

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

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ABSTRACT

Total petroleum hydrocarbons (TPHs) in coastal seawater and marine sediments have attracted much interest because of their potential danger to aquatic ecosystems and human health. This research evaluated seasonal variations and ecological and human health concerns associated with total petroleum hydrocarbon (THPs) in surface water and sediment samples from the northern Gulf of Suez in 2019. Spectrophotometric detection was used to determine the concentration of TPHs in water and sediment samples that had been extracted using liquid-liquid and sonication extraction techniques, respectively. TPHs seasonal means in the water and sediment samples collected from the studied area showed that summer has the lowest levels of this pollutant, and ordered by winter (20.35) > autumn (17.51) > spring (11.55) > summer (17.88 µg/l), and winter (77.64) > summer (64.54 µg/g) for water and sediment samples, respectively. There was a highly significant correlation between physical parameters and TPHs in water and sediments. Evaluation of ecological risk using the pollution index (PI) and Nemerow pollution index (NPI) recorded moderate and low pollution, respectively. The non-carcinogenic risk of TPHs calculated in the water column and sediments by dermal absorption were lower than the target value (< 1) for adults and children. For human health risk, the carcinogenic dermal risk of TPHs was more than the acceptable limits (1×10^{-4} - 1×10^{-6}) for adults. In contrast, the carcinogenic dermal risk of TPHs was lower than the acceptable limits (1×10^{-4} - 1×10^{-6}) for children. Therefore, it follows that the body of water is contaminated and that the necessary measures should be taken to rein in all the sources of pollution there.

Keywords: Total petroleum hydrocarbon, Ecological risk, Human risk, Seawater, Sediments, Gulf of Suez, Egypt

INTRODUCTION

There was a release of several anthropogenic contaminants into the aquatic environment including polycyclic aromatic hydrocarbons (PAHs) since rapid industrialization and urbanization (Brusseau *et al.*, 2019). Carbon and hydrogen, with trace quantities of other

elements, including sulfur, nitrogen, and oxygen, are the primary building blocks of petroleum, which is found in nature (Dembicki Harry, 2017). Anthropogenic activities, such as petroleum extraction and burning fossil fuels, also release hydrocarbons into the environment (Zeneli *et al.*, 2019; Brusseau *et al.*, 2019). Ukpaka

et al. (2020) pointed out how this devastatingly impacts marine ecosystems, fisheries resources, and the diversity of biological communities. In recent years, oil spills have become a major hazard to marine life and human health. Depending on the organism's metabolism, they may be acutely or chronically hazardous (Quintana-Rizzo *et al.*, 2015; Bo *et al.*, 2017). They have limited biodegradability, are carcinogenic, teratogenic, and mutagenic, and cause considerable damage to organisms. As a result of swallowing small amounts or because of direct contact, petroleum hydrocarbons may bioaccumulate in bigger animals through trophic transfer (Kuppusamy *et al.*, 2020). As a result, they cause disorder in food webs, destroy ecosystems, and threaten human health (Muniz *et al.* 2015; Ukpaka *et al.* 2020). Various studies have confirmed the presence of total petroleum hydrocarbon in surface waters and sediments, emphasizing their toxicity and the risks they pose to human health in various regions around the world (Adeniji *et al.*, 2019 a; Du, 2019; Akinola *et al.*, 2019; Anyanwu *et al.*, 2020). Depending on the goals of protection, management, and mitigation, a risk assessment may be conducted in the context of an ecological risk assessment or a human health risk assessment (Pinedo *et al.*, 2012; Tian *et al.*, 2020). These substances' potential carcinogenicity or non-carcinogenicity is assumed mechanically in the risk evaluation of human exposure to contaminants (Chiesa *et al.*, 2019).

The primary source of pollution in Egypt's Red Sea is oil, making the Gulf of Suez the most contaminated location in the country (El-Agroudy *et al.*, 2017). It is a key international trade route and a strategic site near the mouth of the Suez Canal. In addition, the Gulf of Suez's distinctive maritime environment makes it valuable

for fishing stocks and tourism (Mahmoud *et al.*, 2020). Oil terminals and oil refineries, ships, petroleum industries, and other ports that serve as (transit zone, loading/unloading, and fishing operations) are all possible causes of oil pollution. It is thus of local concern that the waters of the Gulf of Suez have a deleterious effect on marine habitats and human health (Abdelmongy & El-Moselhy, 2015; Ibrahim *et al.*, 2019).

This research aimed to assess and identify the concentrations and sources of total petroleum hydrocarbon in surface water and sediments in the northern part of the Gulf of Suez in 2019. In addition, assessment the ecological dangers of TPHs in the investigated locations by using the pollution index approach and the risk assessment method. Also, to investigate the possible hazardous impacts of TPHs on the dermal health of children and adults.

MATERIALS AND METHODS

1. Study area:

The investigated area is located at the northern part of the Gulf of Suez and at the south entrance of the Suez Canal, which extends from Port-Tawfik to El-Sokhna with an average length of nearly about 54 km, and a mean depth is (10-18 m). The northern part of the Gulf of Suez has been subjected to different industrial and human activities covering the coastline (Hamed, 1992). Seven stations were selected depending on the main activities (I: Port-Tawfik; II: El-Zaiytia; III: El-Kabanon; IV: Attaka; V: El-Adabyia; VI: Tourist villages and VII: El-Sokhna) (Fig., 1).

