

PAH QUANTIFICATION AND ESTIMATED CARCINOGENIC RISKS AT SELECTED FUEL STATIONS IN TAMALE METROPOLIS, GHANA

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ABSTRACT

The research aim was to quantify PAHs levels and assess their associated carcinogenic health risks in fuel filling stations located in both high and low vehicular traffic areas in the Tamale Metropolis of Ghana. Dust particles were collected from ten fuel filling stations and extracted using an Accelerated Solvent Extractor (ASE). PAH analysis was conducted using a GC-MS system. Cancer risk assessment was done using the Incremental Life-time Cancer Risk (ILCR) model. The average PAH concentration recorded in the high and low vehicular traffic areas were $158,080 \pm 102,650 \mu\text{g/kg}$ and $370,220 \pm 218,140 \mu\text{g/kg}$ respectively. Benzo(a)Anthracene was the most occurring PAH congener recorded in both traffic areas. Dermal contact as a route of PAH exposure recorded the highest potential cancer risk for both adults and children. Amongst the fuel filling stations, Gab Energy recorded the highest carcinogenic risks of 2.50 (adults) and 2.67 (children). In all, the average cancer risk values for adults (1.02) was slightly higher than that of the children (9.26×10^{-1}) with no significant difference. The mean Σ ILCR values for both populations indicated a high to very high potential carcinogenic risk to the people living or working in and around the study areas.

Keywords: PAHs, fuel filling stations, dust particles, incremental lifetime cancer risk, Benzo(a)Anthracene.

Introduction

Polycyclic aromatic hydrocarbons (PAHs) are a group of organic compounds occurring naturally in resources such as crude oil, gasoline and coal which are introduced into the environment especially through incomplete combustion of organic substances (CDC, 2009). PAHs can be commonly identified in every

part of the environment; from soil media, air mixtures, and water sources, and with their traces found in various food products, thereby characterising them as ubiquitous chemicals in the environment (Ofori *et al.*, 2020).

PAHs are described as organic chemicals having a range of environmental properties which are determined by the amount of

benzene rings and the molecular weight they possess (Stogiannidis & Laane 2015). PAH congeners having four or more benzene rings are categorised as High Molecular Weight (HMW) PAHs, whereas those possessing two to three benzene rings are categorised as Low Molecular Weight (LMW). Some examples of LMW PAHs include Fluorene, Anthracene, Naphthalene, Phenanthrene, while other congeners such as Indeno[1,2,3-cd]pyrene, Chrysene, Pyrene, Benzo[a]pyrene, and Benz[a]anthracene and are described as HMW PAHs (Stogiannidis & Laane, 2015; Ofori *et al.*, 2020).

Law *et al.* (2002) noted that the LMW PAHs stay less longer in the environment due to their high volatility, and their low ability to dissolve in water. On the other hand, HMW PAHs are identified to be persistent in the environment as a result of their low volatile ability and their high resistance to undergo oxidation (Stogiannidis & Laane, 2015). They are also described as being more water-insoluble when present as a mixture with alkyl substituent groups, making them strongly carcinogenic and mutagenic (Nkansah, 2012).

The ubiquitous and carcinogenic nature of PAHs in both indoor and outdoor environments are indicated to constitute a significant risk to public health as their exposure could lead to their uptake into the human body via inhalation, ingestion of food or water, soil, and/or via the skin. Public exposure to PAHs could lead to short- and long-term health effects, particularly associated with breathing and cardiovascular symptoms or ailments (Błaszczuk *et al.*, 2016).

In Africa, PAH emissions from Nigeria and the Democratic Republic of Congo ranked 4th and 8th respectively considering the worldwide view in a report provided by Zhang and Tao (2009) in 2004. One of the reasons that put Nigeria as one of the top countries contributing to the global PAH pollution was

the frequent incidents of oil pollution occurring in the Niger Delta (Ofori *et al.*, 2020). In Africa, a high number of studies have been done in assessing the occurrence and concentration levels of PAHs in various environmental samples as well as the risks they pose to public health. Most of the studies have focused on PAH pollution in agricultural soils, street soils, smoked fish and meat, food items, indoor and outdoor air, water bodies, among others (Ofori *et al.*, 2020). However, only one study has been identified to focus on PAH pollution in fuel filling stations by assessing the levels of PAH pollution in the Calabar Metropolis of Nigeria (Nganje *et al.* 2007). In Ghana, no such study had yet been conducted.

