

## Letter to the editor:

### RECENT STUDIES ON RESVERATROL AND ITS BIOLOGICAL AND PHARMACOLOGICAL ACTIVITY

Yong Joo Kim<sup>1</sup>, Sun Ok Chung<sup>1</sup>, Jae Kwang Kim<sup>2\*</sup>, Sang Un Park<sup>3\*</sup>

<sup>1</sup> Department of Biosystems Machinery Engineering, Chungnam National University, 99 Daehak-ro, Yuseong-gu, Daejeon, 34134, Korea

<sup>2</sup> Division of Life Sciences and Convergence Research Center for Insect Vectors, Incheon National University, Incheon 22012, Korea

<sup>3</sup> Department of Crop Science, Chungnam National University, 99 Daehak-ro, Yuseong-gu, Daejeon, 34134, Korea

\* Corresponding authors

Dr. Jae Kwang Kim: Phone: +82-32-835-8241, E-mail: [kjkpj@inu.ac.kr](mailto:kjkpj@inu.ac.kr)

Dr. Sang Un Park: Phone: +82-42-821-5730, E-mail: [supark@cnu.ac.kr](mailto:supark@cnu.ac.kr)

<http://dx.doi.org/10.17179/excli2017-253>

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0/>).

Dear Editor,

Resveratrol (3,5,4'-trihydroxy-trans-stilbene) is a stilbenoid and polyphenolic compound. It is found naturally, especially in plants sources such as peanuts, grapes, and some berries (Diaz-Gerevini et al., 2016). In plants, resveratrol is produced via the phenylpropanoid pathway. It is derived from *p*-coumaric acid, which is an intermediate formed during lignin production. The 4-coumarate:coenzyme A (CoA) ligase converts *p*-coumaric acid to coumaroyl-CoA, along with three units of malonyl-CoA, which are then condensed to form resveratrol by resveratrol synthase or stilbene synthase (Deng et al., 2016; Zhang et al., 2015; Zheng et al., 2015; Kim et al., 2013).

Resveratrol is the most widely studied plant-derived polyphenol. Several studies have been reported on its numerous biological and pharmacological effects. These include neuroprotective (Su et al., 2016), antiobesity (Chang et al., 2016), antiviral (Abba et al., 2015), hepatoprotective (Faghihzadeh et al., 2015), anti-inflammatory (Liu et al., 2015), cardioprotective (Raj et al., 2015), anticancer (Kumar et al., 2015), anti-atherogenic (Riccioni et al., 2015), antidiabetic (Szkudelski and Szkudelska, 2015), and antioxidant (Yun et al., 2014) activities. Resveratrol shows a wide range of biological activities and health benefits in humans, which makes it a beneficial substance for use in the pharmaceutical, food, and cosmetic industries. We present a review of the most recent studies on the benefits of resveratrol, especially its biological and pharmacological activities (Table 1).

