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RESEARCH ARTICLE

Investigation of the Combined Effects of Temperature and Nutrients on Chlorophyll-a Concentration in the Tigris River, Baghdad

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ABSTRACT

Rivers are exposed to global warming and eutrophication, which in turn affect phytoplankton biomass in rivers. This study investigates the changes in Chl-a concentration, an indicator of phytoplankton biomass. A microcosm experiment was conducted, with water sampling from the Tigris River in Baghdad during the wet season. Temperature increase was based on raising the temperature by 3 °C higher than the ambient temperature, while nutrient enrichment was based on the mean values of soluble reactive NO_{3-1} and PO_{4-3} verified in the Tigris River. Variations in Chl-a concentration are associated with changes in WT and nutrient enrichment; NO_{3-1} and PO_{4-3} . Our results showed that the Chl-a concentration was 0.8 $\mu\text{g/l}$ in the control microcosm. Temperature increase led to an increase in Chl-a of Temperature microcosm 1 $\mu\text{g/l}$, enrichment of nutrient, in turn, led to an increase in Chl-a of NO_{3-1} and PO_{4-3} microcosms to 1.2 and 0.8 $\mu\text{g/l}$, respectively, so the temperature increase and nutrient enrichment together led to a greater increase in Chl-a of T + NO_{3-1} and T + PO_{4-3} microcosms 1.2 and 1.1 $\mu\text{g/l}$, respectively, than if it were a single effect, which constitutes the highest concentration of Chl-a during the experiment. The effects of temperature and/ or nutrient can change the concentration of Chl-a. Nutrient enrichment. NO_{3-1} and PO_{4-3} , were identified as key contributors to Chl-a concentration, while warming influences the growth of Chl-a by enhancing metabolic and photosynthetic processes under nutrient-enriched conditions.

Keywords: Chlorophyll-a, Nutrient enrichment, Phytoplankton, Temperature increase, Tigris river

Introduction

Eutrophication is the systematic response of a fresh-water ecosystem to processing factors, particularly nutrients from soil and the discharge of wastewater into these water sources. Previous studies have identified different factors that promote eutrophication, primarily chlorophyll-a, nutrients (NO_{3-1} and PO_{4-3}), and water temperature.^{1,2} Chlorophyll-a, an indicator of phytoplankton biomass,³ is directly related to the availability of (NO_{3-1} and PO_{4-3}).⁴

Tigris River, the study site, is one of the longest rivers in the Middle East. With the country's growing population and expanding economic activities, the demand for water for various uses has increased. This

has been due to the significant pressure on Iraq's water resources over the past twenty years, particularly in terms of the available water supply.⁵ Iraqi researchers focused on different ways to assess water quality,^{6,7} while using Chl-a phytoplankton as a biological indicator.⁸

Water temperature (WT) is a major factor affecting the growth rates of phytoplankton, species distribution, and the nutrient cycle in the water system.⁹ For this study, WT in the microcosms was changed to reflect the warming scenario as projected by the IPCC.¹⁰ As for nutrients, nitrate (NO_{3-1}) and phosphate (PO_{4-3}), are vital for the growth of phytoplankton and affect chlorophyll-a,¹¹ excessive nutrients from runoff cause eutrophication

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and harmful algal blooms,¹² algal bloom can alter aquatic food webs, which can be harmful to the life of freshwater.¹³ The enrichment of phosphorus in the water poses a risk of future eutrophication.¹⁴

Based on previous research and available data for the Tigris River, water temperature (WT) and nutrients (NO_3^- and PO_4^{3-}) were hypothesized to control chlorophyll-a.¹⁵ To understand the combined effects of temperature and nutrients on chlorophyll-a, we employed a microcosm experiment to investigate this topic. The aim of the study was to elucidate the impact on Chl-a in the Tigris River in response to increasing warming and eutrophication, which are assumed to enhance competitive interactions between different phytoplankton taxa, leading to an increase in phytoplankton biomass. So it is predicted that the effects of combined factors may be greater than the sum of the individual effects, which forms a future scenario in terms of temperature and nutrient content.