In fuel filling stations, where there are countless records of both entries and exits of various vehicles for the purchase of different fuel products, the situation of PAH pollution appears to be common. Higher levels of PAH pollution in this area could be recorded due to observed instances of frequent fuel leakages from fuel dispensers or from faulty vehicular fuel tanks, as well as emissions of exhaust fumes from vehicles that visit or are located in the vicinity of the fuel filling stations. The risk of PAH exposure in the Tamale Metropolis is aggravated by the fact that, some heavy-duty vehicles use some of these stations as parking lots. This presents a high likelihood for fuel leakage and emissions from these vehicles which could thereby be classified as another contributor of PAH pollution into the vicinity. It has also been reported that automobiles, particularly heavy-duty vehicles in the Tamale Metropolis release high concentrations of PAHs into the atmosphere through their exhaust fumes (Obiri *et al.*, 2011). These incidents of PAH pollution collectively create health hazards for workers, customers and people living closer to the fuel filling stations. The limited study in PAH pollution in fuel filling stations in the African continent has creat

ed a knowledge gap in literature which needs to be tackled with. Moreover, in view of the need to promote public education on the potential health risks associated with PAH pollution, it was found necessary to conduct this research by assessing the concentration levels and potential carcinogenic health risks associated with the exposure to PAHs at fuel filling stations located in the Tamale Metropolis of Ghana.

Experimental

Study area

The study was conducted in Tamale Metropolis which is located in the Northern Region of Ghana. The Metropolis is situated at the central

part of the region, sharing boundaries with the Sagnarigu District (West and North direction), Mion District (East direction), and East Gonja (South-West direction). It has a total land size of about 646.9 km² and a geographical location between latitude 09°16 and 09°34 North and longitude 00°57 West. The population of the Metropolis as at 2010 was 233,252 (49.7% males and 50.3% females), corresponding to 9.4% of the regional population with majority being young people (almost 36.4% of the population is below 15 years) (Ghana Statistical Service, 2014).

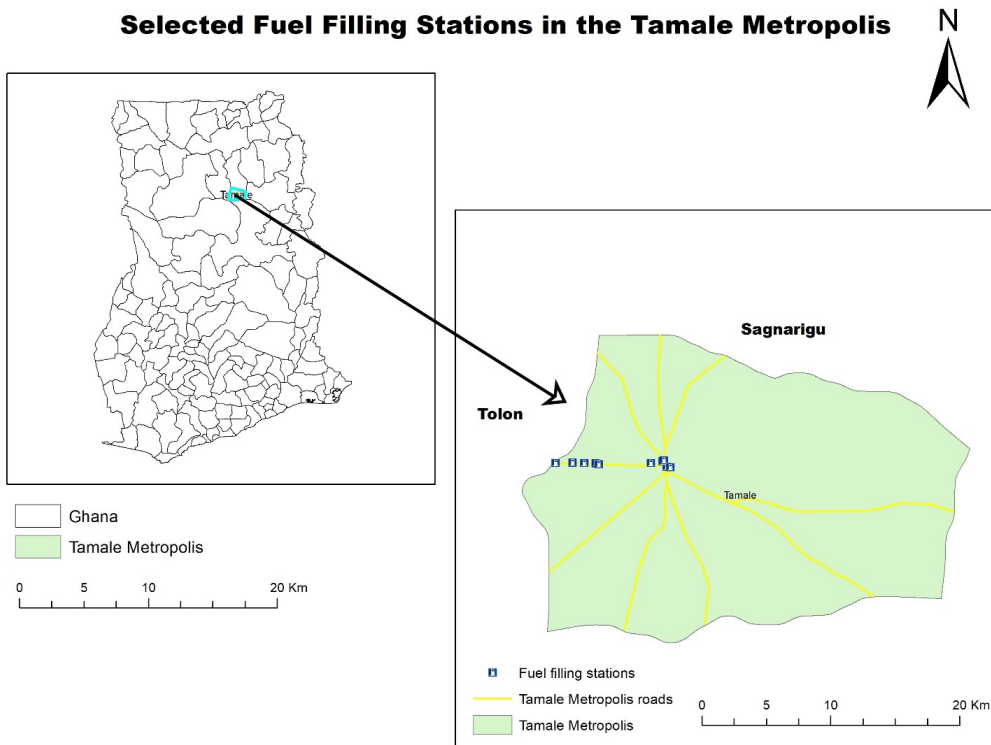


Fig. 1: Tamale Metropolis Showing Selected Fuel Filling Stations.

