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MOLECULAR CHARACTERIZATION OF THE EMERGING SUBLINEAGE IND2001E OF FOOT-AND-MOUTH DISEASE VIRUS SEROTYPE O IN BUFFALO IN MANIKGANJ, BANGLADESH

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ABSTRACT

Foot-and-mouth disease (FMD) is a major economic infectious disease in endemic regions like Bangladesh, affecting cloven-hooved animals and hindering livestock development. Good farm practices, including vaccination, is the only effective control method and continuous monitoring is necessary to detect vaccine effectiveness. Buffalo, a significant livestock resource in the southern coastal region of Bangladesh, requires more disease management efforts than cattle. This study aimed to isolate and make molecularly characterize of the VP1 coding region of FMDV in clinically suspected buffaloes. Tongue epithelial tissues were collected from two buffalos suspected of FMD in June-2018 in the Manikganj district of Bangladesh. Both samples were identified as FMDV serotype O using RT-PCR testing and were subsequently cultured in BHK-21 cell line. Phylogenetic analysis of VP1 region revealed that both isolates clustered into a distinct sublineage Ind2001e, within the dominant lineage Ind2001 of Middle East-South Asia (ME-SA) topotype of FMDV serotype O, which has been circulating in Bangladesh since 2012. Analysis of nucleotide (NT) similarities revealed that 92.38% of NTs are homologous between isolates, whereas similarities with the Indian sequences was 99.91%-99.99%, and with the Nepal sequences was 99.90%-99.99%. The amino acid mutations within the VP1 sequence between the vaccine strain (O/India/R2/75) and the BAN450 isolates was 3.75% (8; 96.25% homology), specifically in the B-C loop and C-terminal region. In conclusion, buffalos get infected with the emerging novel sublineage Ind2001e. The effectiveness of vaccination in Bangladesh with the potential mutational trend poses significant concern regarding the transmission between multiple species.

Keywords: buffalo, Foot-and-mouth-disease, Ind2001e sublineage, molecular characterization, serotype O

INTRODUCTION

Foot-and-mouth disease (FMD) is a transboundary, airborne, acute, and highly

contagious animal disease characterized by high fever and anorexia, followed by vesicular lesions in the oral mucosa and interdigital spaces of cloven-hooved animals, such as buffalo, cattle, goats, pigs, sheep, and various wild species (1, 2). Consequently, the animal undergoes weight loss, resulting in reduced production, particularly in milk and meat production. FMD can also cause abortion, death in calves, international trade restrictions, and high treatment costs (2, 3). It has a significant global economic impact which is estimated between 6.5 and 21 billion US dollars. In endemic countries it can reach higher than 1.5 billion dollars annually (3, 4).

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A study by Giasuddin et al. (5) projected an annual economic impact of 2.22 billion dollars in Bangladesh. Foot-and-mouth disease virus (FMDV) is the etiological agent comprising seven serotypes characterized by distinct genetic and antigenic variations: O, A, Asia 1, C, SAT-1, SAT-2, and SAT-3. Currently, only serotypes O, A, and Asia 1 are prevalent in Bangladesh, with serotype O predominating in 82% of the outbreaks (6). Serotype O is divided into 11 topotypes of which the Middle-East-South Asia (ME-SA) topotype was identified in Bangladesh. To date, four lineages within the ME-SA topotype have been identified: Ind2001, Ind2011, PanAsia, and PanAsia-2 (7, 8). The Ind2001 lineage subsequently developed into five sublineages: Ind2001a, b, c, d, and e (9, 10). A newly identified sublineage Ind2001BD1, was reported to be the predominant sublineage in Bangladesh which was later classified as Ind2001e by the World Reference Laboratory for Foot-and-mouth disease (WRLFMA) (11, 12). However, to prevent FMDV in cattle and buffaloes, the Government of Bangladesh has adopted a national strategic plan to control FMD through early diagnosis, strategic mass vaccination, zonal control measures, and implementation of the OIE FMD control model.