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

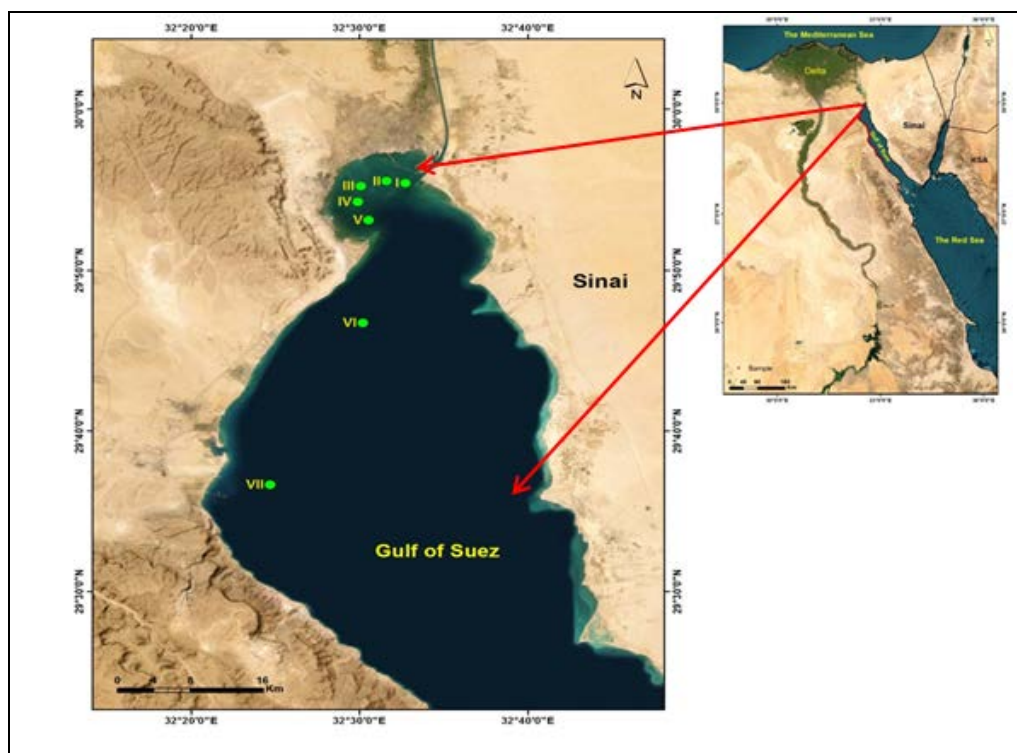


Fig. (1). Map of the northern part of the Gulf of Suez showing the sampling stations.

2. Sampling collection and preparation:

Seven monitoring stations were set up to reflect the varying environmental conditions of the offshore regions of the northern part of the Gulf of Suez in 2019. A total of 84 samples of seawater and 42 samples of surface sediments were collected. According to (UNEP and PERSGA, 1997), subsurface water samples were taken at 50 centimeters; the seawater was acidified to pH 2 using 10% HCl to protect against bacterial activity in dark bottles. The sediment samples were taken using a grab and wrapped in pre-cleaned aluminum foil. The samples were kept at 4°C until they could be sent to the lab for examination.

3. Sampling Analysis:

After adding dichloromethane to a water sample of known volume in a

separator funnel, we gave it a good shake for 2 minutes. After letting the organic layer settle to the bottom of the vial, the water was filtered through 5 g of sodium sulfate-coated filter paper. The sediment sample weighed 10 g, and 30 mL of dichloromethane (the extraction solvent) was added before the vial was sonicated for 15 minutes. The extracted mixture is filtered into the vial when the solid phase has settled, using filter paper with 5 g of sodium sulfate already on it. The water and sediment samples yielded findings in $\mu\text{g/L}$ and $\mu\text{g/g}$, respectively.

4. Reagents, Solvents, and Standards:

The high-performance liquid chromatography grade solvents (dichloromethane, acetone, and hexane) utilized in sample preparation and analysis were purchased from Merck. Absorbent

silica gel (100-200 mesh) made from reagent-grade anhydrous sodium sulfate (Merck).

5. Quality control:

All stages of production were subjected to stringent quality checks. Before taking any samples, we cleaned the equipment with sterile water and laboratory-grade detergent. The accuracy of all instruments was checked. The calibration of the system was checked and confirmed. Ensure there are at least three linear concentration points in the calibration range. Samples were prepared for colorimetric analysis by evaporating oil from produced water; the concentration of samples was determined by the density of crude oil in the range of 100-250 mg/L, and the method was validated by comparing it to other methods for determining oil in water, such as gas chromatography and flame ionization detection. Light crude oil extracted with a low relative error of 31.504 and heavy crude oil extracted with a high relative error of 7.025 was studied. Oil in generated water was successfully quantified using the suggested approach.

6. Pollution assessment

6.1. Ecological Risk

The northern part of the Gulf of Suez water and sediments quality was evaluated using the standard index of single-factor assessment. The pollution index (PI) approach is often used for assessing the pollution status of petroleum hydrocarbons (Zhu *et al.*, 2020; Cao *et al.*, 2020), with the formula for doing so being:

$$Pi = Ci/Ci_o \quad (1)$$

Pi, Ci, and Ci_o represent the

evaluated results, the actual measured data, and the evaluation standard of petroleum hydrocarbon, respectively. TPH is measured at 0.05 mg/L in surface seawater and 500 mg/kg in surface sediments (Pawar *et al.*, 2002; Kwak *et al.*, 2018). PI class is the environment and aquatic products into three categories: The non-polluted type when PI < 0.5; the moderately polluted environment and aquatic products when it varies between 0.5-1; and the highly polluted environment and aquatic products when PI is > 1.