Selection of fuel filling station and sampling technique

This study was conducted in fuel filling stations. For the purpose of this study, fuel filling stations were categorised under High-Vehicular Traffic Areas (H-VTA) and Low-Vehicular Traffic Areas (L-VTA) based on the following parameters:

- Fuel filling stations are close to or far from traffic light system(s)
- Fuel filling stations are close to or far from areas with continuous vehicular traffic
- Fuel filling stations are located in the centre or outskirts of Metropolis

Thus, an area where traffic light system(s) were present was categorised as a H-VTA, and the area without a traffic light system was categorised as a L-VTA. Also, an area where one could find recurrent vehicular traffic jams was categorised as a H-VTA, whereas an area without any occurrence of vehicular traffic jams was categorised as a L-VTA. Moreover, an area with more business activities involving high vehicular movements and located in the centre of the Metropolis was categorised as a H-VTA, while an area with relatively less busy nature and located away from the Metropolis centre was categorised as L-VTA.

In this view, ten fuel filling stations (five from each category) which were able to fully meet the criteria for selection were used in this study (Table 3).

Sample collection

The samples were collected in the dry season from 19th to 21st February 2018, between the hours of 6 pm and 8 pm. This was the best time to find as much dust particles accumulated in the day. Frequent incidences of fuel leakages coupled with the accumulation of dust particles as blown by the dry winds towards the fuel dispensing area gave an indication of a major occurrence of PAH pollution in that area. Therefore, at each fuel filling station, two

samples of 20 g each were collected mainly from the fuel dispensing area by sweeping dust particles with a pre-cleaned brush unto a pre-cleaned dustpan. The fuel dispensing area is described as the space where drivers stop with their vehicles in order to be served, as well as the area around the fuel dispensing machine. Samples were then collected into 15 mL amber glass vials which had already been washed with de-ionised water and covered with aluminium foil. To prevent cross contamination of samples collected from each fuel filling station, the brush and dustpan had to be pre-cleaned before sampling in each fuel filling station. The glass vials were later labelled and then enfolded with aluminium foil and preserved at an average temperature of 4 °C in an ice chest and transported to the laboratory for analyses.

Sample Preparation and extraction

Sample preparation and extraction was done according to protocol recommended by El-Kady et al. (2018). Sample pairs collected from each fuel filling station were added as a composite sample and homogenised by sieving them through a 250 µm mesh to remove stones and unwanted particles present. A sample mixture was prepared with a 3:1 ratio of 30 g of each sample composite to 10 g of diatomaceous earth (DE) to effect homogeneity and dryness. Extraction was carried out using the Dionex Accelerated Solvent Extractor (ASE) 350 with a solvent mixture comprising of 60 mL each of methanol and acetone (v/v; 1:1) per sample extract. The method and sequence of extraction was programmed according to the following order:

- Temperature; 100°C, Static time; 5m, Cycles; 3, Rinse volume; 20%, Purge; 30s.

Sample clean-up and analysis

The clean-up process was done using a rotary evaporator. Sample extracts were concentrated

to a volume of 1 mL at a temperature between 40 - 45 °C under a gentle stream of nitrogen gas. All samples were analysed under the same instrumental conditions defined by USEPA standards (USEPA, 1999).