Although most of the research focuses on FMDV in cattle, there is less synthesized information on monitoring of FMDV in buffaloes, despite being the second biggest livestock population in Bangladesh. Buffalo, referred to as “Black Gold,” holds significant value and is predominantly located in the southern coast area of Bangladesh (13, 14). The buffalo management system remains reliant on traditional practices and lacks modern housing, biosecurity measures, nutritional management, and disease control strategies. Farmers possess misconceptions regarding the necessity of vaccinations and deworming for buffaloes. Consequently, buffaloes were silently affected by many infectious diseases, including FMD, yet remained underreported (15). Therefore, the study aimed to isolate and molecularly type serotypes, and characterize FMDV serotype O, including amino acid mutations in the VP1 coding region, in comparison to the vaccine strain identified in buffaloes from the Manikganj district of Bangladesh. Based on the previous reports of genetic diversity of FMDV circulating in Bangladesh, we hypothesized that the FMDV serotype isolated from buffaloes in Manikganj will carry notable mutations in VP1 coding region which may reduce the match with the vaccine strain.

MATERIAL AND METHODS

Sample collection

Two tongue epithelial tissue samples were aseptically collected into virus transfer medium from two different buffalo farms suspected of having FMD in June 2018, located in the Manikganj District (coordinates: 23.85°N 90.01°E) of Bangladesh (Fig. 1). Clinical signs in buffaloes included absense of appetite, fever, drooling of saliva, vesicles on the tongue and gums. The animals were not vaccinated and were mixed with cattle that have recently recovered from FMD. Samples were immediately transferred to the laboratory in a cooler with ice and stored at -80 °C in a freezer until subsequent laboratory procedures.

Detection of FMDV serotypes by RT-PCR test

The tongue epithelial samples were grinded and macerated in pestle and mortar with sterile sand. A 10% tissue suspension was made with phosphate-buffered saline. Total RNA was extracted from an aliquot of suspension samples utilizing the QIAamp® Viral RNA kit (QIAGEN, Hilden, Germany) following the manufacturer's instructions. The VP1 gene-based PCR diagnostic assay was used for the confirmation of FMDV, adopting the primers and RT-PCR protocols developed by Vangrysterre and De Clercq (16), with minor modifications including the QIAGEN® One-Step RT-PCR Kit (QIAGEN, Hilden, Germany) as the RT-PCR master mix. Additionally, three serotypes A, O, and Asia 1 of FMDV circulating in Bangladesh were examined using the primers and procedures designed by Vangrysterre and De Clercq (16) and Callens and De Clercq (17). The list of primers is described in Table 1, and the RT-PCR temperature profile is described in Table 2.

Isolation of FMDV

The tissue inoculums were inoculated onto a pre-established monolayer of BHK-21 cells (widely used cell line for FMDV culture) using Eagle's medium supplemented with trypticase soy broth and antibiotics, incubated at 37 °C in a 5% CO₂ condition with 10% fetal bovine serum (Gibco, USA). The cells were monitored consistently until the third passage, and after four days post-inoculation. Characteristic cytopathic effects (CPE) became apparent, including cell rounding and flattening, formation of multinucleated giant cells, disruption of intracellular connections, and cell death (18).

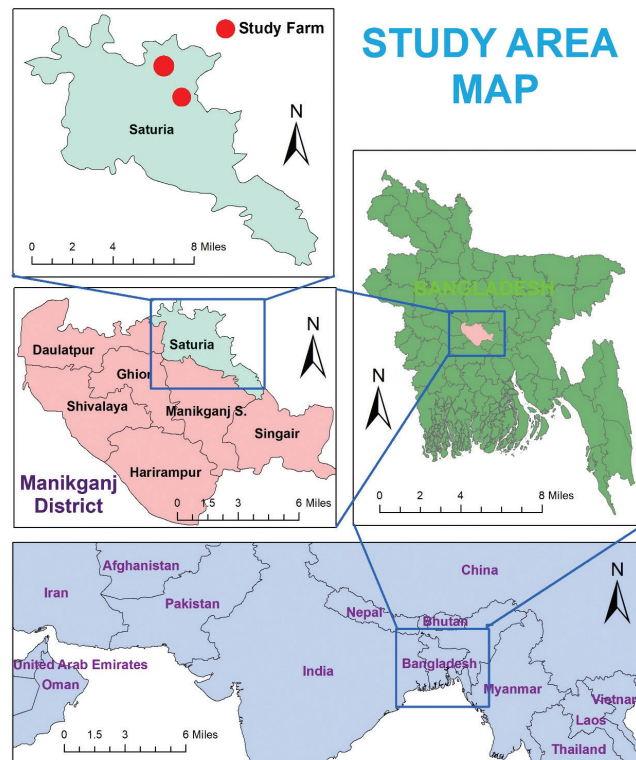


Figure 1. Location of the buffalo farms included in the current study.

