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<tr>
<td>Citation</td>
<td>Japanese Journal of Veterinary Research, 64(3): 171-182</td>
</tr>
<tr>
<td>Issue Date</td>
<td>2016-08</td>
</tr>
<tr>
<td>DOI</td>
<td>10.14943/jjvr.64.3.171</td>
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<tr>
<td>Doc URL</td>
<td><a href="http://hdl.handle.net/2115/62765">http://hdl.handle.net/2115/62765</a></td>
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<td>Type</td>
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<td>File Information</td>
<td>64-3 003.p171-182 FP FARAG.pdf</td>
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Alleviative effect of some phytochemicals on cyadox-induced oxidative damage in rabbit erythrocytes

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Received for publication, February 22, 2016; accepted, May 23, 2016

Abstract
This study was carried out to evaluate the effects of different concentrations of cyadox (CYA), cinnamaldehyde (CIN) and resveratrol (RES) on rabbit isolated erythrocytes in two experiments. Experiment 1 evaluated the effects of different concentrations (2.5, 5, 10, 20, 40, 60, 80 and 100) μg/ml of CYA and μM of both CIN and RES separately on isolated erythrocytes in order to make the concentration response curve. Results indicated that CYA caused a significant depletion of reduced glutathione (GSH) content compared to control at 40 μg/ml and the depletion increased by increasing the concentration. On the other hand, both RES and CIN showed a highly significant elevation of GSH content at 40 μM being more effective by increasing concentrations. Experiment 2 investigated the potential benefits of using CIN and RES separately or in combination on CYA induced alterations in isolated rabbit erythrocytes (the used concentrations were based on the results of experiment 1). Results revealed that exposure to CYA caused a significant decrease in superoxide dismutase (SOD) and catalase (CAT) activities and reduced glutathione (GSH) and total protein (TP) contents, CYA also elevated extracellular hemoglobin (Hb) and adenosine triphosphate (ATP), increased the malonaldehyde (MDA) and protein carbonyl (PrC) contents with increasing caspase3 and caspase8 activities suggesting CYA pro-oxidant effect. Both CIN and RES were able to inverse these hazardous effects of CYA. However, CIN was more effective than RES, and their combination showed a positive synergistic effect in protecting cells against oxidative injury caused by CYA.

Key Words: Cyadox, phytochemicals, oxidative stress, response curve, erythrocytes.

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doi: 10.14943/jjvr.64.3.171
Introduction

Quinoxaline 1,4-dioxides (QdNOs) derivatives are one of the most important synthetic antimicrobial agents used worldwide as feed additives to promote growth and feeding behaviors of different animal species at sub therapeutic levels since the use of high doses for long periods usually causes hazardous effects for both animals and human (Carta et al., 2005). Olaquindox (OLA), carbadox (CBX), mequindox (MEQ), quinocetone (QCT), and cyadox (CYA) are the known members of QdNOs class. QdNOs derivatives were reported to have in vitro mutagenic potential, induce DNA damage and increase the percent of micro-nucleated cell culture (Jin et al., 2009), carcinogenic, developmental and reproductive toxicities and adrenal toxicity in male rats due to oxidative injury (Huang et al., 2009). Compared with other QdNOs derivatives, CYA has been found to be less toxic in mutagenicity tests (Huang et al., 2008). However, CYA showed different signs of toxicity in piglets (Nabuurs et al., 1990) and had a teratogenic potential and reproductive toxicity in rats (Wang et al., 2011). CYA has caused sub-chronic toxic effects in dogs (Wang et al., 2015). Some primary metabolites of CYA showed significant decrease in body weight and change in clinical serum biochemistry in addition to pathological changes in liver of rats (Huang et al., 2016). These undesirable effects could be attributed to the generation of reactive oxygen species (ROS) as a result of QdNOs reduction during their metabolism (Liu et al., 2009).

