

Available Online at: https://www.scholarzest.com Vol. 5 No 08 August 2024 ISSN: 2660-5570

# STUDY OF THE BLOOD CHARACTERISTICS OF PEOPLE EXPOSED TO SMOKE FROM GRILLED MEAT IN SOME AREAS WITHIN AL-HAWIJA DISTRICT IN KIRKUK GOVERNORATE, IRAQ

Sameer saleh hazeem<sup>1</sup> Awaz B Mohammed<sup>2</sup>

1,2, Department of Biology, College of Science, Kirkuk University, Iraq

Corresponding author: <u>scbhm013@uokirkuk.edu.iq</u>

	cell counts, hemoglobin levels, viscosity, and platelet counts were significantly			
	elevated in individuals exposed to barbecue smoke compared to controls.			
Keywords: Blood characteristics, Exposure to grilling smoke, grilled meat				

## INTRODUCTION

Polycyclic aromatic hydrocarbons (PAHs) are widely distributed around the world, owing primarily to long-term anthropogenic pollution sources (Patel *et al.*, 2020). Petroleum hydrocarbon contamination, forest fires and volcanic eruptions can contribute to PAHs emissions (Imam *et al.*, 2022). Additionally, it is possible to combine dripping fat from grilled meat with coals, which produces smoke containing PAHs (Babaoğlu 2023). Polycyclic aromatic hydrocarbons are organic pollutants that consist of two or more fused aromatic rings of carbon and hydrogen atoms. They are typically colorless, white, or pale-yellow solid substances (Abdel-Shafy and Mansour, 2016; Suman *et al.*, 2016). There are allegedly 200 distinct PAHs chemicals compound in this group (Sahin *et al.*, 2020). They have the ability to contaminate meat by directly pyrolyzing food components and by depositing smoke from incomplete combustion of organic materials (Hamidi *et al.*, 2016). Thus, smoked or barbecued meats frequently have significant levels of PAHs (Duedahl-Olesen and Ionas, 2022).

Furthermore, alterations in blood cells may represent the harmful effects of PAHs. Since all types of blood cells in peripheral human blood are susceptible to PAHs exposure due to the highly lipid-solubility and hence easily absorbed by the body (Liu et al., 2021). Studies have linked alterations in CBC cell as a result to exposure to BaP, including elevated neutrophils, reduced lymphocytes, and a decrease in the number of white blood cells (WBCs). According to Wang et al. 2019, found that PAHs impair hematopoiesis, reducing white blood cells, eosinophils, monocytes, and lymphocytes, and affecting red blood cells over two years. There is also a strong link between increased PAHs exposure and raised blood lipids, showing the indirect consequences of inflammation and oxidative stress (Zhou et al., 2023). PAHs can trigger the destruction of red blood cells and the tendency to alter the patterns of leukocytes if inhaled or swallowed in significant quantities (Abdel-Shafy and Mansour, 2016; Fanali et al., 2018). The production of PAHs during charcoal-grilling may be influenced, either directly or indirectly, by the distinct chemical properties of the different varieties of charcoal (white, black, and extruded charcoals), (Kim et al., 2021). PAHs levels are also influenced by various elements, such as the food's distance from the heat source, the fuel source, the degree of processing, the length of time and type of cooking, and other cooking-related parameters. Other methods and processes like the storage enhance the number of PAHs in processed meats and meat products (Adeyeye and Ashaolu, 2022). This study aims to evaluate the content of polycyclic aromatic hydrocarbons (PAHs) in different grilled meat types and chicken. Moreover, the study explores the connection between polycyclic aromatic hydrocarbons (PAHs) and grilled meat and chicken, focusing on their impact on human blood cells.