Table 1. List of primer (forward and reverse) sequences used for detection of FMDV and serotyping of FMDV by RT-PCR

Serotype	Primer name	Primer sequence (5' to 3')	Location	PCR products (bp)	Reference
Universal	1F	GCC TGG TCT TTC CAG GTC T	5'UTR	328	
	1R	CCA GTC CCC TTC TCA GAT C	5'UTR		
O	P38	GCTGCCTACCTCCTTCAA	1D	402	(16)
	P33	AGCTTGTACCAGGGTTTGGC	2B		
Asia-1	P74	GACACCACTCAGGACCGCCG	1D	292	
	P33	AGCTTGTACCAGGGTTTGGC	2B		
A	P110	GT(G:A:T:C)ATTGACCT(G:A:T:C) ATGCA (G:A:T:C) AC (G:A:T:C) CAC	1D	732	(16, 17)
	P33	AGCTTGTACCAGGGTTTGGC	2B		

Table 2. One-Step RT-PCR thermal conditions

Step	Stage	Name of the primers	
		1F and 1R	P33, P38, P74, P110
1	Reverse transcription	50 °C, 30 min	50 °C, 30 min
2	Initial denaturation	95 °C, 15 min	95 °C, 15 min
3	Amplification (30×Denature, Anneal)	94/55 °C, 1 min	35×(94/55 °C, 1 min)
4	Extension	72 °C, 2 min	72 °C, 30 min
5	Final extension	72 °C, 7 min	60 °C, 6 min
6	Hold	4 °C, 4 min	4 °C, 4 min

Sequencing of VP1 coding region

Total RNA was extracted from the third passages of both isolates using the QIAamp® Viral RNA kit. The VP1 gene coding region of FMDV was amplified using three sets of primers and PCR methods as described by Knowles et al. (19) for FMDV serotype O. The amplified PCR products were analyzed using electrophoresis on a 1.5% agarose gel, alongside a 1kb DNA ladder stained with SYBR Safe DNA Gel Stain (ThermoFisher, Waltham, MA), and a UV transilluminator confirmed the amplified band. Later on, the QIAquick® Gel Extraction Kit (QIAGEN®, Hilden, Germany) was used to purify the amplified products. The products were quantified and sent to the commercial source for cycle sequencing reaction of amplified FMD serotype O products. The forward and reverse sequences of the VP1 coding region were subjected to quality assessment, editing, and trimming. They were assembled into a consensus sequence by BioEdit version 7.2.

Phylogeny analysis

A total of 56 VP1 coding sequences were retrieved from the GenBank database through NCBI BLAST (20) search, sourced from neighboring countries and Bangladesh, for an analysis of the evolutionary divergence of FMDV serotype O isolated in this study. The sequences were assembled and aligned using the ClustalW algorithm (21) using MEGA11 software (22). Finally, the phylogenetic tree with different lineages and sublineages was constructed using MEGA11. The best substitution model Tamura-Nei (23), with discrete Gamma distribution (TN93+G) in the Maximum Likelihood method, with 1,000 times bootstrap replications, was applied based on nucleotide alignment data.

Mutational analysis

The VP1 coding amino acid sequence of the BAN 450 (O/BAN/BLRI/450/2018) isolate was compared with the common vaccine strain O/India/R2/75 for checking the mutational spectrum between them by using BioEdit 7.2. The amino acid sequence was translated from the nucleotide sequence based on the standard genetic code after codon-based alignment using MEGA11 software (22).

Ethical approval

The research protocol was approved by the Animal Experimentation Ethics Committee, Bangladesh Livestock Research Institute, Savar, Dhaka 1341, Bangladesh, under reference-BLRI0006.

