

Egyption Journal For Specialized Studies

Quarterly Published by Faculty of Specific Education, Ain Shams University



Board Chairman Prof. Osama El Saved Vice Board Chairman **Prof**. Dalia Hussein Fahmy Editor in Chief Dr. Eman Saved Ali Editorial Board **Prof. Mahmoud Ismail** Prof. Ajaj Selim **Prof. Mohammed Farag Prof. Mohammed Al-Alali Prof. Mohammed Al-Duwaihi Technical Editor** Dr. Ahmed M. Nageib Editorial Secretary **Dr. Mohammed Amer** Laila Ashraf **Usama Edward** Zeinab Wael **Mohammed Abd El-Salam**

<u>Correspondence:</u> Editor in Chief 365 Ramses St- Ain Shams University, Faculty of Specific Education **Tel**: 02/26844594 Web Site : https://ejos.journals.ekb.eg

Email : egyjournal@sedu.asu.edu.eg

ISBN : 1687 - 6164 ISNN : 4353 - 2682

Evaluation (July 2023) : (7) Point Arcif Analytics (Oct 2023) : (0.3881) VOL (12) N (43) P (3) July 2024

Advisory Committee

Prof. Ibrahim Nassar (Egypt) Professor of synthetic organic chemistry Faculty of Specific Education- Ain Shams University

Prof. Osama El Sayed (Egypt) Professor of Nutrition & Dean of Faculty of Specific Education- Ain Shams University

Prof. Etidal Hamdan (Kuwait) Professor of Music & Head of the Music Department The Higher Institute of Musical Arts – Kuwait

Prof. El-Sayed Bahnasy (Egypt) Professor of Mass Communication Faculty of Arts - Ain Shams University

Prof. Badr Al-Saleh (KSA) Professor of Educational Technology College of Education- King Saud University

Prof. Ramy Haddad (Jordan) Professor of Music Education & Dean of the College of Art and Design – University of Jordan

Prof. Rashid Al-Baghili (Kuwait) Professor of Music & Dean of The Higher Institute of Musical Arts – Kuwait

Prof. Sami Taya (Egypt) Professor of Mass Communication Faculty of Mass Communication - Cairo University

Prof. Suzan Al Qalini (Egypt) Professor of Mass Communication Faculty of Arts - Ain Shams University

Prof. Abdul Rahman Al-Shaer (KSA) Professor of Educational and Communication

Technology Naif University **Prof. Abdul Rahman Ghaleb** (UAE)

Professor of Curriculum and Instruction – Teaching Technologies – United Arab Emirates University

Prof. Omar Aqeel (KSA) Professor of Special Education & Dean of Community Service – College of Education King Khaild University

Prof. Nasser Al- Buraq (KSA) Professor of Media & Head od the Media Department at King Saud University

Prof. Nasser Baden (Iraq) Professor of Dramatic Music Techniques – College of Fine Arts – University of Basra

Prof. Carolin Wilson (Canada) Instructor at the Ontario institute for studies in education (OISE) at the university of Toronto and consultant to UNESCO

Prof. Nicos Souleles (Greece) Multimedia and graphic arts, faculty member, Cyprus, university technology

Management Information Systems a Decision Support System		ت المصرية.		المنعة الرئيسية	
ISSN-C السنه نقاط المجله	ISSN-P	اسم الجهه / الجامعة	اسم المجلة	القطاع	م
7 2023 2682-43	53 1687-6164	جامعة عين شمس، كلية التربية النوعية	المجلة المصرية للدراسات المتخصصة	Multidisciplinary عام	1
2023/10/8 التاريخ: 2023/10/8 التاريخ: 2023/177ARCIF التاريخ: المقابرة، مصر، قد نجعت في لي هذه المعايير يمكنكم الدخول هي النفة الوسطى، مع العلم أن في النسخة الورقية لمجلتكم إلى مشكورين. أ.د. سامي الخزندار 'ارسيف المالير	حد مبادرات قاعدة بيا ، كلية التربية النوعية .) معياراً، وللاطلاع ، ي ضمن الغنة (03) و ماعي، وكذلك الإشارة ميف "، التواصل معن	صصة المحترم العربية (ارسيف – ARCIF)، أ 2023. أن الصادرة عن جامعة عين شمس، ر العالمية، والتي يبلغ عددما (32 م). بر أو على مواقع التواصل الاجتم بة خاصة بنجاحكم في معامل " ان يقبول فائق الاحترام والتقدير	لمجلة المصرية للدراسات المتخ التربية النوعية، القاهرة، مصر شهادات المرجعية للمجلات العلمية لتقرير السنوي الثامن للمجلات للعام ارسيف Arcif المتوافقة مع المعاي <u>http://e-marefa.n</u> <u>http://e-marefa.n</u> <u>م</u> العلوم التربوية من إجمالي عدد الم فصص كان (0.511). ي مجلتكم. ي مجلتكم. و تفضلوا و تفضلوا	معادة أ. د. رئيس تحرير المعة عين شمس، كلية حية طيبة وبعد،،، مر معامل التأثير والاست الم معامل التأثير والاست نم معامل التأثير والاست نمايير اعتماد معامل ال معايير اعتماد معامل ال والم التالي: et/arcif(riteria/ معامل ارسيف لهذا الت معامل ارسيف لهذا ال الكم الإعلان عن هذه ال الكم الإعلان عن هذه ال معامل المناتيم. الم معامل المناتيم المحاصل الكم الإعلان عن هذه ال معامل المناتيم. الم معامل المناتيم المحاصل الكم الم الم معامل المائيم الم معامل الم معامل المائيم المائيم معامل الم معامل المائيم المائيم معامل الم معامل المائيم المائيم معامل الم معامل المائيم المائيم المائيم معامل الم معامل المائيم المائيم معامل المائيم المائيم المائيم المائيم المائيم معامل المائيم المائيم المائيم المائيم المائيم المائيم المائيم معامل المائيم المائيم المائيم المائيم المائيم المائيم المائيم المائيم المائيم معامل المائيم المائيم المائيم المائيم المائيم المائيم المائيم المائيم معامل المائيم ال	مع ج ج ت حقيق ويسرين ت تحقيق ويسرين ويامك متوسم ختام

Amman - Jordan 2351 Amman, 11953 Jordan 8





محتويات العدد

بحوث علمية محكمة باللغة العربية:

	تابع محتويات العدد	
	بحوث علمية محكمة باللغة الإنجليزية :	*
•	Dietary Importance of Spirulina and its Efficacy	
	Against sodium arsenite Toxicity in Rats	25
	Dr. Sara A.A. MAhmod	20

Dietary Importance of Spirulina and its Efficacy Against sodium arsenite Toxicity in Rats

Dr. Sara A.A. Mahmod ⁽¹⁾

⁽¹⁾ Lectuerer Nutrition and Food Science Department, Faculty of Home Economics, Helwan University

Dietary Importance of Spirulina and its Efficacy Against sodium arsenite Toxicity in Rats

Dr. Sara A. A. Mahmod

Abstract

This research examined the value of spirulina (sp) in diets and how well it protected rats from the damaging effects of arsenic. Thirty albino Sprague-Dawley strain rats weighing roughly 150±5 gm were randomly assigned into five main groups, six of each When equated with the +ve control group, the groups who were fed SP experienced greater gains in body weight than did the +ve group. When equated with the group serving as the positive control, levels of liver enzymes which include AST, ALT, as well as urea nitrogen and creatinine, reduced significantly in groups three, four and five, respectively. As a result, the spirulina powder had a positive influence on the toxic effects of sodium arsenite toxicity in rats.

