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Research Article

HEALTH AND BODY COMPOSITION OF COSMETOLOGY PROFESSIONALS IN YENAGOA, SOUTHERN NIGERIA

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Abstract

Cosmetologists, typically employed in retail or home-based salons, offer a range of beauty services such as hair shampooing, styling, manicures, pedicures, and facial treatments. With millions employed worldwide, hairdressers and cosmetologists (HC) face regular exposure to chemicals in various hair products, including shampoos, permanent wave solutions, hair dyes, and sprays. Some of these chemicals have been linked to reproductive toxic effects. This study, conducted in Yenagoa Local Government Area (YELGA), Bayelsa State, Nigeria, aimed to assess the health impact of these chemical exposures on cosmetologists. Yenagoa is located in the southern part of Nigeria, between Latitude 4°15' North and 5°23' South, and Longitude 5°22' West and 6°45' East. The study was approved by the Ethical Committee of Niger Delta University, Wilberforce Island, Bayelsa State, and adhered to the principles of the Helsinki Declaration of 1975 (revised in 2008). Informed consent was obtained from all participants, who understood the reasons for providing blood and urine samples. A total of 25 urine and blood samples were randomly collected from cosmetologists working in Yenagoa.

The study focused on common chemicals such as nitrosamines in hair dyes, toluene in nail polish, and formaldehyde in both hair dye and nail polish. Despite the exposure to these chemicals, the study found that the health status of the cosmetologists in YELGA did not show significant abnormalities. The biomarkers examined were not sensitive enough to detect health issues that may result from long-term chemical exposure. The abnormalities observed may be attributed to individual factors such as overall health status, genetic predisposition, lifestyle choices, and personal hygiene, rather than direct chemical exposure.

Keywords: Cosmetology, Anthropometry, Bicarbonate, Formaldehyde, Nitrosamine, Toulene

INTRODUCTION

Cosmetologists are generally defined as individuals who work in retail- or home-based salons and provide a wide range of beauty services, including hair shampooing and styling, manicures, pedicures, and scalp and facial treatments. Hairdressing and cosmetology are common occupations, and several million individuals are employed as hairdressers and cosmetologists (HC) worldwide (European Agency for Safety and Health at Work 2014). Workers in the hairdressing and cosmetology professions are predominantly women, and many of these women are of childbearing age (Halliday-Bell et al., 2009) and begin working before considering family planning (Baste et al. 2008). Therefore, this situation raises concerns that these women of reproductive age could be susceptible

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to the effects of exposure to potential reproductive toxins. Several studies of HC have suggested that their work might adversely affect their reproductive health (Herdt-Losavioet al., 2009; Ronda et al., 2010; Jørgensenet al., 2013; Quachet al., 2014), although various studies have reported conflicting findings. For example, several studies have reported that HC have an increased risks of infertility (Baste et al., 2008), a time to pregnancy of >12 months (Kersemaekers et al. 1997), spontaneous abortion (Ronda et al., 2010), low birth weight (Halliday-Bell et al., 2009; Herdt-Losavioet al., 2009), and preterm delivery (Halliday-Bell et al., 2009), compared to women in other occupations or in the general population. However, other studies have found little or no evidence of an increased reproductive health risk among female hairdressers (Hougaardet al., 2006; Gallicchioet al., 2011).

As a profession, cosmetology is predominantly female inclined, most of whom are of reproductive age. There are more than one million women registered and licensed as cosmetologists in the United States and roughly several million more work as hair stylists. Among cosmetologists, hairdressers and nail technicians make up a large part of the working population. Many cosmetologists begin their careers before reproductive age and before family planning, which may put them at higher risk for reproductive health effects from exposure to workplace cosmetology chemicals. The most common chemicals mentioned in select studies about hair dye and nail polish were nitrosamines in hair dye, toluene in nail polish, and formaldehyde in both hair dye and nail polish.

NITROSAMINES

The primary ingredient in hair dyes is aromatic amines, which are precursors of nitrosamines (McCall et al., 2005). Nitrosamines require bioactivation and have shown mutagenicity in vitro and carcinogenic properties in vivo (Holly, Bracci, Hong, Mueller, & Preston-Martin, 2002). The actual reproductive risk of nitrosamines is unclear due to limited data (Kersemaekerset al., 1995).

TOLUENE

The organic solvent toluene, a common ingredient in nail polish and among the most common exposures in the workplace, has been linked to less fetal growth and shorter pregnancy duration (Hannigan and Bowen, 2010). Rapidly absorbed through the lungs, toluene vapors are distributed to highly perfused and fatty tissues. Organic solvents such as toluene have an affinity for lipid-rich tissues and readily cross the placental barrier (Bukowski, 2001). Little is known about the mechanisms of action for toluene; it is unclear how it is absorbed and distributed through the body (Hannigan and Bowen, 2010).

