

Medicinal Properties of *Njavara* Rice (*Oryza Sativa* L.) cv.

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Abstract— A study was conducted to find out the therapeutic value of medicinal rice (*Oryza sativa* L.) cv. *Njavara*. *Njavara* rice for the study was procured from Rice Research Station, Moncompu.

For assessing the efficacy of *Njavara* on the blood sugar levels, a feeding trial for 3 months was conducted among five subjects who were diabetic and willing to participate but not on medication. Blood sugar levels were monitored during 0, 45 and 90th day of supplementation.

The results revealed that for all subjects' blood sugar levels decreased after supplementation study.

Diphenyl picryl hydrazyl (DPPH) radical scavenging activity, hydroxyl radical scavenging activity, superoxide anion radical scavenging activity and Vitamin E level were also ascertained.

The findings revealed that after the supplementation of *Njavara*, the DPPH scavenging activity, hydroxyl radical activity, superoxide anion-radical scavenging activity and vitamin E level of the blood samples of all the five subjects under study have increased.

Keywords— *Njavara*, *Rice*, *Antioxidants*, *supplementation*.

I. INTRODUCTION

Rice has been used as a medicine by traditional healers from time immemorial. Kerala has an immense wealth of medicinal rice cultivars. Among the various medicinal rices, *Njavara* is a unique grain plant in the *Oryza* genus indigenous to Kerala, widely used in the Ayurvedic system of medicine, especially in Panchakarma treatment. Documents show that it has been under cultivation in Kerala for about 2500 years since the time of Susruta.

Njavara rice, with a distinct gene pool and medicinal properties, can be exploited as nutraceutical rice (Sulochana and Bakiyalakshmi, 2011).

Studies related to therapeutic value of *Njavara* rice are rather limited. So the present study is an attempt to investigate the above said parameters.

II. MATERIALS & METHODS

Njavara rice was collected from Rice Research Station of Kerala Agriculture University, Moncompu

In order to assess the therapeutic value of *Njavara* rice, supplementation study was carried in which *Njavara* rice in grits form was prepared in the laboratory and was given to selected human volunteers with diabetes mellitus.

Conduct of case studies

For the conduct of the case studies, five human subjects who were diabetic but not on medication in the age group of 40-50 years and who were willing to participate were purposively selected through personal interview. After the selection process, preliminary information regarding their socio-economic profile, health status, dietary and life style pattern and nutritional status were collected through a suitably structured questionnaire.

1) Socio-economic profile

In order to elicit information on socio-economic profile of the respondents details regarding age of the subjects, family income, type and size of the family, religion, educational status, money spent on food and health care etc. were collected using the questionnaire.

Using the pre-tested questionnaire, the subjects' food habits and dietary pattern were collected. In life style pattern, the subjects' personal habits like consumption of alcohol, smoking, stress and strain in the daily life, habit of doing exercise etc. were also collected.

2) Nutritional status

Nutritional status of the selected respondents was assessed through anthropometry. Anthropometric measurements relevant to the study include height, weight and BMI and Waist-Hip ratio.

3) Conduct of feeding trail

The selected subjects were supplied the *Njavara* grits for a period of three months based on their individual calorie requirement and disease condition.

4) Assessment of the efficacy of the *Njavara* grits

The feeding trail was conducted for a period of 3 months to assess the efficacy of the *Njavara* grits on the blood glucose levels of the subjects. Blood glucose level of the subjects were taken initially (before the feeding trial), intermittently (after 45 years) and finally (after the feeding trial).

5) Antioxidant properties

a. Diphenyl Picryl Hydrazyl (DPPH) radical scavenging activity

Free radical scavenging activity of sample to characterize antioxidant activity was estimated as suggested by Blois (1958). Different amount of the blood samples were taken (with the help of a lab technician) and DPPH (0.1 mM dissolved in methanol) was mixed together and the reaction mixture was left in dark room for 20 minutes. The absorbance was measured at 517 nm against the blank prepared by mixing DPPH and methanol. The antioxidant activity was expressed in terms of per cent inhibition of DPPH free radicals using the following equation:

$$DPPH\text{radicalscavengingactivity}(\%) = \frac{Abs_{\text{control}} - Abs_{\text{sample}}}{Abs_{\text{control}}} \times 100$$

Where, Abs_{control} = absorbance of DPPH solution (blank) and Abs_{sample} = absorbance of sample.

b. Hydroxyl radical scavenging activity

In order to assess the hydroxyl free radical scavenging activity of the methanolic extracts of the rice samples, the deoxyribose method was used, as described by Halliwell et al. (1987), with some slight modifications. The reaction mixture contained phosphate buffer (20 mM, pH 7.4), 60 mM deoxyribose, 10mM Hydrogen peroxide, 1 mM ferric chloride, 1.04 mM EDTA, different amount of blood samples and the final 2mM ascorbic acid. The reaction mixtures were incubated for 1hr. at 37°C, after which 17 mM trichloro acetic acid (TCA) was added. The mixture was then boiled for 15 minutes, ice cooled and measured for absorbance at 532 nm. Distilled water in lieu of extract was utilized as blank and the sample solution without added deoxyribose was used as a sample blank.

c. Superoxide anion radical scavenging activity

Superoxide anion scavenging activity was measured based on the method described by Robak and Gryglewski (2001). Superoxide radicals were generated in a PMS-NADH system via the oxidation of NADH and then assayed by the reduction of nitro blue tetrazolium (NBT). The superoxide radicals were generated in reaction mixture containing sodium phosphate buffer (100 mM, pH 7.4) containing 150 μ M NBT, 468 μ M NADH solution in sodium phosphate buffer and different concentrations of blood samples. To this 60 μ M phenazine metho sulphite (PMS) solution was added. The reaction mixture was incubated for 5 minutes at 25°C and the absorbance was measured at 560 nm.

d) Vitamin E

Chemiluminescence method was used for the analysis of Vitamin E in blood samples.

III. RESULTS & DISCUSSION

Five subjects having diabetes in the age group of 40-50 years and not on medication were selected for the case study (Subject A-E). Among the five subjects, two were male (A and B) and three were females (Subject C to E). All the subjects were Hindu and were from nuclear family (Table 1).

Table 1 shows that the monthly income of the subjects ranged from 30,000 to 2, 00,000. Their educational qualification ranged from plus two to PG and above, with occupation business (subject A), Govt. employee (B and C), subject D was a housewife whereas subject E was a bank employee.