6.2. Nemerow pollution index (NPI)

The NPI approach considers both the mean and the maximum PI value, provides a more nuanced response to the pollution level, and can emphasize the influence of pollutants using TPH. According to the formula (Nemerow, 1974):

$$NPI = \frac{\sqrt{(Pi)^2 + (Pimax)^2}}{2} \quad (2)$$

Where, Pi is pollution concentration for each station; Pimax is the maximum pollution concentration, NPI, pollution levels are high when NPI > 3, pollution levels are moderate when 2 < NPI ≤ 3.0; pollution levels are low when 1 < NPI ≤ 2.0; NPI pollution levels are low when 0.7 < NPI ≤ 1.0, and finally safe when NPI ≤ 0.7.

7. Evaluation of potential human health

Human health risk assessment is all about assessing the risks that carcinogenic and non-carcinogenic chemicals pose to human health. The risk assessment procedure consists of four steps: calculating toxicity (dose-response), measuring exposure, describing risks, and identifying hazards (USEPA, 2015).

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

7.1. Non-carcinogenic hazard (THQ)

The possible carcinogenic and non-carcinogenic effects of exposure to the TPHs over a particular time were calculated to determine the human health risk (USEPA, 2015; Titilawo *et al.*, 2018). Since this body of water is intended for recreational use, the study was focused only on dermal contact with the contaminants (EOH CES, 2016). The target hazard quotient (THQ) is the ratio of the exposure dose to the reference dose (RfD) (Table 1); THQ was calculated using the formula of Eqs. (3 & 4) in mg/kg/day.

$$(THQ)_{\text{Dermal - water}} \text{ (mg/kg/day)} = \frac{C_s \times S_A \times C_F \times E_T \times E_F \times E_D}{B_w \times T_A \times R_f D} \dots \dots \dots (3)$$

$$(THQ)_{\text{Dermal -Sediment}} \text{ (mg/kg/day)} = \frac{C \times C_F \times S_A \times A_F \times A_B \times E_F \times E_T \times E_D}{B_w \times A_T \times R_f D} \dots \dots \dots (4)$$

Where THQ (Target Hazard Quotients) ingestion and dermal of TPHs (mg/kg/day); C_s stands for the concentration of TPHs in the water sample (mg/L); E_F is the exposure frequency both ingestion and dermal absorption; E_D is the exposure duration; B_w represents the average body weight; A_T means the average time; S_A stands for the exposed

skin area; E_T is the exposure time of shower and bathing; and C_F represents the unit conversion factor and $R_f D$, which represents dermal reference dose (Table 1). Guidelines values provided by the Department of Environmental Affairs and USEPA were used for the estimations (USEPA, 2015; DTSC, 2014; Feng *et al.*, 2016; Wang *et al.*, 2018). Hazard quotient (HQ) ≥ 1 is high risk; $0.1 \leq (HQ) < 1$ is medium risk, and $(HQ) < 0.1$ is low risk.

7.2. Cancer risk assessment

The cancer slope factor (CSF) is used to convert the ratio of the exposure dose of TPHs over a lifetime of exposure to the risk of an individual developing cancer (USEPA, 2015). Potential carcinogen exposure is part of a carcinogenic risk assessment that estimates a person developing cancer over a lifetime. The incidence of cancer was estimated using Eq (5) according to USEPA guidelines (Wei *et al.*, 2015; USEPA, 2015).

$$CSR = ADD \times CSR \dots \dots \dots (5)$$

Where CSF represents the cancer slope factor for Benzo (a) Pyrene is 7.3 mg/kg/day (USEPA, 2015). A risk value $> 1.0 \times 10^{-4}$ indicates carcinogenic effects, according to the US Environmental Protection Agency (USEPA, 2015) (Table 1).

Table (1). Exposure parameters used for the health risk assessment through different exposure pathways (Adeniji *et al.*, 2019a; USEPA, 2015)

Parameters	Values		
	Unit	Child	Adult
Body weight (B _w)	Kg	15	70
Exposure frequency (E _F)	Days/year	365	
Exposure Duration (E _D)	Years	6	30
Stands for the exposed skin area (SA)	Cm ²	18000	6600
The exposure time of shower and bathing (ET)	h/day	0.58	1
RfD (USEPA, 2009)	mg/kg/day	Dermal = 0.04	
Cancer Slop Factor CSF	mg/kg/day	7.3	
Conversion factor (CF)	mg/ng	10 ⁻⁶	
Dermal Absorption from Sediment ABS		0.13	
Adherence Factor from Sediment to Skin AF	mg/cm ²	0.07	0.2
Average Time		Days/year	
For Carcinogenic		365 x 70 = 25550	
For non-Carcinogenic		365 x E _D	365 x E _D

8. Data Analysis

The statistical analysis of the collected dataset was performed using XL-state, 2020. Pearson's correlation coefficient (CM) analysis was performed to identify the relationship between total petroleum hydrocarbons in seawater and sediments and physical parameters. In addition, TPH content in seawater and sediment samples and physical parameters were subjected to one-way ANOVA. A probability of 0.05 was considered a significance level (Stephan *et al.*, 2019).

RESULTS AND DISCUSSION

1. Physical parameters:

Table (2) shows the physical characteristics measured in this work for water samples taken from 7 locations in the

northern part of the Gulf of Suez in 2019. Water temperature, pH, and salinity varied between (18.33 - 30.22 °C), (7.78 - 8.20), and (39.93 – 42.33 ‰), respectively. The present results matched the previous study by Khedr *et al.* (2019).