Keywords: Spirulina, sodium arsinate toxicity, Antioxidant enzyme, liver and kidney function

ملخص: العنوان : أهمية المتناول من الاسبير ولينا وفاعليته ضد التسمم بالزئبق في الفئر ان المؤلفون: سارة عاطف على محمود يهدف هذا البحث إلى معرفة قيمة السبيرولينا في النظام الغذائي وتأثيره على حماية الفئر ان من الآثار الضارة للزرنيخ. تم استخدام ثلاثون فار من سلالة سبر أجو داولى يبلغ وزنها حوالي (150±5 جم) تم تقسيمها عشوائيًا إلى خمسُ مجموعات رئيسية، ستة فئر ان في كل منها أظهرت النتَّائج زيادة فيَّ زيادة وزن الجسم في المجموعات التي تم تغذيتها على السبير ولينا مقارنة مع المجموعة الضابطة الموجبة. إنزيمات الكبد مثل (ALT, AST) بالإضافة إلى نيتر وجين اليوريا والكرياتينين انخفضت بشكل ملحوظ في المجموعات (3، 4، 5) على التوالي مقارنة مع المجموعة الضابطة الموجبة ، توصلت هذه الدر اسة إلى استنتاج أن مسحوق السبير ولينا له تأثير إيجابي على الصحة من خلال تقليل التأثير ات السامة للمعادن الثقيلة، وخاصبة الزر نيخ.

الكلمات الدالة : سبير ولينا، سمية أرسينات الصوديوم، إنزيم مضادات الأكسدة، وظائف الكبد والكلي.

Introduction

Arsenic has the potential to spread widely throughout the plant and animal kingdoms by entering the food chain. Arsenic is a naturally and element that is not vital to life that has a glossy, grey appearance. The most poisonous types of arsenic are those that are inorganic (FAO and WHO, 1983). One of the most ubiquitous and dangerous metalloids in the environment is arsenic. Arsenic in its inorganic form poses a danger to millions of people throughout the world who consume tainted food and water. Metalloid arsenic (As) can be found in water and soil in both organic and inorganic forms (Khatun *et al.*, 2020 and Silbergeld *et al.*, 2008). A major public health issue involving millions of individuals is arsenic toxicity. Arsenic is distributed widely in the environment and comes from both natural and artificial sources (Amalendu *et al.*, 2014 & Erbanova *et al.*, 2008).

Spirulina (SP) is a non-toxic species of cyanobacteria and blue-green algae. It is a dietary supplement that includes important fatty acids, vitamins, and minerals. In alkaline waters in tropical regions, the Spirulina (SP) family naturally grows as microalgae and the nation's manufacturing output is always expanding. It is an excellent source of nourishment and has a high percentage of protein, vitamins (B6, A, B2, Vitamin B12 and E), enzymes, minerals (Ca, Fe, Mg, P, Cu, Zn, Mn, K, Na, Cr, and Se), essential amino acids, essential fatty acids, & other nutrients. It was selected as ((generally recognized) product by the FDA in 2011(Guldas *et al.*, 2022; Gutierrez-Salme *et al.*, 2015 and Kent *et al.*, 2015).

The Pacific Ocean in Japan and Hawaii, as well as sizable freshwater lakes in Africa, Mexico, South America and North America, are habitats where spirulina grows extensively. Spirulina (SP) maxima and Spirulina platensis are the two varieties of Spirulina that are most frequently utilised in dietary supplements (**Ghaeni**, & Roomiani, 2016; Khan *et al.*, 2005). Free-floating filamentous cyanobacteria known as SP are originate in tropical also subtropical lakes in Asia, Africa, Central America and South in addition Europe. They are distinguished by their Central America, and South America, in addition to Europe composed of several cells arranged in a left-handed open helix (**El-Sheekh** *et al.*, **2014; Vonshak 1997**). Spirulina has a number of therapeutic benefits, including lowering blood cholesterol, preventing some cancers, increasing intestinal lactobacilli, boosting the immune system, protecting against radiation, lowering hyperlipidemia, reducing nephrotoxicity from drugs and heavy metals, and reducing obesity (**Ghaeni** *et al.*, **2014; Jimenez** *et al.*, **2003**). The research aimed to determine the nutritional value of SP in the diet and its effectiveness in protecting rats from arsenic toxicity.



Materials and methods

Materials:

El-Gomhoria Company in Cairo, Egypt provided the minerals, cellulose, and arsenic that were purchased. The Imtenan Company sold spirulina, which was purchased. At the Helwan Farm for Experimental Animals in Cairo, Egypt, a total of thirty albino Sprague-Dawley strain rats that weighed approximately 150 ± 5 grammes each were acquired for the purpose of conducting experiments. Kits for conducting biochemical analysis were available for purchase from the Biodiagnostic Company for Pharmaceuticals and Chemicals in Dokki, Egypt. In Cairo,

Egypt, at the Morgan Chemical Factory, vitamins, casein, cellulose, minerals, flour and choline were purchased.

Experimental Animal Design:

After an adaptation period, thirty Sprague-Dawley albino rats, weighing 150 ± 5 grams, were randomly assigned into two main groups (6 rats). The 1st group being kept on the basal diet as the negative control (-ve), and the 2nd group24 rats being subjected to sodium arsenite exposure (10 mg/kg) daily in intake water for 6 weeks and divided into 4 groups. The first group was used as a positive control (+ve), while the other 3 were given basal diet with dried SP added at levels of 25, 50, and 75 mg/kg respectively for a period of six weeks.

The animals were housed in the Animal House of the Agricultural Research Center, Giza, Egypt, under hygienic conditions at a temperature of $25\pm2^{\circ}$ C with a moderate humidity of 56%. Ad libitum access to basic diet and water was provided.