Duration of the diseases revealed that subject A, C and D were having diabetes for the past 2 years, whereas subject B too was having diabetes for the last nine months and Subject E for the past 3 years. Except subject B and C, all others were not having any diabetic history.

The two male subjects were found to have no exercise habit and both of them were non-vegetarians, whereas female subjects were found to have the habit of doing exercise and all of them were vegetarians.

The anthropometric data of the respondents (Table 2) revealed that subject A and E were coming under Obese grade I category, with a waist-hip ratio of 1.05 and 0.99 respectively. Whereas the other three subjects i.e. subject B, C and D were of normal category, with a waist hip ratio of 0.96, 0.89 and 0.94 respectively.

Price et al. (2006) is of the opinion that waist hip ratio (WHR) has been found to be a more efficient predictor of mortality than BMI.).

The National Institute of Diabetes, Digestive and Kidney Diseases (NIDDK) stated that women with waist-hip ratios of more than 0.8, and men with more than 1.0, are at increased health risk because of their fat distribution (Marlowe et al., 2005). The author also stated that a WHR of 0.7 for women and 0.9 for men has been shown to correlate strongly with general health. Women within the 0.7 range have optimal levels of estrogen and are less susceptible to major diseases such as diabetes, cardiovascular disorders and ovarian cancers. Men with WHRs below 0.9, similarly, have been shown to be healthier.

Efficacy of *Njavara* on the blood sugar levels

The subjects were given *Njavara* grits for a period of three months based on their requirement.

Table 3 shows that all the subjects have shown a decrease in the blood sugar level. The initial blood sugar levels of subject A,B,C, D and E were 193 mg/dl, 140mg/dl, 140 mg/dl, 150mg/dl and 250 mg/dl and their final blood sugar level was 173 mg/dl, 90 mg/dl, 110mg/dl, 109 mg/dl and 160 mg/dl respectively.

The present investigation revealed that *Njavara* has glucose lowering effect. The exact factor for this effect is not known but there can be a number of factors like amino acids, soluble fibre, antioxidants, phenolic compound, minerals and B-complex vitamins.

Antioxidant properties

a) DPPH radical scavenging activity

Free radical scavenging is one of the known mechanisms by which antioxidants inhibit lipid peroxidation. The results of the DPPH scavenging activity in the blood samples of the subjects revealed that after supplementation of *Njavara* rava, their DPPH radical activity has increased on a dose dependent manner (Figure. 1-5).

In a study done by Chrzczanowicz et al. (2008) mean values of DPPH radical scavenging activity obtained in human serum (healthy subjects) were 11.2+/-3.3 per cent.

b) Hydroxyl radical scavenging activity

Hydroxyl and superoxide radicals are the two most representative free radicals. In cellular oxidation reactions, superoxide radical is normally formed first, and its effects can be magnified because it produces other kinds of cell damaging free radicals and oxidizing agents. However, the damaging action of the hydroxyl radical is the strongest among free radicals.

In the present investigation, the results revealed that after the supplementation of *Njavara* grits for three months the hydroxyl scavenging activity of the serum has increased when compared to the activity level before supplementation (Figure. 6-10).

c) Superoxide anion-radical scavenging activity

Superoxide radical is known to be very harmful to cellular components as a precursor of more reactive species.

The superoxide scavenging activity of serum of subjects under study of the present investigation has shown an increase after the feeding trial, which is also on a dose dependent manner (Figure. 11-15).

d) Vitamin E

Vitamin E is a fat-soluble antioxidant that stops the production of ROS formed when fat undergoes oxidation. Scientists are investigating whether, by limiting free-radical production and possibly through other mechanisms, vitamin E might help prevent or delay the chronic diseases associated with free radicals (Dietrich et al., 2006).

In the present investigation, the vitamin E status of the subjects under study revealed that there is an increase in their vitamin E levels after the supplementation of *Njavara* rava from 23.00 to 25.62µg/ml, 9.80 to 13.35 µg/ml, 19.96 to 22.65µg/ml, 24.00 to 25.47 µg/ml and 16.00 to 18.90 µg/ml respectively for subjects A, B, C, D and E.

IV. CONCLUSION

Medicinal plants are a source of antioxidants and bioactive compounds present in them are responsible for the prevention of oxidative stress that leads to many degenerative diseases and conditions (Smitha et al., 2012). The results of the study revealed that *Njavara* was having glucose lowering effect and good antioxidant properties.

Table.1: Socio economic profile of the selected respondents

Particulars	Subjects				
	A	B	C	D	E
Age (yrs.)	47	43	48	49	48
Gender	M	M	F	F	F
Family income (Rs.)	30,000-40,000	40,000-45,000	>2,00,000	50,000-60,000	70,000-80,000
Educational status	Degree	Degree	PG and above	Plus two	PG

Table.2: Anthropometric parameters of the selected subjects

Body measurements	Subject A	Subject B	Subject C	Subject D	Subject E
Ht. (cms)	165	170	157	150	162
Wt. (Kg)	73	65	60	53	74
BMI	26.81	22.49	24.34	23.56	28.20

Table.3: Fasting Blood sugar levels of subjects before and after supplementation

Monitoring intervals	Blood sugar levels (mg/dl)				
	Subject A	Subject B	Subject C	Subject D	Subject E
Initial (Fasting)	193	140	140	150	250
Final	173	90	110	109	160