The Agilent Technologies GC-MS system (GC 7890B and MS 7000C) was used to identify the presence and concentrations of PAH congeners in the sample extracts. A 2.0 µL of each sample extract was injected into the GC-MS system using the Agilent Split/Splitless inlet with a split time of 1 min after injection and at an inlet temperature set at 280°C. The oven temperature programme was initially held at 80°C for 0.5 min, then increased to 230°C at 80°C/min, and finally increased to 280°C at 5°C/min (held at 5 min) with a run time of 33.648 min. Helium was used as the carrier gas with a column flow rate of 1 mL/min. The MS mode of operation was in the electron impact ionisation mode at 70 eV with selected ion monitoring at a temperature of 230°C. After all semi-volatile target compounds were eluted from the GC, the MS data acquisition was terminated, and data files were stored on the data system storage device.

The resulting limit of detection (LOD), which is calculated as three times the standard deviations of the blank sample, was recorded as 1.00 µg/kg for all the 16 PAHs that were present in the samples.

Health risk assessment

Mutagenicity and carcinogenicity of PAHS

The MEQ and TEQ levels for the sum of the PAHs were calculated using defined values of MEF and TEF with the following equations;

$$MEQ = \sum_{i=1}^n (\text{PAH})_i \times MEF \quad (1)$$

$$TEQ = \sum_{i=1}^n (\text{PAH})_i \times TEF \quad (2)$$

Where MEQ is Mutagenic Equivalent, TEQ is Carcinogenic Equivalent, MEF is Mutagenic

Equivalence Factor and TEF is Carcinogenic Equivalence Factor.

Cancer risk assessment

The incremental lifetime cancer risk (ILCR) model (USEPA, 1989) developed for quantitative estimation of exposure risks to PAHs was adopted in this study using the following assumptions:

- i. Exposure to PAHs in the fuel filling stations is via ingestion of dust particles, dermal contact and direct inhalation of particulate matter,
- ii. Overall cancer risk could be estimated by adding up the individual risks calculated for the respective exposure pathways,
- iii. The carcinogenic risk is evaluated based on exposure in reference to the type of land use pattern (i.e. fuel filling stations) over the entire lifetime.

The assessment models recommended by the Risk Assessment Guidance of USEPA (1989) and adapted by Yang *et al.* (2015) and Bandowe and Nkansah (2016) were used to evaluate the ILCR in terms of ingestion, dermal contact and inhalation for both children and adults:

$$ILCRs_{(\text{inhalation})} =$$

$$\frac{CS \times \left(CSF_{\text{inhalation}} \times \sqrt[3]{\frac{BW}{70}} \right) \times IR_{\text{inhalation}} \times EF \times ED}{BW \times AT \times PEF} \quad (3)$$

$$ILCRs_{(\text{dermal})} =$$

$$\frac{CS \times \left(CSF_{\text{dermal}} \times \sqrt[3]{\frac{BW}{70}} \right) \times SA \times AF \times ABS \times EF \times ED}{BW \times AT \times 10^6} \quad (4)$$

$$ILCRs_{(\text{ingestion})} =$$

$$\frac{CS \times \left(CSF_{\text{ingestion}} \times \sqrt[3]{\frac{BW}{70}} \right) \times IR_{\text{ingestion}} \times EF \times ED}{BW \times AT \times 10^6} \quad (5)$$

Where CSF is carcinogenic slope factor (mg/kg/day); $CSF_{\text{Ingestion}}$, CSF_{Dermal} and $CSF_{\text{Inhalation}}$ of B(a)P were addressed as 7.3, 25, and 3.85 re-

spectively, determined by the cancer-causing ability of B(a)P. CS ($\mu\text{g}/\text{kg}$) is the sum of the PAHs concentrations of the samples based on Toxic Equivalents (TEQ) of B(a)P using the Toxic Equivalency Factor (TEF) (Bandowe & Nkansah, 2016).

Bandowe & Nkansah (2016) provides a qualitative description of the ILCR as follows; a

value which is $\leq 10^{-6}$ represents a very low cancer risk, a value ranging from 10^{-6} to 10^{-4} represents a low cancer risk, a value ranging from 10^{-3} to 10^{-1} represents a high cancer risk, and a value $\geq 10^{-1}$ represents a very high risk.

The Table below provides the parameters used in the incremental lifetime cancer risk (ILCR) assessment.