The seasonal water temperature distribution showed summer > autumn > spring > winter. Various factors could affect the water temperature, such as variations in climatic conditions, latitude, height, season, wind, depth, and waves (Abdelmongy & El-Moselhy, 2015). The pH value declined more in the summer than in the other seasons. In which the volume of sewage released into urbanized regions and the photosynthetic activity of algal biomass significantly impact the pH value (Abdel-Halim & Aly-Eldeen, 2016). The lowest annual pH level (7.87) was

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

observed at station (III) El-Kabanon. This might be related to high sewage discharges and increased microbial activity in the maritime environment (Abo El-Khair *et al.*, 2016; Mahmoud *et al.*, 2020). The annual mean of salinity ranged between (40.02 – 42.14 ‰) for Tourist villages and

El-Zaityia stations, respectively. Salinity variations were affected by temperature and wastewater discharge (Mahmoud *et al.*, 2020). The current results agreed within acceptable limits with coastal water quality guidelines tailored to the marine environment (DSME, 2004),

Table (2): Seasonal and annual means of physical parameters measured in water samples collected from the northern part of the Gulf of Suez during (2019).

Stations		Seasons	Temp °C	pH	Salinity ‰
I	Port-Tawfik	Winter	18.69	8.14	41.33
		Spring	23.74	8.10	41.35
		Summer	29.88	8.00	41.89
		Autumn	23.87	8.11	41.54
Annual Mean			24.05 ± 4.58	8.09 ± 0.06	41.53 ± 0.26
II	El-Ziaytia	Winter	18.33	8.19	41.89
		Spring	23.34	8.13	42.15
		Summer	29.27	8.01	42.33
		Autumn	23.55	8.15	42.20
Annual Mean			23.62 ± 4.47	8.12 ± 0.08	42.14 ± 0.18
III	El-Kabanon	Winter	18.39	7.99	39.93
		Spring	23.86	7.84	39.95
		Summer	29.55	7.78	40.35
		Autumn	23.85	7.88	40.09
Annual Mean			23.91 ± 4.56	7.87 ± 0.09	40.08 ± 0.19
IV	Attaka	Winter	19.09	8.18	40.17
		Spring	24.50	8.10	40.18
		Summer	30.22	8.06	40.16
		Autumn	24.35	8.13	40.21
Annual Mean			24.54 ± 4.55	8.12 ± 0.05	40.18 ± 0.02
V	El-Adabyia	Winter	18.22	8.20	40.29
		Spring	24.78	8.17	40.33
		Summer	29.88	8.11	40.31
		Autumn	25.50	8.18	40.33
Annual Mean			24.60 ± 4.81	8.17 ± 0.04	40.32 ± 0.02
VI	Tourist villages	Winter	18.50	8.20	39.97
		Spring	24.80	8.16	40.01
		Summer	30.10	8.10	40.07
		Autumn	24.22	8.13	40.02
Annual Mean			24.41 ± 4.74	8.15 ± 0.04	40.02 ± 0.04
VII	El-Sokhna	Winter	18.90	8.10	40.89
		Spring	25.26	8.01	41.23
		Summer	29.70	7.95	41.27
		Autumn	24.10	8.00	41.11
Annual Mean			24.49 ± 4.44	8.02 ± 0.06	41.13 ± 0.17

2. Total petroleum hydrocarbons in water and sediments:

The presence of petroleum hydrocarbons in marine ecosystems is rapidly rising to the top of the world's most environmental concerns. It helped us recognize the human impacts on marine ecosystems and shed light on the dramatic shifts that have occurred in recent years (Beiras, 2018). The quantity, origin of pollutants, the temporal and spatial patterns of TPHs in water may be assumed from monitoring (Ahmed *et al.*, 2015). Marine sediments act as a reservoir for petroleum hydrocarbon components. Therefore, surface sediment may provide insight into the contamination level of sediments today (Angela *et al.*, 2012). The present study analyzed all samples from the northern Gulf of Suez for total petroleum hydrocarbons (TPH) in the sediments and surface water (Table 3 and Figs. 2 – 4).

TPH concentration recorded the highest annual mean (28.53 $\mu\text{g/l}$ and 122.04 $\mu\text{g/g}$) at El-Zaiytia station (II) for water and sediments, respectively. It can be attributed to its proximity to numerous oil pollution sources, including loading and unloading activities at El-Zaiytia harbor and the effluents of oil refineries owned by El-Nasr and Suez petroleum companies (Diab, 2017). TPHs level observed in sediments is higher than in water, which may be attributed to TPHs being less affected by biological or photochemical oxidation in bottom sediments. Also, TPHs may stay long and build up to dangerous levels in sediments (Guzzella & Paolis, 1994). According to these findings, Neff (1985) stated that the petroleum hydrocarbons in sediments are more than

those observed in the water column by a factor of 1000 or more.

The present results showed the spatial pattern of TPHs mean concentrations in water samples in the investigated area fall in the following order El-Zaiytia > El-Sokhna > Attaka > El-Adabyia > Port-Tawfik > Tourist villages > El-Kabanon (28.53, 21.40, 17.21, 15.39, 14.19, 13.56 and 7.50 $\mu\text{g/l}$, respectively). While, in sediment samples was El-Zaiytia > Port-Tawfik > Attaka > El-Adabyia > El-Sokhna > Tourist villages > El-Kabanon (122.04, 97.76, 85.22, 55.11, 41.63 and 23.36 $\mu\text{g/g}$, respectively). Ports activities highly contribute to the TPH's value in the marine ecosystem. This conclusion was consistent with the previous results of (Eed & Kaiser, 2016), which suggested that marine-based activities may be the most effective variables in the distribution patterns and concentrations of TPHs at severely contaminated stations. According to Adeniji *et al.* (2017), all stations had TPH concentrations below the EU's maximum allowable level for hydrocarbons in harbor basin water of 300 $\mu\text{g/L}$. In comparison, our data showed that the sediments of the present study revealed slightly to moderately polluted (Table 3).