Methods and materials

Microcosm experiment design

In our study, a microcosm experiment was conducted Fig. 1. Water sampling from the (AL-Gherai'at site) was on January 8th 2025. It consisted of 18 outdoor insulated polyethylene microcosms; each microcosm had a capacity of 20 L. Water sampling from the Tigris River included three replicates of the 6 treatments. The temperature increase was based on raising the temperature 3 °C above the ambient temperature. The nutrient enrichment was based on the mean values of soluble reactive and verified in the Tigris River, to stimulate nutrient loading, solutions of NaNO_3 and Na_2HPO_4 , (5 mg/l and 1 mg/l, respectively) were used. Each gram of NaNO_3 contains 0.72 grams of nitrate, so we added 100 mg of this compound to each NO_3^- and T + NO_3^- treatment. Each gram of Na_2HPO_4 contains 0.66 grams of phosphate, so we added 20 mg of this compound to each PO_4^{3-} ,

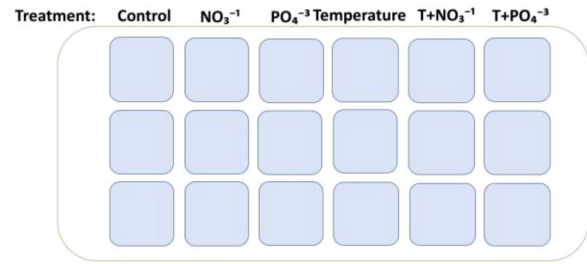


Fig. 1. Schematic representation of the experimental design.

and T + PO_4^{3-} treatment on the 1st day of the experiment. We adjusted the pH of the medium after adding the nutrient salts to minimize the potential effect of sodium salts on pH and salinity, ensuring pH stability across all treatments. Salinity remained within the normal range for our study, and the effect of sodium from the added salts was minimal.

Microcosms were performed as follows: (a) Ambient temperature microcosms: Control treatment (no nutrient enrichment), NO_3^- treatment, and PO_4^{3-} treatment. (b) Heated microcosms: Temperature treatment (no nutrient enrichment), T + NO_3^- treatment, and T + PO_4^{3-} treatment.

Analytical methods

Triplicate subsamples were collected and analysed eight times during the study period from January 8th to February 1st 2025, subsamples were taken to estimate Physico-chemical variables (WT, pH, EC, Salinity, DO, CO_2) in the field, while NO_3^- , PO_4^{3-} , DOC, and biological analysis of phytoplankton chlorophyll-a (Chl-a, proxy phytoplankton biomass) were analyzed in the laboratory, according to the methods mentioned in Table 1.

Statistical analyses

Statistics were based on microcosm-specific means and standard deviations (Mean $\pm$ SD) of measured response variables. A two-way ANOVA test was used to

Table 1. Methods of Physico-chemical and biological variables.

Variable	Method	Equation	Unit	Reference
WT	Water Quality Multimeter (Yieryi, China)	-	°C	-
pH	Water Quality Multimeter (Yieryi, China)	-	-	-
EC	Water Quality Multimeter (Yieryi, China)	-	$\mu\text{S}/\text{cm}$	-
Salinity	Calculated from EC	Salinity ‰ = $\text{EC } (\mu\text{S}/\text{cm}) \times 0.00064$	ppt or ‰	16
DO	Dissolved Oxygen Analyzer (Yieryi)	-	mg/l	-
CO_2	Titrimetric by using Na_2CO_3 and HCl	$\text{CO}_2 = (\text{A} + \text{B} \times 4.4) \times 10$	μM	17,18
NO_3^-	UV-Visible Spectrophotometric	-	mg/l	19
PO_4^{3-}	UV-Visible Spectrophotometric	-	mg/l	20
DOC	DOC Determination by Filtration (0.45 μm)	The final value of TOC after filtering = DOC	mg/l	21
Chl-a	Acetone Extraction Spectrophotometer	$\mu\text{g Chl-a} = 11.9 [2.43 (\text{Db-Da})] (\text{V/L}) (1000/\text{S})$	$\mu\text{g/l}$	22

Table 2. Summary statistics (Mean $\pm$ SD) of the physicochemical and biological variables were measured throughout the study period for the microcosm experiment.