TABLE 1

Parameters used in ILCR Assessment

Exposure variable	Unit	Adult	Child	Reference
Exposure frequency (EF)	day year ⁻¹	350	350	USEPA (2001)
Exposure duration (ED)	year	30	6	USEPA (2001)
Body weight (BW)	kg	70.8	15	Shirazu et al., 2017; Vuvur and Harrison (2017), Bandowe and Nkansah (2016)
Dust ingestion rate ($IR_{\text{ingestion}}$)	$\text{mg}\cdot\text{day}^{-1}$	100	200	USEPA (2001)
Inhalation rate ($IR_{\text{inhalation}}$)	$\text{m}^3\cdot\text{day}^{-1}$	20	10	USEPA (2001)
Dermal adherence factor (AF)	$\text{mg}\cdot\text{cm}^{-2}$	0.07	0.2	USEPA (2001)
Dermal exposure area (SA)	cm^2	5700	2800	USEPA (2001)
Particle emission factor (PEF)	$\text{m}^3\cdot\text{kg}^{-1}$	1.36×10^9	1.36×10^9	USEPA (2001)
Dermal adsorption fraction (ABS)	unitless	0.13	0.13	USEPA (2001)
Averaging life span (AT)	day	$70\times 365=25550$	$70\times 365=25550$	USEPA (2001)

Results and discussion

16 PAH congeners were identified using GC/MS as presented in Table 2. Amongst the 16 PAHs identified, Benzo(a)Anthracene recorded the highest concentrations in both H-VTAs (56.27%) and L-VTAs (53.62%) (Table 2). This agrees with the results of Essumang *et al.* (2016) and Franco *et al.* (2017) which indicate Benzo(a)Anthracene as the commonest PAHs in most environments. The prevalence of Benzo(a)Anthracene presents a carcinogenic risk to the health of the people living or working in the vicinity of the fuel filling stations since Benzo(a)Anthracene is listed by the International Agency for Research on Cancer (IARC) as a Class 2B carcinogenic compound (JRC-IRMM, 2011). The National Cancer Institute Thesaurus (2019) confirms the common sources of Benzo(a)Anthracene production to be wood and soot smoke, gasoline and diesel exhaust, as well as coal tar and coal tar pitch. These common sources are nowhere exclusive from certain human activities including the combustion of gasoline and diesel in vehicles, the burning activities of sawmills, bush burning activities and the singeing activities of abattoirs, all occurring in neighbouring communities to some of the sampling sites.

In the study conducted by Obiri *et al.* (2011) on PAH pollution in road dusts sampled from the Tamale Metropolis, higher mean PAH concentration were recorded in H-VTA as compared to L-VTA. In the current study, it was identified that fuel filling stations in L-VTA rather recorded higher mean PAH concentration than those of the H-VTA (Table 2). Since this study deals with fuel filling stations, a number of factors may come into play to influence the total concentration of PAHs produced. For instance, Gab Energy which is located in the L-VTA recorded the highest concentration of PAHs

amongst all the other fuel filling stations. It was observed that this station was visited more often by heavy-duty vehicles like trucks and buses, and directly served as a parking lot for them overnights. Also, many vehicular users chose to purchase their fuels there because it was newly opened, and the fuel products sold there were relatively cheaper to the public. Moreover, wood and grass combustion could also be a contributing factor to the higher PAH concentrations in the L-VTA since Gab Energy and Star Oil in particular were located near local communities where bush burning was common during the dry seasons.

The mean B(a)P concentration levels recorded in this study were $74,735.22 \pm 41,038.98$ $\mu\text{g}/\text{kg}$ for L-VTA fuel filling stations and $3,890.56 \pm 1,688.48$ $\mu\text{g}/\text{kg}$ for H-VTA fuel filling stations. A study conducted by Nganje *et al.* (2007) on PAH pollution in surface soil present in four fuel filling stations located in the Calabar Metropolis of Nigeria recorded a mean B(a)P concentration ranging from 4 to 680 $\mu\text{g}/\text{kg}$. Moreover, studies have been conducted on PAH pollution in street dust by Obiri *et al.* (2011) in the Tamale Metropolis (Ghana), Essumang *et al.* (2006) in the Kumasi Metropolis (Ghana), and Singh (2014) in Delhi (India) with recorded average B(a)P concentration levels of 10,900 $\mu\text{g}/\text{kg}$, 27,900 $\mu\text{g}/\text{kg}$ and 365.7 $\mu\text{g}/\text{kg}$ respectively. A comparison of the B(a)P concentration levels recorded in this study to those of the aforementioned studies indicated that the B(a)P pollution levels at fuel filling stations in the Tamale Metropolis is higher (Table 2). B(a)P is a common human carcinogen (WHO, 2010). Consequently, its high levels in fuel filling stations of the Tamale Metropolis should be of great concern.