The Basal Diet Preparation:

The basal diet was made in accordance with **Reeves** *et al.*, (1993). Twenty percent protein, 10 % sucrose, 4.7 % maize oil, 2 % choline chloride, 3.5 % mixture of salts, 1 % mixture of vitamins, and 5 % fibers makes up the ingredient list. Corn starch made up the remaining 1% to the full 100%.

Gain in overall body mass was computed at the conclusion of the trial (6 weeks). All rats were given a brief ether anaesthesia while they fasted overnight. Blood was drawn into a dry, plastic centrifuge tubes. After centrifuging blood samples, serum was extracted and kept at -20°C in a clean, well-sealed vial until analysis.

Biochemical Analysis:

Alanine (ALT) amino transeferase enzyme was identified in serum to the technique of *Sherwin (1984)*. Aspartate amino transeferase(AST) was identified as the enzyme along with *Young (1990)*. The serum alkaline phosphatase (ALP) level was measured consistent with the process designated by **Roy** (1970). The level of serum urea nitrogen was measured by the technique of *Fossati et al.*, (1980). Creatinine was evaluated consistent with the technique designated by *Henry* (1974). The concentration of serum arsnic was measured by the method of <u>Gora et al.</u>, (2014). Malondialdehyde (MDA) levels in the blood were measured according of *Draper & Hadly* (1990). The serum Catalase CAT activity in tissue homogenate was evaluated consistent with *Aebi* (1984). Superoxide Dismutase (SOD) was dignified as stated by *Nishikimi et al.*, (1972).

Statistical Analysis:

Mean and standard error were used to represent the results (SE). The Dunk'test multiple range post-hoc test was used to conduct the statistical analysis using SPSS, PC statistical software (Verion 18.0 SPSS Inc., Chicago, USA). One-way analysis of alteration was functioned to analyse the data (ANOVA). At P under 0.05, the values were considered substantially different (*Snedecor & Cochran, 1980*).

Results and Discussion

	Nutrients (100 g)		
	Energy	25kcal	
Nutrients	Protein	6 g	
(g)	Fat	0.45 g	
	Carbohydrate	3.5 g	
Minanala	Calcium	11 mg	
(ma)	Iron	2.99 mg	
(mg)	Sodium	100 mg	
Vitamins	Vitamin C	1mg	
(mg)	Vitamin A	60 IU	

 Table (1): Chemical Composition of 100 gm Spirulina

As shown in Table 1, aspirin contains 25 kcal of energy, 6g of protein, 3.5g of carbohydrate, 0.45 g of lipids, 11mg of calcium, 2.99mg of iron, 100 mg of sodium, 1 mg of vitamin C, and 60 IU of vitamin A. Based on the outcomes of the research performed by **Ali**, (2017), the lipids as well as the ash content of SP platensis were, respectively, 5.3 percent & 6.9 percent, which

were fewer than our findings. On the other hand, the protein and fibers content were, respectively, 61.8 & 9.5%, which were greater than our outcomes. In our research, Spirulina platensis revealed a higher percentage of protein in comparison to SP platensis obtained in Nomayos & Chad, which respectively accounting for 37.55% & 50.24% of the total protein content. According to the findings of our tests, the concentration of protein was much lower in comparison to that which was harvested in Switzerland & Burkina Faso, where it stretch to 65 as well as 61.3% respectively (Branger et al., 2003). This divergence raises the possibility that SP platensis' protein content shifts during the course of a cultivation cycle in response to the amount of photons (sunlight) available to the plant. (AFAA, 1982). The trial that create this variation was published in 1982. The amounts, quantity, in addition accessibility of the protein's amino acids are what determine the quality of the protein, which is why it is necessary to evaluate the quality of the protein in addition to the quantity.

Table (2): Effect of A spirulina on feed intake, body weight gain% and feed efficiency ratio of rats treated with sodium arsenite

Groups	Body weight Gain (%)	Feed Intake (g)	Feed Efficiency Ratio
Control (-Ve)	120.60±4.037c	15.06±0.321b	8.02±0.164c
Control (+Ve)	113.60±5.413d	12.40±1.056c	9.20±0.980a
Spirulina (25mg/kg) + sodium arsenite	127.00±2.121b	16.00±0.400b	7.96 ±0.297c
Spirulina (50mg/kg) + sodium arsenite	145.00±1.871a	16.44±0.699a	8.80±0.418b
Spirulina (75mg/kg) + sodium arsenite	149.20±8.167a	16.14±0.924a,b	9.26±0.747a

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p \leq 0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

Results in table (2) showed that body weight gain % (BWG), feed consumption was significantly decreased for the +ve

group as compared to with the -ve control group but feed Efficiency ratio was significantly rise for +ve group Compared to the -ve control group. Furthermore, body weight gain, (FI) for rats in groups three, four, besides five presented significant increase in body weight gain, FI when equated with the +ve control rats. The influence of the taste, odour, and deliciousness of the rats' food may be the cause of the increases in BWG and feed intake along with an upsurge in the levels of dehydrated SP.

During the whole study period, the treated rats did not exhibit any clinical symptoms or lesions, however most of the groups did notice a minor rise in body weight. When compared to rats given arsenic with spirulina (T2), animals in the arsenictreated group (T1) had lower body weights, but these differences were not very significant. In light of this, it may be concluded that feeding rats NaAsO2 (4 mg/kg BW) will not result in chronic arsenic toxicity, however it may take longer or involve a larger dose of As for symptoms to appearing (**Hasan** *et al.*, 2015).

 Table (3): Protective effect of aspirulina on weight of some organs in rats treated with sodium arsnic.

Parameters	Liver	Kidney
Groups	(%)	(%)
Control (-Ve)	8.24 ±0.207c	$7.96 \pm .451b$
Control (+Ve)	9.26±0.611a	8.88±.597a
aspirulina (25mg/kg) + sodium arsenite	9.18±0.192a	8.10±.158a
aspirulina (50mg/kg) + sodium arsenite	8.96±0.391b	8.24±.114a
aspirulina (75mg/kg) + sodium arsenite	8.80±0.339b	7.96±.358b

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p \leq 0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

Outcomes in table (3) presented that organs weight/ body weight % (liver & kidney) was significantly enlarged for the +ve control group equated with the -ve control one. Additionally, organs weight (liver & kidney) for rats in groups 3,4,5 respectively exposed significant decreased in liver, kidney when compared with the positive control rats.