Seasonally, TPHs in water varied from (8.90 – 35.46), (7.22 – 30.19), (5.52 – 19.03), and (8.35 – 29.43) $\mu\text{g/L}$ throughout the winter, spring, summer, and autumn seasons, respectively, and from (26.36 – 138.98) and (20.35 – 105.11) $\mu\text{g/g}$ during the winter and summer, respectively, in the sediments. In particular, the TPHs distribution confirmed that summer has the lowest level of these pollutants in the studied area, with average means of (20.39 > 17.87 > 17.49 > 11.55 $\mu\text{g/l}$) and (77.64 > 64.54 $\mu\text{g/g}$) for water and sediments,

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

respectively. High temperatures accelerate the breakdown of petroleum hydrocarbons

due to increased microbial activity (El-Agroudy *et al.*, 2017; Kottb *et al.*, 2019).

Table (3). Seasonal and annual mean for TPHs in water and sediment samples collected from the northern part of the Gulf of Suez during 2019.

		Surface water samples $\mu\text{g/l}$					MAC for TPHs for Seawater = 0.05 m (ATSDR 2013) (Omayma <i>et al.</i> , 2015), wastewater = 0.5 mg/l; DPR (2015) = 10 mg/l
		Stations	Winter	Spring	Summer	Autumn	
Water	I	Port-Tawfik	17.47	14.03	10.02	15.22	14.19
	II	El-Zaiytia	35.46	30.19	19.03	29.43	28.53
	III	El-Kabanon	8.90	7.22	5.52	8.35	7.50
	IV	Attaka	20.78	19.15	10.71	18.19	17.21
	V	El-Adabyia	18.56	15.05	11.32	16.61	15.39
	VI	Tourist villages	16.21	13.95	9.97	14.09	13.56
	VII	El-Sokhna	25.08	22.99	14.29	23.24	21.40
	Seasonal Mean			20.35 ± 8.27	17.51 ± 7.42	11.55 ± 4.19	17.88 ± 6.78
		Sediment samples $\mu\text{g/g}$					
Sediment	Station		Winter	Summer		Annual Mean	
	I	Port-Tawfik	101.97	93.55		97.76	
	II	El-Zaiytia	138.98	105.11		122.04	
	III	El-Kabanon	26.36	20.35		23.36	
	IV	Attaka	92.50	77.93		85.22	
	V	El-Adabyia	59.61	50.62		55.11	
	VI	Tourist villages	45.98	37.29		41.63	
	VII	El-Sokhna	78.06	66.96		72.51	
	Seasonal Mean			77.64 ± 37.71	64.54 ± 30.45		71.09 ± 33.94
Guideline for sediment	Unpolluted		10 – 15 mg/kg		(Adeniji <i>et al.</i> , 2017)		
	Slightly polluted		15 – 50 mg/kg				
	Moderately polluted		50 – 200 mg/kg				
	Heavily polluted		> 200 mg/kg				

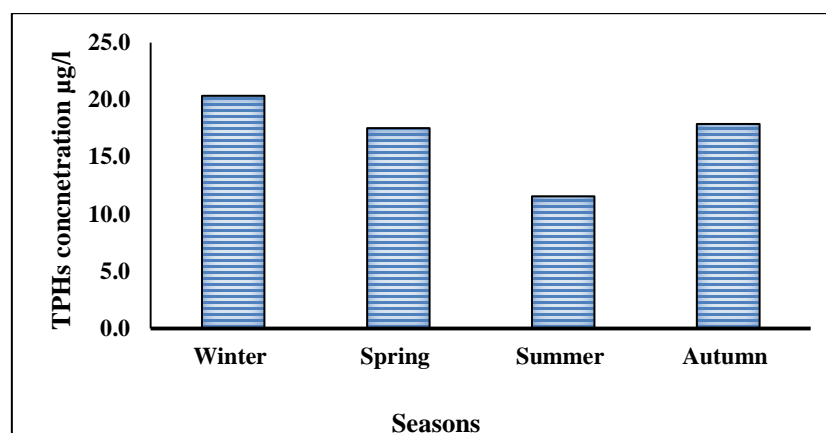


Fig. (2). Seasonal mean for TPHs in the water samples from the investigated area during 2019

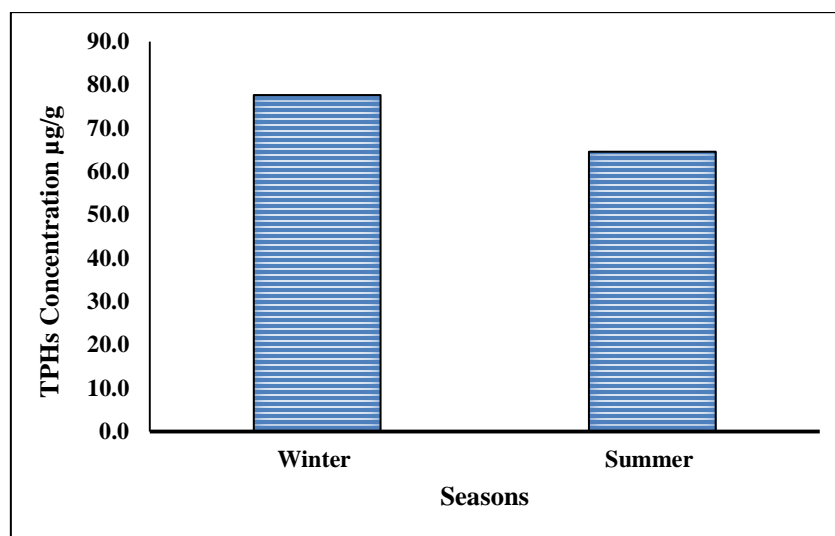


Fig. (3). Seasonal mean for TPHs in the sediment samples from the investigated area during 2019