Treatment Variable	Ambient Temperature Microcosms			Heated Microcosms		
	Control	NO ₃₋₁	PO ₄₋₃	Temperature	T + NO ₃₋₁	T + PO ₄₋₃
WT (°C)	16.3 $\pm$ 0.9	15.8 $\pm$ 0.8	15.6 $\pm$ 0.7	18.7 $\pm$ 0.9	19.4 $\pm$ 1.3	20.2 $\pm$ 0.6
pH	8.2 $\pm$ 0.3	7.8 $\pm$ 0.6	8.3 $\pm$ 0.2	8.3 $\pm$ 0.1	8.3 $\pm$ 0.1	8.3 $\pm$ 0.2
Salinity (‰)	0.3 $\pm$ 0.06	0.3 $\pm$ 0.05	0.3 $\pm$ 0.05	0.4 $\pm$ 0.08	0.4 $\pm$ 0.08	0.5 $\pm$ 0.1
EC (μ S/cm)	576.5 $\pm$ 89.4	589.7 $\pm$ 90	569.9 $\pm$ 87.6	638.2 $\pm$ 125	669 $\pm$ 122.4	804.1 $\pm$ 261.1
DO (mg/l)	7.9 $\pm$ 0.4	7.7 $\pm$ 0.4	7.8 $\pm$ 0.4	7.8 $\pm$ 0.3	7.6 $\pm$ 0.5	7.2 $\pm$ 0.7
CO ₂ (μ M)	176.1 $\pm$ 13.8	181.8 $\pm$ 26.3	178.1 $\pm$ 12.1	187.7 $\pm$ 19.8	192.3 $\pm$ 21.5	209.7 $\pm$ 25.8
NO ₃₋₁ (mg/l)	1.7 $\pm$ 1.1	3.5 $\pm$ 2.3	0.5 $\pm$ 0.2	1.7 $\pm$ 0.8	4.7 $\pm$ 1.1	1.2 $\pm$ 0.5
PO ₄₋₃ (mg/l)	0.3 $\pm$ 0.5	0.2 $\pm$ 0.3	1.9 $\pm$ 0.6	0.06 $\pm$ 0.02	0.2 $\pm$ 0.5	3.2 $\pm$ 0.9
DOC (mg/l)	7.6 $\pm$ 3.2	6.4 $\pm$ 3	7.1 $\pm$ 3	7 $\pm$ 2.8	8 $\pm$ 3.4	5.9 $\pm$ 2.7
Chl-a (μ g/l)	0.8 $\pm$ 0.5	1.2 $\pm$ 2	0.8 $\pm$ 0.5	1 $\pm$ 0.8	1.2 $\pm$ 0.8	1.1 $\pm$ 1

Table 3. Two-way ANOVA analysis showing the effects of Time, Treatments, and the interactions between them on the studied environmental variables.

Variable	Time		Treatment		Time $\times$ Treatment	
	F-value(df)	p-value	F-value(df)	p-value	F-value(df)	p-value
WT	59.31(6,84)	<0.0001	439.56(5,84)	<0.0001	3.57(30,84)	<0.0001
pH	22.28(6,84)	<0.0001	9.57(5,84)	<0.0001	3.28(30,84)	<0.0001
Salinity	46.21(6,84)	<0.0001	22.31(5,84)	<0.0001	1.91(30,84)	0.01
EC	42.20(6,84)	<0.0001	22.03(5,84)	<0.0001	1.81(30,84)	0.01
DO	94.62(6,84)	<0.0001	26.10(5,84)	<0.0001	1.91(30,84)	0.01
CO ₂	42.11(6,84)	<0.0001	23.00(5,84)	<0.0001	1.65(30,84)	0.03
NO ₃₋₁	1.27(6,84)	0.27	40.32(5,84)	<0.0001	0.89(30,84)	0.62
PO ₄₋₃	1.67(6,84)	0.13	127.95(5,84)	<0.0001	0.68(30,84)	0.87
DOC	10.42(6,84)	<0.0001	2.02(5,84)	0.08	1.19(30,84)	0.25
Chl-a	2.88(6,84)	0.001	0.90(5,84)	0.65	2.29(30,84)	0.02

Significance level: <0.05.