When contrasted with the normal control group, the arsenic control group demonstrated a statistically significant and apparently decreased end body weight alongside an increased liver weight. When compared to the arsenic control group, the SP platensis- treated group indicated a significant increase in both the final body weight and liver weight. This was because arsenic toxicity was not present in the Spirulina platensis- treated group. When compared to those of the arsenic control group, the ultimate body and liver weights of the SP platensis/arsenic-treated group did not significantly vary from those of the arsenic group. In this study, the exposure to arsenic had no impact on the weight of the kidneys. When compared to the arsenic control group, the SP platensis- treated group demonstrated a significant increase in both the final body weight & liver weight. This was caused by the absence of any adverse effects from arsenic. However, the final body besides liver masses were unaffected. In contrast, there was no alteration between the SP platensis/arsenic-treated group & the arsenic control group in terms of the ultimate weights of the body and liver. Arsenic exposure in this experiment had no effect on kidney weight. (Bashandy et al., 2018).

Table (4): Protective effect of aspirulina on liver enzymes in
rats treated with sodium arsnic.

Parame	eters	ALT	AST	ALP
Groups		(U/L)		
Control (-Ve)		39.40a±1.673d	49.40a±2.881d	843.60±0.817c
Control (+Ve)		72.00±2.121a	109.00±5.390a	861.20±2.683a
Spirulina (25mg/kg) + sod arsenite	lium	57.40±1.817b	60.20±2.864b	853.80±2.588b
Spirulina (50mg/kg) + sod arsenite	lium	53.40±1.140b	55.40±1.140c	849.20±1.789b
Spirulina (75mg/kg) + sod arsenite	lium	44.60±1.140c	53.80±1.304c	847.80±1.304b,c

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p \leq 0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

Table 4 shows the impact of SP powder on serum Alanine Aminotransferase serum activity. The +ve group exposed to

Arsnic toxicity had higher ALT activity, as compared to the negative control group $(72.00\pm2.121 \text{ vs. } 39.40\pm1.673 \text{ u/l})$ while the Spirulina-fed groups showed significant reductions, as compared to the positive control group. Group 5 rats fed SP powder had the lowest ALT activity with a mean value of $44.60\pm1.140 \text{ u/l}$.

Table (3) also indicated the impact of Spirulina powder on activity of AST. serum levels The of Aspartate the Aminotransferase in the group of rats subjected to Arsnic toxicity (positive control group) were considerably higher (109.00±5.390 u/l) than those in the group of rats not exposed to Arsnic toxicity $(49.40\pm2.881 \text{ u/l})$. When compared to the positive control group, the outcomes established that the serum activity of Aspartate Aminotransferase was decreased in the rats fed a diet containing SP powder, regardless of the amount consumed. Serum AST activity was observed to be 53.80±1.304u/I lower in the group of rats administered Spirulina powder (group 5).

Also, Table (4) presented the influence of Spirulina powder on Alkaline Phosphatase. When rats were subjected to arsnic toxicity, the activity of ALP in serum elevated considerably, with a mean value of 861.20±2.683 u/L in comparison to the negative control group's value of 843.60±1.817 u/L. The distinction was statistically When contrasted with the positive control $(861.20\pm 2.683 \text{ U/L})$, the results demonstrated that the mean value of Alkaline Phosphatase in serum decreased significantly in a group of mice nurtured on SP powder at three levels in diet as 853.80±2.588U/L, value 849.20±1.789 U/L. & mean 847.80±1.304 U/L, respectively. These values were found to be significantly lesser than the positive control's value.

The data in this table revealed that the mean values of AST, ALT and ALP increased in the positive control group compared to the negative control group, while treating rats with spirulina at the three levels improved liver enzymes.

When rats received both SP and arsenic, these metrics improved. This rise is a sign of cellular absence & a breakdown in the integrity of liver cell membranes' functional integrity. The liver is a significant target organ of toxicity owing to its distinct metabolic processes. This toxicity is known to be linked with oxidative stress. (Korany *et al.*, 2019; Chowdhury *et al.*, 2016).

biochemical analysis in our trial revealed a The considerable increase in liver enzymes (AST & Alanine Aminotransferase), & these findings were comparable with those of previous research (Ahmed et al., 2019; Flora et al., 1997; Saved et al., 2015). The precise process causing the increase of these enzymes is still unknown. According to several researchers, this impact could be brought on by cellular injury or improved plasma membrane absorbency. In addition, other variables could be at play, like elevated synthesis or diminished enzyme degradation (Ahmed et al., 2019; Sayed et al., 2015). In contrast, the co-administration of Sp in the current study considerably reduced the levels of aspartate aminotransferase and ALT to levels that were close to control levels. This study also shown that A- treatment caused a noticeable rise in serum creatinine levels, indicating impaired renal function.

Table (5): Protective Effect of Spirulina on Serum KidneyFunctions of Rats with sodium arsenite

Parameters	Urea	Creatinine	Uric acid
Groups		mg/dl	
Control (-Ve)	44.80±1.483c	0.78±0.110c	4.02±0.110c
Control (+Ve)	54.00±2.915a	1.54 ±0.134a	5.54±0.114a
Spirulina (25mg/kg) + sodium arsenite	$52.40 \pm 2.074a$	1.32±0.130a	4.18 ±0.084b
Spirulina (50mg/kg) + sodium arsenite	49.00±0.707b	1.16±0.055b	4.38±0.084b
Spirulina (75mg/kg) + sodium arsenite	45.80±0.837c	1.12±0.084b	4.14±0.261b, c

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p \leq 0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

Table (5) demonstrated the impact that Spirulina powder has on kidney functioning as measured by serum concentrations of creatinine, urea nitrogen, and uric acid. The concentration of serum levels of urea nitrogen was dramatically raised with a mean value of 54.00 ± 2.915 mg/dl when rodents were subjected to arsnic toxicity. This was in comparison to the -ve control group, which had levels that were 44.80 ± 1.483 mg/dl. When compared to the positive control group, which had a serum level of urea nitrogen that was 54.00 ± 2.915 mg/dl, groups 3, 4, and 5 had concentrations of urea nitrogen in the serum that ranged from 52.40 ± 2.074 mg/dl to 45.80 ± 0.837 mg/dl, respectively.

Table (5) Relative to the negative control group, which had a creatinine concentration that was 0.78 ± 0.110 mg/dl on average, the +ve group, which had been exposed to arsenic toxicity, had a concentration of creatinine that was 1.54 ± 1.34 mg/dl higher than the negative control group. When compared with the positive control group, the groups of rats who were fed Spirulina powder at whatever amount of intake demonstrated a lower concentration of creatinine in their serum. The optimal level was determined in group 5, with a mean value of 1.12 ± 0.84 mg/dl; this group is considered to be superior to the negative control group.

In addition, table (5) exhibited the reduction of uric acid by SP powder. Rats supplementing their diet with spirulina powder had lower serum uric acid activity compared to the positive control group at all doses. Group 5 rats administered SP powder had the lowest serum uric acid activity, with a mean value of 4.14 ± 0.261 mg/dl.

