, as shown by increasing TEER and reducing FD-4 diffusion, in both untreated and DON-treated cells, by almost the same extent. Similarly, pre-treatment of the cell monolayers with OXY, was also found to reduce DON-induced bacterial translocation. All of these suggest a possible protective role of OXY to mitigate barrier dysfunction caused by DON.

The gut barrier is formed to a large extent by tight junctions (TJ), which are multi-protein complexes that link adjacent epithelial cells near their apical border. The presence of TJ that can seal the paracellular space between neighboring cells accounts for the main physical barrier to bacterial translocation. Previous studies have shown the correlation between decreased barrier function and reduced expression of the tight junction protein, particularly claudin-4 in DON-treated intestinal cells (Pinton et al., 2009; Pinton et al., 2010). Such a decrease in claudin-4 protein expression was found to be associated with an increase in permeability of porcine intestinal epithelial cells to pathogenic E. coli (Pinton et al., 2009).

In this sense, claudin-4 apparently is a key protein in the preservation of barrier function by establishing a paracellular seal in the intestinal cells (Pinton et al., 2010). As previously observed, some polyphenols increased TJ protein

Figure 8. Levels of (A) p38 phosphorylation, (B) JNK phosphorylation and (C) ERK phosphorylation in IPEC-J2 cells treated with or without OXY in the presence of DON. Proteins were extracted 30 min after DON administration. Data are mean ± S.D. for n = 3 in each group. Asterisk indicates a significant pairwise difference between treatments, P < 0.05.
gene expression in the Caco-2 and T84 cells (Amasheh et al., 2008; Rogoll et al., 2010). Similarly, our laboratory has recently demonstrated that RES fortified intestinal barrier and protected against DON induced damage via modulation of claudin-4 expression. In the present study, the same protocols were followed through for OXY to characterize its effects on intestinal barrier function and to compare with RES. OXY acted similarly as RES which was shown to enhance the claudin-4 expression and protected against DON-induced decreased claudin-4 expression. Regulation of TJ structure and function involves the MAPK signaling molecules (Matter et al., 2003). MAPK cascades participate in an array of cellular responses, it was shown that p38 and JNK cascades function in stress responses, including inflammation and apoptosis (Schaeffer et al., 1999). The exact cellular responses are cell-type or stimuli specific (Oshima et al., 2008). Previously, DON was shown to activate MAPKs in the Caco-2 and IPEC-1 cells (Pinton et al., 2010; Sergent et al., 2006). It was suggested that DON impaired intestinal barrier function by decreasing the expression of TJ protein such as claudin-4 through a MAPK-dependent mechanism (Pinton et al., 2009). Polyphenols such as OXY, on the other hand, can modulate a series of signaling cascades, including MAPKs, by directly binding to their ATP binding sites, a property determined by the number and the location of hydroxyl groups on the B ring. OXY can therefore penetrate into the cytoplasm and may have a direct modulation effect on MAPK molecules (Chao et al., 2008). Our results clearly demonstrated that OXY significantly reduced DON-induced phosphorylation for p38 and JNK. Therefore, it indicated that OXY was a prominent inhibitor to DON-induced MAPK activation, which may protect against DON-induced barrier dysfunction by preventing decreased expression of claudin-4. Nevertheless, further research is needed to disen- tangle the underlying mechanisms.

Owing to the fact that RES and OXY are relatively non-polar, they are lipophilic enough to permeate the GI tract and absorb readily into the blood (Huang et al., 2008). This suggests that they may exert their effects in the intestine very quickly after oral administration. The solubility of RES is, rather limited in aqueous solutions, which provides difficulties for drug formulation. However, OXY is better water soluble, which may contribute to its ability to exert a protective effect over a wider range of concentrations than RES (Lorenz et al., 2003). In addition, OXY is a natural analogue of RES and can easily be obtained from natural sources. All these properties make OXY a promising candidate for pharmaceutical formulations.

Taken together, due to the structural similarity of OXY and RES, they may share similar health benefits by enhancing epithelial barrier functions and protecting against DON-induced intestinal damage. Chronic exposure of DON could result in intestinal damage, which may have severe health consequences in animals and humans in terms of sepsis and inflammation. Polyphenols, such as RES and OXY, may provide a novel intervention strategy to restrict the consequences of DON ingestion. Based on the results presented, it will be of great interest to compare RES and OXY in an in vivo model for protection against DON-induced intestinal damage. Given that OXY represents only one member out of an enormous pool of RES derivatives, it is anticipated that other members may confer similar, or even more potent, effects to intestinal barrier function and may be used as a novel dietary intervention strategy to reduce DON toxicity in humans/animals.

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