**Table 1:** The biological and pharmacological properties of resveratrol reported by recent studies

Key findings	Reference
Resveratrol helped to improve blood supply to the femoral head in a rabbit model of osteonecrosis, which might help to protect the vascular endothelium and reduce thrombosis through an anti-inflammatory effect.	Zhai et al., 2016
Dietary intake or therapeutic administration of resveratrol affects the skeletal muscle phenotype in a muscle-specific manner. Therefore, resveratrol supplementation could be a vital approach for promoting the generation of fatigue-resistant myosin heavy chain (type I) isoform, specifically if its expression is suppressed as a result of a long-term high-fat or sugar diet.	Hyatt et al., 2016
The findings of the study emphasize new mechanistic insights that dihydro-resveratrol can protect against oxidative damage in the pancreas. In addition, the findings support the potential of dihydro-resveratrol for the remedy of acute pancreatitis, especially in patients who are not responsive to trans-resveratrol because they lack the necessary microbiota strains.	Tsang et al., 2016
Resveratrol chemosensitized cancer cells to doxorubicin by protecting against cell proliferation and breast cancer. Additionally, co-treatment with resveratrol and doxorubicin induced apoptosis through the suppression of chronic inflammation and autophagy.	Rai et al., 2016
Resveratrol influences the differentiation of preadipocytes by hindering insulin signaling, mitochondrial biogenesis, lipogenesis, and lipid storage, which contribute to weight loss in both humans and animals.	Li et al., 2016a
Resveratrol has beneficial effects on redox balance in bovine mammary epithelial cells. Therefore, it could be a vital therapeutic agent against oxidative insults in lactating animals.	Jin et al., 2016
Resveratrol has the ability to ameliorate sporadic hypoxia-induced anxiety. Furthermore, it causes spatial learning deficits through its effects on hippocampal oxidative pathways, which may directly involve decreasing the expression of the p47Phox subunit of nicotinamide adenine dinucleotide phosphate (NADPH) oxidase. In addition, resveratrol has a potential therapeutic role in obstructive sleep apnea, which should be investigated in further studies.	Abdel-Wahab et al., 2016
Resveratrol acts as a blocker of signal transduction, as well as an activator of the transcription of STAT3 and STAT5. Moreover, through these mechanisms, resveratrol may exert a potential growth inhibitory effect on renal carcinoma cells.	Kim et al., 2016
Combined therapy using resveratrol and an enriched environment evidently improved the neuroprotective effect of resveratrol, compared to monotherapy using either treatment. Therefore, this combination therapy could be researched for treating ischemic brain injury.	Su et al., 2016
It was reported that resveratrol showed a potent immune-enhancing activity in immunosuppressed mice. The authors indicated that the possible mechanism underlying the observed action was the activation of nuclear factor-kappa B.	Lai et al., 2016
Resveratrol influenced the behavior of rats undergoing cocaine withdrawal. In addition, it was indicated that oxidative stress, inflammation, apoptosis, and silent information regulator 1 signaling pathways in the hippocampus, prefrontal cortex, or in both might be involved in mediating the observed effects of resveratrol.	Hu et al., 2016
Resveratrol modified dendritic morphology in the prefrontal cortex, hippocampus, and dentate gyrus, which explained the potential therapeutic effect of resveratrol in both aging and Alzheimer's disease (AD).	Montserrat Hernández-Hernández et al., 2016
Resveratrol advanced the vasoprotective effect of captopril on aortic remodeling and fibrosis during renovascular hypertension. This effect was mediated through the synergetic antioxidant and nitric oxide generation actions of resveratrol.	Natalin et al., 2016
Resveratrol showed an anti-obesity effect through the induction of cytotoxicity when it was used at a high dosage. However, at lower concentrations, it influenced preadipocyte differentiation and lipolysis in mature adipocytes.	Chang et al., 2016
Low-intensity exercise training combined with supplementation with resveratrol and piperine increased the mitochondrial capacity of the forearm skeletal muscle.	Polley et al., 2016