#### **PRIMARY OBJECTIVE:**

The present study aimed to evaluate the complete blood picture in individuals exposed to barbecue smoke. **MATERIALS AND METHODS** 

The study involved the collection of 105 blood samples from workers in Hawija city restaurants and 20 control samples from healthy individuals. The sampling period spanned from the beginning of November 2023 to the end of March 2024. After obtaining verbal consent from each participant, blood was drawn using a medical syringe in a 3.0 ml volume and placed into an EDTA tube for complete blood count

#### **COMPLETE BLOOD COUNT**

This analysis is considered one of the most important tests that provides a complete picture of the main components in the blood. Levels of some blood indicators were measured in the studied groups, including hemoglobin (Hb), hematocrit (HCT), red blood cell count (RBC), white blood cell count (WBC), and platelets (PLT). This was done using a Swelab hematology analyzer, which automatically analyzes each sample and provides readings for 22 parameters of the blood picture.

Whole blood was collected into an EDTA tube and mixed thoroughly. The EDTA tube was placed in the designated position of the analyzer for sample aspiration. Subsequently, 13 microliters of whole blood and 20 microliters of prediluted blood were aspirated. Red blood cells were lysed, releasing hemoglobin which subsequently reacted to produce a colored solution. Concurrently, white blood cells underwent cytoplasmic lysis, causing the nuclear and granular components to shrink. Due to the electrical inaccessibility of the cells, the hematology analyzer categorized white blood cells into groups such as neutrophils, lymphocytes, and monocytes by detecting changes in electrical resistance as particles suspended in an electrolyte solution passed through an aperture.

#### **RESULTS AND DISCUSSION**

The results presented in Table (1) indicated that there was no significant difference in white blood cell count between individuals exposed to barbecue smoke and healthy controls. While a slight increase in white blood cell count was observed in the exposed group (8.17 X10<sup>9</sup>/L) compared to the control group (7.73 X10<sup>9</sup>/L), red blood cell count showed a significant difference, with values of 5.190 X10<sup>12</sup>/L and 4.60 X10<sup>12</sup>/L for the exposed and control groups, respectively. Hemoglobin levels and viscosity also exhibited significant differences, with values of 14.92 g/dl and 14.31 g/dl, and 48.31% and 46.58%, respectively, for the exposed and control groups. Platelet counts also showed significant differences between the exposed and control groups, with values of 235.2X10<sup>9</sup>/L and 211.3X10<sup>9</sup>/L, respectively.

The results of this study can be attributed to the presence of harmful compounds in barbecue smoke, which can affect blood cell functions and induce changes in blood component composition. The findings of the present study align with those of Liu (2019) in China, which reported an increased white blood cell count in both men and women exposed to barbecue smoke. Similarly, Duhl, Tefft, and TerAvest (2018) in the United States found that exposure to barbecue smoke was associated with elevated red blood cell counts and hemoglobin levels in both genders. Furthermore, Aparicio et al. (2017) in South Korea reported a positive correlation between barbecue smoke exposure and platelet counts in both men and women. However, these findings contrast with those of D. J. Lee *et al.* (2016) in Japan and Ballas (2014) in Germany, who found no significant association between barbecue smoke exposure and white blood cell count, red blood cell count, hemoglobin levels, or platelet counts in either gender.

Groups Variables	Control	Exposure
WBC	7.73 a	8.17 a
LYM%	27.30 b	30.27 a
MON%	6.36 b	6.84 a
GRA%	63.31 a	62.93 b
LYM#	1.93 b	2.50 a
MON#	0.42 b	0.51 a
GRA#	4.98 b	5.45 a
RBC	4.60 b	5.190 a
HGB	14.31 b	14.92 a
HCT	46.58 b	48.31 a
MCV	91.05 a	89.06 b
MCH	26.85 b	28.76 a
MCHC	29.71 b	32.22 a
RDW-CV	11.44 b	12.74 a
RDW-SD	40.90 b	50.37 a
PLT	211.3 b	235.2 a
MPV	8.15 b	11.03 a
PCT	0.18 b	0.20 a
PDW	14.49 a	14.21 b