TABLE 2
Mean concentrations of individual PAHs in low and high-vehicular traffic area fuel filling stations

PAHs	Low-Vehicular Traffic Area			High-Vehicular Traffic Area		
	Total Conc. (µg/kg)	Mean Conc. (µg/kg)	Standard Deviation (µg/kg)	Total Conc. (µg/kg)	Mean Conc. (µg/kg)	Standard Deviation (µg/kg)
Naphthalene	787.10	157.42	81.88	38.70	7.74	3.31
Acenaphthene	43,787.20	8,757.44	4,855.22	5,623.30	1,124.66	474.52
Acenaphthylene	421.70	84.34	49.85	144.30	28.86	16.07
Fluorene	305.10	61.02	33.98	62.40	12.48	4.67
Anthracene	8,892.90	1,778.58	937.99	1,658.40	331.68	144.43
Phenanthrene	8,955.10	1,791.02	954.77	1,704.10	340.82	149.76
Fluoranthene	447,821.60	89,564.32	51,392.37	82,482.20	16,496.44	8,952.67
Pyrene	1,782,923.20	356,584.64	220,423.35	945,255.70	189,051.14	93,200.59
Benzo(a)Anthracene	3,176,397.80	635,279.56	278,810.95	1,423,277.80	284,655.56	154,237.67
Chrysene	53,532.80	10,706.56	6,364.05	47,903.90	9,580.78	5,303.57
Benzo(a)Pyrene	373,676.10	74,735.22	41,038.98	19,452.80	3,890.56	1,688.48
Benzo(k)Fluoranthene	1,731.50	346.30	196.29	143.20	28.64	15.95
Benzo(b)Fluoranthene	15,318.20	3,063.64	1,751.32	623.20	124.64	76.86
Dibenzo[a,h]anthracene	989.30	197.86	90.48	233.30	46.66	29.19
Indeno[1,2,3,cd]pyrene	957.10	191.42	91.92	249.60	49.92	30.75
Benzo[g,h,i]perylene	7,035.00	1,407	978.48	451.50	90.30	39.89
Mean concentration (µg/kg)	370,220.73			158,081.53		

The estimated average TEQs and MEQs for the various fuel filling stations were 86,074.267 µg/kg and 77,836.93 µg/kg respectively (Table 3). In comparison to the average TEQs recorded in urban soils from Xi'an (1,150 µg/kg), Guangzhou (2,763 µg/kg), and Tianjin (4,554 µg/kg) (Bandowe and Nkansah, 2016) which are known to be megacities in China, it provided an indication that PAH concentration

levels in dust particles at fuel filling stations in Tamale Metropolis presented significantly higher potential carcinogenic effects. It is however important to note that since samples in this study were directly collected from fuel filling stations, there is a greater probability of recording high average values of TEQs and MEQs due to a larger contribution of localised source of pollution.

Amongst the selected fuel filling stations, Gab Energy (483,651.14 $\mu\text{g}/\text{kg}$ and 452,219.52 $\mu\text{g}/\text{kg}$) and Star Oil (213,162.58 $\mu\text{g}/\text{kg}$ and 188,536.77 $\mu\text{g}/\text{kg}$) of the L-VTAs recorded the highest estimates of TEQs and MEQs respectively. This could be attributed to the fact that the two fuel filling stations served as parking lots for heavy-duty vehicles, and were also located close to local communities which oftentimes practiced bush burning during the

dry seasons, leading to the high accumulation of PAHs in their vicinity. Among the H-VTA fuel filling stations, Goil [Melcom Branch] (124,849.21 $\mu\text{g}/\text{kg}$ and 104,008.07 $\mu\text{g}/\text{kg}$) recorded the highest TEQ and MEQ values respectively. The high TEQ and MEQ values presented by Goil [Melcom Branch] could be as a reason of its location in the centre of the Tamale Business District, serving the greater portion of the public.