Similarly, arsnic-treated rats showed considerably increased levels of oxidative stress and kidney arsenic buildup; however, resveratrol pretreatment prevented these alterations from occurring. Additionally, creatinine, the blood urea nitrogen, in addition to renal tubular epithelial cell necrosis levels were lower before starting resveratrol therapy. The reactivity of arsenic, which results in the production of ROS and compounds containing sulfur, is connected to its toxicity. (**Zhang** *et al.*, **2014**; **Hughes** *et al.*, **2011**; **Lu** *et al.*, **2012**). ROS, which are implicated in the pathogenic mechanisms of diseases, are particularly prone to causing harm to the kidney (**Zhang** *et al.*, **2014**; **Rodrigo and** **Rivera**, **2002**). Our investigation also confirmed that arsenicinduced kidney oxidative damage was caused by ROS.

Spirulina platensis indicated a significant positive improvement in both kidney & liver functions through its modulation of the oxidative stress parameters in addition to the inflammatory cytokines, regardless of whether it was given alone (to ascertain any problems) or in combination with sodium arsenite intoxicated rats. This improvement appeared through its modulation of the oxidative stress parameters as well as the inflammatory cytokines. The purpose of our experimental investigation is to enhance atherosclerotic indices, & this improvement in the previously indicated parameters, in particular the levels of uric acid and IL-6, reflected in the much lower level of MCP-1. SP may be employed as an inexpensive, safe, and natural adjuvant therapy for atherosclerosis in the future rather than expensive, traditional medication therapy with several adverse effects. (Bashandy et al., 2018; Jensen et al., 2016).

Table (6): Influence of Spirulina on Serum Lipid Profile ofRats with sodium arsenite

Parameters	TC	TG	HDL-c	LDL-c	VLDL-c	
Groups		(mg/dl)				
Control (-Ve)	141.60±8.678d	106.80±1.789d	38.40±1.517c	81.84±9.910b	21.36±0.358c	
Control (+Ve)	254.20±7.855a	193.80±1.789a	53.00±2.828a	162.44±8.764a	38.76±0.358a	
Spirulina (25mg/kg) + sodium arsenite	151.20±1.643b	139.80±1.483b	48.60±1.517b	74.64±2.613c	27.96±0.297b	
Spirulina (50mg/kg) + sodium arsenite	146.20±1.643c	134.60±2.702b	45.80±0.837b	73.48±2.492c	26.92±0.540b	
Spirulina (75mg/kg) + sodium arsenite	144.20±4.438c	113.60±3.362c	40.80±0.837c	80.68±4.793b	22.72±0.672c	

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p≤0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

As stated in table (6), the levels of total cholesterol, total fat, very low-density lipoprotein cholesterol, in addition to low density lipoprotein cholesterol all amplified significantly in the positive control group, with P values more than 0.05. However, as compared to the group serving as a positive control, the HDL-C serum level was statistically significantly reduced (P above 0.05).

The mean values of the serum lipid profile exhibited a statistically significant drop (P over 0.05) after SP treatment. On the other hand, when compared to the control group with the negative result, the serum HDL-C level enlarged significantly (P above 0.05). As opposed to the positive control group, the lipid profile of the arsnic-toxic rats that were administered spirulina displayed significant improvement. These alterations point to the significance of consuming a diet rich in SP either for the treatment of arsnic toxicity or the prevention of its occurrence.

Spirulina's hypolipemic effects were also demonstrated in mice that had been given alloxan (250 mg/kg body weight), which was used to experimentally cause diabetes. Hepatic triacylglycerols decreased after receiving 5% SP. Improvement in blood HDL levels and decreases in serum LDL and VLDL were also seen. Li et al discovered that when fed a high-fat diet, spirulina for 8 weeks increased HDL-C and decreased LDL-C, TG, and TC levels (**DiNicolantonio** *et al.*, 2020 and Li *et al.*, 2018). Similar to prior trials, it was also demonstrated to normalise hepatic steatosis with enhancements in liver function tests, including free fatty acids, transaminases & total lipid profile.

The outcomes were distinguished by a marked decrease in TC during the first four weeks of SP supplementation, which recovered to baseline once it was stopped. These TC alterations were proportionate to the levels of serum and dietary TC. No adjustments were made to TG, HDL, or body weight (**DiNicolantonio** *et al.*, 2020; Nakaya *et al.*, 1988). Spirulina supplementation at a dose of 2 to 8 grammes per day has been shown to enhance lipid profiles, especially by lowering TG, TC, & LDL-c in addition to raising HDL-c; increasing apolipoprotein A1 and decreasing apolipoprotein B; and promoting weight reduction and lowering BMI.

Table (7): Protective Effect of Spirulina on SerumMalondialdehyde and Glutathione of Rats with sodium
arsenite

	Parameters	Malondialdehyde	Glutathione
Groups		(ng/m	l)
Control (-Ve)		67.60±2.074d	4.82±0.192a
Control (+Ve)		158.80±3.271a	3.32±0.148b
Spirulina (25mg/kg) + sodium arsenite		97.40±1.673b	4.16±0.089a
Spirulina (50mg/kg) + sodium arsenite		83.80±2.387c	3.34±0.207b
Spirulina (75mg/kg) + sodium arsenite		87.40±1.949c	3.44±0.114b

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p \leq 0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

Outcomes in Table (7) revealed the effect that varying doses of SP powder had on serum levels of antioxidant activity like malonaldehyde & glutathione. Results indicate that there was a significant reduction in the level of serum glutathion in the positive control group, compared to the negative control group with a man value of 3.32 ± 0.148 (ng/ml) and 4.82 ± 1.92 (ng/ml). On the other hand, the results showed that the concentrations of malondahyde in serum increased substantially when contrasted with the positive control. On the other hand, the findings demonstrated that the level of malondialdehyde was much higher in the group that served as the positive control as opposed to the group that served as the negative control. When compared with the group serving as the positive control, the serum MDA levels in groups 3, 4, and 5 were shown to have reduced significantly (P under 0.05). Additionally, the highest value of MDA was found in group 4, which was observed.

The blood MDA levels were elevated in the current investigation by As feeding and reached their maximum levels in the experimental groups, increasing the trend from the beginning to the completion of the trial. This finding suggests that As feeding increased lipid peroxidation in the treated rat, indicating an increase in the formation of free radicals. Our results support a prior study that found elevated blood levels of lipid peroxidation to be related to high as exposure. In comparison to the control and As-treatment groups, serum MDA levels were dramatically reduced by both the (agro-based spirulina) Ab-Sp and (commercially accessible spirulina) Com-Sp treatments. This result demonstrates that lipid peroxidation was reduced during the experiment and reverted to the control level with Ab-Sp and Com-Sp treatments alone (**Khair** *et al.*, 2021; Hu *et al.*, 2020).