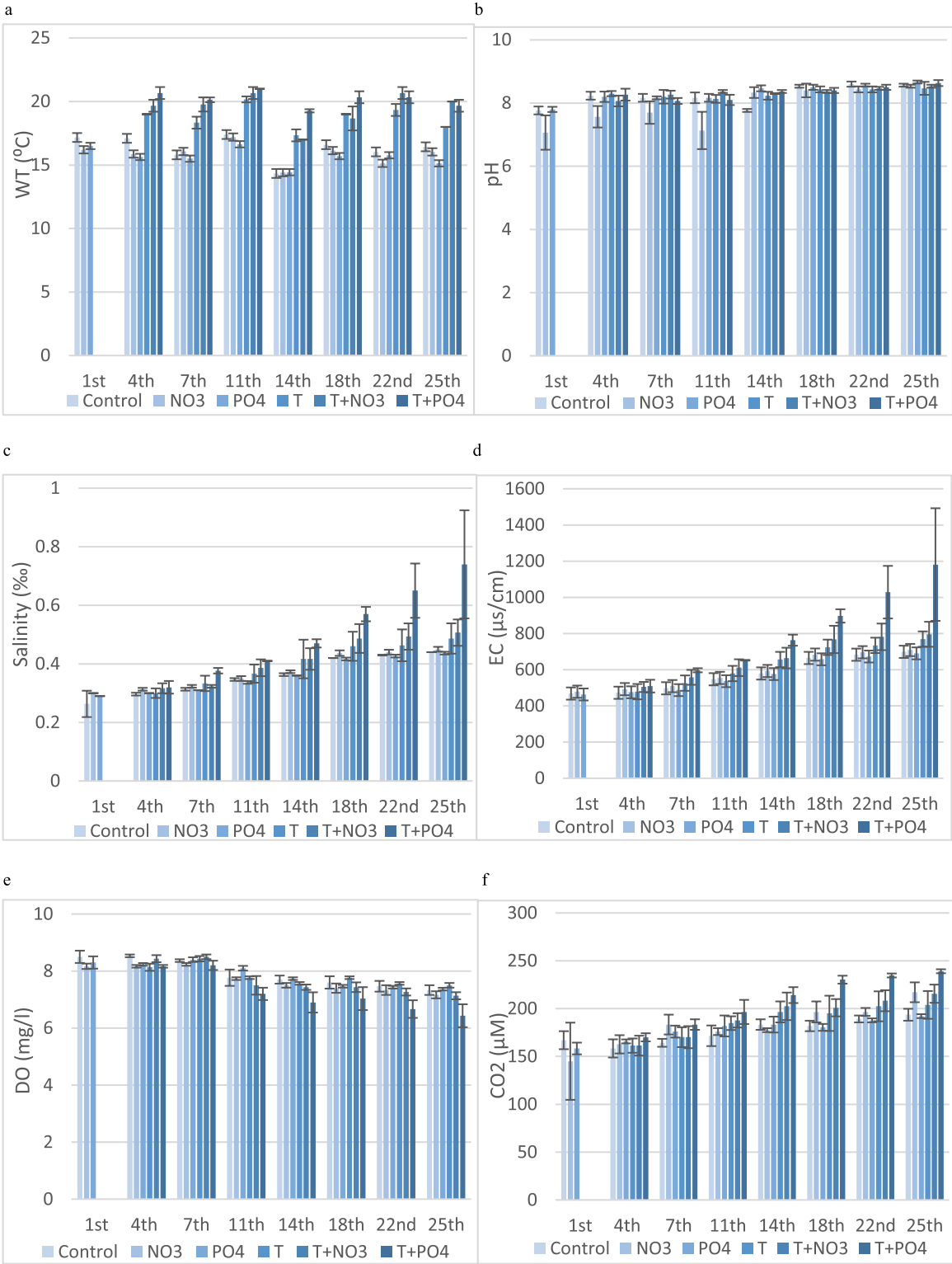
show if the variables changed significantly between time, treatments, and interactions between them. A standard t-test was used to test the differences in water chemistry between the control treatment and (NO₃₋₁, PO₄₋₃, Temperature, T + NO₃₋₁, and T + PO₄₋₃ treatments). Further, correlations between Chl-a and variables were calculated by using the Pearson correlation coefficient (r).

Results

Statistics (Mean $\pm$ SD) of the physicochemical and biological variables measured throughout the study period for the microcosm experiment are presented in Table 2. These variables changed significantly during the study period Table 3.

Water temperature ranged between 14.3 $\pm$ 0.28 °C in the control microcosm on the 14th day and 21 $\pm$ 0 °C in the heated PO₄₋₃ microcosm on the 11th day Fig. 2-a. WT value was affected by temperature increase and/or nutrient enrichment Table 4. pH ranged between 7 $\pm$ 0.5 in the NO₃₋₁ microcosm on the first day, and 8.6 $\pm$ 0.04 in PO₄₋₃ microcosm on the last day Fig. 2-b. pH value was only affected by the

individual effect of NO₃₋₁ enrichment, but not affected by the individual effects of PO₄₋₃, temperature, the combined effect of T + NO₃₋₁, and T + PO₄₋₃ Table 4. Salinity ranged between 0.26 $\pm$ 0.04 ‰ in the control microcosm on the first day and 0.74 $\pm$ 0.18 ‰ in the T + PO₄₋₃ microcosm on the last day Fig. 2-c. Salinity was affected by individual effect of NO₃₋₁, temperature, the combined effect of T + NO₃₋₁, and T + PO₄₋₃, but not affected by individual effect of PO₄₋₃ Table 4. Electrical conductivity (EC) ranged between 463.3 $\pm$ 0.94 μ S/cm in the PO₄₋₃ microcosm on the first day and 1181.3 $\pm$ 311.1 μ S/cm in the T + PO₄₋₃ microcosm on the last day Fig. 2-d. EC is affected by temperature increase and/or nutrient enrichment Table 4. Dissolved oxygen (DO) ranged between 8.5 $\pm$ 0.04 mg/l in the control microcosm on the fourth day and 6.4 $\pm$ 0.4 mg/l in the T + PO₄₋₃ microcosm on the last day Fig. 2-e. DO was affected by the individual effect of NO₃₋₁, combined effects of T + NO₃₋₁, and T + PO₄₋₃, but not affected by the individual effect of PO₄₋₃ and Temperature Table 4. CO₂ ranged between 145 $\pm$ 40.3 μ M in the NO₃₋₁ microcosm on the first day and 239 $\pm$ 2 μ M in the T + PO₄₋₃ microcosm on the last day Fig. 2-f. CO₂ was