FORMALDEHYDE

Formaldehyde is found in both hair dye and nail polish. Formaldehyde is considered a human carcinogen by the IARC, the U.S. National Toxicology Program (NTP), the U.S. Environmental Protection Agency (EPA), and the Occupational Safety and Health Administration (OSHA). Formaldehyde has been associated with nasal cancers in workers exposed in occupational settings (Agency for Toxic Substances and Disease Registry [ATSDR, 1999; EPA, 2010; NTP, 2010). Formaldehyde has been named a Group 1 carcinogen by the IARC, meaning that "there is sufficient evidence in humans for the carcinogenicity of formaldehyde" (WHO and IARC, 2006). The NTP classified formaldehyde as "reasonably anticipated to be a human carcinogen," although there is currently a proposal to reclassify formaldehyde as "known to be a human carcinogen" (NTP, 2010). Reproductive and developmental toxicity have been speculated to be associated with formaldehyde for some time, but this has not been confirmed. Although a small number of human studies have suggested that formaldehyde exposure may

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cause reproductive toxicity, the current understanding of the mechanisms of action is limited. Currently, formaldehyde's mechanisms of action are proposed to induce reproductive and developmental toxicity via genotoxicity, oxidative stress, disruption of the activity of proteins, enzymes, and hormones important for the maturation of the male reproductive system, apoptosis, and DNA methylation (Duong et al., 2011). These mechanisms are hypothetical and require validation, particularly for reproductive system effects (Duong et al., 2011).

The majority of studies showing occupational reproductive effects among hairdressers and nail technicians suggest that chemical exposure is the probable cause of these findings (Ronda et al., 2009). However, the effects of exposures to mixtures of chemicals, such as those found in salons, are largely unknown (Hougaardet al., 2006). The evidence is inconclusive regarding hair dyes and potential human carcinogenicity. Regarding carcinogenicity, the primary concern is chemical absorption through human skin (McCallet al., 2005). However, the level of absorption depends on the extent of dermal contact (Kersemaekerset al., 1995). Hair dyes are found in many forms (i.e., liquids, creams, gels, shampoos, and rinses), and the method of application may affect exposure. As an example, permanent cream dyes are commonly applied with a brush, whereas other dyes are more often worked into the hair by hand. The major route of entry for hair dye chemicals is cutaneous absorption (Kersemaekerset al., 1995). The main concern with nail polish is inhalation exposure. Working in close proximity to multiple agents exposes nail technicians to potential sensitizers and respiratory irritants.

(Reutmanet al., 2009). Nail technicians commonly inhale and breathe harmful vapors, dusts, or mists, and can get the product on their skin or in their eyes or can swallow the product if it is accidentally transferred onto food or cigarettes (OSHA, 2013). These exposures can accumulate if the products are used daily or if poor ventilation exists in salons (OSHA, 2013). Chemical exposure over time is a concern. Several other potentially hazardous chemicals can affect workers in nail salons. Acetone (nail polish remover) can cause headaches, dizziness, and eye, skin, and throat irritation. Acetonitrile (fingernail glue remover) can cause nose and throat irritation, breathing problems, nausea, and vomiting. Ethyl methacrylate (artificial nail liquid) can cause asthma, eye, skin, nose, and mouth irritation, and reproductive effects for the fetus if exposure occurs during pregnancy (OSHA, 2013).

MATERIALS AND METHODS Study Area

The study was conducted in Yenagoa (YELGA), Bayelsa State. Bayelsa State is a cosmopolitan state in the southern part of Nigeria, which is geopolitically located within Latitude 415 North, 523 South, Latitude 522 West and 645 East. It has an area of 706km. It shares boundaries with Delta State on the North, Rivers State on the East with the Atlantic Ocean on the West and South. The official language is English language but the major language spoken is the Izon language.

Ethical clearance

The study which got the ethical approval from the Ethical Committee of Niger Delta University, Wilberforce Island, Bayelsa state, Nigeria. It was carried out with compliance to the principle of Helsinki declaration of 1975 as revised in 2008. Informed consent was obtained from all the recruited volunteers who were made to know the reasons their blood and urine samples were needed for this research.

Sample collection

A total number of twenty-five (25) Urine and Blood samples were randomly selected from cosmetologist in yenagoa. Sample Size Calculation

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Taro Yamane formula with 95% confidence level according to Yamane, 1973, was used to determine the sample size of this research (Yamane, 1973). Calculation of sample size using the Taro Yamane method n = N + N(e)

Where n = sample size required N = Population size e = Allowable error which is between 0.01-0.05 Assuming N=25 and e =0.1, therefore n = 100 k+100(0.1)2 Therefore, n = 25

Experimental design: Venous blood and Urine were collected from 13 males and 12 females making a total of 25 adult individuals. The blood was used to assay for Packed Cell Volume (PCV) and urine was used for urinalysis. **Materials, Equipment and Reagent**

Materials: Capillary tubes, sealant, Microhaematocrit reader, EDTA anticoagulated container, plain universal container, methylated spirit, cotton wool,5ml syringe, and Combi 11.

