

Cite this article as: Khan RA, Khan MR Khan A. Comparative antioxidant scavenging and lipid peroxidation activity of rutin and gallic acid. Bangladesh J Pharmacol. 2015; 10: 637-638.

A Journal of the Bangladesh Pharmacological Society (BDPS)

Journal homepage: www.banglajol.info

Abstracted/indexed in Academic Search Complete, Agroforestry Abstracts, Asia Journals Online, Bangladesh Journals Online, Biological Abstracts, BIOSIS Previews, CAB Abstracts, Current Abstracts, Directory of Open Access Journals, EMBASE/Excerpta Medica, Global Health, Google Scholar, HINARI (WHO), International Pharmaceutical Abstracts, Open J-gate, Science Citation Index Expanded, SCOPUS and Social Sciences Citation Index ISSN: 1991-0088; DOI: 10.3329/bjp.v10i3.23362

Letter to the Editor

Comparative antioxidant scavenging and lipid peroxidation activity of rutin and gallic acid

Sir.

Reactive oxygen species (ROS) are generated in the normal metabolism of living organisms, and beside of their beneficial role in signal transduction, they are involved in dispersion of several degenerative diseases like malignant tumors, rheumatic joint inflammation, cataract, Parkinson's disease, Alzheimer's disease, hypertension, diabetes, oxidative stress, tissue damages and atherosclerosis (Halliwell and Gutteridge, 1984). Natural products and secondary plant metabolites play a key role in the prevention of various chronic diseases and improvement of health. Rutin is used as a chemotherapeutic agent in addition to food supplement. The present study was designed to evaluate the antioxidant of free radical and lipid peroxidation activities of various concentrations of rutin and gallic acid.

To evaluate the antioxidant efficacy of both natural compounds various free radical scavenging assays are used.

During these characterizations 1, 1-diphenyl-2-picrylhydrazyl (DPPH), ABTS (2, 20-azinobis-(3-ethylbenzothiazoneline-6-sulphonic acid, 7.4 mM), ammonium molybdate, β-carotene, nitroblue tetrazolium, 2-deoxyribose, hydrogen peroxide and FeCl₂.4H₂O solution are free radicals providers. DPPH assay was performed according to the procedure as reported by Gyamfi et al. (1999). ABTS radical cation (Re et al., 1999), phosphomolybdenum method (Prieto et al., 1999), β-carotenelinoleate (Sun and Ho, 2005), superoxide radical scavenging activity (Nishikimi et al., 1972), hydroxyl radicals (Nagai et al., 2005), hydrogen peroxide (Ruch et al., 1989) and the chelating ability of ferrous ions by various fractions were estimated by the method of Dinis et al. (1994). After % activity of free radical inhibition, IC₅₀ of various assays was determined using Graph pad software.

The scavenging activities of various concentrations of rutin and gallic acid were determined using free radicals of 1, 1-diphenyl 1-2-picrylhydrazyl (DPPH). Results showed that rutin (IC₅₀ 6.7 \pm 0.1 μ g/mL) possessed highest antioxidant activity as compared to gallic acid (19.0 \pm 0.0 μ g/mL) (Table I; Figure 1).

Table I Extracts of rutin and gallic acid for various antioxidant system		
	Rutin	Gallic acid
DPPH activity	6.7	19.0
	$(0.1)^a$	$(0.0)^{b}$
ABTS radical inhibition	52.7	122.1
	(3.2)a	(5.2)b
Phosphomolybdenum assay	58.3	123.0
	$(1.8)^{a}$	(3.1) ^b
β-Carotene bleaching	61.6	190.2
	$(2.4)^{a}$	(2.8)b
Chelating activity	75.3	92.5
	(3.2)a	(3.3)b
Superoxide scavenging	68.6	220.7
	(2.3)a	(7.8) ^b
Nitric oxide scavenging	68.5	145.0
	$(4.2)^{a}$	(3.2) ^b
Lipid peroxidation	57.4	112.2
	$(3.1)^a$	$(2.4)^{b}$
Hydrogen peroxide scavenging	86.3	100.7
	$(4.0)^{a}$	(3.8) ^b
Hydroxyl radical	93.2	98.5
	(2.6)a	$(0.6)^{b}$

Each value is represented as mean (SD); n=3; Means not sharing the same letter are significantly different (LSD) at p<0.01 probability level in each column

The ABTS radical scavenging activity order of bioactive constituents are: rutin>gallic acid with IC50 values of $52.7 \pm 3.2 \,\mu\text{g/mL}$ and $122.1 \pm 5.2 \,\mu\text{g/mL}$, respectively (Table I). The results showed that rutin possessed significantly higher ABTS radical scavenging activity (p<0.01) as compared to gallic acid.

Table I showed the reduction of Mo (VI) to Mo (V) by administration of rutin (IC₅₀ 58.3 \pm 1.8 μ g/mL) and gallic acid (IC₅₀ 123.0 \pm 3.1 μ g/mL).

The absorbance of β -carotene was found to be decreased in the presence of 50-250 µg/mL of the various ascorbic acid, gallic acid and rutin. Various concentrations of these compounds effectively inhibited the oxidation of linoleic acid and subsequent bleaching of β -carotene. Rutin (IC₅₀ 61.6 \pm 2.4 μ g/mL) was more potent then that of gallic acid (IC₅₀ 190.2 \pm 2.8 μ g/mL).

Scavenging activity for superoxide radicals exhibited by



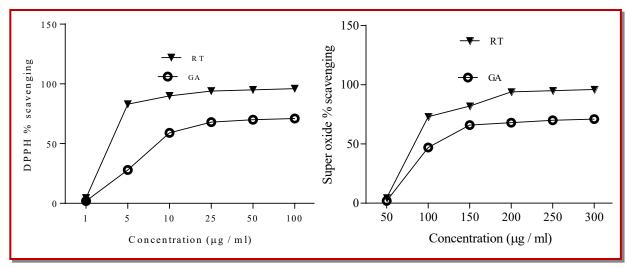


Figure 1: DPPH and superoxide scavenging activity rutin (RU) and gallic acid (GA)

rutin (IC $_{50}$ 68.6 ± 2.3 $\mu g/mL$) and gallic acid (IC $_{50}$ 220.7 ± 7.8 $\mu g/mL$).

Table I shows the hydroxyl radical scavenging activities of rutin (IC₅₀ 93.2 \pm 2.6 μ g/mL) and gallic acid (IC₅₀ 98.5 \pm 0.6 μ g/mL).

Scavenging effect of rutin and gallic acid indicated that rutin possessed (p<0.01) highest hydroxyl radical scavenging effect and was most potent.

Various concentrations of compounds showed an ability to chelate iron (II) ions in a dose-dependent manner. Among these rutin had potent iron chelating activity. The highest activity was observed for rutin (p<0.01) and was more potent in inhibition of lipid peroxidation than gallic acid. Overall, the rutin showed the highest nitric oxide scavenging (p<0.01) capability compared to other compound.

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