TABLE 3
Carcinogenicity and mutagenicity of identified PAHs

Area Category	Name of Fuel Filling Station	Average PAHs ($\mu\text{g}/\text{kg}$)	TEQ ($\mu\text{g}/\text{kg}$)	MEQ ($\mu\text{g}/\text{kg}$)
High-Vehicular Traffic Area	Goil (Melcom Branch)	192,454	124,849.21	104,008.07
	Total (CM Branch)	587,067	37,158.94	32,016.17
	Goil (HR. Branch)	5,635	507.50	420.04
	Total (HR. Branch)	11,658	1,134.51	933.13
	Shell (Aboabo Branch)	396	0.78	0.91
Low-Vehicular Traffic Area	Gab Energy	4,147,852	483,651.14	452,219.53
	Star Oil	1,772,030	213,162.58	188,536.77
	Compass Oleum	1,243	111.53	94.20
	Naagamni	1,808	164.78	138.38
	Blanko Oil	597	1.71	2.11

Depending on the TEQs of the fuel filling stations as well as some relevant parameters mentioned in Table 1, a potential risk assessment framework was designed to estimate the risks posed to the people (adults and children) living and working in the vicinity of the fuel filling stations via exposure routes of inhalation, ingestion and dermal contact (Table 4). For adults, the estimated cancer risk via inhalation route of exposure ranged between $2.57\text{E}-10$ and $1.60\text{E}-04$, that of dermal contact ranged between $5.89\text{E}-06$ and 3.65, and that of ingestion between $3.32\text{E}-06$ and 2.06. For children, the estimated cancer risk via inhala-

tion route of exposure ranged between $7.24\text{E}-11$ and $4.49\text{E}-05$, that of dermal contact ranged between $4.65\text{E}-06$ and 2.89, and that of ingestion between $3.73\text{E}-06$ and 2.32. Of both populations, PAH exposure via inhalation route recorded the lowest value of cancer risk, while dermal route of exposure recorded the highest value of cancer risk. These findings agree with Yang *et al.* (2015) in China, Bandowe and Nkansah (2016) in Ghana, Tarafdar and Sinha (2018) in India, and Ghosh and Maiti (2019) in India, who stated that PAH exposure in urban soils via inhalation route posed the least potential cancer risk whereas exposure via der-

mal contact recorded the highest risk for both adults and children. This is explained by the lower concentrations of the LMW PAHs such as Naphthalene, Phenanthrene, Anthracene and Fluorene recorded in this study. Although these PAH congeners are normally present in the atmosphere because of their high volatility, they are however present in lower levels (Law *et al.*, 2002). In contrast, the HMW PAHs such as Pyrene, Benz[a]anthracene, and Benzo[a]pyrene which contributed greatly to the total PAH concentrations recorded in this study are less volatile with low mobility in organic matter (e.g. B[a]P log K_{oc} = 6.6–6.8) (WHO, 2010; Stogiannidis and Laane, 2015).

A cancer risk value ranging from 10^{-3} to 10^{-1} represents a high cancer risk, while a value $\geq 10^{-1}$ represents a very high risk (Bandowe and Nkansah, 2016). The recorded cancer risk values in this study for both populations via ingestion and dermal pathways of exposure do explain that the associated population are potentially exposed to a high to very high cancer risk. The exposure of both groups to the identified PAHs can be linked with an increased risk of cancer development in different tissues of the body, including the lung, bladder, stomach, and skin (including the scrotum), depending on the mode of exposure and the type of PAH they are exposed to (CCME, 2010). The higher carcinogenic risk values recorded by the adults via both inhalation and ingestion exposure pathways could be associated with their longer exposure duration especially during working hours. However, children

are at a higher risk of acquiring cancer than the adult population due to their smaller body weight and their less developed immune system necessary to detoxify toxic chemicals. Once the PAHs are able to get into the body system of the children, they become more active in their systems and thus can interfere with organ development and proper operation of the central nervous system (Iwegbue *et al.*, 2016). Moreover, the WHO (2010) stated that the incidental ingestion of soil by adult males is estimated to be of the order of a few milligrams per day, but that of the children were of the order of 100 mg/day, re-emphasising the high risks the children population may be exposed to.