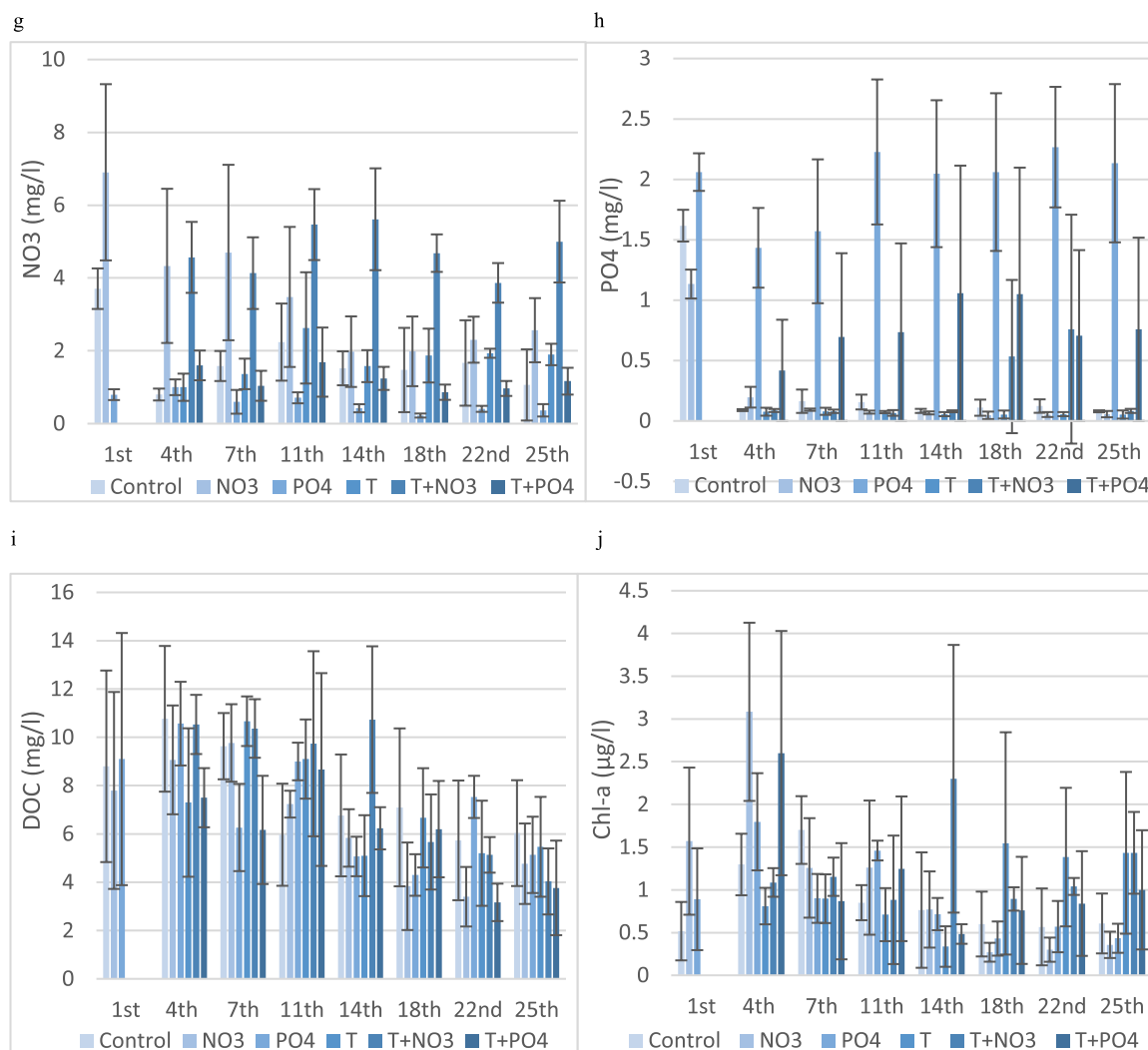


Fig. 2. Physico-chemical and biological variables throughout the microcosm experiment.