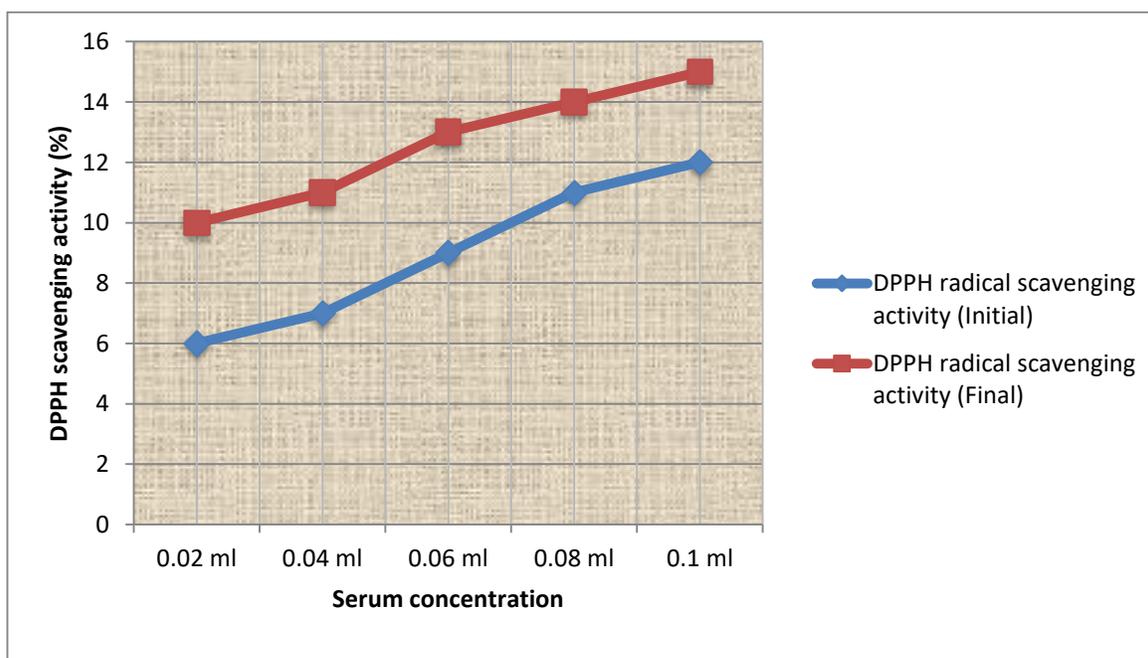


Fig.1: DPPH radical scavenging activity of subject A (before and after supplementation)

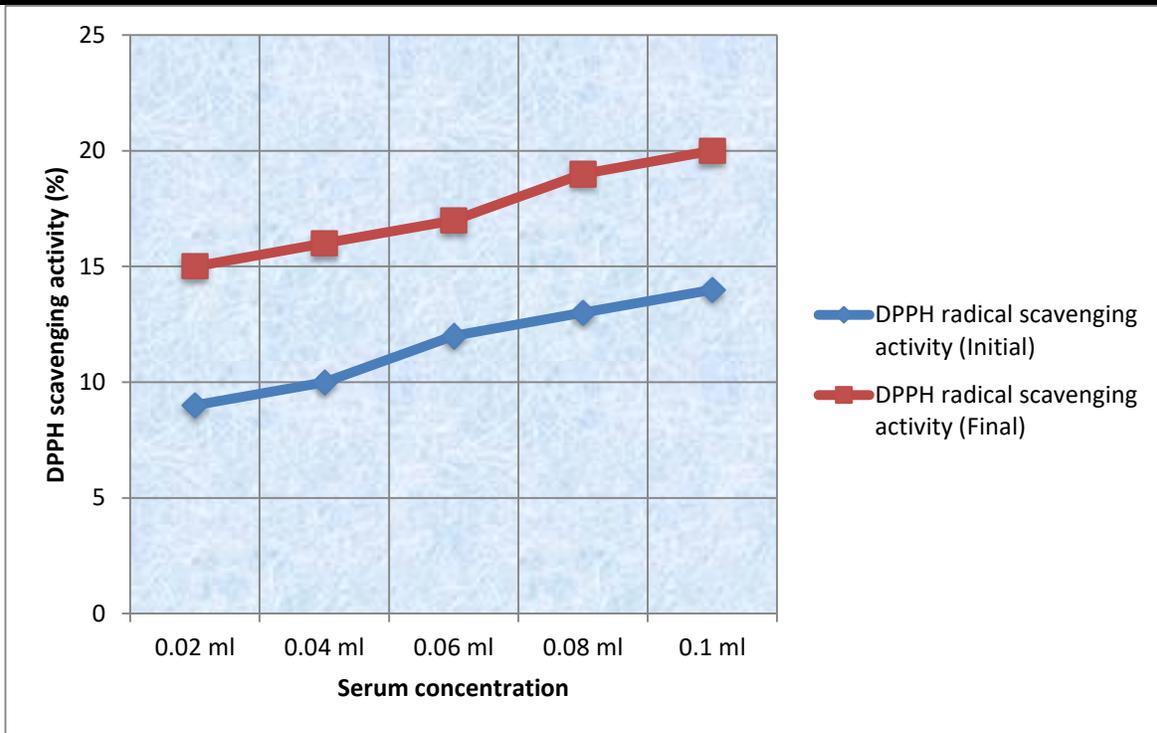


Fig.2: DPPH radical scavenging activity of subject B (before and after supplementation)

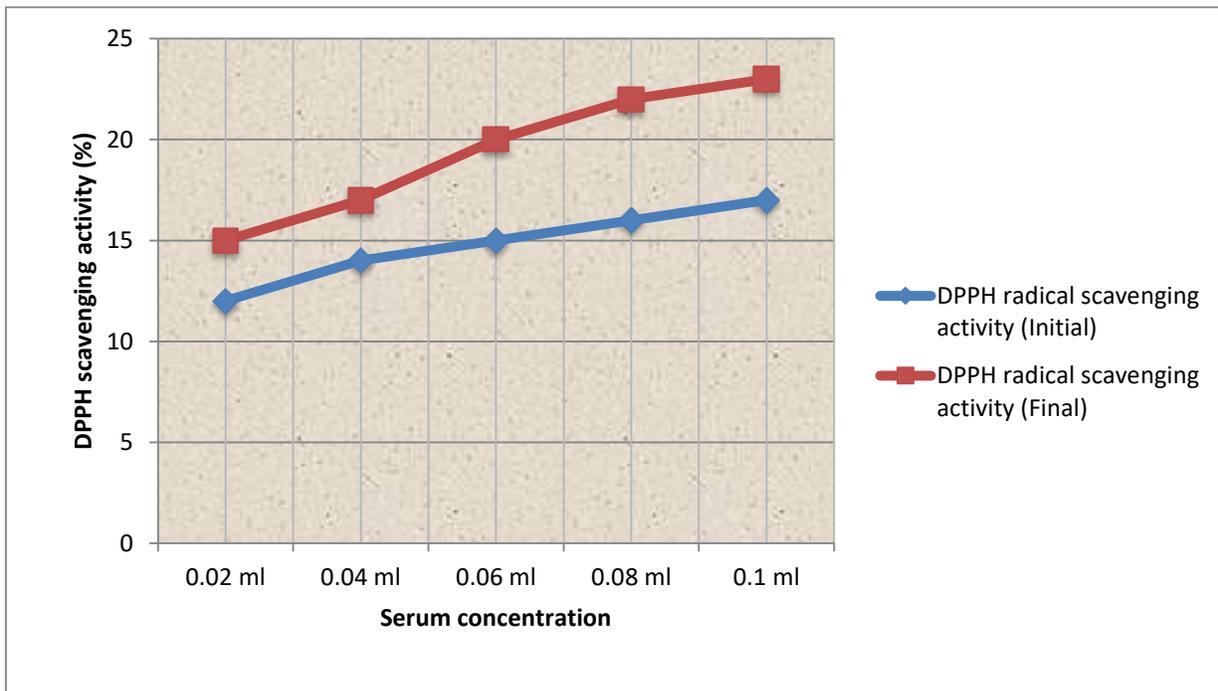


Fig.3: DPPH radical scavenging activity of subject C (before and after supplementation)