Table 1: Complete Blood Count in Individuals Exposed to Barbecue Smoke

Table 2 demonstrated a significant difference in white blood cell count based on the duration of exposure to barbecue smoke. Individuals exposed for 8 hours exhibited a slightly higher white blood cell count ( $8.83 \times 10^9/L$ ) compared to those exposed for 6 and 12 hours ( $7.83 \times 10^9/L$  and  $7.80 \times 10^9/L$ , respectively). Red blood cell count was highest in individuals exposed for 12 hours ( $5.33 \times 10^{12}/L$ ), followed by 8 hours ( $5.20 \times 10^{12}/L$ ) and 6 hours ( $5.07\times 10^{12}/L$ ). Hemoglobin levels and viscosity did not show significant differences based on exposure duration. However, the highest hemoglobin levels were observed at 8 hours (15.13 g/dl), followed by 12 hours (15.01 g/dl) and 6 hours (14.66 g/dl). Viscosity showed significant differences, with higher values at 12 hours (47.35%), 8 hours (46.67%), and 6 hours (45.92%). Platelet counts did not show significant differences based on exposure duration, with the highest values observed at 8 hours ( $245.2 \times 10^9/L$ ), followed by 12 hours ( $223.9 \times 10^9/L$ ), while exposure to barbecue smoke for longer durations did not significantly affect hemoglobin levels or platelet counts, it led to a slight increase in white blood cell count and a gradual increase in red blood cell count.

Groups Variables	6	8	12
WBC	7.83 b	8.83 a	7.80 a
LYM%	30.47 a	29.83 a	30.55 a
MON%	6.92 a	6.84 a	6.66 a
GRA%	63.56 a	62.81 a	62.23 a
LYM#	2.30 a	2.56 a	2.34 a
MON#	0.474 b	0.572 a	0.486 b
GRA#	5.33 a	5.85 a	5.10 a
RBC	5.07 b	5.20 ab	5.33 a
HGB	14.66 a	15.13 a	15.01 a
HCT	45.92 a	46.67 a	47.35 b
MCV	89.48 a	89.19 a	88.33 a
MCH	28.82 a	29.07 a	28.29 a
MCHC	32.03 ab	32.67 a	31.94 b
RDW-CV	12.64 a	12.96 a	12.60 a
RDW-SD	49.80 ab	52.40 a	48.63 b
PLT	223.9 a	245.2 a	234.7 a
MPV	8.60 a	8.58 a	8.47 a
PCT	0.189 a	0.210 a	0.194 a
PDW	14 a	13.89 a	14.20 a

Table 3 revealed a significant correlation between the age of individuals exposed to barbecue smoke and their white blood cell counts. A slight increase in white blood cell count was observed in the 30-39 age group, reaching a value of  $8.29 \times 10^9$ /L. Conversely, the highest red blood cell count was found in individuals aged 40 and above, with a value of  $5.30 \times 10^{12}$ /L. Hemoglobin levels showed significant differences, with the highest values observed in the 30-39 and below 20 age groups, reaching 15.01 g/dl. Viscosity also exhibited significant variations, with the highest values recorded in the below 20 age group at 47.73%. Platelet counts showed significant differences as well, with the highest values observed in the above 40 age group, reaching 276.4  $\times 10^9$ /L.

The observed elevation in white blood cell count can be attributed to an inflammatory response of the immune system against the harmful substances in barbecue smoke, particularly benzo[a]pyrene (Jin et al., 2024). The increased red blood cell count is likely due to a compensatory increase in hemoglobin levels to counteract the hypoxia caused by the respiratory effects of benzo[a]pyrene (Chandrashekar, 2022). The elevated blood viscosity may be a result of increased protein levels, especially fibrinogen, as an inflammatory response to barbecue smoke (Qigang et al., 2023). The elevated platelet count could be attributed to increased platelet production as a compensatory mechanism for potential microbleeding caused by the effects of benzo[a]pyrene on blood vessels (Levin and Lilis, 2008)

Table 3: Mean effects of age and the interaction between age and exposure on complete blood count

Groups(year) Variables	< 20	20-29	30-39	>20
WBC	7.87 b	8.18 a	8.29 a	8.16 a
LYM%	31.18 a	29.17 c	30.75 b	31.12 a
MON%	7.38 a	6.56 d	6.80 c	7.15 b
GRA%	61.94 c	64.47 a	62.93 b	59.52 d
LYM#	2.21 d	2.35 c	2.48 b	3.32 a
MON#	0.513 c	0.485 d	0.517 b	0.577 a
GRA#	4.99 d	5.59 b	5.61 a	5.13 c
RBC	5.22 b	5.13 d	5.20 c	5.30 a
HGB	15.01 a	14.80 c	15.01 a	14.92 b
HCT	47.73 a	45.72 d	46.66 c	47.51 b