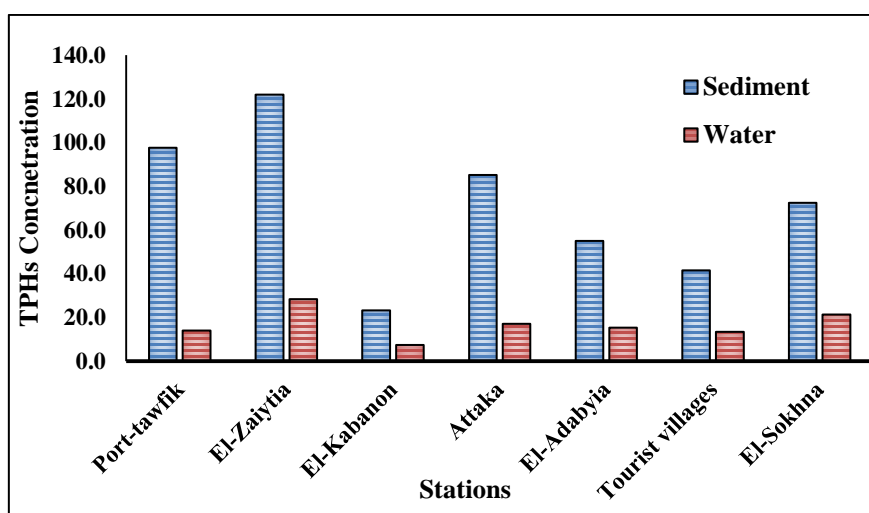


Fig. (4). Annual mean for TPHs in water and sediment samples from the investigated area during 2019

3. Ecological risk assessment:

Coastal ecosystem health may be estimated by assessing pollutants such as petroleum contaminates in seawater and sediments (Khudur *et al.*, 2018). Each sample's PI was used to determine how much pollution was present in the area of investigation and the results are shown in Table (4) and Figure (5). Seasonal PI for

TPHs in water and sediments ranged between (0.18 – 0.71), (0.14 – 0.60), (0.11 – 0.38) and (0.17 – 0.59) for winter, spring, summer, and autumn, respectively, and from (0.053 – 0.278) and (0.041 – 0.201) for winter and summer, respectively. The order of the mean values of PI for water showed winter (0.41) > autumn (0.36) > spring (0.35) > summer (0.23), whereas in

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

sediment samples was winter (0.155) > summer (0.129). Seasonal PI was classified as a non-polluted type (<0.5) for water and sediment samples. Regionally, the annual mean values ranged (0.15 – 0.57) with an average (0.34 ± 0.13) and (0.047 – 0.244) with an average (0.142 ± 0.07) for water and sediments, respectively. Only El-Zaiytia station recorded PI (0.57) higher than the lower pollution limit in water, and was classified as a moderately polluted station.

Whereas, NPI values ranged from 0.51 to 0.64 with an average of 0.56 ± 0.04 and 0.199 – 0.262 with an average of 0.224 ± 0.022 for water and sediments, respectively. The seven water monitoring stations revealed the following

geographical distribution trend: El-Zaiytia (II) > El-Sokhna (VII) > Attaka (IV) > El-Adabyia > Port-Tawfik (I) = Tourist villages (VI) > El-Kabanon (III). While in sediments was El-Zaiytia (II) > Port-Tawfik (I) > Attaka (IV) > El-Sokhna (VII) > El-Adabyia > Tourist villages (VI) > El-Kabanon (III). According to the NPI values, the current investigated stations were classified as a safe pollution area. Consequently, the northern part of the Gulf of Suez is classified as non-polluted and safe even though shipping operations and land-based wastewater discharges are the primary causes of oil pollution in the present study area (Liu *et al.*, 2019).

Table (4). Pollution indices of the measured TPHs in water and sediment samples collected from the northern part of the Gulf of Suez for aquatic life utilizations.

		Pi= Ci/Ci _o					NPI	
Stations		Winter	Spring	Summer	Autumn	Annual Mean		
Water	I	Port-Tawfik	0.35	0.28	0.20	0.30	0.28	0.54
	II	El-Zaiytia	0.71	0.60	0.38	0.59	0.57	0.64
	III	El-Kabanon	0.18	0.14	0.11	0.17	0.15	0.51
	IV	Attaka	0.42	0.38	0.21	0.36	0.34	0.56
	V	El-Adabyia	0.37	0.30	0.23	0.33	0.31	0.55
	VI	Tourist villages	0.32	0.28	0.20	0.28	0.27	0.54
	VII	El-Sokhna	0.50	0.46	0.29	0.46	0.43	0.59
	Average		0.41 ±0.17	0.35 ±0.15	0.23 ±0.08	0.36 ±0.14	0.34 ±0.13	0.65 ±0.04
Sediment	Stations		Winter		Spring		Annual Mean	NPI
	I	Port-Tawfik	0.204		0.187		0.196	0.240
	II	El-Zaiytia	0.278		0.210		0.244	0.262
	III	El-Kabanon	0.053		0.041		0.047	0.199
	IV	Attaka	0.185		0.156		0.170	0.231
	V	El-Adabyia	0.119		0.101		0.110	0.211
	VI	Tourist villages	0.092		0.075		0.083	0.205
	VII	El-Sokhna	0.156		0.134		0.145	0.222
Average		0.155± 0.08		0.129 ± 0.06		0.142 ±0.07	0.224± 0.05	