As a result of therapy, considerably higher amounts of MDA have been identified in blood in contrast to the control group, while co-treatment with SP platensis resulted in lower MDA levels. Rats exposed to as had lower blood GSH levels, and when Spirulina platensis was also administered, the serum antioxidant capacity rose (GSH). MDA concentrations may rise as a result of arsenic. Because of the increased creation of ROS and oxidative stress brought on by the lipid peroxidation of membranes, arsenic exposure speeds up the destruction of phospholipids, which in turn causes cellular degeneration. Increased ROS production and oxidative stress, which can lead to an increase in glutathione consumption, may be the cause of the glutathione level being depleted following exposure to arsenic. As the original line of antioxidant resistance against arsenic-induced damage, glutathione has been demonstrated to be a major indicator of oxidative stress. Reduced tissue GSH levels signify oxidative injury (Reda et al., 2019; Ahangarpour et al., 2018).

Table (8): Protective Effect of Spirulina on Different LevelsSerum Concentrations of arsnic Intoxicated Rats.

Groups	Parameters	Arsnic blood (ug)
Control (-Ve)		0.68±0.17e
Control (+Ve)		5.38±0.249a
Spirulina (25mg/kg) + sodium arsenite		3.26±0.182b
Spirulina (50mg/kg) + sodium arsenite		2.46±0.089c
Spirulina (75mg/kg) + sodium arsenite		1.88±0.148d

The data were presented as mean \pm S.E.

(* *) = highly significant at p< 0.01, (*) = significan a p≤0.05, (-) = Non significant p>0.05.

Acco ding o "D ncan" he mean val es we e a anged in a descending o de f om "a": "d"

When contrasted to the negative control group, the findings indicated that the blood level of arsnic was significantly elevated when rats were subjected to arsnic toxicity (positive control group), with a mean value of 5.38 ± 0.249 ug. This was in contrast to the negative control group, which had a value of 0.68 ± 0.17 ug. When compared to the positive control group, the arsnic concentration in the serum of rats that were administered aspirina at varying levels of intake resulted in a substantial drop across all groups, with a mean value of 3.26 ± 0.182 ug; 2.46 ± 0.089 ug; and 1.88 ± 0.148 ug, respectively.

Arsenic feeding to the rats increased As loads in the lung, liver and kidney and caused hepatic and renal insufficiencies; and separate treatments of SP and selenium significantly reduced tissue loads and recovered the hepatic and renal insufficiencies. Researchers from a variety of institutions (Balaji et al., 2014) looked at the bioremediation capability of SP as it relates to the deletion of heavy metal ions from industrial effluents. According to Ghosh et al., (2014) research, giving goat's arsenic while also giving them spirulina appears to offer a protective mechanism against the toxicity caused by the arsenic. Based on the results of the research, Hossain et al., (2013), using SP in conjunction with vitamin A as a treatment for chronic arsenicosis in rats is an effective strategy. According to Vedi M. et al., (2013) research, it also has a protective influence contrary to galactosamine-induced hepatotoxicity in mice. In arsenic-induced toxicities in ducks, spirulina has been presented to be advantageous for toxic symptoms as well as body weight & hematological parameters (Islam et al., 2009).

Reference

- Aebi, H. (1984): Catalase in vitro. In: Methods of Enzymology; 105: 121–126.
- Ahangarpour, A.; Zeidooni, L.; Samimi, A.; Alboghobeish, S. and Khorsandi, S. (2018): Chronic exposure to arsenic and high fat diet additively induced cardiotoxicity in male mice. Res Pharmaceut Sci.; 13(1): 47-56.
- Ahmed, A.; Korany, S.; Halawany, A. and Ahmed, S. (2019): *Spirulina platensis* alleviates arsenic-induced toxicity in male rats: biochemical, histopathological and immune-histochemical studies. Adv Anim Vet Sci.; 7(8): 701-710.
- Ali, O. (2017): Effects of dietary supplementing of spirulina platensis and chlorella vulgaris microalgae on hematologic parameters. J Pediatri Hematol Oncol. 7(3):163-9.
- Amalendu, G.; Abdul, A.; Ali, K.; Shahi, A.; Shahi, G.; Shafiqul, I. and Mahfuzul, B. (2014): Effects of spirulina in arsenic poisoning in the Black Bengal goat. Turkish Journal of Veterinary & Animal Sciences.; 38(1): 11.
- Association française pour l'algologie appliquée (AFAA). (1982): Actes du premier symposium sur la spiruline Spirulina platensis(Gom.) Geitler de l'AFAA.
- Balaji, S.; Kalaivani, T. and Rajasekaran, C. (2014): Biosorption of Zinc and nickel and its effect on growth on different Spirulina species, clean-Soil Air Water 42: 507-512.
- **Bashandy, S.; Amin, M. and Morsy, F. (2018):** Spirulina platensis reduced liver and kidney injuries induced by Sodium arsenite. International Journal of PharmTech Research.; 11(1):35-48.
- Branger, B.; Cadudal, J.; Delobel, M.; Ouoba, H.; Yameogo, P. and Ouedraogo, D. (2003): Spiruline as a food supplement in case of infant malnutrition in Burkina-Faso. Archives de pediatrie: organe officiel de la Societe francaise de pediatrie. 10(5):424-31
- Chowdhury, U.; Islam, S.; Akter, R.; Khaleda, L. and Rahman, Z. (2016): A study on the effect of arsenic on tissue histology and its deposition pattern in various organs of Wistar albino rat. European J Pharmaceut Medic Res.; 3(5): 580-587.
- **DiNicolantonio, J.; Bhat, G. and OKeefe, J. (2020):** Effects of spirulina on weight loss and blood lipids a review. *Open Heart.*; 7(1003): 2018-001003.
- Draper, H. and Hadley, M. (1990): Malondialdehyde determination as index of lipid peroxidation. Methods Enzymol. 186, 421-431.