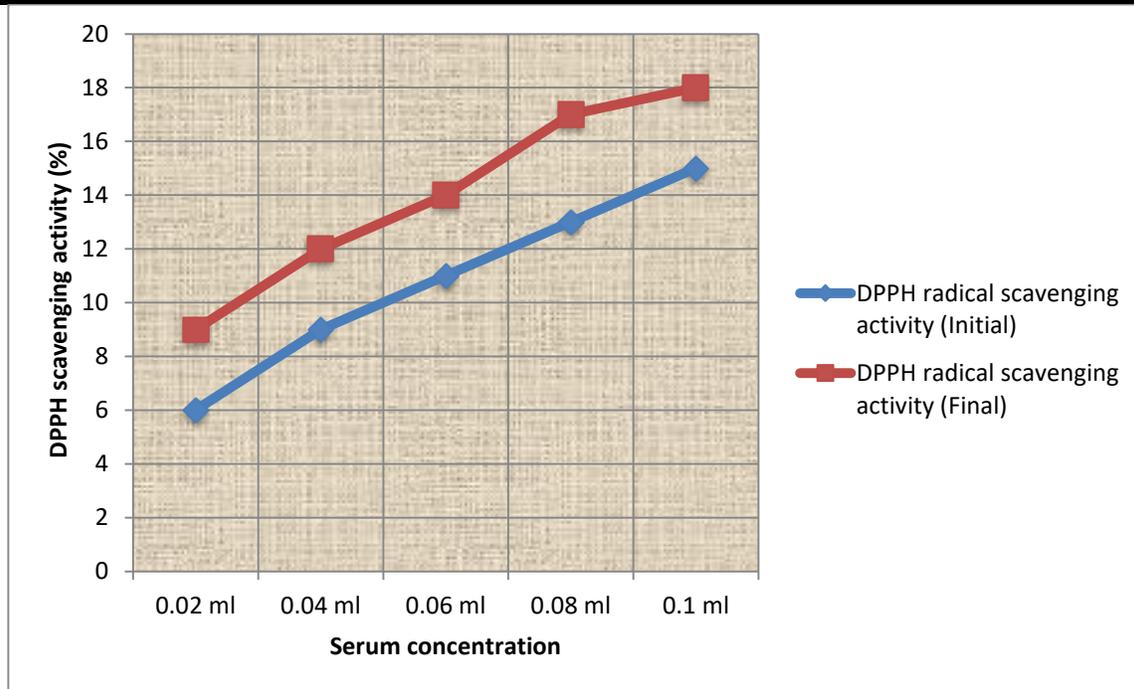


Fig.4: DPPH radical scavenging activity of subject D (before and after supplementation)

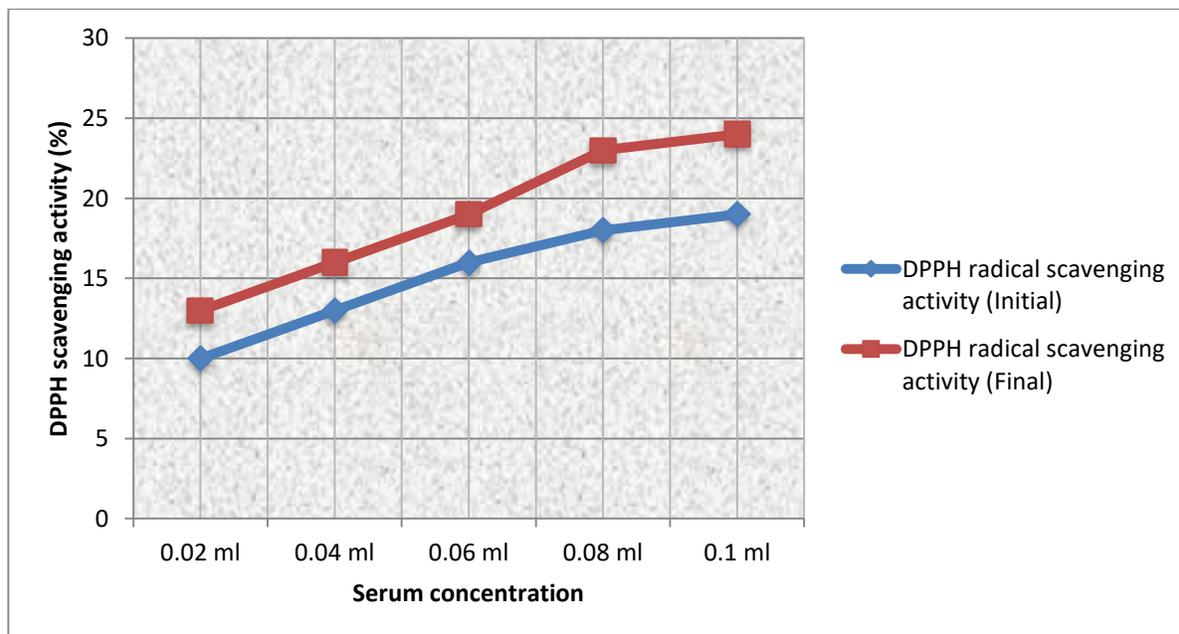


Fig.5: DPPH radical scavenging activity of subject E (before and after supplementation)