Key findings	Reference
Co-treatment with diclofenac and resveratrol may represent a novel approach to reducing the dosage and gastrointestinal side effects of diclofenac.	Bedada et al., 2016
Metabolic and cardiac functions improved independence of sex through dietary supplementation with resveratrol in Sprague-Dawley rats. This study showed the therapeutic role of resveratrol in the male and female rat offspring that experienced prenatal hypoxia.	Shah et al., 2016
Resveratrol had beneficial effects in the liver by attenuating oxidative stress and downregulating the expression of receptor for advanced glycation end products.	Khazaei et al., 2016
Resveratrol played a positive role in suppressing pancreatic cancer progression induced by hypoxia-driven reactive oxygen species (ROS) by inhibiting the hedgehog signaling pathway. This result suggests that resveratrol may be a potential candidate for the chemoprevention of cancer.	Li et al., 2016b
It was demonstrated that resveratrol serves as a clinically and economically feasible therapeutic agent for reducing the global burden of iron-overload cardiomyopathy at its early and chronic stages.	Das et al., 2015
Resveratrol induced the apoptosis of human leukemic K562 cells via the mitochondrial signaling pathway. This finding necessitates in vivo studies on the potential use of resveratrol as an anticancer agent.	Wang et al., 2015
It was demonstrated that resveratrol is effective against ischemia-induced apoptosis in the rat hippocampus through its neuroprotective and antioxidant effects.	Meng et al., 2015
Resveratrol inhibited glucose uptake and had a significant antineoplastic effect in a preclinical mouse model of ovarian cancer. In addition, treatment with resveratrol suppressed tumor regrowth after therapy with cisplatin, suggesting that resveratrol has the potential to prolong disease-free survival.	Tan et al., 2016
Resveratrol exerted both anti-apoptotic and anti-catabolic effects on compression injury in the skeletal muscle of rats. However, these effects required the action of silent mating type information of 2 homolog 1.	Sin et al., 2015
The authors demonstrated the potential therapeutic use of resveratrol to prevent the development of proliferative vitreoretinopathy by targeting retinal pigment epithelial cells during their epithelial to mesenchymal transition.	Ishikawa et al., 2015
It was shown that resveratrol has a considerable antiepileptic effect. In addition, it has a neuroprotective effect and counters epileptic depression. The findings indicated that resveratrol has the potential to be a new antiepileptic drug; therefore, more studies are needed to further elucidate its therapeutic potential.	Lu and Wang, 2015
Resveratrol alleviates hepatic steatosis and inflammation in choline-deficiency-induced non-alcoholic steatohepatitis (NASH) through autophagy. Therefore, resveratrol may be useful for inhibiting accumulation of lipids and the inflammatory processes associated with NASH.	Ji et al., 2015
It was reported that habitual dietary intake of resveratrol is associated with a low risk of developing frailty syndrome in community-dwelling elderly people. This effect was observed during the first 3 years of follow-up but not after longer follow-up periods.	Rabassa et al., 2015
Intake of fructose for a long period may have different metabolic and vascular effects on male and female rats; however, these effects were observed to be modified by resveratrol.	Pektaş et al., 2015
It was documented that resveratrol is a compound for the potential treatment of thrombovascular diseases through its antiplatelet effect. This effect was observed to occur via the inhibition of ROS production induced by NADPH oxidase and the subsequent oxidative inactivation of SHP-2.	Jang et al., 2015
This recent study provided direct electrophysiological evidence of the inhibitory effect of resveratrol on pyramidal neurons. This effect was mediated, at least in part, by the reduction of evoked neural activity.	Meftahi et al., 2015
It was demonstrated that attenuation of inflammatory responses might be involved in the mechanisms underlying the cardioprotective effect of resveratrol. The findings showed the response of resveratrol to myocardial ischemia/reperfusion injury.	Dong et al., 2015