- El-Sheekh, M.; Hamad, S. and Gomaa, M. (2014): Protective Effects of Spirulina on the Liver Function and Hyperlipidemia of Rats and Human. Braz. Arch. Biol.; 57 (1): 77-86.
- Erbanova, L.; Novak, M.; Fottova, D. and Dousova, B. (2008): Export of arsenic from forested catchments under easing atmospheric pollution. Environ Sci Technol.; 42: 7187–7192.
- Flora, J.; Pant, C.; Malhotra, R. and Kannan, M. (1997): Biochemical and histopathological changes in arsenic-intoxicated rats coexposed to ethanol Alcohol. 14(6):563-568.
- Food and Agricultural Organization (FAO) and World Health Organization (WHO). (1983). WHO Food Addit, Ser. 18.
- Fossati, P.; Prencipe, L. and Berti, G. (1980): Enzymatic colorimetric method of determination of urea in serum. Clin. Chem. 6(18) 499-502.
- Islam, M.; Awal, M.; Mostofa, M.; Begum, F.; Khair, A. and Myenuddin, M. (2009): Effect of spirulina on toxic signs, body weight and hematological parameters in arsenic induced toxicities in ducks. Int. J. Poult. Sci.,8(1): 75-79.
- Ghaeni, M. and Roomiani, L. (2016): Review for Application and Medicine Effects of Spirulina, Spirulina platensis Microalgae. Journal of Advanced Agricultural Technologies.; Vol. 3, No. 2.
- Ghaeni, M.; Roomiani, L. and Masomozadeh, Z. (2014): Review for uses and therapeutic effects of spirulina, Spirulina platensis microalgae. *Global Journal on Advances in Pure and Applied Sciences [Online].*; 4: 365-371.
- Ghosh, A.; Abdul, A.; Khan, A.; Alam, G.; Islam, S. and Bari, A. (2014): Effects of spirulina in arsenic poisoning in the Black Bengal goat. Turkish Journal of Veterinary and Animal Sciences, 38(1): 63-72
- <u>GORA</u>, R.; <u>BAXLA</u>, S.; <u>KERKETTA</u>, P.; <u>PATNAIK</u>, S. AND <u>ROY</u>, B.(2014): HEPATOPROTECTIVE ACTIVITY OF TEPHROSIA PURPUREA AGAINST ARSENIC INDUCED TOXICITY IN RATS.; 46(2): 197-200.
- Guldas, M.; Gurbuz, O.; Cakmak, B. and Yildiz, E. (2022): Effects of honey enrichment with Spirulina platensis on phenolics, bio-accessibility, antioxidant capacity and fatty acids. LWT - Food Science and Technology.; 153: 112-461.
- Gutierrez-Salme, G.; Fabila-Castillo, L. and Chamorro-Cevallos, G. (2015): Nutritional and toxicological aspects of Spirulina (Arthrospira). Nutricion Hospitalaria.; 32(1): 34-40.

- Hasan , M.; Awal, M.; Ara, A.; Rashid, M.; Azam, M. and Ali, M. (2015): Study on Prevention of Induced Arsenic Toxicity in Rats by using Spirulina and Thankuni. Scholars Journal of Applied Medical Sciences (SJAMS).; 3(2D):828-832.
- Henry, R. (1974): Creatinine measurement with colorimetric method. In clinical Chem., Principles and technics. Second edition, Haper and Row publishers. hepatocytes. Cancer Lett, 97: 61-67.
- Hossain, F.; Hossain, M.; Kabir, M. and Fasina, F. (2013): Effectiveness of combined treatment using Spirulina and vitamin A against chronic arsenicosis in rats. 7(20): 1260-1266.
- Hughes, F.; Beck, D.; Chen, Y.; Lewis, S. and Thomas, J. (2011): Arsenic exposure and toxicology a historical perspective. Toxicol Sci.; 123:305-32.
- Hu, Y.; Li, J.; Lou, B.; Wu, R.; Wang, G. and Lu, C. (2020): The role of reactive oxygen species in arsenic toxicity. Bio molecules.; 10(2):240.
- Jensen, S.; Drapeau, C.; Lenninger, M. and Benson, F. (2016): (Spirulina) platensis Results from a Randomized, Double-Blind, Placebo-Controlled Study with a Focus on Anticoagulant Activity and Platelet Activation. *Journal of Medicinal Food*.;19(7): 645–653.
- Jimenez, C.; Cossio, B.; Labella, D. and Xavier Niell, F. (2003): The feasibility of industrial production of spirulina in southern Spain, Aquaculture.; 217: 179-190.
- Kent, M.; Welladsen, M.; Mangott, A. and Li, Y. (2015): Nutritional evaluation of Australian microalgae as potential human health supplements. PloS One.;10(2): 118-985.
- Khan, M.; Varadhara, S.; Gansesa, L.; Shobha, J.; Naidu, M. and Parmandi, N. (2005): "C-Phycocyanin protects against ischemia- reperfusion injury of heart through involvement of p38 and ERK signaling. Am. J. Physiol. Heart Circ. Physiol.; 290 (5): 136-145.
- <u>Khair</u>, A.; <u>Abdul Awal</u>, M.; <u>Islam</u>, M.; <u>Islam</u>, M. and <u>Damanna</u>, <u>R. (2021)</u>: Potency of spirulina (*Spirulina platensis*) on arsenicinduced lipid peroxidation in rat. Journal of advanced veterinary and animal research.; 8(2): 330–338.
- Khatun, M.; Hasan1, M.; Islam, R.; Sarkar, S. and Haque, M. (2020): Effect of spirulina (*Spirulina platensis*) and vitamin E on arsenic induced toxicity in Quail. Asian J. Med. Biol. Res., 6 (1):93-98.
- Korany, R.; Ahmed, K.; El Halawany, H. and Ahmed, K. (2019): Effect of long-term arsenic exposure on female albino rats with

special reference to the protective role of spirulina platensis. Explore Anim Med Res.; 9 (2) 125-136.

- Li, T.; Liu, Y. and Wan, Z., (2018): Regulatory Efficacy of the Polyunsaturated Fatty Acids from Microalgae *Spirulina platensis* on Lipid Metabolism and Gut Microbiota in High-Fat Diet Rats. *Int J Mol Sci.*; 19 (9): 20-18.
- Lu, H.; Lee, J.; Huang, L.; Lai, C.; Ho, L.; Chang, S. and Chi, W. (2012): Synergistic apoptosis-inducing antileukemic effects of arsenic trioxide and mucuna macrocarpa stem extract in human leukemic cells via a reactive oxygen species-dependent mechanism. Evid Based Complement Alternat.; 20(12):921-430.
- Nakaya, N.; Homma, Y. and Goto, Y. (1988): Cholesterol lowering effect of spirulina. *Nutrition reports international.*; 37: 1329–37.
- Nishikimi, M.; Appaji, N. and Yagi, K. (1972): The occurrence of superoxide anion in the reaction of reduced phenazine methosulphate and molecular oxygen. Biochem. Biophys. Res. Commun.; 46(2): 849–854.
- Reda, M.; Khaled, S.; Hanaa A. and Kawkab, A. (2019): Effect of long-term arsnic exposure on female albino rats with special reference to the protective role of spirulina platensis. Exploratory Animal and Medical Research.; 9, (2):125-136.
- Reeves, P.; Nielsen, F. and Fahmy, G. (1993): Purified diets for laboratory rodents: Final report of the American Institute of Nutrition writing committee on the reformulation of the AIN- 76 a rodent diet. J. Nutr. 123(51): 1939-1951.
- Rodrigo, R. and Rivera, G. (2002): Renal damage mediated by oxidative stress: a hypothesis of protective effects of red wine. Free Radic Biol Med.; 33:409-22.
- **Roy, E. (1970):** Colorimetric determination of Co. St Louis. Toronto. Princeton; 1088-1273.
- Sayed, A.; Gofur, R.; Khair, A. and Awal, A. (2015): Protective role of *spirulina* and vitamin e against arsenic toxicity in rats. Asian J Anim Sci.; 9 (6): 330-340.
- Sherwin, J. (1984): Liver Function. In Kaplan LA, Pesce AJ, eds. Clinical chemistry, theory, analysis, and correlation. St Louis: Mosby 55(25):420-438.
- Snedecor, G. and Cochran, W. (1980): Statistical methods.,7th Ed., Iowa State University Press, Ames, USA (90).
- Silbergeld, E.; Graham, J. and Price, L. (2008): Industrial food animal production, antimicrobial resistance, and human health. Annu. Rev. Public Health.; 29:151-169.