In an attempt to reduce the oxidative stress induced by different types of chemical feed additives, herbal plants and their extracts are used to provide animals with antioxidant polyphenols to improve their health. Among the most abundant beneficial compounds that are used as feed additives for animal and poultry to reduce free radicals are resveratrol (3, 5, 4’-trihydroxy-trans-stilbene; RES), a stilbenes type aromatic phytoalexin which was predominantly found in grapes, peanuts, berries, Yucca schidigera and turmeric (Alagawany et al., 2015) and cinnamaldehyde (3-phenyl-2-propenal; CIN) which presents in cinnamon (Faix et al., 2009). Numerous in vitro and in vivo studies described different biological effects of resveratrol, including antioxidant, cardio-protective, anti-aging, anticancer, anti-inflammatory, immunomodulatory, antioxidants and metabolic modifying activities (Hao et al., 2011).

Similarly, CIN has been demonstrated to have an antimicrobial, anti-platelet aggregating, vasodilatory, anticancer and immunomodulatory effects and it could inhibit the induction of nitric oxide synthase and nitric oxide in a dose-dependent manner (Vangalapati et al., 2012).

The present study aims at exploring the benefits of using CIN, RES and their combination on CYA induced alterations in isolated rabbit erythrocytes where rabbits are one of the most important animals exposed to treatment by CYA as antimicrobial agent and the oxidative damage of could result in some pathological conditions like anaemia and cardiovascular diseases especially with using CYA for long periods. Erythrocytes were used as it is very sensitive to the peroxidation process owing to its high oxygen tension, membrane concentration of polyunsaturated fatty acids and redox active Hb molecules as recently reported by Ahmed and Beg (2013). Moreover, GSH was used for studying the concentration response curve as it is known to provide the primary antioxidant defense for the stored erythrocytes, protect membrane lipids and proteins and keep the stability of the membrane skeleton and survival of erythrocytes against oxidative damage.

Materials and methods

Chemicals: Cyadox (CYA, C₁₂H₉N₅O₃, molecular weight 271.23 gmol, CAS No: 65884-46-0, purity 98%) was obtained from Hangzhou Uniwise International Co., Ltd. (Zhejiang, China (Mainland). Resveratrol (3, 5, 4’-trihydroxy-trans-stilbene,
purity of 99%) and pure cinnamaldehyde (3-phenyl-2-propenal, purity \( \geq 98\% \)) were purchased from Oxford Laboratory Mumbai, India. All other chemicals were purchased from Sigma (St. Louis, MO, USA). All other reagents used were of analytical grade. Caspase3 and caspase8 assay kits were purchased from Biovision Inc. (MountainView, CA, USA). LDH Cytotoxicity Detection Kit (plus) (LDH) (Roche Applied Science, Mannheim, Germany).

Animals and care: Male New Zealand White rabbits obtained from rabbit farm of the faculty of agriculture of Zagazig University (aged 3 months with initial body weight of 2.00 ± 0.05 kg) were used.

**DPPH** free Radical-Scavenging Activity of CIN and RES: The electron donation ability of cinnamaldehyde and resveratrol was measured by bleaching of the purple colored solution of 2, 2-diphenyl-1-picrylhydrazyl (DPPH) according to the method of Hanato et al. (1988).

Preparation of erythrocytes: Five ml of blood was collected aseptically from the ear vein of all animals in a graduated test-tube containing heparin to avoid coagulation. Blood was centrifuged at 3000 rpm for 10 min at 4°C in graduated tubes and the plasma and buffy coat were carefully removed. The erythrocytes were harvested by centrifugation after washing once with 0.9% NaCl solution and two times with ice-cold phosphate buffered saline (PBS: 145 mMNaCl, 1.9 mM NaH₂PO₄, 8.1 mM Na₂HPO₄) and finally suspended in PBS to obtain cell suspensions at 10 % hematocrit to be used for incubations (Yang et al., 2006).