Equipment: Microhaematocritcentrifuge (Vanguard V6000).

Packed Cell Volume (PCV) Principle of test

Blood specimen is centrifuged in a sealed capillary tube and Packed Cell Volume is determined by a special haematocrit reader and gives the result as percentage.

Materials provided with the kit:

- Microhaematocrit reader
- Capillary tubes
- Sealant
- Marker pen

Materials required but not provided:

Non

Storage of Test Kits and Instrumentation Urinalysis Principle of test

The Urinalysis Reagent Strips (Urine) are firm plastic strips onto which several separate reagent areas are affixed. The test is for the qualitative and semi-quantitative detection of one or more of the following analytes in urine: Specific Gravity, pH, Leukocytes, Nitrite, Protein, Glucose, Ketone Bodies, Urobilinogen, Bilirubin, Blood, and Ascorbic Acid.

Reagents

Materials provided with the kit:

Medi Test Combi 11

Materials required but not provided:

Non

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Statistical Analysis

All the values were expressed as mean \pm standard deviation of triplicate determinations. The data obtained from the analysis was subjected to one way Analysis of variance (ANOVA) and t-test, the means were separated and compared at 95% confidence level.

RESULTS

Table 4.1: Socio-demographic distribution

	TEST	CONTROL
SEX		
Male	13	0
Female	12	
Sub-total	25	
MARITAL STATUS	23	
Single	02	18
Married	25	25
Sub-total	01	21
TYPE OF DIET	02	04
Carbohydrate	22	25
Protein		02
	25	00
Carbohydrate/protein	02	23
Sub-total	01	25
WORK HOURS	00	00
7am – 9pm	02	21
8am – 4pm	18	01
8am – 5pm	02	00
8am – 8pm	25	03
8am – 9pm		00
8am – 10pm		25
Sub-total		

Table 4.2: general descriptives statistics of test against control category. Results are expressed as mean \pm standard deviation and statistical significance set at P \leq 0.05, which is calculated at 95% confidence interval.

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		TEST	CONTROL	P-VALUE	REMARK
AGE	24.08±3.65	27.92±11.96	0.001	S	
PCV	36.88±9.06	34.68 ± 7.60	0.121	NS	
BLOOD PI	RESSURE	115/72±11.54	119/72±8.67	0.225	NS
BMI	21.94±3.01	25.83±5.68	0.225	NS	
PULSE RA	TE 72.4	8±13.38 75.24±	13.86 0.11	11 NS	
HEIGHT	5.48 ± 0.24	5.44 ± 0.46	0.291	NS	
WEIGHT	60.96±10.88	8 68.64±11.48	0.254	NS	

Table 4.3: gender based analytics of test category. Results are expressed as mean \pm standard and statistical significance set at P \leq 0.05, which is calculated at 95% confidence interval.

deviation

	MAL	E	FEMALE	P-VALU	JE REN	IARK	
AGE	25.46±	±4.21	22.58±2.23	0.217	NS		
PCV	39.15±	<u> </u>	34.42±10.11	0.118	NS		
BLOC	D PRI	ESSURE	117/75±13.79	113.	.69±10.73	0.255	NS
	22.75± ±13.90	±2.40 0.411	21.06±3.44 NS	0.121	NS	PULSE RAT	ΓE 69.54±12.71
HEIG	HT	5.56±0.24	5.40±0.22	2	0.191	NS	
WEIG	HT	66.00±10.	13 55.50±9.1	14	0.001	S	

Table 4.3: Urine analysis demographics of test against control category.

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COLOUR		
Amber	22	
Straw	00	21
Pale	02	01
Amber/cloudy	01 25	03
Sub-total	00	
LEUCOCYTE	20	25
Negative		25
(+) 00	00	
(++) 05	00	
Sub-total	25	25
NITRITE		
Negative 25	25	

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Positive	05		
Sub-total	00 25	5	
PH	25		
5	14		12
6	06		08
6.5 7	01	00	
8	02	03	
Sub-total	02		02
BLOOD	25 25		
Negative	25	22	
Trace	24	23	
(+)	00 01	01 01	
Sub-total	25	25	
PROTEIN		23	20
Negative	23		20
Normal	00		02
(+)	02		02
(++)	00		01
Sub-total	2525		
GLUCOSE	25		21
Negative	00		02
Normal	00		02
(+)	25		02
Sub-total	25 25		
ASCORBIC ACID	24		25
Negative	00	00	23
(+)	01	00	
(++)	25		25
Sub-total	25		23
KETONE	00		02