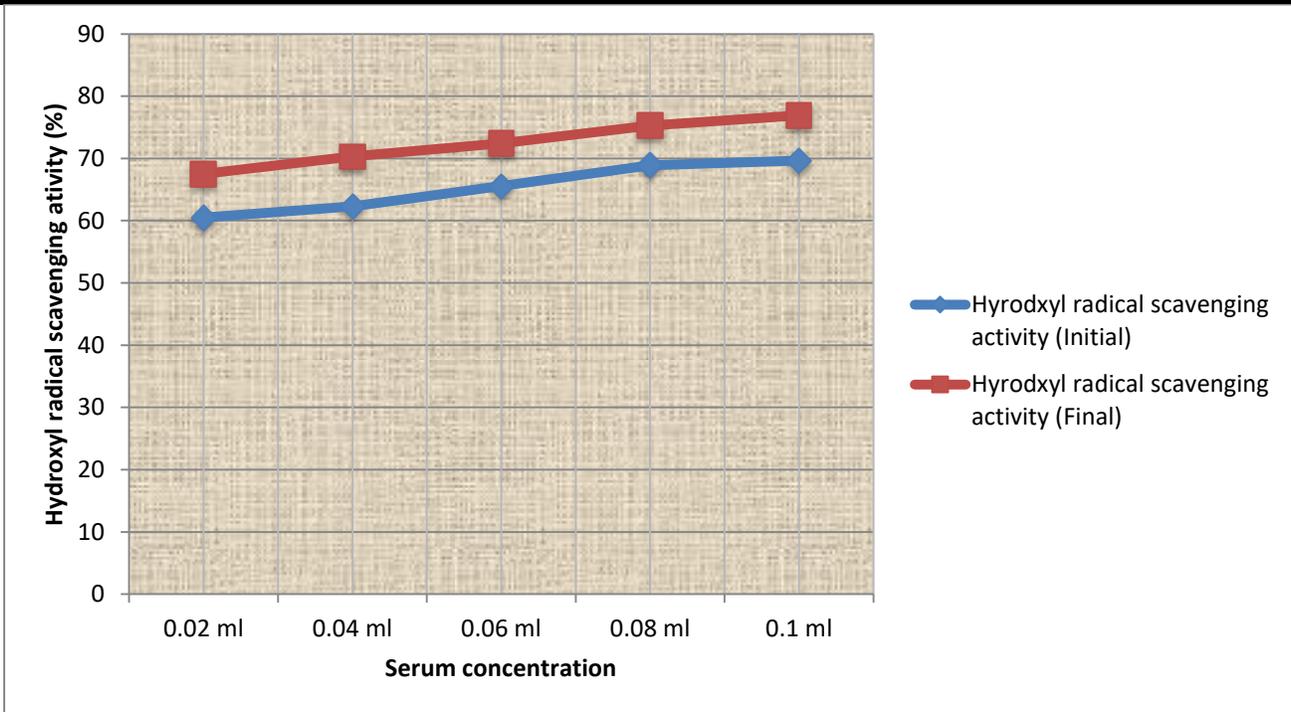


Fig.6: Hydroxyl radical scavenging activity of subject A (before and after supplementation)

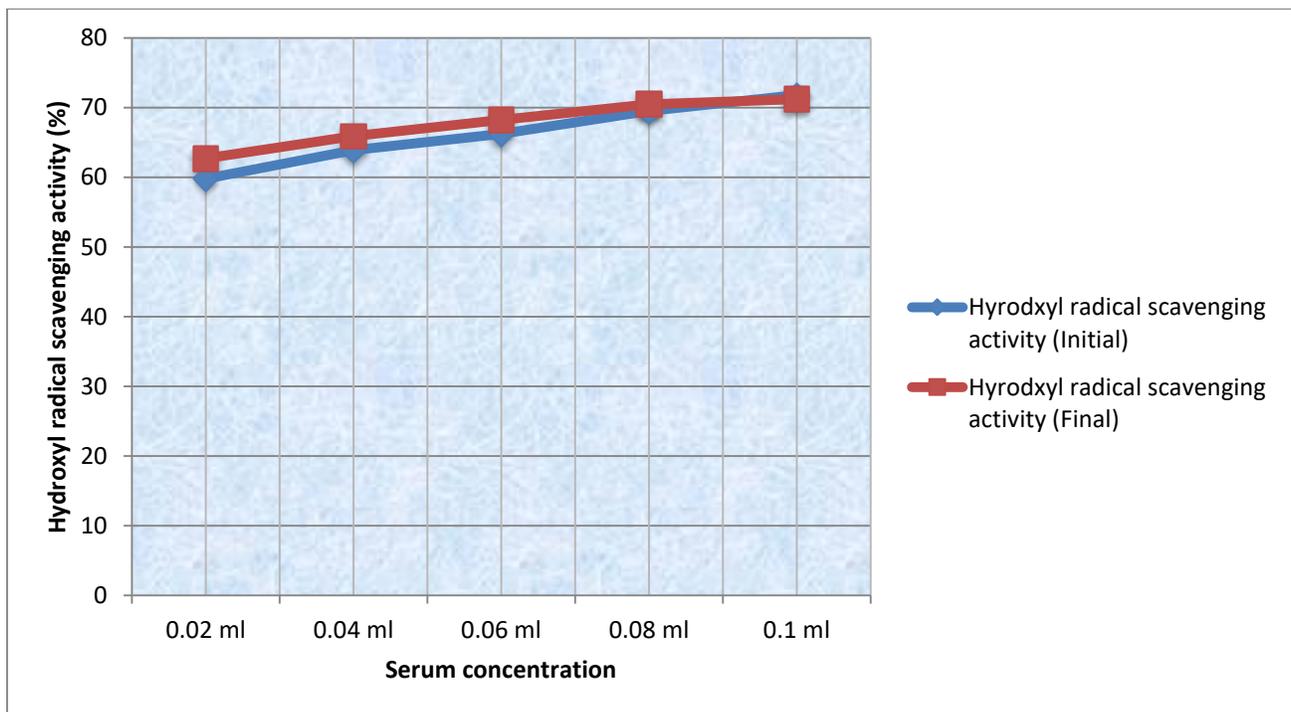


Fig.7: Hydroxyl radical scavenging activity of subject B (before and after supplementation)