**Treatment of erythrocytes:**

**Experiment 1:** Concentration response curve: CYA, CIN and RES were primarily solubilized in a small amount of DMSO (not exceed 0.1%) and further diluted in PBS to get different concentrations (2.5, 5, 10, 20, 40, 60, 80 and 100) \( \mu \)g /ml of CYA and \( \mu \)M of both CIN and RES, where each concentration was incubated with erythrocyte suspension individually in groups of 5 samples at 37°C for 3 h and compared with control group that was treated with PBS with DMSO (0.1% v/v) in order to make the concentration response curve.

**Experiment 2:** This experiment was carried out based on the data obtained from Experiment 1. Seven groups (\( n = 5 \)) of erythrocytes suspension were used. In particular, the 1\(^{st} \) group kept as control treated with PBS with DMSO, 2\(^{nd} \) group was subjected to CYA (40 \( \mu \)g /ml), 3\(^{rd} \) group was incubated with CIN (40 \( \mu \)M), 4\(^{th} \) group were subjected to RES (40 \( \mu \)M) while the 5\(^{th} \) group was co exposed to CYA (40 \( \mu \)g /ml) and CIN (40 \( \mu \)M). The 6\(^{th} \) group was co exposed to CYA (40 \( \mu \)g /ml) and RES (40 \( \mu \)M). The 7\(^{th} \) group was exposed to CYA in combination with both CIN and RES at the same indicated concentrations. The reaction mixtures of different groups were incubated at 37°C for 3 h with gentle shaking every 15 min.

Assessment of antioxidant activities of erythrocytes: Antioxidant indices: For measuring the antioxidant parameters, after incubation the mixtures were stored at \(-20°C\) and thawed one day later for RBCs lysis by osmotic pressure, then they were centrifuged and supernatants obtained and used for measuring the activities of superoxide dismutase (SOD) according to Misra and Fridovich, (1972), catalase (CAT) according to Aebi (1984), total reduced glutathione (GSH) according to Ellman, (1959).

Lipid peroxidation: Lipid peroxidation was assessed by determining erythrocyte MDA levels using the thiobarbituric acid method (Bartosz, 2004) and the reaction product was measured spectrophotometrically at 535 nm.

Protein oxidation: Protein carbonyl content was measured as an index of protein oxidation as described by Uchida and Stadtman, (1993).
Hemoglobin (Hb) and protein determination: The hemolysis process of erythrocytes was monitored by hemoglobin (Hb) release. After incubation time periods, 50 ml of cell suspension was added to 1 ml of PBS and centrifuged (1000 g, 7 min). The Hb content of supernatants was measured by absorbance at 540 nm using the Varian Cary 50 UV-vis spectrophotometer. While total protein (TP) concentrations in the hemolysates was determined spectrophotometrically by standard kits based on the colorimetric biuret method.

Measurement of erythrocyte release of ATP: The procedure for measurement of the ATP content in the reaction mixture was based on the reactions described by Adams, (1963).

Membrane toxicity assay: The membrane toxicity can be rated by quantifying the liberation of the intracellular enzyme lactate dehydrogenase (LDH) into the supernatant using cytotoxicity detection Kit (plus) (LDH) according to the manufacturer’s instruction.

Determination of caspase activities: The proteolytic activities of caspase3 and caspase8 were evaluated in erythrocyte lysates using the fluoro-metric substrates DEVD-AFC (caspase-3 substrate) and IETD-AFC (caspase-8 substrate), following the protocols of the caspase activity assay kits.

Statistical analysis: The experiment was carried out as a completely randomized design. Data were statistically analyzed using GLM procedure SAS (SAS Institute Inc., 2001). Orthogonal polynomial contrasts were used to test the linear and quadratic effects of the increasing levels of supplements to isolated rabbit erythrocytes. The differences among means were determined using the post-hoc Tukey’s test. Statements of statistical significance are based on P < 0.05, unless otherwise stated.

