INTRODUCTION

Painting is an age-long profession. Painters are often occupationally exposed to substances in paints (such as solvents and metals) which may be deleterious to their health due to chronic poisoning [1,2]. Paints are a mixture of solvents, pigments, organic compounds, and other additives. Most of these compounds are volatile and easily spread into the environments as paint mists during spraying activities [3]. Different paints of varying chemistry have been in use for both domestic and industrial purposes. Paints and its constituents such as organic solvents, including aromatic hydrocarbons (mainly toluene), aliphatic hydrocarbons, alcohols, ketone and esters, and metals such as aluminum, cobalt, chromium, titanium, and lead may cause unfavorable cellular effects to occupationally exposed individuals [4,5]. Despite the fact that the International Agency for Research on Cancer has not considered some of these compounds contained in paints as being carcinogenic, some individual metals or their mixture may contribute to an increased risk of cancer in exposed individuals hence the classification of paints as a Group 1 carcinogen [6-8]. Furthermore, exposure to these organic solvents, metals, and other potentially mutagenic compounds in paints (such as phthalic acids and chlorophenols) is known to be capable of inducing DNA damage and chromosomal changes sometimes manifesting as micronuclei [1,9].

The examination of exfoliated buccal epithelial cells using the buccal cell micronuclei (MN) test as a biomarker serves as a minimally invasive method for bio-monitoring of human exposure to potentially genotoxic environmental pollutants [5]. In experimental animal models, MN assay has also been employed to monitor the damage of chromatin materials caused by exposure of animals to chemicals. A previous study documented an increase in the percentage frequency of micronucleus in animals treated with arsenite, which indicates chromosomal damage [10]. Another study also employed the in vivo micronucleus assay in bone marrow cells of male and female Sprague-Dawley rats to evaluate genotoxicity [11]. MN are extra-nuclear bodies that contain damaged chromosome fragments and/or whole chromosomes that were not incorporated into the nucleus after cell division [12]. An increased number of MN in exfoliated buccal cells is often an indication of genetic damage [13]. The presence of MN can be seen as an early warning sign for the potential risk of developing long-term health problems [14].

Car spray workers in Enugu metropolis, Nigeria constitute a major workforce with a considerable population. In the course of their work, these painters are exposed to paint mist or vapor primarily through inhalation, dermal absorption, and ingestion [15]. Mishandling of paints, inhalation of paint materials, and inadequate use of personal protective equipment (PPE) observed among some car spray painters have increased their risk of exposure to these harmful chemicals which are potential threats to health [16]. In view of this, the present study was, therefore, designed to evaluate cytogenetic damage of exfoliated cells of the buccal mucosa of car spray painters by determination of MN frequency using the MN test and binucleated cells.

METHODS

Ethical consideration

Approval for the study was obtained from the Health Research and Ethics Committee of the University of Nigeria Teaching Hospital Ituku-Ozalla,
with approval number:-[NHREC/05/01/2008B-FWA00002458-1RB00002323]. Before recruitment, the details of procedures involved in the study were explained to the study participants, volunteers who consented to participate in the study signed an informed consent form.

Study area and design
The study was conducted in Enugu, the capital city of Enugu state, Nigeria, and lasted for 6 months from June 2018 to December 2018. The study adopted a cross-sectional design.

Study participants
The study participants were adult males between the ages of 18 and 65 years, recruited from Enugu metropolis, South East, Nigeria. A total of 352 individuals participated in the study, 200 of the participants were car spray painters directly exposed to paints while the other 152 neither worked as car spray painters nor were exposed to paints and served as control. A structured questionnaire was used to obtain information such as age, smoking habit, alcohol consumption, exposure to radiation, and occupational exposure characteristics of the selected participants.

Inclusion criteria
Adult males between the ages of 18 and 65 years, who were apparently healthy with no oral lesion that has worked as car spray painters for at least 6 months and consented to participate in the study.

Exclusion criteria
Individuals outside the age range of 18–65 years, those with oral lesions and had been exposed to X-ray radiations in the past 1 year and non-consenting individuals were excluded from the study.

Sample collection, staining, and evaluation
Before sample collection, the participating individuals thoroughly rinsed their mouth with water to remove any unwanted debris. Buccal cells were obtained by scraping the inside of both cheeks gently with a sterile wooden spatula; the sample was suspended in a labeled universal container containing 95% ethanol which served as a fixative. The samples were prepared in a laboratory by transferring the sample into centrifuge bottles and centrifuging at 5000 rpm for 5 min, the supernatants were decanted, and the sediments were smeared on poly-L-lysine charged grease free slides. They were allowed to air dry and stained with hematoxylin and eosin staining technique and then examined using a light microscope at x40 magnification to identify and record the number of buccal cells with MN and BNC. At least 1000 intact buccal epithelial cells per individual were examined and scored using the criteria described by Tolbert et al. [17] and the MN and BNC frequencies recorded. MN were counted by two independent observers who evaluated the slides twice at different intervals to minimize intra- and inter-observer bias.