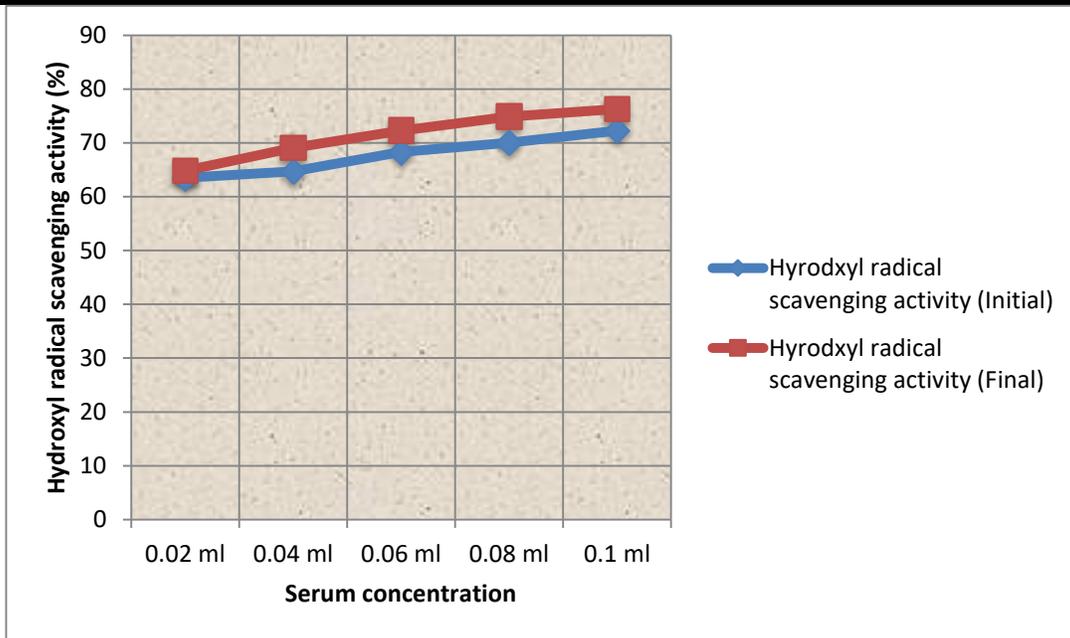


Fig.8: Hydroxyl radical scavenging activity of subject C (before and after supplementation)

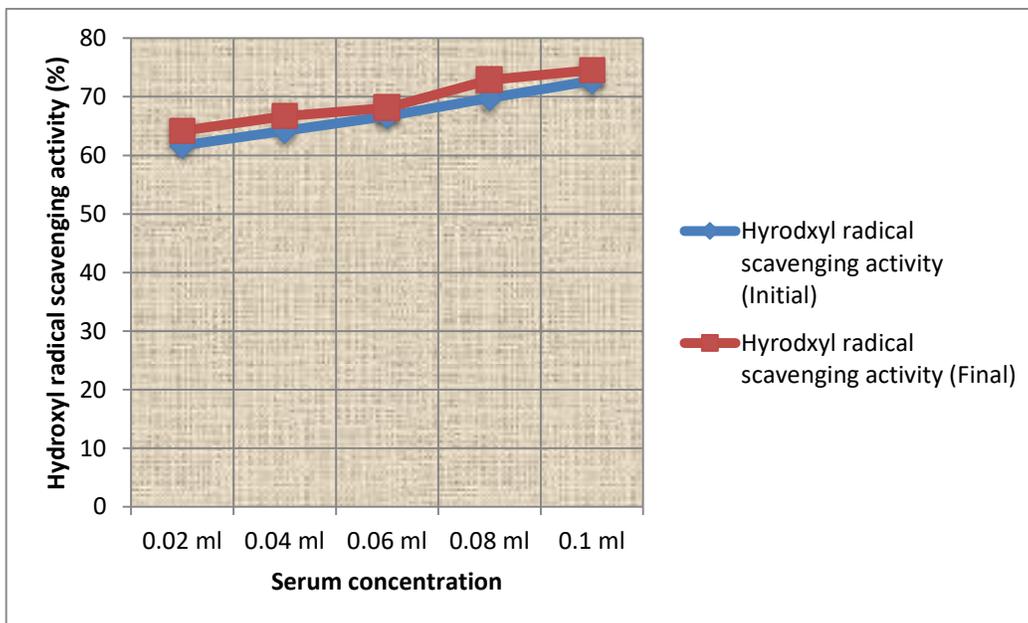


Fig. 9: Hydroxyl radical scavenging activity of subject D (before and after supplementation)

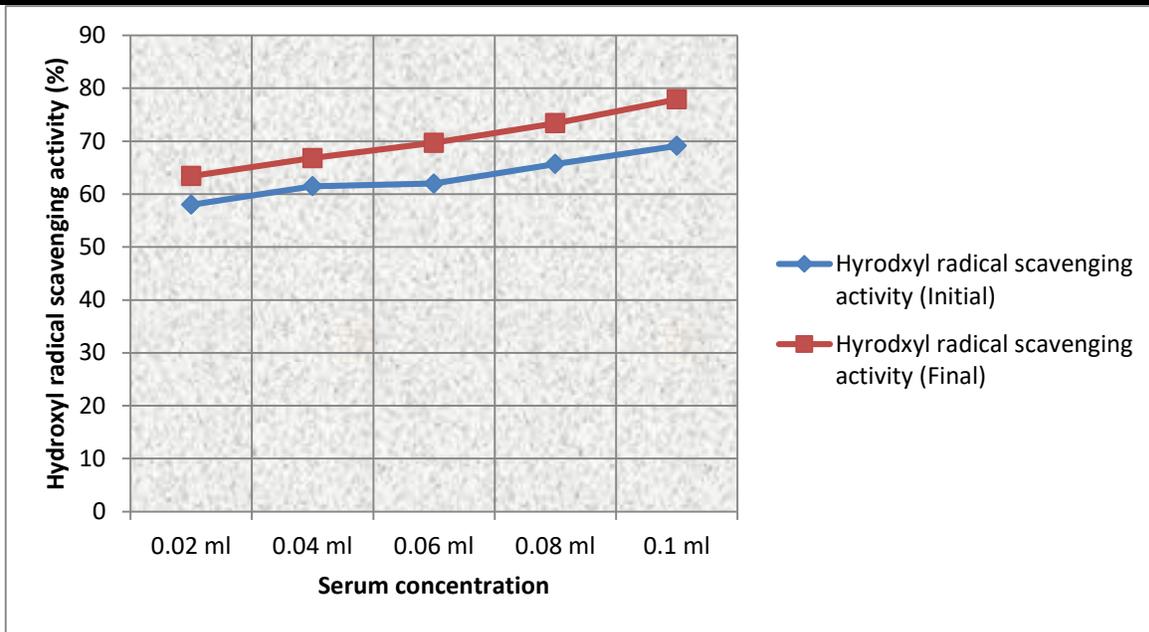


Fig.10: Hydroxyl radical scavenging activity of subject E (before and after supplementation)

