



A Cross Sectional Serologic and Epidemiological Study of Dengue Virus Infection in North Central Area of Trinidad and Tobago

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Authors' contributions

This work was carried out in collaboration between all authors. Authors KK and PEA designed the study wrote the protocol and wrote the first draft of the manuscript. Authors KK, CU and RS managed the analyses of the study. Authors KK, PEA and CU managed the literature searches. All authors read and approved the final manuscript.

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ABSTRACT

Aims: This study was carried out to determine the observed serological and significant epidemiological risk factors for dengue fever infection in a cross-section of the population in Trinidad and Tobago.

Study Design: This was an observational cross sectional study.

Place and Duration of Study: The study was carried out in the department of Paraclinical Sciences of the University of the West Indies, St. Augustine Campus, Trinidad and Tobago, over a period of 10 months, October 2016 to July 2017.

Materials and Methods: Over 450 individuals from a cross section of the population residing in the northern part of Trinidad Island were surveyed. These included individuals suspected of having

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dengue fever that presented to the healthcare facilities with complaints of fever along with some other symptoms suggestive of dengue viral illness. There was no age, gender or ethnic bias. A standardized questionnaire was used to obtain epidemiological data. Blood samples taken from consented participants were analyzed using rapid immune chromatographic tests (ICTs) – Panbio, SD Bioline and Enzyme Linked Immunosorbent Assays (ELISA). The samples were also tested for baseline blood parameters such as platelets and haemoglobin. The epidemiological data were analysed using SPSS version 21.

Results: Analysis of 380 individuals who fulfilled study criteria revealed that there were no demographic characteristics (age, gender, locality, etc.) that showed statistical significance with having a dengue infection. Retro-orbital pain, headaches and respiratory symptoms (e.g., cough, cold) showed differences that were significant with those having a dengue infection. No statistical significance was found in any comorbidity (diabetes, hypertension and asthma) factors considered and patients with dengue infections. Evaluation of platelet counts showed that only 5.4% samples had abnormal range, while 80% of those that tested positive were not significant either. Monitoring of platelet levels is still very important, but it showed that it is not an indicator of worsening dengue because 95.3% of the positive cases were within normal levels.

Conclusions: Except for nonspecific symptoms observed among patients suspected of dengue fever, there were no other significant factors that were exclusive in identifying dengue infection among the subjects studied. Platelet monitoring may not be the only parameter to use in determining deteriorating dengue patients. Vector eradication activities should be intensified with other efforts such as education program.

Keywords: Dengue fever; ELISA, epidemiology; serology; panbio; Trinidad and Tobago.

1. INTRODUCTION

Dengue is a global public health problem and in the last decade has increased substantially due to human travel and changing suitability for the mosquito vector [1,2,3]. Dengue is endemic in more than 100 countries with an estimated 50 – 100 million infections annually [4,5]. Dengue fever is an acute manifestation of this arthropod borne viral infection belonging to the *Flaviviridae* family. The dengue virus is transmitted by female mosquito *Aedes aegypti*. Four serotypes of the virus are known to exist DEN-1-4 [6], and a recently documented fifth serotype appears to have been emerged [7]. Classic dengue fever is usually self-limiting, especially in children. Dengue infection is characterized by sudden onset of a headache, retro-orbital pain, high fever, joint pain and rash. More serious manifestations of dengue virus infection includes the dengue haemorrhagic fever (DHF) and dengue shock syndrome (DSS) [2]. Dengue haemorrhagic fever is associated with re-infection, characterized by the defects in homeostasis and plasma leakage into interstitial spaces associated with increased levels of vasoactive cytokines [8]. This leads to life-threatening shock (DSS) in some cases.

The severe syndromes occur in patients with passively acquired or pre-existing, non-neutralizing, heterologous antibody caused by

previous infection with a different serotype of the virus [9]. The antibodies from the previous infection bind to the new infecting serotype and facilitate viral entry via Fc-receptor binding cells, so the number of antigen-presenting cells is increased at secondary infection [8]. In 2016, there were a recorded 1,801 probable cases alone in Trinidad and Tobago out of the total 9,993 probable cases in the non-Latin (English, French and Dutch) Caribbean [10]. This is a significant decrease in the number of reported probable cases when compared to 2014; with 5,157 probable dengue cases. As was noted in a prospective seroepidemiological study from Trinidad and Tobago, many dengue infections do not produce severe symptoms, and the number of reported cases underestimates the actual prevalence of dengue in the population [11, 12].

The aim of this study was to serologically confirm the frequency of dengue virus infection and determine epidemiological risk factors associated with dengue infections among patients suspected of having dengue fever and attending health care facilities in the north-central region of Trinidad and Tobago.