MCV	91.48 a	88.53 b	88.98 b	87.90 b
MCH	28.97 a	28.95 b	28.68 c	28.15 d
MCHC	31.31 d	32.59 a	32.34 b	31.94 c
RDW-CV	12.53 d	12.73 c	12.79 b	12.89 a
RDW-SD	46.49 d	52.23 a	51.28 b	47.05 c
PLT	233.6 c	221.7 d	235.8 b	276.4 a
MPV	8.37 b	8.73 b	15.67 a	8.32 b
PCT	0.194 c	0.190 d	0.200 b	0.219 a
PDW	14.92 a	14.33 b	13.83 d	14.04 c

Table 4 demonstrates a significant difference in white blood cell count among individuals exposed to barbecue smoke, healthy controls, and diseased individuals. White blood cell counts exhibited a slight increase in both exposed and diseased groups compared to the control group, with values of 8.126 X10<sup>9</sup>/L, 8.38 X10<sup>9</sup>/L, and 7.725 X10<sup>9</sup>/L, respectively. Red blood cell counts also showed significant differences between exposed healthy individuals, exposed diseased individuals, and the control group, with values of 5.31 X10<sup>12</sup>/L, 5.17 X10<sup>12</sup>/L, and 4.60 X10<sup>12</sup>/L, respectively. Hemoglobin levels and viscosity exhibited significant differences among the three groups, with values of 14.88 g/dl, 15.12 g/dl, and 14.31 g/dl, and 46.43%, 47.29%, and 48.31%, respectively, for the exposed diseased, exposed healthy, and control groups. Platelet counts showed significant differences among all three groups, with values of 228.5X10<sup>9</sup>/L, 267.3 X10<sup>9</sup>/L, and 211.3 X10<sup>9</sup>/L, respectively.

Groups(veat)	Exposure		
Variables	Healthy	Patients	Control
WBC	8.383 a	8.126 b	7.725 c
LYM%	30.54 a	30.21 a	27.30 b
MON%	7.239 a	6.759 b	6.355 c
GRA%	60.91 b	63.35 a	63.31 a
LYM#	3.01 a	2.39 b	1.93 c
MON#	0.578 a	0.498 b	0.420 c
GRA#	5.39 b	5.46 a	4.98 c
RBC	5.31 a	5.17 b	4.60 c
HGB	15.12 a	14.88 b	14.31 c
HCT	47.29 b	46.43 c	48.31 a
MCV	87.25 c	89.44 b	91.05 a
MCH	28.06 b	28.91 a	26.85 c
MCHC	32.20 b	32.23 a	29.71 c
RDW-CV	12.93 a	12.70 b	11.44 c
RDW-SD	50.38 a	50.37 b	40.90 c
PLT	267.3 a	228.5 b	211.3 c
MPV	8.38 b	11.58 a	8.15 c
PCT	0.216 a	0.194 b	0.180 c

Table 4: Mean effects of age and the interaction between age and exposure on complete blood count

PDW 13.57 c 14	4.34 b 14.49 a
----------------	----------------

The results presented in Table (5) indicated that there was no significant difference in white blood cell count between individuals exposed to barbecue smoke, regardless of smoking status. While a slight increase in white blood cell count was observed in smokers  $(8.17 \times 10^9/L)$  compared to non-smokers  $(7.73 \times 10^9/L)$ , red blood cell count showed a significant difference, with values of  $5.190 \times 10^{12}/L$  and  $4.975 \times 10^{12}/L$  for the exposed and control groups, respectively. Hemoglobin levels and viscosity also exhibited significant differences, with values of 14.92 g/dl and 14.31 g/dl, and 46.58% and 48.31%, respectively, for the exposed and control groups. Platelet counts also showed significant differences between the exposed and control groups, with values of  $235.2\times 10^9/L$  and  $211.3\times 10^9/L$ , respectively. These findings can be attributed to the presence of harmful compounds in barbecue smoke, which can affect blood cell function and induce changes in blood component composition.

