CHARACTERISATION OF Plasmodium falciparum PFMDR1 GENE MUTATIONS IN MALARIA PATIENTS FROM SELECTED HOSPITALS IN OTA, OGUN STATE, NIGERIA

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AUGUST, 2024

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BY

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DISSERTATION **SUBMITTED** TO **SCHOOL OF** THE POSTGRADUATE STUDIES IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THE AWARD OF MASTER OF SCIENCE (M. Sc) DEGREE IN MICROBIOLOGY IN THE DEPARTMENT OF SCIENCES, COLLEGE **OF** BIOLOGICAL **SCIENCE AND** TECHNOLOGY, COVENANT UNIVERSITY, OTA, OGUN STATE, **NIGERIA**

AUGUST, 2024

ACCEPTANCE

| ACCEITANCE | | |
|---|------------------------|--|
| This is to attest that this dissertation is accepted in partial fulfilment of the | e requirements for the | |
| award of the degree of Master of Science in Microbiology in the Depa | artment of Biological | |
| Sciences, College of Science and Technology, Covenant University, Ota, Nigeria | | |
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DECLARATION

I, ARIRIGUZOH, VICTORIA-GRACE ONYEKACHI (22PCQ02456) declare that this research was carried out by me under the supervision of Professor Olayemi O. Akinnola of the Department of Biological Sciences, College of Science and Technology, Covenant University, Ota, Nigeria. I attest that the dissertation has not been presented either wholly or partially for the award of any degree elsewhere. All sources of data and scholarly information used in this dissertation are duly acknowledged.

ARIRIGUZOH VICTORIA-GRACE ONYEKACHI

Signature and Date

CERTIFICATION

We certify that this dissertation titled "CHARACTERISATION OF Plasmodium falciparum PFMDR1 GENE MUTATIONS IN MALARIA PATIENTS FROM SELECTED HOSPITALS IN OTA, OGUN STATE, NIGERIA" is an original research work carried out by ARIRIGUZOH, VICTORIA-GRACE ONYEKACHI in the Department of Biological Sciences, College of Science and Technology, Covenant University, Ota, Ogun State, Nigeria under the supervision of Professor Olayemi O. Akinnola. We have examined and found this work acceptable as part of the requirements for the award of Master of Science in Microbiology.

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DEDICATION

For my dear mom, without whom my academic achievements would not have been possible. Your unwavering support, encouragement, and sacrifices have been the driving force behind my success. Thank you for always believing in me and pushing me to be my best self. This project is dedicated to you with all my love and gratitude.

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ABSTRACT

Malaria continues to be a significant global health problem, especially in tropical and

subtropical areas. The 2023 World Malaria Report by the World Health Organization reveals

that there were 249 million cases of malaria and 608,000 fatalities in 2022. A significant

obstacle in malaria control is the rise and spread of resistance to antimalarial drugs, which

endangers the efficacy of current therapies and undermines worldwide endeavours to eradicate

the illness. The efficacy of routinely used antimalarial medicines, such as artemisinin-based

combination treatments (ACTs), is increasingly endangered by the formation and spread of

multi-drug resistance (MDR). The Plasmodium falciparum multidrug resistance 1 gene

(*Pfmdr1*) is frequently linked to multi-drug resistance. This study was conducted to detect the

presence and prevalence of the *Pfmdr1* gene in *Plasmodium falciparum* isolated from malaria-

positive blood samples to understand its role in anti-malaria drug resistance within Ota, Ogun

State, Nigeria. Ethical approval was obtained from the Covenant Health Research Ethics

Committee, Ogun state, Nigeria, before the commencement of the study. Three hundred and

fifty-six blood samples were obtained from outpatients attending Covenant University Medical

Centre and the General Hospital, Ado-Odo Ota, Ogun State. Microscopy was used to identify

P. falciparum in all samples. The prevalence of malaria was recorded at 66%, with the age

group of children ≤ 10 having the highest prevalence at 80%. DNA extraction was performed

to isolate P. falciparum from 50 malaria-positive blood samples. Nested PCR was performed

to identify and amplify the *Pfmdr1* gene and determine its mutations. The prevalence of the

Pfmdr1 was recorded at 66% among samples amplified using two-step Nested PCR. Sanger

Sequencing was carried out on 10 samples to provide a detailed genetic characterization of the

Pfmdr1 gene and three mutations of the Pfmdr1 gene the F74L, Y184F and S164S were

identified.

KEYWORDS: Malaria, Multidrug resistance, Gene Mutations, Prevalence

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