Key findings	Reference
Resveratrol improved the hepatic and peripheral insulin resistance induced by free fatty acids. Therefore, it may help to mitigate the health consequences of obesity.	Pereira et al., 2015
The genotoxic potential of resveratrol might be attributed to its sulfate- and glucuronide-phase II metabolites through the inhibition of topoisomerase II activity. In addition, resveratrol-3-sulfate might serve as a pool for the parent compound by deconjugation at its target site.	Schroeter et al., 2015
Diseases involving pathological neovascularization could be managed based on the therapeutic value of resveratrol.	Lee et al., 2015
Resveratrol modified kidney histology and leptin expression level in diabetic rats through its antioxidant and antidiabetic effects. Therefore, resveratrol might be useful in the prevention of kidney damage caused by long-term hyperglycemia.	Yaylali et al., 2015
Resveratrol performed more favorably than famotidine did because resveratrol prevents long-term methotrexate toxicity but does not inhibit gastric acid secretion.	Arslan et al., 2015
Resveratrol exhibited a therapeutic potential against myeloproliferative neoplasms, which is evident of its aberrant activation of the janus kinase 2 pathway.	Trung et al., 2015
Resveratrol emerged as a potential agent for the treatment of conditions associated with androgen excess, such as polycystic ovarian syndrome. However, further studies are required to evaluate the efficacy of resveratrol for the treatment of gynecological conditions.	Ortega and Duleba, 2015
Resveratrol was recognized as a safe compound since no significant toxic effects were observed after several concentrations were tested. In addition, it was considered as an effective anti-atherogenic agent.	Riccioni et al., 2015
Resveratrol was found to be a potential agent for reducing human rhinovirus replication and virus-induced cytokine/chemokine production.	Mastromarino et al., 2015
It was considered that NADPH oxidase might be a potential target in AD treatment and that resveratrol may be used as a natural product with a therapeutic potential against AD.	Yao et al., 2015
From histopathological and immunohistochemical analyses that were performed, it was observed that resveratrol ameliorates cisplatin-induced oxidative injury in the rabbit kidney.	Cigremis et al., 2015
Resveratrol might be considered a potential natural product for development as a medicine or dietary supplement for the prevention and treatment osteoporosis.	Zhao et al., 2015
Resveratrol ameliorated spatial learning and memory deficit in a rat model of sub-clinical hypothyroidism. The mechanism involved in this effect might modify the hyperactive hypothalamic-pituitary-thyroid axis and upregulate the hippocampal hypoxexpression of synaptotagmin 1 and brain-derived neurotrophic factor.	Ge et al., 2015

### Acknowledgements

This research was supported by Agriculture, Food and Rural Affairs Research Center Support Program, Ministry of Agriculture, Food and Rural Affairs.

### Conflict of interest

The authors declare no conflict of interest.

### REFERENCES

Abba Y, Hassim H, Hamzah H, Noordin MM. Antiviral activity of resveratrol against human and animal viruses. *Adv Virol.* 2015;2015:184241.

Abdel-Wahab BA, Abdel-Wahab MM. Protective effect of resveratrol against chronic intermittent hypoxia-induced spatial memory deficits, hippocampal oxidative DNA damage and increased p47Phox NADPH oxidase expression in young rats. *Behav Brain Res.* 2016; 305:65-75.

Arslan A, Ozcicek F, Keskin Cimen F, Altuner D, Yarali O, et al. Protective effect of resveratrol against methotrexate-induced oxidative stress in the small intestinal tissues of rats. *Int J Clin Exp Med.* 2015;8: 10491-500.

Bedada SK, Yellu NR, Neerati P. Effect of resveratrol treatment on the pharmacokinetics of diclofenac in healthy human volunteers. *Phytother Res.* 2016;30: 397-401.