RESULTS

The VP1-based RT-PCR test confirmed that the two tongue epithelium samples obtained from both buffalo farms were positive for FMDV. Subsequent serotyping via RT-PCR revealed that both were confirmed as FMDV serotype O. The VP1 coding sequences of positive isolates were submitted to the NCBI GenBank database with accession IDs PQ468693 (O/BAN/BLRI/450.5/2018) and PQ468694 (O/BAN/BLRI/450.6/2018) (October 2024). The VP1 sequence data analysis for FMDV using the NCBI BLAST database revealed 100% nucleotide query coverage with FMDV serotype O. The 26 sequences of FMDV serotype O, including the two sequences of the current study, were clustered within the emerging novel sublineage Ind2001e from lineage Ind2001 under the Middle East-South Asia (ME-SA) topotype (Fig. 2). Phylogenetic analysis showed that the isolates identified in this study clustered within the same group, corresponding to FMDV serotype O strains previously reported in Bangladesh between 2012 and 2018.

The VP1 coding region sequences had a nucleotide similarity of 92.38%. The VP1 coding region sequences of FMDV serotype O obtained from NCBI GenBank had nucleotide similarity of 99.10% to 99.97% with various Bangladesh isolates, 99.91% to 99.99% with Indian isolates, and 99.90% to 99.99% with Nepalese sequences (Fig. 3).

Eight mutations were identified among the isolates and the vaccine strains BAN 450 and O/India/R2/75, with the majority occurring in the G-H loop (D138E, S140A, S140H, I144V) in the VP1 regions of buffalo isolates. Two mutations were seen in the C-terminus (N197E and E198Q), while no mutations were found in the B-C loop (Fig. 4). Cytopathic effects were observed in the cell line (Fig. 5), and the presence of the virus was subsequently confirmed by RT-PCR.

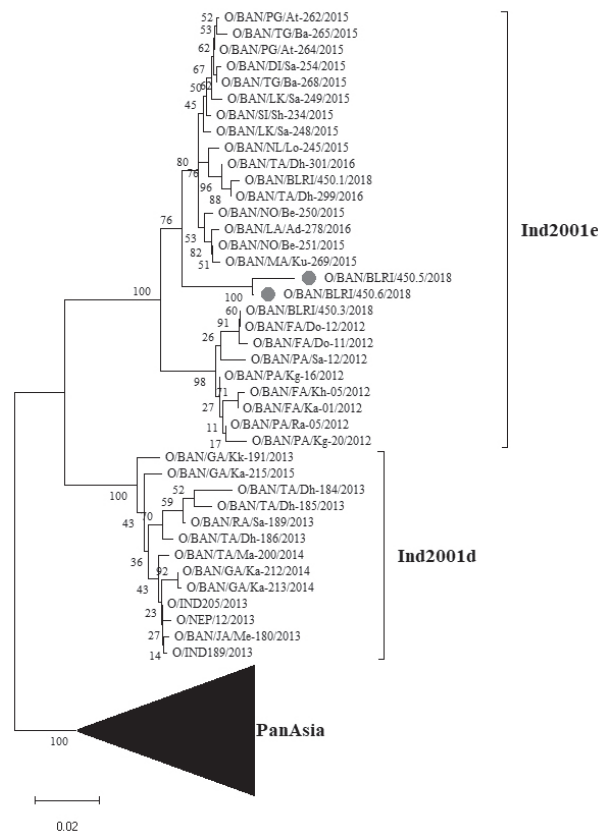
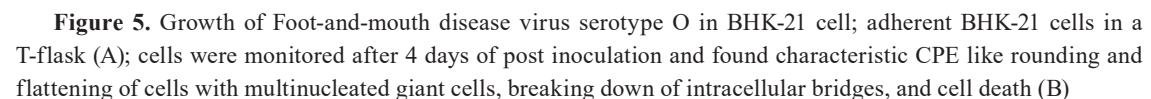
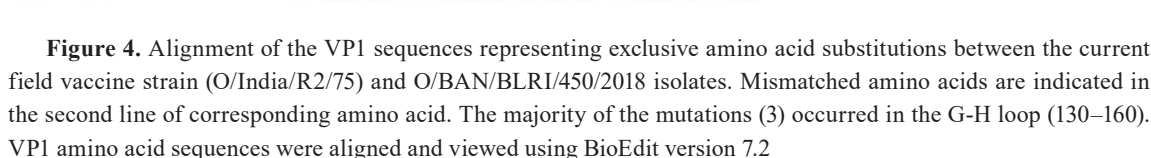