Table 5: Mean Effects of the Interaction between Exposure and Smoking on Complete Blood Count

Groups Variables	smoking	non smoker
WBC	8.17 a	7.73 b
LYM%	30.27 a	27.30 b
MON%	6.84 a	6.36 b
GRA%	62.93 b	63.31 a
LYM#	2.50 a	1.93 b
MON#	0.512 a	0.420 b
GRA#	5.446 a	4.975 b
RBC	5.190 a	4.595 b
HGB	14.92 a	14.31 b
НСТ	46.58 b	48.31 a
MCV	89.06 b	91.05 a
MCH	28.76 a	26.85 b
MCHC	32.22 a	29.71 b
RDW-CV	12.74 a	11.44 b
RDW-SD	50.37 a	40.90 b
PLT	235.2 a	211.3 b
MPV	11.03 a	8.15 b
PCT	0.198 a	0.180 b
PDW	14.21 b	14.49 a

#### REFERENCES

- 1. Abdel-Shafy, H. I., & Mansour, M. S. (2016). A review on polycyclic aromatic hydrocarbons: source, environmental impact, effect on human health and remediation. *Egyptian journal of petroleum*, *25*(1), 107-123.
- 2. Adeyeye, S. A. O., & Ashaolu, T. J. (2022). Polycyclic aromatic hydrocarbons formation and mitigation in meat and meat products. *Polycyclic Aromatic Compounds*, *42*(6), 3401-3411.
- 3. Amadou, A., Praud, D., Coudon, T., Deygas, F., Grassot, L., Faure, E., ... & Fervers, B. (2021). Risk of breast cancer associated with long-term exposure to benzo [a] pyrene (BaP) air pollution: Evidence from the French E3N cohort study. *Environment international*, *149*, 106399.
- Aparicio, María et al. 2017. "What Are Families Most Grateful for after Receiving Palliative Care? Content Analysis of Written Documents Received: A Chance to Improve the Quality of Care." BMC palliative care 16(1): 47.
- 5. Babaoğlu, A. S. (2023). Assessing the formation of polycyclic aromatic hydrocarbons in grilled beef steak and beef patty with different charcoals by the quick, easy, cheap, effective, rugged, and safe (QuEChERS) method with gas chromatography–mass spectrometry. *Food Science of Animal Resources*, *43*(5), 826.
- 6. Ballas, Juan Pablo Estévez. 2014. "Moragas Manouver Correlation Clinically Arthroscopy, in the Diagnosis of Lateral Meniscal Injury." Orthopaedic Journal of Sports Medicine 2(4 Suppl).