- Chang CC, Lin KY, Peng KY, Day YJ, Hung LM. Resveratrol exerts anti-obesity effects in high-fat diet obese mice and displays differential dosage effects on cytotoxicity, differentiation, and lipolysis in 3T3-L1 cells. *Endocr J.* 2016;63:169-78.
- Cigremis Y, Akgoz M, Ozen H, Karaman M, Kart A, Gecer M, et al. Resveratrol ameliorates cisplatin-induced oxidative injury in New Zealand rabbits. *Can J Physiol Pharmacol.* 2015;93:727-35.
- Das SK, Wang W, Zhabyeyev P, Basu R, McLean B, Fan D, et al. Iron-overload injury and cardiomyopathy in acquired and genetic models is attenuated by resveratrol therapy. *Sci Rep.* 2015;5:18132.
- Deng N, Chang E, Li M, Ji J, Yao X, Bartish IV, et al. Transcriptome characterization of *Gnetum parvifolium* reveals candidate genes involved in important secondary metabolic pathways of flavonoids and stilbenoids. *Front Plant Sci.* 2016;7:174.
- Diaz-Gerevini GT, Repossi G, Dain A, Tarres MC, Das UN, Eynard AR. Beneficial action of resveratrol: How and why? *Nutrition.* 2016;32:174-8.
- Dong W, Yang R, Yang J, Yang J, Ding J, Wu H, et al. Resveratrol pretreatment protects rat hearts from ischemia/reperfusion injury partly via a NALP3 inflammasome pathway. *Int J Clin Exp Pathol.* 2015;8:8731-41.
- Faghihzadeh F, Hekmatdoost A, Adibi P. Resveratrol and liver: A systematic review. *J Res Med Sci.* 2015;20:797-810.
- Ge JF, Xu YY, Li N, Zhang Y, Qiu GL, Chu CH, et al. Resveratrol improved the spatial learning and memory in subclinical hypothyroidism rat induced by hemi-thyroid electrocauterization. *Endocr J.* 2015;62:927-38.
- Hu P, Zhu W, Zhu C, Jin L, Guan Y, Guan X. Resveratrol fails to affect cocaine conditioned place preference behavior, but alleviates anxiety-like behaviors in cocaine withdrawn rats. *Psychopharmacology (Berl).* 2016;233:1279-87.
- Hyatt JP, Nguyen L, Hall AE, Huber AM, Kocan JC, Mattison JA, et al. Muscle-specific myosin heavy chain shifts in response to a long-term high fat/high sugar diet and resveratrol treatment in nonhuman primates. *Front Physiol.* 2016;7:77.
- Ishikawa K, He S, Terasaki H, Nazari H, Zhang H, Spee C, et al. Resveratrol inhibits epithelial-mesenchymal transition of retinal pigment epithelium and development of proliferative vitreoretinopathy. *Sci Rep.* 2015;5:16386.
- Jang JY, Min JH, Wang SB, Chae YH, Baek JY, Kim M, et al. Resveratrol inhibits collagen-induced platelet stimulation through suppressing NADPH oxidase and oxidative inactivation of SH2 domain-containing protein tyrosine phosphatase-2. *Free Radic Biol Med.* 2015;89:842-51.
- Ji G, Wang Y, Deng Y, Li X, Jiang Z. Resveratrol ameliorates hepatic steatosis and inflammation in methionine/choline-deficient diet-induced steatohepatitis through regulating autophagy. *Lipids Health Dis.* 2015;14:134.
- Jin X, Wang K, Liu H, Hu F, Zhao F, Liu J. Protection of bovine mammary epithelial cells from hydrogen peroxide-induced oxidative cell damage by resveratrol. *Oxid Med Cell Longev.* 2016;2016:2572175.
- Khazaei M, Karimi J, Sheikh N, Goodarzi MT, Saidijam M, Khodadadi I, et al. Effects of resveratrol on receptor for advanced glycation end products (rage) expression and oxidative stress in the liver of rats with type 2 diabetes. *Phytother Res.* 2016;30:66-71.
- Kim C, Baek SH, Um JY, Shim BS, Ahn KS. Resveratrol attenuates constitutive STAT3 and STAT5 activation through induction of PTP $\epsilon$  and SHP-2 tyrosine phosphatases and potentiates sorafenib-induced apoptosis in renal cell carcinoma. *BMC Nephrol.* 2016;17(1).
- Kim YB, Thwe AA, Kim Y, Yeo SK, Lee C, Park SU. Characterization of cDNA encoding resveratrol synthase and accumulation of resveratrol in tartary buckwheat. *Nat Prod Commun.* 2013;8:1571-4.
- Kumar A, Dhar S, Rimando AM, Lage JM, Lewin JR, Zhang X, et al. Epigenetic potential of resveratrol and analogs in preclinical models of prostate cancer. *Ann N Y Acad Sci.* 2015;1348:1-9.
- Lai X, Pei Q, Song X, Zhou X, Yin Z, Jia R et al. The enhancement of immune function and activation of NF- $\kappa$ B by resveratrol-treatment in immunosuppressive mice. *Int Immunopharmacol.* 2016;33:42-7.
- Lee CS, Choi EY, Lee SC, Koh HJ, Lee JH, Chung JH. Resveratrol inhibits hypoxia-induced vascular endothelial growth factor expression and pathological neo-vascularization. *Yonsei Med J.* 2015;56:1678-85.
- Li S, Bouzar C, Cottet-Rousselle C, Zagotta I, Larmarche F, Wabitsch M, et al. Resveratrol inhibits lipogenesis of 3 T3-L1 and SGBS cells by inhibition of insulin signaling and mitochondrial mass increase. *Biochim Biophys Acta.* 2016a;1857:643-52.