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	Research A	rticle		
Negative			25	
Normal			25	
Sul	o-total		22	25
UROBILINOGI	EN		03	00
Normal			25	25
(+)				
Su	ıb-total			
BILIRUBIN No	egative 22	23 Normal 0	00	
(+) 02	00			
(++) 01 25	00	Sub-total 25		

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DISCUSSION

From this study, blood and urine samples were collected from 25 individuals in the cosmopolitan city of yenagoa. From the 25 individuals, 13 were males and 12 werefemales, 23single and 2 married. 1 individual was on a carbohydrate diet, 2 were on a protein diet, and 22 were on a carbohydrate/protein diet. Out of the 25 individual sampled, 2 worked from 7am-9pm, 1 worked from 8am-4pm, 2 worked from 8am-8pm, 18 worked from 8am9pm and 2 worked from 8am10pm.

From table 4.2 It was observed that the p-value of the average age(25.46 ± 4.21 years) of the individuals is 0.001 which is significant to the study as statistical significance was set at P \leq 0.05. It was also observed that P-value of the average Packed Cell Volume ($36.88\pm9.06\%$) is 0.121, Blood pressure($115/72\pm11.54$ mmHg) is 0.225, Body Mass Index (21.94 ± 3.01 kg/m2) is 0.225, Pulse rate(72.48 ± 13.38 bpm) is 0.111, Height(5.48 ± 0.24 ft) is 0.291 and Weight(60.96 ± 10.88 kg) is 0.254, all of which were seen to be statistically insignificant based on the study carried out.

Table 4.3 shows gender-based analytics of the test group. It was observed that statistical significance was evident only in weight for both males and females which was expressed as 0.001 but was not observed for other groups which were all statically insignificant with a $P \le 0.05$.

Table 4.4 shows the urine analysis demographic of the test category and results were obtained for different analytes. Out of the Twenty-five (25) urine samples, 22 were amber colored, 2 were pale colored and 1 was amber/cloudy. (20) reacted negative for leucocyte and 5 reacted positive (++) for leucocyte which is suggestive of an ongoing Urinary tract infection (UTI) in those individuals. 20 samples reacted negative for nitrite and 5 reacted positive for nitrite which is also suggestive of a UTI. 14 samples had a pH of 5.0, 6 samples had a pH of 6.0, 1 sample had a pH of 6.5, 2 samples had a pH of 7.5, and 2 samples had a pH of 8.0. 24 samples reacted negative for blood and 1 reacted positive (+) which may be due to an underlying health condition. 23 samples reacted negative for protein and 2 samples reacted positive (+) for protein.

25 samples reacted negative for glucose. 24 samples reacted negative for ascorbic acid and 1 reacted positive (++) which suggest an intake of vitamin C before the sample was collected. 25 samples reacted negative for ketones. 22 samples had normal urobilinogen and 3 had abnormal (+) urobilinogen. 22 samples reacted negative for bilirubin 2 reacted positive (+) and one reacted positive (++) for bilirubin.

CONCLUSION

Cosmetologists are individuals who work in retail or home-based salons and provide a wide range of beauty services, including hair shampooing and styling, manicures, pedicures, and scalp and facial treatments. This exposes them to different chemicals and toxic substances which can be hazardous to their reproductive health. The most common chemicals mentioned in this study are nitrosamines in hair dye, toluene in nail polish, and formaldehyde in both hair dye and nail polish. This study, shows that the health status of cosmetologists residing in yenagoa local government area are not affected by the chemicals they are exposed to, due to the fact that the parameters worked on are unable to pick out certain abnormalities that may determine the hormonal and electrolytes levels of the individuals, which are a major pointer to determine hormones (oestrogen and testosterone) which are responsible for reproduction and electrolytes (Potassium, Sodium, Chloride and Bicarbonate) that are essential for basic life functioning. The abnormalities observed from this study may arise from an individual's overall health status, genetic predisposition, lifestyle and personal hygiene.

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RECOMMENDATION

- Cosmetologist should wear protective gear (latex gloves, nose mask, face shields, hair nets etc.) to prevent exposure to toxic chemicals.
- Working hours should be adjusted as long working periods stresses the body and may affect overall health and reproductive status.
- Another study should be carried out and should include hormonal assay of the individuals to ascertain their hormonal status.
- Further studies should be carried out to better understand the effect of different chemicals (toluene, formaldehyde, nitrosamines) on reproductive and health status.

COMPETING INTEREST

No competing interest.

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