affected by the individual effect of Temperature, the combined effect of T + NO₃₋₁, and T + PO₄₋₃, but not affected by the individual effect of NO₃₋₁ and PO₄₋₃ Table 4. NO₃₋₁ ranged between 0.22 ± 0.05 mg/l in PO₄₋₃ microcosm on the 18th day, and 6.9 ± 2.41 mg/l in NO₃₋₁ microcosm on the 14th day Fig. 2-g. NO₃₋₁ was affected by the individual effect of NO₃₋₁, PO₄₋₃, and the combined effect of T + NO₃₋₁, but not affected by the individual effect of Temperature, and the combined effect of T + PO₄₋₃ Table 4. PO₄₋₃ ranged between 0.04 ± 0.03 mg/l in NO₃₋₁ microcosm on the 18th day, and 3.76 ± 0.75 mg/l in T + PO₄₋₃ microcosm on the last day Fig. 2-h. PO₄₋₃ was affected by the individual effect of NO₃₋₁, PO₄₋₃, Temperature, and the combined effect of T + PO₄₋₃, but not affected by the combined effect of T + NO₃₋₁ Table 4. Dissolved organic carbon (DOC) ranged between 3.1 ± 0.7 mg/l in the T + PO₄₋₃ microcosm on the 22nd day and 10.7

± 3 mg/l in the control microcosm on the fourth day Fig. 2-i. DOC was not affected by temperature and/or nutrient enrichment at all Table 4.

Chlorophyll-a ranged between 0.27 ± 0.1 μg/l in the ambient temperature NO₃₋₁ microcosm on the 18th day and 3.08 ± 1 μg/l in the ambient temperature NO₃₋₁ microcosm on the fourth day Fig. 2-j. Our results showed the concentration of Chl-a 0.8 ± 0.5 μg/l in the control microcosm. Temperature increase led to an increase in Chl-a in the Temperature microcosm (1 ± 0.8 μg/l). Nutrient enrichment, in turn, led to an increase in Chl-a in the NO₃₋₁ and PO₄₋₃ microcosms (1.2 ± 2 and 0.8 ± 0.5 μg/l, respectively), so the temperature increase and nutrient enrichment together led to a greater increase in Chl-a of T + NO₃₋₁ and T + PO₄₋₃ microcosms, 1.2 ± 0.8 and 1.1 ± 1 μg/l, respectively, than if it were a single effect, which constitutes the highest concentration of

Table 4. Summary statistics (T-test); Effects of Physico-chemical and biological variables measured throughout the microcosm experiment.

Variable	Control vs. Treatment	t-value	p-value
WT	NO ₃ ⁻¹	3.57	0.0007
	PO ₄ ⁻³	5.61	<0.0001
	Temperature	-15.16	<0.0001
	T + NO ₃ ⁻¹	-14.05	<0.0001
	T + PO ₄ ⁻³	-24.46	<0.0001
pH	NO ₃ ⁻¹	2.78	0.005
	PO ₄ ⁻³	-1.58	0.06
	Temperature	-0.38	0.35
	T + NO ₃ ⁻¹	-0.82	0.20
	T + PO ₄ ⁻³	-0.64	0.26
Salinity	NO ₃ ⁻¹	-3.4	0.001
	PO ₄ ⁻³	-0.10	0.45
	Temperature	-2.82	0.005
	T + NO ₃ ⁻¹	-4.78	<0.0001
	T + PO ₄ ⁻³	-4.89	<0.0001
EC	NO ₃ ⁻¹	-4.98	<0.0001
	PO ₄ ⁻³	3.92	0.0003
	Temperature	-2.81	0.005
	T + NO ₃ ⁻¹	-5.24	<0.0001
	T + PO ₄ ⁻³	-4.84	<0.0001
DO	NO ₃ ⁻¹	3.74	0.0005
	PO ₄ ⁻³	0.47	0.31
	Temperature	0.08	0.46
	T + NO ₃ ⁻¹	3.42	0.001
	T + PO ₄ ⁻³	7.61	<0.0001
CO ₂	NO ₃ ⁻¹	-1.40	0.08
	PO ₄ ⁻³	-0.91	0.18
	Temperature	-3.29	0.001
	T + NO ₃ ⁻¹	-4.57	<0.0001
	T + PO ₄ ⁻³	-8.07	<0.0001
NO ₃ ⁻¹	NO ₃ ⁻¹	-3.79	0.0004
	PO ₄ ⁻³	4.77	<0.0001
	Temperature	-1.23	0.11
	T + NO ₃ ⁻¹	-9.49	<0.0001
	T + PO ₄ ⁻³	1.12	0.13
PO ₄ ⁻³	NO ₃ ⁻¹	2.17	0.02
	PO ₄ ⁻³	-10.17	<0.0001
	Temperature	3.57	0.0009
	T + NO ₃ ⁻¹	-1.07	0.14
	T + PO ₄ ⁻³	-14.14	<0.0001
DOC	NO ₃ ⁻¹	1.40	0.08
	PO ₄ ⁻³	0.69	0.24
	Temperature	0.38	0.35
	T + NO ₃ ⁻¹	-0.61	0.27
	T + PO ₄ ⁻³	1.73	0.05
Chl-a	NO ₃ ⁻¹	-3.44	0.001
	PO ₄ ⁻³	-3.66	0.001
	Temperature	-2.49	0.01
	T + NO ₃ ⁻¹	-3.07	0.004
	T + PO ₄ ⁻³	-2.46	0.01

