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- Ching-Shu Lai, Shiming Li, Yutaka Miyauchi, Michiko Suzawa, Chi-Tang Ho and <u>Min-Hsiung</u> <u>Pan\*</u>. Potent anti-cancer effects of citrus peel flavonoids in human prostate xenograft tumors. *Food & Function*. 2013, 4 (6), 944 - 949.
- <u>Pan, Win-Hsiung</u>, Lai, Ching-Shu, Wang, Hong, Lo, Chin-Yu, Ho, Chi-Tang, Li, Shiming. Black tea in chemoprevention of cancer and other human diseases. *Food Science and Human Wellness*, 2013, 2, 13-22.
- Yi-Shiou Chiou, Shengmin Sang, Chi-Tang Ho, Ying-Jan Wang\* and <u>Min-Hsiung Pan\*</u>. Peracetylated (-)-epigallocatechin-3-gallate (AcEGCG) potently prevents skin carcinogenesis by suppressing the PKD1-dependent signaling pathway in CD34+ skin stem cells and skin tumors. *Carcinogenesis*. 2013. PMID: 23385063 [PubMed - as supplied by publisher] (SCI IF= 5.702. rank: 23/194=11.86% in ONCOLOGY, JCR 2011).(Corresponding author)
- <u>Min-Hsiung Pan</u><sup>\*</sup>, Ching-Shu Lai, Jia-Ching Wu and Chi-Tang Ho. Epigenetic and diseases targets by polyphenols. *Current Pharmaceutical Design*. 2013. (Review) Accepted. (Co-Corresponding author)
- Ming-Fen Lee, Min-Lung Liu, An-Chin Cheng, Mei-Ling Tsai, Chi-Tang Ho, Wen-Shiung Liou, <u>Min-Hsiung Pan\*</u>. Pterostilbene inhibits dimethylnitrosamine -induced liver fibrosis in rats. *Food Chemistry*. 2013, 138, 802-807 (SCI IF= 3.655. rank: 3/71=4.23% in CHEMISTRY, APPLIED, JCR 2011). (Corresponding author)
- Ming-Fen Lee, Mei-Ling Tsai, Pei-Pei Sun, Ling-Lung Chien, An-Chin Cheng, Nianhan Jia-Lin Ma, Chi-Tang Ho, <u>Min-Hsiung Pan\*</u>. Phyto-power dietary supplement potently inhibits dimethylnitrosamine-induced liver fibrosis in rats. *Food & Function.* 2012. 4 (3):470-5. (SCI IF= 1.179. rank: 64/128=50% in Food Science & technology, JCR 2011).

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Int J Food Microbiol. 2014 Jul 16;182-183:1-8. doi: 10.1016/j.ijfoodmicro.2014.05.001. Epub 2014 May 9.

Biocontrol activity of an alkaline serine protease from Aureobasidium pullulans expressed in Pichia pastoris against four postharvest pathogens on apple.

Banani H<sup>1</sup>, Spadaro D<sup>2</sup>, Zhang D<sup>1</sup>, Matic S<sup>1</sup>, Garibaldi A<sup>1</sup>, Gullino ML<sup>3</sup>.

Author information

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Malus/microbiology\*

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Abstract Ginseng and lingzhi (Ganoderma lucidum) both traditional medicines in many Asian countries. I contains a lot of bioactive compounds such as fermentation medium of G. lucidum to produce fermentation products were analysed during fer K (CK) significantly increased, especially Rd, w ginsenoside in the final fermentation product. F Copyright © 2013 Elsevier Ltd. All rights reserv KEYWORDS: Biotransformation; Ganoderma lucid	are valuable traditional Chinese medicines a However, massive quantity of ginseng residue ginsenosides. The goal of this study was to re bioactive ginsenoside enriched biotransforma rmentation. Our results showed that after 30 d thile other ginsenosides (Re, Rb1 and Rc) dec furthermore, the biotransformation of ginsenou red.	and have been extensively utilised in functional foods and e is produced after extraction of ginseng which still euse the American ginseng extraction residue as the ation products. The changes of ginsenosides in the days of fermentation, ginsenoside Rg1, Rd, and compound creased during fermentation. Ginsenoside Rd is the major sides was the major reaction in this fermentation process. -文章識別號	Related citations in PubMed Bioconversion of ginsenosides in the american ginseng ( xī yár [J Tradit Complement Med. 2013] Highly selective microbial transformation of major ginsenoside Rb1 to gyper [J Appl Microbiol. 2012] Transformation of ginsenosides Rb2 and Rc from Panax ginseng by food mi [Biol Pharm Bull. 2005] Review Comparison of the pharmacological effects of Panax ginse [Acta Pharmacol Sin. 2008] Review Biotransformation of ginsenosides by bydebezing the (Appl Microbiol Biotechnol. 2010)
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Oral Rg1 supplementation strengthens antioxidant defense system against exercise-induced oxidative stress in rat skeletal muscles.