- Li W, Cao L, Chen X, Lei J, Ma Q. Resveratrol inhibits hypoxia-driven ROS-induced invasive and migratory ability of pancreatic cancer cells via suppression of the Hedgehog signaling pathway. *Oncol Rep.* 2016b;35:1718-26.
- Liu FC, Tsai YF, Tsai HI, Yu HP. Anti-Inflammatory and Organ-Protective Effects of Resveratrol in Trauma-Hemorrhagic Injury. *Mediators Inflamm.* 2015;2015:643763.
- Lu S, Wang X. The role and potential mechanism of resveratrol in the prevention and control of epilepsy. *Future Med Chem.* 2015;7:2005-18.
- Mastromarino P, Capobianco D, Cannata F, Nardis C, Mattia E, De Leo A, et al. Resveratrol inhibits rhinovirus replication and expression of inflammatory mediators in nasal epithelia. *Antiviral Res.* 2015;123:15-21.
- Meftahi G, Ghotbedin Z, Eslamizade MJ, Hosseinnardi N, Janahmadi M. Suppressive effects of resveratrol treatment on the intrinsic evoked excitability of CA1 pyramidal neurons. *Cell J.* 2015;17:532-9.
- Meng Z, Li J, Zhao H, Liu H, Zhang G, Wang L, et al. Resveratrol relieves ischemia-induced oxidative stress in the hippocampus by activating SIRT1. *Exp Ther Med.* 2015;10:525-30.
- Monserrat Hernández-Hernández E, Serrano-García C, Antonio Vázquez-Roque R, Díaz A, Monroy E, Rodríguez-Moreno A, et al. Chronic administration of resveratrol prevents morphological changes in prefrontal cortex and hippocampus of aged rats. *Synapse.* 2016;70:206-17.
- Natalin HM, Garcia AF, Ramalho LN, Restini CB. Resveratrol improves vasoprotective effects of captopril on aortic remodeling and fibrosis triggered by renovascular hypertension. *Cardiovasc Pathol.* 2016;25:116-9.
- Ortega I, Duleba AJ. Ovarian actions of resveratrol. *Ann N Y Acad Sci.* 2015;1348:86-96.
- Pektaş MB, Sadi G, Akar F. Long-term dietary fructose causes gender-different metabolic and vascular dysfunction in rats: modulatory effects of resveratrol. *Cell Physiol Biochem.* 2015;37:1407-20.
- Pereira S, Park E, Moore J, Faubert B, Breen DM, Oprescu AI, et al. Resveratrol prevents insulin resistance caused by short-term elevation of free fatty acids in vivo. *Appl Physiol Nutr Metab.* 2015;40:1129-36.
- Polley KR, Jenkins N, O'Connor P, McCully K. Influence of exercise training with resveratrol supplementation on skeletal muscle mitochondrial capacity. *Appl Physiol Nutr Metab.* 2016;41:26-32.
- Rabassa M, Zamora-Ros R, Urpi-Sarda M, Bandinelli S, Ferrucci L, Andres-Lacueva C, et al. Association of habitual dietary resveratrol exposure with the development of frailty in older age: the Invecchiare in Chianti study. *Am J Clin Nutr.* 2015;102:1534-42.
- Rai G, Mishra S, Suman S, Shukla Y. Resveratrol improves the anticancer effects of doxorubicin in vitro and in vivo models: A mechanistic insight. *Phytomedicine.* 2016;23:233-42.
- Raj P, Zieroth S, Neticadan T. An overview of the efficacy of resveratrol in the management of ischemic heart disease. *Ann N Y Acad Sci.* 2015;1348:55-67.
- Riccioni G, Gammone MA, Tettamanti G, Bergante S, Pluchinotta FR, D'Orazio N. Resveratrol and anti-atherogenic effects. *Int J Food Sci Nutr.* 2015;66:603-10.
- Schroeter A, Groh IA, Del Favero G, Pignitter M, Schueller K, Somoza V, et al. Inhibition of topoisomerase II by phase II metabolites of resveratrol in human colon cancer cells. *Mol Nutr Food Res.* 2015;59:2448-59.
- Shah A, Reyes LM, Morton JS, Fung D, Schneider J, Davidge ST. Effect of resveratrol on metabolic and cardiovascular function in male and female adult offspring exposed to prenatal hypoxia and a high-fat diet. *J Physiol.* 2016;594:1465-82.
- Sin TK, Yung BY, Yip SP, Chan LW, Wong CS, Tam EW, et al. SIRT1-dependent myoprotective effects of resveratrol on muscle injury induced by compression. *Front Physiol.* 2015;6:293.
- Su Q, Pu H, Hu C. Neuroprotection by combination of resveratrol and enriched environment against ischemic brain injury in rats. *Neurol Res.* 2016;17:1-9.
- Szkudelski T, Szkudelska K. Resveratrol and diabetes: from animal to human studies. *Biochim Biophys Acta.* 2015;1852:1145-54.
- Tan L, Wang W, He G, Kuick RD, Gossner G, Kueck AS, et al. Resveratrol inhibits ovarian tumor growth in an in vivo mouse model. *Cancer.* 2016;122:722-9.
- Trung LQ, Espinoza JL, An DT, Viet NH, Shimoda K, Nakao S. Resveratrol selectively induces apoptosis in malignant cells with the JAK2V617F mutation by inhibiting the JAK2 pathway. *Mol Nutr Food Res.* 2015;59:2143-54.
- Tsang SW, Guan YF, Wang J, Bian ZX, Zhang HJ. Inhibition of pancreatic oxidative damage by stilbene derivative dihydro-resveratrol: implication for treatment of acute pancreatitis. *Sci Rep.* 2016;6:22859.