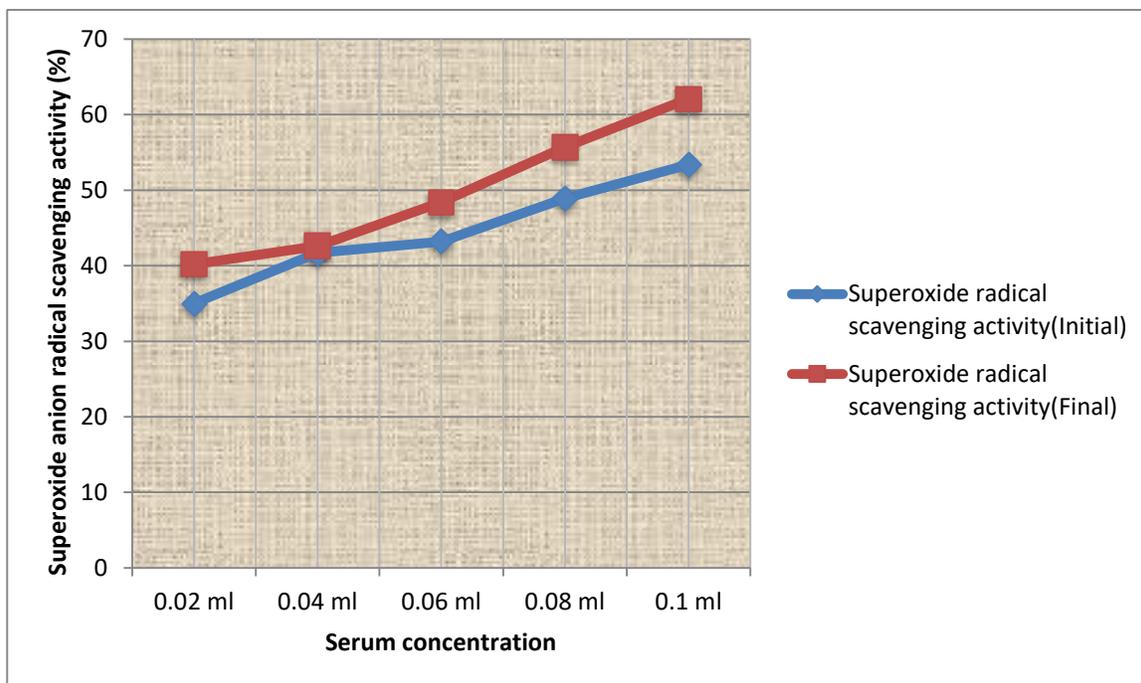


Fig.11: Superoxide anion radical scavenging activity of subject A (before and after supplementation)

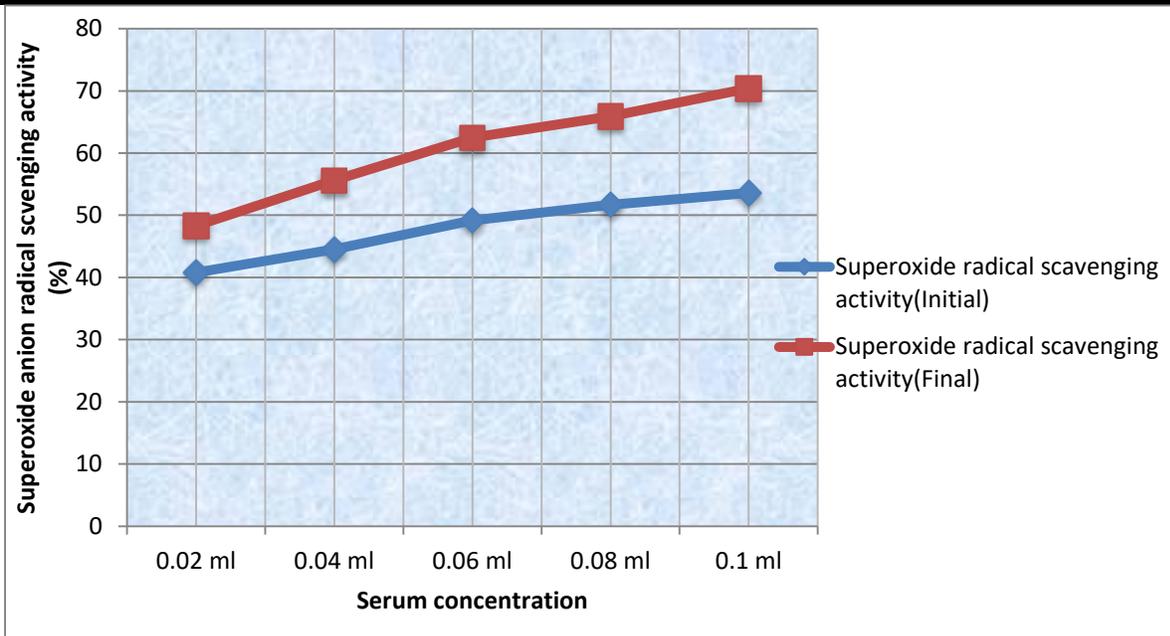


Fig. 12: Superoxide anion radical scavenging activity of subject B (before and after supplementation)