Yu SH1, Huang HY, Korivi M, Hsu MF, Huang CY, Hou CW, Chen CY, Kao CL, Lee RP, Lee SD, Kuo CH.

Author information

#### Abstract

BACKGROUND: Previous studies reported divergent results on nutraceutical actions and free radical scavenging capability of ginseng extracts. Variations in ginsenoside profile of ginseng due to different soil and cultivating season may contribute to the inconsistency. To circumvent this drawback, we assessed the effect of major ginsenoside-Rg1 (Rg1) on skeletal muscle antioxidant defense system against exhaustive exerciseinduced oxidative stress.

METHODS: Forty weight-matched rats were evenly divided into control (N = 20) and Rg1 (N = 20) groups. Rg1 was orally administered at the dose of 0.1 mg/kg bodyweight per day for 10-week. After this long-term Rg1 administration, ten rats from each group performed an exhaustive swimming, and

remaining rats considered as non-exercise control. Tibialis anterior (TA) m exercise rats.

RESULTS: Exhaustive exercise significantly (p<0.05) increased the lipid p (MDA) levels. The increased oxidative stress after exercise was also confid (GSH/GSSG ratio) in control rats. However, these changes were complete (GPx) activities were significantly (p<0.05) increased by Rg1 in non-exerci reductase (GR) and glutathione S-transferase (GST) activities were signific

CONCLUSIONS: This study provide compelling evidences that Rg1 supple and completely attenuate the membrane lipid peroxidation induced by exh supplement to buffer the exhaustive exercise-induced oxidative stress.

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### Steroid-like compounds in Chinese medicines promote blood circulation via inhibition

Chen RJ1, Chung TY, Li FY, Yang WH, Jinn TR, Tzen JT.

#### Author information

### Abstract

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AIM: To examine if steroid-like compounds found in many Chinese medicinal products conventionally used for the act as active components via the same molecular mechanism triggered by cardiac glycosides, such as ouabain.

METHODS: The inhibitory potency of ouabain and the identified steroid-like compounds on Na(+)/K(+)-ATPase act Molecular modeling was exhibited for the docking of these compounds to Na(+)/K(+)-ATPase.

RESULTS: All the examined steroid-like compounds displayed more or less inhibition on Na(+)/K(+)-ATPase, with equivalent to ouabain) exhibiting significantly higher inhibitory potency than the others. In the pentacyclic triterpeno oleanolic acid were moderate inhibitors of Na(+)/K(+)-ATPase, and their inhibitory potency was comparable to that high inhibitory potency of ursolic acid or oleanolic acid was due to the formation of a hydrogen bond between its ca in the deep cavity close to two K(+) binding sites of Na(+)/K(+)-ATPase. Moreover, the drastic difference observed bufalin, ginsenoside Rh2, and pentacyclic triterpenoids is ascribed mainly to the number of hydrogen bonds and pa interaction between the compounds and residues around the deep cavity of Na(+)/K(+)-ATPase.

CONCLUSION: Steroid-like compounds seem to contribute to therapeutic effects of many cardioactive Chinese me such as Prunella vulgaris L, rich in ursolic acid, oleanolic acid and their glycoside derivatives may be adequate sou inhibition on Na(+)/K(+)-ATPase.

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The inhibition of $Na(+)/K(+)$ -ATP ase by versatile steroid-like compounds contributes to the putative therapeutic	effects of many Chinese medicin	al				
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