Immunocytochemical (ICC) staining and evaluation for p53 and Ki-67
ICC staining of buccal smears was carried out according to previously described methods [18,19]. Monoclonal antibodies Ki-67 and p53 were employed. Expose Mouse and Rabbit Specific Horseradish Peroxidase/Diaminobenzidine detection immunohistochemistry kit was employed for immunostaining while detection of immunoreactivity was performed according to the manufacturer’s instruction.

Procedure
Smears were hydrated by passing through 50% ethanol for 15 s and then to distilled water for 15 s. Slides were arranged in slide racks and treated in protein block and biotin block solutions for 25 min in each solution. Thereafter, they were arranged on a staining rack and flooded with phosphate buffer saline (PBS) solution to prevent drying. The smears were drained afterward, the exact portions of smears on slides were carefully ringed with a hydrophobic pen, and diluted antibodies (1:100) (anti-Ki-67 and p53) were applied onto smears with the aid of Pasteur pipette and allowed to incubate at room temperature for 1 h. After incubation in the primary antibodies, the smears were washed with PBS, flooded with a secondary antibody for 25 min, washed with PBS, drained and diaminobenzidine DAB were applied for 5 min. Finally, the smears were washed with PBS; counterstained in Harris Hematoxylin for 5 min, washed in water and was differentiated by dipping 10 times in 1% acid alcohol. Slides were later washed and blued in tap water, dehydrated by passing through 70%, 90% and two changes of absolute ethyl alcohol for 15 s each, cleared in xylene and mounted in DPX. Ki-67 and p53 positive immune control sections were also stained alongside test and control smears. The ICC staining was semi-quantitatively scored, according to Zlobec et al. [18].

Statistical analysis
Statistical analysis was performed using Statistical Package for the Social Sciences version 20.0. Data obtained from the assay were expressed as the mean ± standard deviations. Student’s t-test (two-tailed) was used to compare nuclear abnormalities among test and control groups. The level of significance was set at *p<0.05. Overall effects of age, exposure duration, alcohol consumption, and smoking were determined using one-way analysis of variance, followed by post-hoc multiple comparisons.

RESULTS
Effects of exposure on buccal cell nuclei
Nuclear damage indices assessed in the buccal cells of study participants were MN and BNC. (Fig. 1a–b) is representative micrographs showing normal buccal cells (Fig. 1a), cells with MN (Fig. 1b) and BNC (Fig. 1c).

Table 1 compared the mean values of the frequencies of MN and BNC in control and exposed car spray painters. Car spray painters were found to have significantly higher MN (p=0.000) and BNC (p=0.000) when compared to the control subjects.

Variations of nuclear abnormalities (MN and BNC) based on demographics, lifestyle, and exposure characteristics
Car spray painters were further subdivided according to age, occupational exposure factors, and lifestyle to assess the influence of these factors on the frequency of MN and BNC (Table 2). With respect to age, the result showed a statistically significant increase in the MN of car spray painters in the age group >35 years when compared with those in the lower age groups <25 years and 26–35 years.

Car spray painters who have worked for 5–10 years, 11–15 years, and >15 years had significantly higher MN (p=0.000) when compared with those who have worked for <5 years. Duration of exposure and use of PPE did not significantly affect the distribution of MN and BNC, though cars spray painters who worked >10 h daily had an increased frequency of MN and BNC though not statistically significant. Cigarette smoking and alcohol consumption significantly increased MN frequency (p=0.000) of car spray painters.

Effects of exposure on immunoreactivities for p53 gene
Buccal cell cytology of both car spray painters and controls participants was negative for p53 immunoreactivities (Fig. 2a and b). The positive stain immune control section, however, showed p53 reactivity indicated using arrows (Fig. 2c).

Effects of exposure on immunoreactivities for Ki-67 gene
There was also no expression of Ki-67 in the buccal cell nuclei of both car spray painters and control participants (Fig. 3a and b). The positive stain immune control section shows expression of Ki-67 (Fig. 3c).

DISCUSSION
Occupational and environmental exposure to hazardous chemical agents poses a great risk to human health and has become a global concern [20,21]. Buccal epithelial cells are often the first to come in contact with airborne pollutants in occupationally exposed individuals, especially when exposure is through inhalation or ingestion. These cells are capable of metabolizing proximate carcinogens into reactive products [22].
In the present study, both car spray painters and the unexposed control group had MN and BNC in their exfoliated buccal epithelial cells. However, the car spray painters had significantly higher frequencies of buccal cell MN and BNC when compared to the unexposed control subjects. The result obtained from the present study corroborates with the findings of a preliminary study conducted on automobile spray painters in the coal camp mechanic village in Enugu [23]. More so, previous studies have documented similar increased levels of MN and other nuclear abnormalities in buccal epithelial cells of workers occupationally exposed to potentially genotoxic substances [9,24-26].