Amongst the fuel filling stations, Gab Energy recorded the highest carcinogenic risks of 5.71 (adults) and 5.21 (children), and then followed by Star Oil with 2.52 (adults) and 2.29 (children). The least carcinogenic risk was recorded at the Shell station with values of $9.21\text{E-}06$ (adult) and $5.33\text{E-}05$ (children). Ultimately, the average ILCR for adults (1.02) amongst all the fuel filling stations in both vehicular areas was slightly greater than that of the children ($9.26\text{E-}01$). An unpaired t-test conducted presented a p-value of 0.9226 at 95% confidence level, indicating that there was no significant difference existing between the mean carcinogenic risk values recorded between adult and children populations. This gives a pressing issue of concern for both age groups, especially the children group who are more prone to effects from exposure to toxic chemicals.

TABLE 4
The potential carcinogenic risks and exposure pathways for each fuel filling station

Area Category	Name of Fuel Filling Station	Inhalation (ILCRs _(inhalation))		Dermal Contact (ILCRs _(dermal))		Ingestion (ILCRs _(ingestion))		Carcinogenic Risk Σ(ILCRs)	
		Adult	Child	Adult	Child	Adult	Child	Adult	Child
High-Vehicular Traffic Area	Goil (M. Branch)	4.12E-05	1.16E-05	5.31E-01	5.98E-01	9.43E-01	7.45E-01	1.47E+00	1.34E+00
	Total (CM Branch)	1.23E-05	3.45E-06	1.58E-01	1.78E-01	2.81E-01	2.22E-01	4.39E-01	4.00E-01
	Goil (HR. Branch)	3.74E-07	4.71E-08	2.16E-03	2.43E-03	3.83E-03	3.03E-03	5.99E-03	5.46E-03
	Total (HR. Branch)	1.67E-07	1.05E-07	4.83E-03	5.43E-03	8.57E-03	6.77E-03	1.34E-02	1.22E-02
	Shell (AB)	2.57E-10	4.49E-05	3.32E-06	3.73E-06	5.89E-06	4.65E-06	9.21E-06	5.33E-05
Low-Vehicular Traffic Area	Gab Energy	1.60E-04	1.98E-05	2.06E+00	2.32E+00	3.65E+00	2.89E+00	5.71E+00	5.21E+00
	Star Oil	7.03E-05	1.04E-08	9.07E-01	1.02E+00	1.61E-00	1.27E+00	2.52E+00	2.29E+00
	Compass Oil	3.68E-08	1.53E-08	4.74E-04	5.34E-04	8.43E-04	6.66E-04	1.32E-03	1.20E-03
	Naagamini	5.44E-08	1.59E-10	7.01E-04	7.89E-04	1.25E-03	9.83E-04	1.95E-03	1.77E-03
	Blanko Oil	5.64E-10	4.49E-05	7.27E-06	8.19E-06	1.29E-05	1.02E-05	2.02E-05	6.33E-05
Average ILCRs		2.84E-05	1.25E-05	3.66E-01	4.13E-01	6.50E-01	5.14E-01	1.02E+00	9.26E-01

Conclusions

The identified PAHs in the dust particles present in the selected fuel filling stations were of high concentrations, especially Benzo(a)Anthracene and Pyrene which are of public health concern. Benzo(a)Pyrene which is identified as a common carcinogen recorded higher concentration levels when compared with similar studies conducted in major cities like Kumasi of Ghana, and Delhi of India. The exposure of associated population to the high PAH concentration levels recorded in this study is more through dermal pathway than that of ingestion or inhalation. The results indicate that there is a high potential risk of cancer to both adults and the children who work, patronise, or live around the fuel filling stations. It is therefore recommended that government agencies like the Environmental Protection Agency (EPA) and the National Petroleum Authority (NPA) take up effective measures to help control the high level of PAH pollution occurring in the fuel filling stations.

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