Figure 2. The phylogenetic tree reconstructed from isolates of the current study (red circles) and collected sequences of Bangladesh, India, and Nepal of diverse lineages of VP1 coding sequences of foot-and-mouth disease virus serotype O by Tamura-Nei model with discrete Gamma distribution (TN93+G) in Maximum Likelihood method and 1,000 times robustness

		Similarity																	
		1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
O/BAN/BLRI/450.1/2018	1		99.99	99.99	99.98	99.986	99.98	99.979	99.98	99.98	99.98	99.979	99.84	99.85	99.91	99.91	99.90	99.94	99.96
O/BAN/TA/Dh-299/2016	2	0.003		99.99	99.99	99.99	99.99	99.982	99.983	99.983	99.99	99.982	99.84	99.86	99.91	99.91	99.91	99.95	99.96
O/BAN/TA/Dh-301/2016	3	0.006	0.003		99.99	99.993	99.99	99.983	99.985	99.985	99.99	99.986	99.85	99.86	99.91	99.91	99.91	99.95	99.96
O/BAN/SL/Sh-234/2015	4	0.015	0.013	0.011		99.99	99.99	99.994	99.993	99.993	99.99	99.99	99.85	99.86	99.92	99.92	99.92	99.96	99.97
O/BAN/NL/Lo-245/2015	5	0.014	0.010	0.007	0.010		99.99	99.984	99.988	99.988	99.99	99.983	99.85	99.86	99.92	99.93	99.92	99.95	99.97
O/BAN/LK/Sa-248/2015	6	0.017	0.014	0.012	0.006	0.011		99.992	99.992	99.992	99.99	99.991	99.86	99.87	99.92	99.92	99.92	99.96	99.97
O/BAN/LK/Sa-249/2015	7	0.021	0.018	0.017	0.006	0.016	0.008		99.993	99.993	99.97	99.984	99.85	99.87	99.91	99.92	99.91	99.95	99.96
O/BAN/NO/Be-250/2015	8	0.015	0.013	0.011	0.008	0.010	0.007	0.014		99.987	99.99	99.993	99.85	99.87	99.92	99.92	99.92	99.95	99.97
O/BAN/DI/Sa-254/2015	9	0.020	0.017	0.015	0.007	0.012	0.008	0.007	0.013		99.98	99.985	99.85	99.87	99.92	99.92	99.91	99.95	99.97
O/BAN/TG/Ba-268/2015	10	0.017	0.014	0.012	0.003	0.011	0.005	0.003	0.010	0.000		99.988	99.86	99.87	99.92	99.92	99.92	99.96	99.97
O/BAN/MA/Ku-269/2015	11	0.021	0.018	0.014	0.010	0.017	0.009	0.016	0.007	0.015	0.012		99.85	99.86	99.92	99.92	99.91	99.95	99.97
O/BAN/BLRI/450.2/2018	12	0.162	0.157	0.152	0.150	0.152	0.145	0.147	0.153	0.146	0.145	0.153		99.88	99.87	99.87	99.87	99.81	99.84
O/BAN/BLRI/450.4/2018	13	0.145	0.141	0.141	0.135	0.137	0.128	0.133	0.134	0.132	0.131	0.143	0.119		99.87	99.87	99.87	99.83	99.85
O/IND189/2013	14	0.095	0.091	0.089	0.078	0.076	0.080	0.086	0.079	0.085	0.080	0.083	0.128	0.129		99.99	99.99	99.90	99.92
O/IND205/2013	15	0.093	0.089	0.087	0.077	0.074	0.078	0.084	0.077	0.083	0.078	0.081	0.127	0.127	0.002		99.99	99.90	99.92
O/NEP/12/2013	16	0.097	0.093	0.091	0.080	0.077	0.082	0.087	0.081	0.087	0.082	0.085	0.130	0.131	0.005	0.003		99.90	99.92
O/BAN/BLRI/450.5/2018	17	0.058	0.054	0.050	0.045	0.045	0.045	0.050	0.046	0.046	0.045	0.047	0.191	0.169	0.099	0.097	0.101		99.99
O/BAN/BLRI/450.6/2018	18	0.043	0.039	0.037	0.032	0.032	0.034	0.039	0.032	0.034	0.034	0.034	0.164	0.146	0.080	0.079	0.082	0.014	