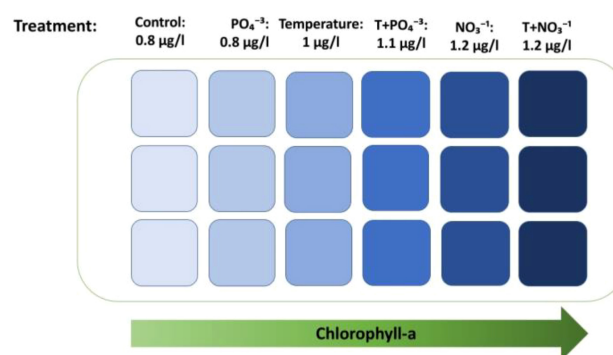
Significance level: <0.05.

Chl-a during experiment **Fig. 3**. Chl-a was affected by temperature increase and/or nutrient enrichment **Table 4**. Chl-a was affected by WT, pH, DO, CO₂, NO₃⁻¹, PO₄⁻³ and DOC, but not affected by salinity and EC **Table 5**.

Table 5. Effects of the Physico-chemical variables on chlorophyll-a.

Chl-a vs. Variable	Pearson correlation coefficient (r)	p-value
WT	0.18	0.03
pH	-0.21	0.01
Salinity	-0.12	0.13
EC	-0.13	0.12
DO	0.19	0.02
CO ₂	-0.18	0.03
NO ₃ ⁻¹	0.29	0.0004
PO ₄ ⁻³	-0.17	0.04
DOC	0.41	<0.0001

Significance level: <0.05.

**Fig. 3.** Schematic representation of our results for Chl-a concentration.

Discussion

Our experiment has provided a remarkably consistent pattern in the short-term response of chlorophyll-a concentrations. The cellular chlorophyll content of phytoplankton was stimulated by both increasing temperature and adding nutrients, an effect that was synergistically enhanced under the combined effects of temperature and nutrient enrichment.

The marked stimulatory effect of nitrate and phosphate additions on phytoplankton biomass confirmed our expectations. However, nutrient enrichment effects on chlorophyll-a concentrations differed between the treatments and between the types of nutrient (NO₃⁻¹, PO₄⁻³), indicating species-specific resource requirements. phytoplankton cells regulate their content of chlorophyll-a dependent on nitrogen availability,²³ but reductions in nitrogen loading at high phosphorus levels may lead to an increase in chlorophyll-a and a decrease in water clarity, resulting in a clear deterioration in water quality,²⁴ combination of high temperatures and phosphorus loading stimulated the maximum biomass built by the phytoplankton community,²⁵ these mainly reflected in nutrient-enriched microcosms, while low chlorophyll-a concentrations are reflected in nutrient-poor microcosms (Control and Temperature microcosms).

The short duration of our experiment may have prevented individual phytoplankton populations and communities from fully adapting to the increased temperatures, thus affecting the phytoplankton biomass's response to warming. Conversely, sudden temperature changes can exacerbate negative growth effects, particularly in the context of global warming, precisely because thermal acclimatization requires longer exposure periods.²⁶

Warm water promotes the growth of phytoplankton,²⁷ and this was mainly reflected in heated microcosms. However, in this study, chlorophyll-a concentration decreased in some heated microcosms as WT increased, possibly because WT exceeded that which is suitable for the growth of most phytoplankton, and high WT began to inhibit phytoplankton photosynthesis.²⁸

An increase was observed for pH value during the experiment; mild alkaline conditions may result in elevated chlorophyll-a.²⁹ An increase was observed for salinity and EC during the experiment. Our studies indicated that salinity and EC have no significant effect on chlorophyll-a, which could be attributed to the complicated interplay between several environmental elements (WT, pH, DO, and carbon sources) with species composition and ecological dynamics.³⁰ A decrease was observed in the DO concentration during the experiment. Although DO was high in the first days of the experiment due to low organism activity under low temperature such as respiration.³¹ An increase was observed for CO₂ concentration during the experiment, high CO₂ leads to increasing phytoplankton biomass.³² DOC influences algal blooms and overall ecosystem health.³³

To isolate the impact of the altered variable on chlorophyll-a concentration, all environmental variables were held constant in this study. As a result, changes in chlorophyll-a concentration primarily represent the responses of phytoplankton and may also be affected by other biotic factors, such as zooplankton grazing and interspecies interactions, which were not examined in our study. Future studies should include the structure and abundance of phytoplankton and zooplankton communities to better illustrate the ecosystem responses.