Results

DPPH· Radical-Scavenging Activity of CIN and RES

The results of DPPH· radical-scavenging activities of CIN and RES are represented in (Fig. 1). The results indicated that CIN and RES exhibited antioxidant activity. The CIN showed relatively high antioxidant activity (93.00%) than RES (90.12%) compared to the radical scavenging activity of TBHQ (98.65%) after 120 min.

Results of experiment 1: Concentration response curve

The results indicated that the first concentration of CYA caused a significant depletion of GSH content compared to control (40 μg/ml) and the degree of depletion increased by increasing concentration. Thus, from the toxicological point of view 40 μg/ml was used in the second experiment to study the toxic effect of CYA. On the other hand, both RES and CIN began to show a highly significant elevation of GSH level at 40 μM being more effective by increasing the concentrations, but from the economical point of view we choose this concentration to study their modulatory effects on CYA exposed erythrocytes (Fig. 2).

Results of experiment 2

The effect of CYA on antioxidant indices

Results in Table 1 showed that CYA at a dose of (40 μg/ml) inhibited the activities of SOD and CAT and decreased GSH concentration linearly and quadratically (P < 0.001) compared to control and other experimental groups. Incubation of erythrocytes with CIN or RES (40 μM) separately significantly (P < 0.001) increased the activities of SOD and CAT as well as the concentration of GSH to levels better than control, where CIN showed the height values followed by RES then control. In addition, CIN+CYA resulted in a linear and quadratic (P < 0.001) improvement in the antioxidant capacity of erythrocytes than CYA+RES in which GSH concentration did not
statistically differ than CYA group. Meanwhile, CYA+CIN+RES showed results comparable to control.

**The effect of CYA on lipid and protein oxidation**

Lipid peroxidation measured by the MDA concentration and protein oxidation represented by PrC content were significantly (\(P < 0.001\)) increased in erythrocytes incubated with CYA compared with control and the other groups (Table 1). Incubation of erythrocytes with CIN or RES (40 \(\mu\)M) individually resulted in a linear and quadratic (\(P < 0.001\)) decrease in the MDA and PrC contents to being lower than control values, where CIN resulted in a much reduction in concentration than RES. CIN+CYA resulted in a linear and quadratic (\(P < 0.001\)) decrease in the MDA and PrC contents than CYA+RES. Meanwhile, CYA+CIN+RES succeeded to minimize the MDA and PrC contents to control values.
The effect of CYA on hemoglobin content

The effect of CYA, RES, CIN and their combinations on Hb release are summarized in Table 2. Incubation of erythrocytes with CYA (40 μg/ml) showed linear and quadratic significant (P < 0.001) increase in Hb release in the reaction mixture compared to control. On the other hand, Hb was linearly and quadratically (P < 0.001) decreased by incubation with CIN or RES. CIN + CYA linearly and quadratically (P < 0.001) decreased the Hb content than CYA + RES. Both CIN and RES showed a synergistic effect against CYA, where treatment of erythrocytes with CYA + CIN + RES reduced the Hb content compared to control.

The effect of CYA on protein content:

Incubation of erythrocytes with CYA alters...
the TP content of isolated erythrocytes \( (P < 0.001) \). On the other hand, TP content was both linearly and quadratically \( (P < 0.001) \) increased by incubation with CIN or RES, where the CIN achieved the highest TP content compared to the other groups. CIN+CYA linearly and quadratically \( (P < 0.001) \) increased the TP content than CYA+RES. CIN and RES exhibited a good synergistic effect by elevating the reduced TP content induced by CYA by increasing its level to be similar to that of control (Table 2).