Figure 3. Comparison of the nucleotide similarities and divergence of the sequences of Bangladesh samples of FMDV with other sequences of this virus



FMD affects all cloven hoof animal species in Bangladesh every year, especially cattle and buffalo. It has been reported that the genetic evolution of FMDV leads to consistently increasing tendency of infection and mortality (24, 25). Buffalo were perceived as having a lower susceptibility to diseases, and farmers considered buffaloes resistant to FMDV (26). Evidence indicates that Bangladeshi

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and to compare the amino acid mutation against the vaccine strain based on the VP1 coding sequence of the genome.

The VP1 sequencing analysis indicated that the lineage Ind2001 has been predominantly circulating in Bangladeshi cattle since 2009 (9, 25). The study identified an emerging novel sublineage Ind2001e (previous name Ind2001BD1) under the ME-SA topotype circulating in both buffalo and cattle. The sublineages Ind2001e and Ind2001BD2 were first identified in samples collected between 2012 and 2016 by Siddique et al. (9). Later, lineage PanAsia along with other two sublineages Ind2001e and Ind2001BD2 were reported in cattle in 2018 from Manikganj (27). However, the present isolates were collected from diverse host buffaloes within the same geographical region and year. The farms were not vaccinated against FMD and practiced mixed farming, housing both cattle and buffalo. Recently, the farm experienced an FMD infection among the cattle. Consequently, there was a high probability of FMDV transmission from cattle to buffalo (28). The lineage Ind2001 has been categorized into four distinct sublineages: Ind2001a, Ind2001b, Ind2001c, Ind2001d, and Ind2001e (8). Sublineage Ind2001e, is the most prevalent and dominant sublineage in Bangladesh since 2009 (9, 12, 25). Recently, another novel sublineage MYMBD21 under SA-2018 was identified in the sample from 2019 to 2021 in Mymensingh, Bangladesh (12). Therefore, five distinct lineages of the FMDV serotype O circulating in the Indian subcontinent: SA-2018, Ind2001, PanAsia, PanAsia-2, and Ind2011, with potential cross-border transmission among neighboring countries (10, 12, 25).

The O/India/R2/75 strain, used in India's national FMD control program as the primary component of the trivalent vaccine, serves as a relevant reference to assess antigenic variation and implications for vaccine efficacy (29). The VP1 region amino acid sequence of vaccine strain (O/India/R2/75) and BAN 450 isolates differed in 3.75% amino acids and BAN 450 isolates had mutations in both B-C loop and C-terminal region. These mutations in the antigenic sites of the BAN 450 isolates may contribute to future vaccine escape by field viruses (30). As a result, the co-circulation of several sublineages with ongoing mutations that differ from the vaccine strains may pose a significant threat to future outbreaks with high mortality rates. Vaccine failures will have a direct effect on Bangladesh's livestock industry. However, regular surveillance of FMD, along with the study of its genomic and antigenic evolution,

and the implementation of an effective national vaccination program, are essential to limit the spread of FMD in Bangladesh.

CONCLUSION

The emerging sublineage Ind2001e of FMDV serotype O is circulating in buffalo in Manikganj district of Bangladesh. There are several mutations in the VP1 amino acid sequences as compared to vaccination strains, which could make the current FMD situation in Bangladesh even more complicated. The emergence of multiple sublineages of FMDV serotype O in multiple susceptible hosts is a concern for FMD control program. Emphasis should be placed on increasing vaccination coverage in buffaloes, similar to the vaccination efforts in cattle, by motivating buffalo farmers and continuously monitoring FMD in buffalo through evaluating the genomic and antigenic variation of FMDV.

CONFLICT OF INTEREST

The authors declare that they have no financial or non-financial conflict of interest regarding authorship and publication of this article.

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AUTHORS' CONTRIBUTIONS

MZA made the conceptualization, investigation, methodology, data analysis, and original draft writing. NT and MAT performed the data analysis, writing, reviewing, and editing; MHR and MMM were included in software support, reviewing, and editing. MG contributed to funding acquisition, supervision, methodology, reviewing, and editing. All authors have read and approved the final version of the manuscript.

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