Conclusion

A study proved that temperature increase and/or nutrient enrichment influence physicochemical variables, which in turn influence chlorophyll-a across a microcosm experiment. Nutrient enrichment; NO₃₋₁ and PO₄₋₃, identified as the main contributors to chlorophyll-a concentration, while

temperature affects chlorophyll-a growth by enhancing metabolic processes and photosynthesis under nutrient-rich conditions. The relationship between elevated temperature and/or nutrient enrichment and chlorophyll-a may be more complex under climate change and eutrophication. Furthermore, the experimental study designs can't accommodate the complexity of freshwater ecosystems. Furthermore, studies are needed to integrate the experimental method and field observation to better understand the impacts of climate change on aquatic ecosystem dynamics.

Authors' declaration

- Conflicts of Interest: None.
- We hereby confirm that all the Figures and Tables in the manuscript are ours. Furthermore, any Figures and images that are not ours have been included with the necessary permission for republication, which is attached to the manuscript.
- No animal studies are present in the manuscript.
- No human studies are present in the manuscript.
- Ethical Clearance: The project was approved by the local ethical committee at the University of Baghdad.

Authors' contributions statements

Y. A. and J. S. A. contributed to the design and implementation of the Microcosm Experiment, to the analysis of the data, to the statistical analysis of the results, and to the writing of the manuscript.

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دراسة التأثيرات المشتركة لدرجة الحرارة والمغذيات على تركيز الكلوروفيل-أ في نهر دجلة، بغداد

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الخلاصة

تتعرض الأنهار لظاهرة الاحتباس الحراري والتخثث، مما يؤثر بدوره على الكتلة الحيوية للعوالق النباتية فيها. تبحث هذه الدراسة في التغيرات التي تطرأ على تركيز الكلوروفيل-أ، وهو مؤشر على الكتلة الحيوية للعوالق النباتية. أجريت تجربة مصغرة، حيث جُمعت عينات مياه من نهر دجلة في بغداد خلال موسم الأمطار. استندت زيادة درجة الحرارة إلى رفعها بمقدار 3 درجات مئوية فوق درجة الحرارة المحيطة، بينما استند إثراء المغذيات إلى متوسط قيم التي تم التحقق منها في نهر دجلة. ترتبط التغيرات في تركيز الكلوروفيل-أ بالتغيرات في درجة حرارة الماء وإثراء المغذيات؛ النتراة والفوسفات. أظهرت نتائجنا أن تركيز الكلوروفيل-أ بلغ 0.8 ميكروغرام/لتر في النظام البيئي المصغر الضابط. أدى ارتفاع درجة الحرارة إلى زيادة تركيز الكلوروفيل-أ في النظام البيئي المصغر ذي درجة الحرارة بمقدار 1 ميكروغرام/لتر، بينما أدى إثراء المغذيات بدوره إلى زيادة تركيز الكلوروفيل-أ في النظامين البيئيين المصغرين ذوي النتراة والفوسفات بمقدار 1.2 و 0.8 ميكروغرام/لتر على التوالي. وبذلك، أدى ارتفاع درجة الحرارة وإثراء المغذيات معاً إلى زيادة أكبر في تركيز الكلوروفيل-أ في النظامين البيئيين المصغرين ذوي النتراة والفوسفات بمقدار 1.2 و 1.1 ميكروغرام/لتر على التوالي، مقارنة بتأثير كل منهما على حدة، وهو ما يمثل أعلى تركيز للكلوروفيل-أ خلال التجربة. أظهرت النتائج أن درجة الحرارة و/أو المغذيات يمكن أن تؤثر على تركيز الكلوروفيل-أ. وقد تم تحديد إثراء المغذيات، وخاصة النتراة والفوسفات، كعوامل رئيسية في زيادة تركيز الكلوروفيل-أ، بينما يؤثر ارتفاع درجة الحرارة على نمو الكلوروفيل-أ من خلال تعزيز عمليات التمثيل الغذائي والتمثيل الضوئي في ظل ظروف غنية بالمغذيات.

الكلمات المفتاحية: الكلوروفيل-أ، إثراء المغذيات، العوالق النباتية، ارتفاع درجة الحرارة، نهر دجلة.