The effect of CYA on ATP release

Incubation of erythrocytes with CYA \( (40 \mu g/ml) \) for 3 h produced linear \( (P = 0.001) \) and quadratic \( (P < 0.001) \) increase in the ATP content to be compared to control (Table 2). Incubation of erythrocytes with CIN or RES \( (40 \mu M) \) in absence of CYA resulted in a linear \( (P = 0.001) \) and quadratic \( (P < 0.001) \) decrease in the amount of released ATP to levels lower than control values however RES showed lower inhibition than CIN. CIN+CYA resulted in a linear and quadratic \( (P < 0.001) \) decrease in the ATP content than CYA+RES. Treatment of erythrocytes with CYA in combination with both CIN and RES showed a linear \( (P = 0.001) \) and quadratic \( (P < 0.001) \) decrease in ATP content better than CYA+CIN or CYA+RES.

The effect of CYA on LDH release

Erythrocytes incubated with CYA for 3 h exhibited marked significant elevation of LDH release \( (P < 0.005) \) compared to the experimental groups. Contrarily, CIN and RES significantly decreased LDH release from erythrocytes, where the lowest measured LDH \( (120 \text{ and } 130 \text{ IU}) \) was obtained after incubation with CIN and RES respectively compared to control and the other groups. CIN+CYA significantly reduced the release of LDH than CYA+RES and both were higher than control. The synergistic effect of CIN and RES against CYA was also observed where treatment of erythrocytes with CYA+CIN+RES reduced the LDH release to the level control (Fig. 3).

The effect of CYA on caspase activity

Exposure of erythrocytes to CYA led to the activation of caspase3 and caspase8. Incubation of erythrocytes with CIN or RES individually or in combination with CYA significantly decreased caspases activities. However, CIN showed advantage over RES in both treatments. The synergistic effect of CIN+RES on erythrocytes treated with CYA was clear on decreasing of both
natural antioxidant effect on erythrocytes

Discussion

This study was conducted to investigate the possible modulatory effects of CIN and RES as natural phytogenic additives against the hazardous impacts of CYA in isolated rabbit erythrocytes. Under normal physiological conditions, internal antioxidant enzymes such as SOD and CAT act as defense mechanisms against intracellular oxidative stress and provide cell protection by their free radical scavenging activities and elimination of ROS. In the present work, CYA caused a significant decrease in SOD and CAT activities in rabbit erythrocytes. This decrease may be returned to the ability of CYA to produce a short and unstable release of ROS, which was the key mediator of QdNOs-induced cell death and this agrees with the study of Liu et al., (2009) who suggested that QdNOs displayed prooxidant activities on adrenocortical cells.

Furthermore, CYA induced depletion of the GSH level which is required to protect important proteins of RBCs against oxidation by maintaining caspases (Fig. 4 and 5).
SH groups in Hb and enzymes in the reduced state, and binding with MDA and other deleterious endogenous substances (Hao et al., 2011).

Depletion of GSH is accompanied with increased MDA content in CYA exposed group indicating the capability of CYA in producing a state of considerable erythrocytic oxidative injury as MDA is a highly reactive bifunctional molecule, that cross-link erythrocyte phospholipids and proteins to alter the functions of cell membrane leading to decreased erythrocytic survival and has been proposed as a general mechanism for cell injury (i.e induce hemolysis) (Banerjee et al., 2008). This comes on line with the results obtained in the present study concerning the effects of CYA on increasing the released Hb and ATP and confirmed by the increased release of LDH which is considered as indicator of membrane toxicity. Moreover, these results suggest the role of CYA in induction of ROS in erythrocytes membrane and peroxidation of membrane lipids leading to hemolysis and altering the protein and lipid contents to different extents. Additionally, the interaction between MDA and CuZn-SOD of erythrocytes leads to the modification of histidine amino acid residues and the production of protein–protein cross-linked derivatives as a result each type of ROS gives a different protein oxidation pattern (Kwon et al., 2000). This could explain the generation of protein carbonyl derivatives which could be also account for the decreased protein content of the rabbit erythrocytes after incubation with CYA.