- Vedi, M.; Kalaiselvan, S.; Rasool, M. and Sabina, E. (2013): Protective effects of blue green algae Spirulina fusiformis against galactosamine-induced hepatotoxicity in mice. Asian J. Pharm. Clin. Res, 6(3): 150-154.
- Vonshak, A. (1997): Spirulina platensis (Arthrospira), Physiology, Cell-biology and Biotechnology.; London: Taylor & Francis.
- Young, D. (1990): Effect of drugs on clinical laboratory tests. Am. J. Clin. Pathol 3(7):6-12.
- Zhang, W.; Liu, Y.; Ge, M.; Jing, J.; Chen, Y.; Jiang, H. and Zhang, Z. (2014): Protective effect of resveratrol on arsenic trioxideinduced nephrotoxicity in rats. *Nutrition Research and Practice.*; 8(2): 220-226.



للدراسات المتخصصة

دورية فصلية علمية محكمة - تصدرها كلية التربية النوعية - جامعة عين شمس

<u>الهيئة الاستشارية للمجلة</u>

أ.د/ إبراهيم فتحي نصار (مصر) استلا الكيمياء العضوية التخليقية كلية التربية النوعية - جامعة عين شمس

أ.د/ أسامة السيد مصطفى (مصر)
استاذ التغذية وعميد كلية التربية النوعية - جامعة عين شمس

أ.د/ اعتدال عبد اللطيف حمدان (الكويت) استاذ الموسيقى ورنيس قسم الموسيقى بالمعهد العالي للفتون الموسيقية دولة الكويت

> أ.د/ السيد بهنسي حسن (مصر) استاذ الإعلام – كلية الآداب – جامعة عين شمس

1.1 / بدر عبدالله الصالح (السعودية) استاذ تكنولوجيا التعليم بكلية التربية جامعة الملك سعود

أ.د/ رامى نجيب حداد (الأردن)
استاذ التربية الموسيقية وعميد علية الفنون والتصميم الجامعة الأردنية

أ.د/ رشيد فايز البغيلي (الكويت)
استاذ الموسيقى وعميد المعهد العالي للفنون الموسيقية دولة الكويت

أ.د/ سامى عبد الرؤوف طايع (مصر) استاذ الإعلام – كلية الإعلام – جامعة القاهرة ورنيس المنظمة الدولية للتربية الإعلامية وعضو مجموعة خبراء الإعلام بمنظمة اليونسكو

1.6 / سوزان القليني (مصر) استاذ الإعلام- كلية الأداب – جامعة عين شمس عضو المجلس القومي للمرأة ورنيس الهينة الاستشارية العليا للإتحاد الأفريقي الأسيوي للمرأة

أ.د/ عبد الرحمن إبراهيم الشاعر (السعودية)
استاذ تكنولوجيا التعليم والاتصال - جامعة نايف

أ.د/ عبد الرحمن غالب المخلافي (الإمارات) استاذ مناهج وطرق تدريس- تقنيات تعليم – جامعة الأمارات العربية المتحدة

> أ.د/ عمر علوان عقيل (السعودية) استاذ التربية الخاصة وعميد خدمة المجتمع كلية التربية - جامعة الملك خالد

1.6 / ناصر نافع البراق (السعودية) استاذ الاعلام ورنيس قسم الاعلام بجامعة الملك سعود

أ.د/ فاصر هاشم بدن (العراق) استاذ تقنيات الموسيقى المسرحية قسم الفنون الموسيقية كلية الفنون الجميلة - جامعة البصرة

Prof. Carolin Wilson (Canada) Instructor at the Ontario institute for studies in education (OISE) at the university of Toronto and consultant to UNESCO

Prof. Nicos Souleles (Greece) Multimedia and graphic arts, faculty member, Cyprus, university technology

(*) الأسماء مرتبة ترتيباً ابجدياً.



رئيس مجلس الإدارة أ.د/ أسامة السيد مصطفى نائب رئيس مجلس الإدارة أ.د/ داليا حسين فهمي رئيس التحرير أ.د/ إيمان سيد على هيئة التحرير i.د/ محمود حسن اسماعیل (مصر) **أ.د/ عجاج سليم** (سوريا) **i**.د/ محمد فرج (مصر) 1.د/ محمد عبد الوهاب العلالى (المغرب) 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1.
 1. المحرر الفني د/ أحمد محمد نجيب سكرتارية التحرير د / محمد عامر محمد عبد الباقي أ/ أسامة إدوارد أ/ليلي أشرف أ/ محمد عبد السلام أ/ زينب وائل المراسلات : ترسل المراسلات باسم الأستاذ الدكتور / رئيس

التحرير، على العنوان التالى ٣٦٥ ش رمسيس – كلية التربية النوعية – جامعة عين شمس ت/ ٤٩٩٤ ٢٢/٢٦٨٤ • الموقع الرسمي: <u>https://ejos.journals.ekb.eg</u> البريد الإلكتروني: egyjournal@sedu.asu.edu.eg

الترقيم الدولي الموحد للطباعة : 6164 - 1687 الترقيم الدولي الموحد الإلكتروني : 2682 - 4353 تقييم المجلة (يونيو ٢٠٢٣) : (7) نقاط معامل ارسيف Arcif (أكتوبر ٢٠٢٣) : (0.3881)

المجلد (١٢). العدد (٤٣). الجزء الثالث

يوليو ۲۰۲٤