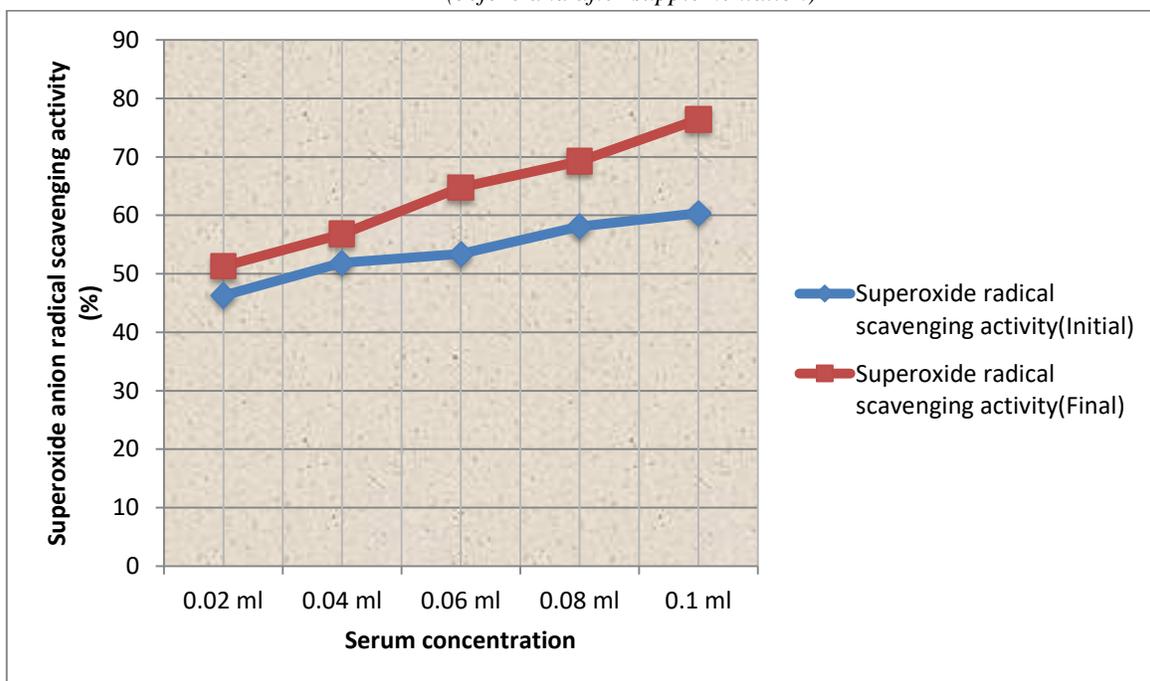


Fig.13: Superoxide anion radical scavenging activity of subject C (before and after supplementation)

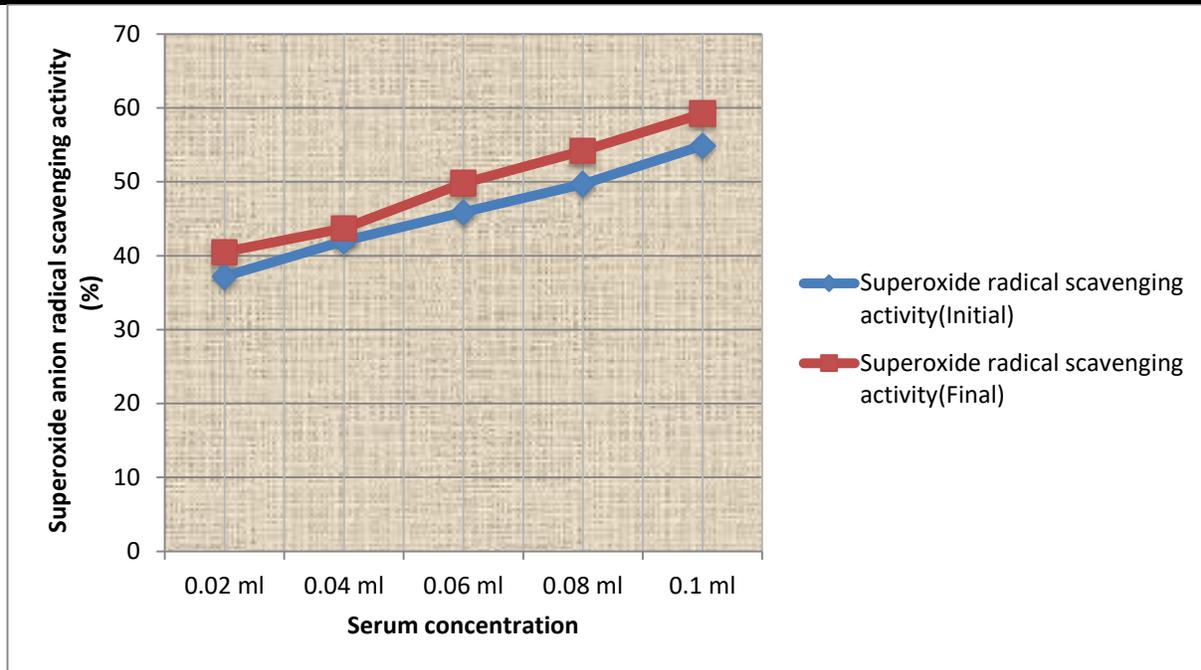


Fig. 14: Superoxide anion radical scavenging activity of subject D (before and after supplementation)

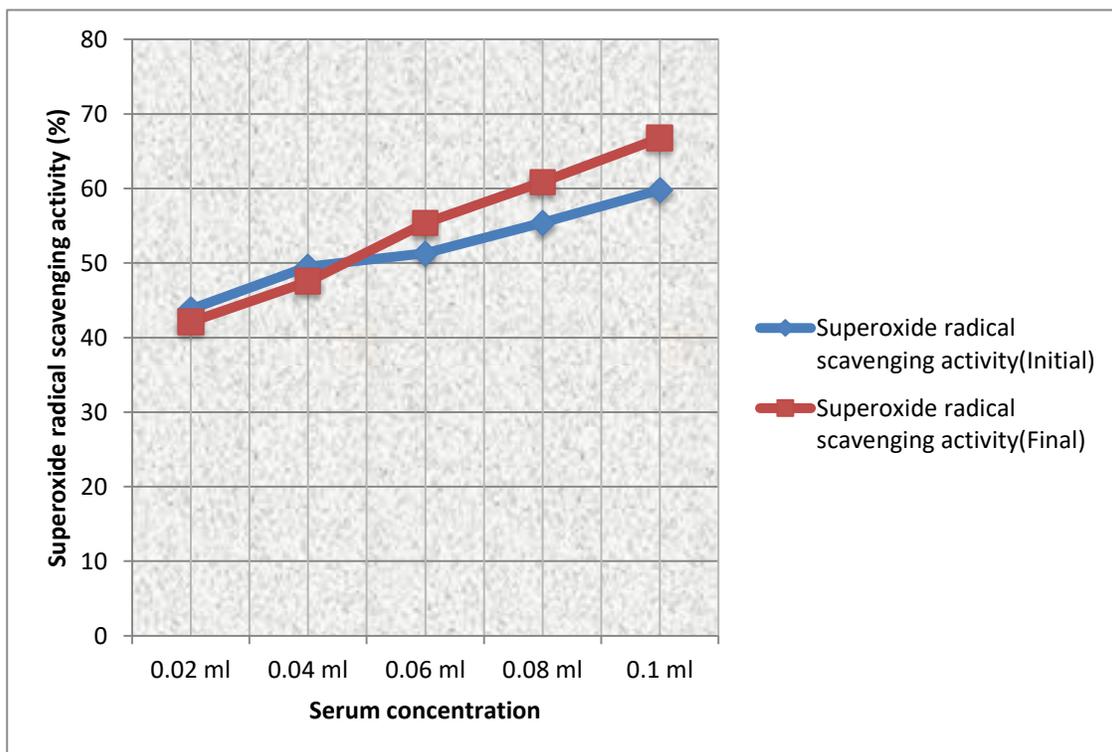


Fig. 15: Superoxide anion radical scavenging activity of subject E (before and after supplementation)

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