Oxidative stress and impairment of antioxidant defense system observed in CYA exposed erythrocytes could be the main cause of activating the caspase8 which is a membrane-bound mediator initiating the cellular cascade for apoptosis and caspase3 that is the effector mediator leading to proteolysis of cellular proteins as reported (Mandal et al., 2012).

Interestingly, incubating erythrocytes with CIN or RES with CYA either separately or in combination resulted in increased concentration of GSH and the activities of SOD and CAT and protein content in the rabbit erythrocytes while decreased the levels of MDA and PrC and the markers of hemolysis (Hb and ATP) and reduced the LDH release and decreased the caspase activities.

The results concerning the effects of RES on erythrocyte antioxidant defense system could be attributed to its DPPH radical scavenging activity observed in the present work and also by Lopez-Velez et al. (2003) who stated that resveratrol could provide cell protection due to antioxidant capacity. RES could act directly as an effective eliminator of free radicals and through balancing of hydroxyl phenolic groups, increasing the activities of antioxidant enzymes including SOD and CAT, glutathione S-transferase and NADPH quinone oxido-reductase, reducing lipid peroxidation or scavenging the generated ROS (Das, 2011). Our results are also in agreement with Mikstacka et al. (2010) where resveratrol prevented the reduction in GSH content and reduced lipid peroxidation in human erythrocytes.

RES has been also suggested to exert its antioxidant protection effects through decreasing the generation of ROS in some in vitro studies like (Vieira de Almeida et al., 2008; Sayin et al. 2011) and in vivo (Liu et al., 2014; Sridhar et al., 2015). RES antioxidant activity is parallel with its DPPH radical scavenging activity observed in the present work which agrees with Ioanna et al. (2015).

In a like manner, CIN could reverse the undesirable effects of CYA by enhancing the activities of SOD and CAT, increasing GSH content and reducing lipid and protein oxidation. These results are in agreement with Subash-Babu et al. (2014) where CIN enhances the activity of antioxidant defense system against ROS produced under hyperglycemic conditions in animal providing protection to pancreatic β-cells. These effects may be returned to the ability of CIN to act as a potential source of antioxidant scavenging free radicals as reported in different in vitro models like DPPH, superoxide, nitric oxide, H₂O₂ scavenging activity and reducing
power (Haripriya et al., 2013). CIN also exhibit strong antioxidant capacity to scavenge free radicals of oxygen and lipids. Additionally, ROS release from lipopolysaccharide (LPS) stimulated J774A.1 macrophages was reduced by CIN (Chao et al., 2008).

Our findings also showed that CIN was more effective than RES as antioxidant. Agents inducing lipid peroxidation cause the release of iron from the biological complexes that enhance peroxidative damage to lipids and consequent hemolysis. It is suggested that the protection against oxidative damage to erythrocyte membrane by CIN may be due to its antioxidant activity, and independently, to intracellular chelation of iron. Intra cellular chelation of iron could be excluded as a mechanism of antioxidant activity of RES where it inhibited Fe (II)-catalyzed lipid peroxidation mainly by scavenging lipid peroxyl radicals within the membrane, and that it cannot change the oxidation state of the metal in a way that would alter the Fe[II]/Fe[III] ratio (Tadolini et al., 2000). This could explain the differences in response to the two phyogenic additives.

In conclusion, the impacts noted in the present study indicated that CYA have hazardous prooxidant effects on body cells. These results may be attributed to the possibility of this type of phytochemicals to generate ROS and a state of oxidative injury. Both, CIN and RES as phytochemical additives could be helpful in reducing the hazardous effects of CYA, keeping the normal function of the body cells and efficiently protecting the cells against oxidative injury suggesting their role in eliminating ROS which are responsible for lipid peroxidation, peroxidative hemolysis and aging of cells. Moreover, it was found that CIN and RES have valuable synergistic effects against the prooxidant activity of CYA.

**Declaration of interest**

The authors report no conflicts of interest.

**References**


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