Wang B, Liu J, Gong Z. Resveratrol induces apoptosis in K562 cells via the regulation of mitochondrial signaling pathways. *Int J Clin Exp Med*. 2015;8:16926-33.

Yao Y, Li J, Niu Y, Yu JQ, Yan L, Miao ZH, et al. Resveratrol inhibits oligomeric A $\beta$ -induced microglial activation via NADPH oxidase. *Mol Med Rep*. 2015;12:6133-9.

Yaylali A, Ergin K, Ceçen S. Effect of resveratrol on leptin and sirtuin 2 expression in the kidneys in streptozotocin-induced diabetic rats. *Anal Quant Cytopathol Histopathol*. 2015;37:243-51.

Yun H, Park S, Kim MJ, Yang WK, Im DU, Yang KR, et al. AMP-activated protein kinase mediates the antioxidant effects of resveratrol through regulation of the transcription factor FoxO1. *FEBS J*. 2014;281:4421-38.

Zhai JL, Weng XS, Wu ZH, Guo SG. Effect of Resveratrol on Preventing Steroid-induced Osteonecrosis in a Rabbit Model. *Chin Med J (Engl)*. 2016;129:824-30.

Zhang E, Guo X, Meng Z, Wang J, Sun J, Yao X, et al. Construction, expression, and characterization of *Arabidopsis thaliana* 4CL and *Arachis hypogaea* RS fusion gene 4CL::RS in *Escherichia coli*. *World J Microbiol Biotechnol*. 2015;31:1379-85.

Zhao L, Wang Y, Wang Z, Xu Z, Zhang Q, Yin M. Effects of dietary resveratrol on excess-iron-induced bone loss via antioxidative character. *J Nutr Biochem*. 2015;26:1174-82.

Zheng S, Zhao S, Li Z, Wang Q, Yao F, Yang L, et al. Evaluating the effect of expressing a peanut resveratrol synthase gene in rice. *PLoS One*. 2015;10(8):e0136013.