With reference to age, the result showed that car spray painters older than 35 years had the highest frequency of MN and BNC. Furthermore, there was a significantly higher MN frequency in the buccal cells of car spray painters who were older than 35 years when compared to those <35 years. This finding suggests that there may be an association between an individual's age and the frequency of MN, as previously documented [26]. In line with this finding, a previous study also reported a significant change in MN frequency due to age in a study involving automobile painters in South India [27]. In contrast; however, a previous study reported no significant change in MN frequency in relation to age among gas station attendants [16].

In the present study, a greater percentage of the exposed participants spent 5–10 h daily in their workplace, this may have also attributed to the significantly higher frequency of MN among those that have worked as car spray painters for 5–10 years, 11–15 years, and above 15 years when compared to those who have worked for <5 years. This suggests a positive correlation between the increased MN frequency and the number of years of exposure/ Frequency of exposure which is similar to the findings documented by a previous study which showed an increased MN frequency with a positive correlation with increased period of exposure in motor garage workers [2]. Car spray painters who have worked for more than 15 years had a higher frequency of BNC when compared to those who have worked for <5 years. Even though the finding was not significant, it is of importance because an increase in BNC frequency of BNC is considered as an indicator of cytotoxicity [1].

With the use of PPE such as goggles, overalls, gloves, and boots did not significantly alter the frequency of MN and BNC in car spray painters even though most of the workers sampled in this study did not strictly adhere to an adequate use of PPE while working. A previous study in Nigeria documented that only a few paint workers use some form of PPE despite public awareness on the use of PPE [2]. Regardless, adequate use of PPE among car spray painters is essential in preventing health hazards associated with occupational exposure.

According to previous studies [22,28], lifestyle factors such as alcohol consumption and smoking habits are considered as contributing factors for increased frequency of nuclear abnormalities in individuals who are exposed to genotoxic substances. The present study showed that alcohol consumption in addition to cigarette smoking habits...
significantly increased the frequency of MN in exposed car spray painters when compared to car spray painters that neither smoked cigarette nor consumed alcohol. This may be attributed to benzene, which is used in the production of paints and also a content of cigarette smoke [1]. Previous epidemiological studies showed a clear relationship between increase in MN frequency and exposure to benzene and its metabolites [29,30]. Cigarette smoking is one of the factors that may influence the rate of cytogenetic damage, such as MN in humans [31]. Alcoholic beverages have been described as containing mutagenic substances [32]. Similar to this finding, a previous study also documented an increase in MN frequency in buccal cells of motor garage workers who consumed alcohol [33]. Another study showed a direct relationship between buccal mucosa nuclear changes and cigarette smoking and alcohol consumption [34].

The expression of tumor suppressor gene p53 and cell proliferation marker, Ki-67 has been widely used to monitor the progression of epithelial dysplasia of the oral cavity [35]. There was no expression of both p53 and Ki-67 genes in the buccal epithelial cells of car spray painters and control subjects. However, there was significantly increased MN frequency in the epithelial buccal cells of car spray painters in the present study, morose, a previous study reported that increased MN frequency in peripheral blood lymphocytes of healthy subjects is a predictive biomarker of cancer risk, 12-15 years after the MN test was performed [36]. Another study documented a gradual increase in MN frequency of exfoliated buccal cells from normal mucosa to precancerous lesions then to oral squamous carcinoma, which suggested a link of this biomarker with neoplastic progression [37]. With the observed increase of MN frequency in buccal epithelial cells of car spray painters in the present study, periodical biological monitoring of these exposed car painters is therefore needful. Similar to our findings, a previous study on ICC staining for p53 and Ki-67 to characterize urothelial cells in urine cytology revealed a negative immunoreactivity for p53 and Ki-67 genes in subjects with normal cytology with median percentage values (first to third quartile) of p53 and Ki-67 being 0 (0–5) and 0 (0–1), respectively. The authors, however, recorded positive immunoreactivities for p53 and Ki-67 genes in subjects with urothelial carcinoma for 30 (10–80) and 20 (10–30), respectively [38].

**CONCLUSION**

The results obtained from the present study showed that car spray paint workers in Enugu metropolis occupationally exposed to genotoxins have increased MN and BNC frequencies, which may pose potential health risks with fatal consequences. There is, therefore, need to provide appropriate training for workers on the importance of using PPE and other safety practices so as to reduce the exposure to genotoxic agents and improve conditions of occupational safety.

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**AUTHOR’S CONTRIBUTIONS**

Anulika Onyemelukwe – conducted the study and drafted the manuscript, Peter Achukwu – designed and supervised the study, Nnukwu Azubuike – performed the statistical analysis, Uzoamaka Madakor – assisted in laboratory investigations, and Okechukwu Onwukwe – critically reviewed the manuscript. All authors read and approved the final manuscript.

**CONFLICTS OF INTEREST**

The authors declare that there are no conflicts of interest for this research.

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