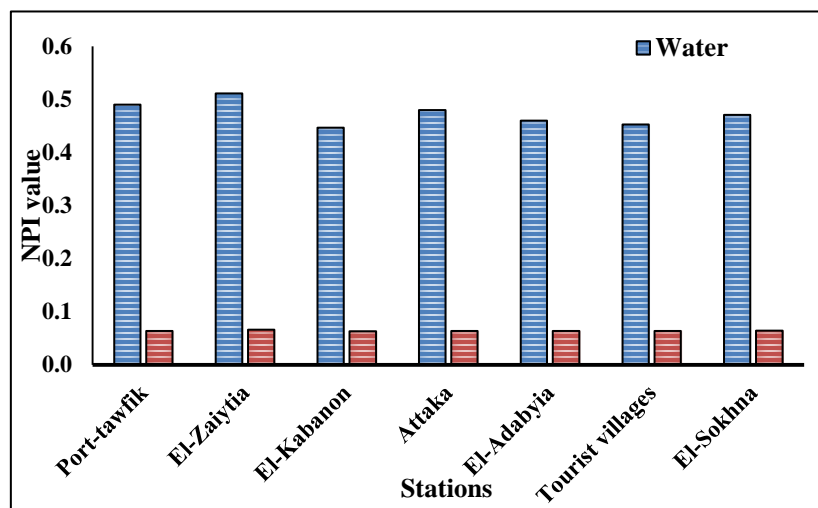


Fig. (4). Annual mean for NPI in water and sediment samples from the area of investigation during 2019

4. Human health risk assessment:

TPHs directly threaten humans in the aquatic environment because of their toxic properties and negative impact on the food chain (ITRC, 2018). In light of the above, the following are the most typical forms of serious risk assessment that include assessing the impact of water bodies on human health risks:

4.1. Non-Carcinogenic risk (THQ)

Exposure to carcinogenic compounds during a certain period was considered in human health risk assessments (USEPA, 2015). The danger to human health from dermal TPHs-contaminated water was calculated in this research using equations (3 and 4). In water, TPHs in the studied area were calculated to have target hazard quotients (THQ) ranging from (4.8×10^{-5} - 1.8×10^{-4}) and (3.1×10^{-7} - 1.2×10^{-6}) for adults and children, respectively, for dermal absorption (Table 5). In sediments, the THQ for TPHs ranged from (3.9×10^{-6} - 2.0×10^{-5}) and (2.3×10^{-10} - 1.2×10^{-9}), respectively. The findings show that the

THQ in the water column and sediments for both age groups was lower than the USEPA's threshold (<1). Therefore, recreational use is unlikely to expose people to non-carcinogenic diseases or other health problems (ATSDR, 2013; Wei *et al.*, 2015; Benson *et al.*, 2017). Consequently, adults are highly susceptible to non-carcinogenic risk than a child.

4.2. Carcinogenic Risk

Using equation (5), the effect of each particular hydrocarbon in water across all exposure routes was calculated independently for children and adults, yielding unique cancer risk assessments. Results from an analysis of the carcinogenic hazards posed by TPHs showed that these compounds have a low risk of causing cancer (Table 5). Therefore, the US Environmental Protection Agency (USEPA, 2015) accepts a cancer risk range of 1×10^{-6} - 1×10^{-4} . According to Adeniji *et al.* (2019 a, b), who stated that if the CSR is $\leq 1 \times 10^{-6}$, the risk is considered acceptable by the USEPA, and if it is between $1 \times 10^{-5} < \text{CSR} < 1 \times 10^{-4}$, the

Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

risk is considered unacceptable. If $CSR \geq 1 \times 10^{-4}$, it can cause serious health effects.

Table (5) shows that the water content of CSRs ranged from ($1.3 \times 10^{-5} - 3.5 \times 10^{-5}$) and ($2.1 \times 10^{-9} - 5.6 \times 10^{-10}$) for dermal in both adults and children, respectively. In comparison, CSR for sediments content varied between ($2.6 \times 10^{-6} - 4.7 \times 10^{-7}$) and ($3.17 \times 10^{-12} - 7.1 \times$

10^{-13}) for dermal in both adults and children, respectively. Adult group risk indices were higher than the limit value of ($1 \times 10^{-4} - 1 \times 10^{-6}$), which is considered unacceptable. At the same time, child group risk indices were lower than the limit value (acceptable limit) (Wei *et al.*, 2015; Wang *et al.*, 2018).

Table (5). THQ_{Dermal} of non-carcinogenic and Cancer Slop Risk (CSR) for adults and children in the area of investigation.

	Stations		Adult	Child	Adult	Child
			THQ _{Dermal}	THQ _{Dermal}	CSR	CSR
Water	I	Port-Tawfik	9.1E-05	5.8E-07	6.6E-06	1.1E-09
	II	El-Zaiytia	1.8E-04	1.2E-06	1.3E-05	2.1E-09
	III	El-Kabanon	4.8E-05	3.1E-07	3.5E-06	5.6E-10
	IV	Attaka	1.1E-04	7.1E-07	8.0E-06	1.3E-09
	V	El-Adabyia	9.9E-05	6.3E-07	7.2E-06	1.1E-09
	VI	Tourist villages	8.7E-05	5.6E-07	6.3E-06	1.0E-09
	VII	El-Sokhna	1.4E-04	8.8E-07	1.0E-05	1.6E-09
Sediment	I	Port-Tawfik	1.6E-05	9.5E-10	2.0E-06	3.0E-12
	II	El-Zaiytia	2.0E-05	1.2E-09	2.6E-06	3.7E-12
	III	El-Kabanon	3.9E-06	2.3E-10	4.9E-07	7.1E-13
	IV	Attaka	1.4E-05	8.3E-10	1.8E-06	2.6E-12
	V	El-Adabyia	9.2E-06	5.3E-10	1.2E-06	1.7E-12
	VI	Tourist villages	7.0E-06	4.0E-10	8.7E-07	1.3E-12
	VII	El-Sokhna	1.2E-05	7.0E-10	1.5E-06	2.2E-12
					CSR= $1 \times 10^{-6} - 1 \times 10^{-4}$	