- 7. Birkett, N., Al-Zoughool, M., Bird, M., Baan, R. A., Zielinski, J., & Krewski, D. (2019). Overview of biological mechanisms of human carcinogens. *Journal of Toxicology and Environmental Health, Part B, 22*(7-8), 288-359.
- 8. Bukowska, B., & Sicińska, P. (2021). Influence of benzo (a) pyrene on different epigenetic processes. *International Journal of Molecular Sciences*, *22*(24), 13453.
- 9. Chandrashekar, Naveenkumar. 2022. "Benzo (a) Pyrene-Induced Oxidative Stress during Lung Cancer and Treatment with Baicalein." In Handbook of Oxidative Stress in Cancer: Mechanistic Aspects, Springer, 787–804.
- 10. Duhl, Kody L, Nicholas M Tefft, and Michaela A TerAvest. 2018. "Shewanella Oneidensis MR-1 Utilizes Both Sodium- and Proton-Pumping NADH Dehydrogenases during Aerobic Growth." Applied and environmental microbiology 84(12).
- 11. Duedahl-Olesen, L., & Ionas, A. C. (2022). Formation and mitigation of PAHs in barbecued meat–a review. *Critical Reviews in Food Science and Nutrition*, *62*(13), 3553-3568.
- 12. Fanali, L. Z., Franco-Belussi, L., Bonini-Domingos, C. R., & de Oliveira, C. (2018). Effects of benzo [a] pyrene on the blood and liver of Physalaemus cuvieri and Leptodactylus fuscus (Anura: Leptodactylidae). *Environmental pollution*, *237*, 93-102.
- 13. Hamidi, E. N., Hajeb, P., Selamat, J., & Razis, A. F. A. (2016). Polycyclic aromatic hydrocarbons (PAHs) and their bioaccessibility in meat: A tool for assessing human cancer risk. *Asian Pacific Journal of Cancer Prevention*, *17*(1), 15-23.
- 14. Imam, A., Suman, S. K., Kanaujia, P. K., & Ray, A. (2022). Biological machinery for polycyclic aromatic hydrocarbons degradation: A review. *Bioresource Technology*, *343*, 126121.
- 15. Jin, Hui et al. 2024. "Effects and Mechanisms of Polycyclic Aromatic Hydrocarbons in Inflammatory Skin Diseases." Science of The Total Environment: 171492.
- 16. Kim, H. J., Cho, J., & Jang, A. (2021). Effect of charcoal type on the formation of polycyclic aromatic hydrocarbons in grilled meats. *Food Chemistry*, *343*, 128453.
- Lee, Daniel J et al. 2016. "Comparative Effectiveness of Targeted Prostate Biopsy Using Magnetic Resonance Imaging Ultrasound Fusion Software and Visual Targeting: A Prospective Study." The Journal of urology 196(3): 697–702.
- 18. Levin, Stephen, and Ruth Lilis. 2008. "Diseases Associated with Exposure to Chemical Substances." *Public Health Preventive Med* 619.
- 19. Liu, Yiting. 2019. "New Ex Vivo Demyelination/Remyelination Models to Defeat Multiple Sclerosis and Neuromyelitis Optica." Neural regeneration research 14(10): 1715–16.
- Liu, C., Wu, M., Fu, M., Wang, H., & Nie, J. (2021). Dose–response relationships between polycyclic aromatic hydrocarbon exposure and blood cell counts among coke oven workers: a sex-stratified analysis. *BMJ* open, 11(12).
- 21. Patel, A. B., Shaikh, S., Jain, K. R., Desai, C., & Madamwar, D. (2020). Polycyclic aromatic hydrocarbons: sources, toxicity, and remediation approaches. *Frontiers in Microbiology*, *11*, 562813.
- 22. Qigang, Nie et al. 2023. "The Effect of Polycyclic Aromatic Hydrocarbon Biomarkers on Cardiovascular Diseases." *Reviews on Environmental Health* (0).
- 23. Sahin, S., Ulusoy, H. I., Alemdar, S., Erdogan, S., & Agaoglu, S. (2020). The presence of polycyclic aromatic hydrocarbons (PAHs) in grilled beef, chicken and fish by considering dietary exposure and risk assessment. *Food Science of Animal Resources*, *40*(5), 675.
- 24. Suman, S., Sinha, A., & Tarafdar, A. (2016). Polycyclic aromatic hydrocarbons (PAHs) concentration levels, pattern, source identification and soil toxicity assessment in urban traffic soil of Dhanbad, India. *Science of the Total Environment*, *545*, 353-360.
- 25. Susanto, A., Yusril, N., Zaini, J., & Nuwidya, F. (2021). Comparison of serum benzo (a) pyrene diol epoxide protein adducts level between kretek cigarette smokers and nonsmokers and the related factors. *J. Nat. Sci. Biol. Med*, *12*, 52.
- 26. Wang, Y., Zhao, H., Wang, T., Liu, X., Ji, Q., Zhu, X., ... & Duan, H. (2019). Polycyclic aromatic hydrocarbons exposure and hematotoxicity in occupational population: a two-year follow-up study. *Toxicology and applied pharmacology*, *378*, 114622.
- Zhou, S., Li, X., Dai, Y., Guo, C., Peng, R., Qin, P., & Tan, L. (2023). Association between polycyclic aromatic hydrocarbon exposure and blood lipid levels: the indirect effects of inflammation and oxidative stress. *Environmental Science and Pollution Research*, *30*(59), 123148-123163.