5. Data analysis

According to the ANOVA analysis, the studied physical parameters and TPHs showed significant spatial and temporal variations (p -value < 0.05). Table (6) displays the correlation matrix of numerous variables in water and sediments across stations. Salinity and total petroleum hydrocarbons in water and

sediments were shown to have very positive associations (0.89 and 0.77, respectively). The correlation between total petroleum hydrocarbons in water and sediments was (0.82). In contrast, there were strong to moderate inverse associations between pH and total petroleum hydrocarbons in water and sediments (- 0.79 and - 0.58).

Table (6). Correlation matrix with different variables in water and sediment samples.

	Salinity	pH	Temp.	TPHs S	TPHs W
Salinity	1				
pH	-0.695	1			
Temp.	-0.272	-0.263	1		
TPHs S	0.885	-0.582	-0.355	1	
TPHs W	0.766	-0.789	-0.234	0.815	1

Conclusion

This study evaluated the spatial and temporal distribution patterns of the total petroleum hydrocarbons in water and sediment samples from the northern part of the Gulf of Suez, Egypt. Overall, TPHs were observed to be modest, and the principal sources of TPHs in the studied area were likely to be different intensive shipping activities associated with the main harbors and effluents discharged from petroleum oil refineries and petroleum production and industry companies. The PI and NPI pollution levels were classified as moderate and low, respectively, for ecological risk. For human health risk, the non-carcinogenic risk of TPHs calculated in the water column and sediments by dermal absorption was lower compared to the target value (< 1) for both adults and children. For adults, the carcinogenic dermal risk of TPHs was more than the acceptable limits (1×10^{-4} - 1×10^{-6}). In contrast, the carcinogenic dermal risk of TPHs was less than the acceptable limits (1×10^{-4} - 1×10^{-6}) for children. Therefore, more precautions have to be followed to protect the body of water from contamination with TPH from different sources.

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Ecological and human health risk assessment of total petroleum hydrocarbons in surface water and sediments from the northern part of the Gulf of Suez, Egypt

تقييم المخاطر البيئية والصحية للإنسان للمواد الهيدروكربونية البترولية الكلية في المياه السطحية والرسوبيات من الجزء الشمالي لخليج السويس، مصر

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المستخلص

اجتذبت المواد الهيدروكربونية البترولية الكلية (TPHs) في المياه الساحلية والرسوبيات البحرية اهتمامًا كبيرًا نظرًا لخطرها المحتمل على النظم البيئية المائية وكذلك على صحة الإنسان. وقد هدفت الدراسة الحالية إلى تقييم الاختلافات الموسمية والمخاطر البيئية والصحية المرتبطة بالمواد الهيدروكربونية البترولية الكلية في عينات من المياه السطحية والرسوبيات من شمال خليج السويس خلال عام ٢٠١٩. حيث تم استخدام الكشف الطيفي لتقدير تركيز المواد الهيدروكربونية البترولية الكلية في عينات المياه والرسوبيات التي تم استخلاصها باستخدام تقنيات الاستخلاص السائل وتقنيات الاستخلاص بالذئذبات الصوتية، على التوالي. أظهرت نتائج المتوسطات الموسمية لـ TPHs في عينات المياه والرسوبيات التي تم جمعها من منطقة الدراسة أن الصيف احتوي على أقل المستويات، وكان ترتيبها كالتالي: الشتاء (٢٠.٣٥) < الخريف (١٧.٥١) < الربيع (١١.٥٥) < الصيف (١٧.٨٨ ميكروجرام/ لتر) للمياه، والشتاء (٧٧.٦٤) < الصيف (٦٤.٥٤ ميكروجرام/جرام) للرسوبيات. كما كانت هناك علاقة طردية عالية بين الخواص الفيزيائية والمواد الهيدروكربونية البترولية الكلية في الماء والرسوبيات. سجل تقييم المخاطر البيئية باستخدام مؤشر التلوث (PI) ومؤشر التلوث Nemerow (NPI) تلوثًا متوسطًا ومنخفضًا، على التوالي. كانت المخاطر غير المسرطنة للمواد الهيدروكربونية البترولية الكلية في عمود الماء والرسوبيات عن طريق امتصاص الجلد أقل من القيمة المستهدفة (> ١) للبالغين والأطفال. بالنسبة للمخاطر على صحة الإنسان، كان خطر الإصابة بالسرطان الجلدي من المواد الهيدروكربونية البترولية الكلية أكثر من الحدود المقبولة (١ × ١٠^{-٤} - ١ × ١٠^{-٦}) للبالغين. على النقيض من ذلك، كان خطر الإصابة بالسرطان الجلدي من المواد الهيدروكربونية البترولية الكلية أقل من الحدود المقبولة (١ × ١٠^{-٤} - ١ × ١٠^{-٦}) للأطفال. وعلى ذلك فإن منطقة الدراسة تتعرض لمخاطر التلوث بالمواد الهيدروكربونية البترولية الكلية خاصة في عمود المياه ويجب اتخاذ الاجراء اللازمة لمعرفة مصادر تلك الملوثات والعمل على الحد من أثارها.