2. MATERIALS AND METHODS

2.1 Study Design, Sites and Population

This was an observational cross sectional study conducted during the period of October 2016 –

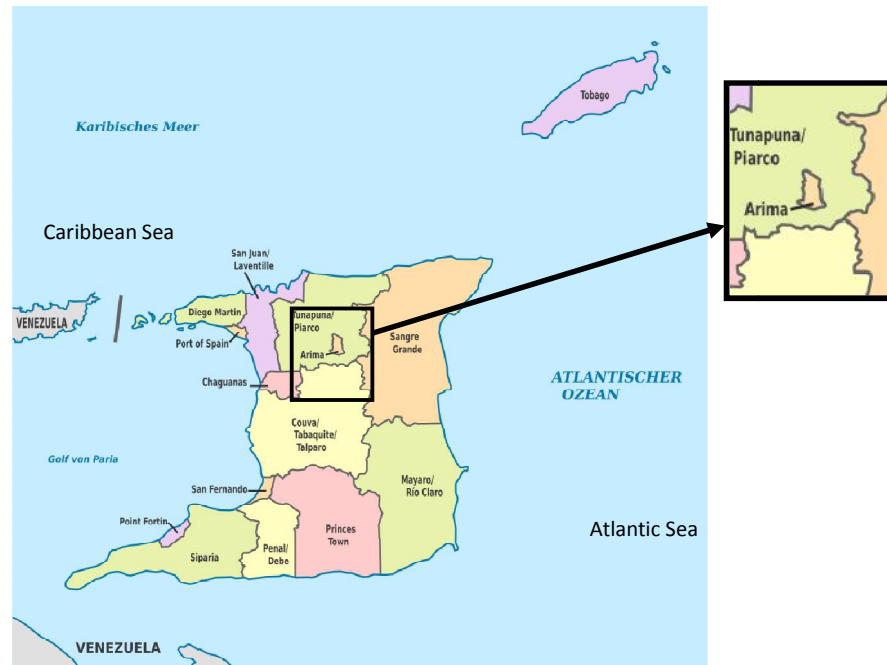


Fig. 1. Geographical map of Trinidad and Tobago showing the locality of individuals surveyed for dengue virus fever in Trinidad and Tobago

July 2017, among patients with suspected dengue infection. The study was carried out at two health care facilities of the North Central Regional Health Authority (NCRHA) in Trinidad of the twin Island, Trinidad and Tobago with catchment areas as indicated in the figure above (Figure 1). This area has a high population density in the country and most dengue cases in the past were localized to this region [13]. This region was chosen as the area of study so as to reassess the current burden of dengue virus infection. This study was carried out among patients who presented to these healthcare facilities with suspected dengue infection. Fever characterizes suspected dengue infection along with the following symptoms - anorexia, rash, aches and pains, vomiting and nausea, abdominal pains and warning signs include positive tourniquet test, leukopenia, thrombocytopenia (platelet count $<150 \times 10^9/L$), abdominal tenderness, clinical evidence of plasma leakage and/or increase in haematocrit [14]. The study enlisted voluntary participants who gave written consent and were systematically randomly selected. Standardized data collection form was used to obtain epidemiological information from all enrolled participants who were seen and physically examined by medical personnel in the study.

2.2 Inclusion Criteria

All patients of all age groups, gender, ethnic groups, social and educational level who presented to these health facilities with suspected dengue infection symptoms as enumerated above and gave written consent or assent were included in the study. Any patient who did not meet the previously mentioned requirements for suspected dengue infection or did not give consent was excluded from the study.

2.3 Collection of Specimen

A standardized questionnaire was used to obtain patient biodata or information and clinical history. This was administered by one of the trained investigators to avoid bias and misinterpretation or misrepresentation of the responses from the participants.

About 10 ml of blood (5 ml each in red and purple top tubes) was obtained through venipuncture and transported to the Department of Paraclinical Sciences, The University of West Indies, St. Augustine Campus; and Pathology Laboratory at the Eric Williams Medical Sciences Complex for further analysis. The blood samples were allowed to clot at room temperature,

centrifuged and separated as soon as possible the same day for the rapid kits (Panbio and SD Bioline). They were then stored at 2-8°C for a maximum of two days or stored frozen at -30°C until complete testing using the ELISA kits that were performed in batches.

2.4 Laboratory Analysis - Complete Blood Count

All samples were subjected to a routine complete blood count as part of the routine services offered to the patients by the healthcare facilities including platelet counts for each patient.

2.5 Rapid Immuno-chromatographic Tests (ICTs)

The samples collected in the red top tubes were subjected to serological analysis using enzyme linked immunosorbent assay - ELISA, (Dengue Virus IgM/IgG capture DxSelect ELISA, Focus Diagnostics, Cypress, PA, USA) for detection of human serum IgM and IgG antibodies in dengue virus (DV) infections. Rapid immuno-chromatographic tests (ICT) kits were used for detection of IgM and IgG antibodies, and non-structural protein 1 (NS1) antigen; of sera collected and the results were recorded. The relative sensitivity and specificity for the Panbio rapid ICT is 96.3% (90.8 -99.0%) and 95.0% (87.7 – 98.6%), respectively. The sensitivity and specificity of the SD Bioline rapid ICT is 92.8% and 98.4%, respectively. The kits were used within one to three months of procuring them from the distributors and manufacturers, while their life span (expiration dates) were still within two to three years.

2.6 Quality Controls

Controls for both the IgM/IgG ELISA kits were provided as follows: Detectable controls (human sera), non-detectable controls (human sera) and cut-off calibrators (human sera). Samples that were collected from asymptomatic and healthy individuals during the time of the study were used as controls for both of the rapid ICT tests. Controls were run every time when procedures were carried out.

2.7 Statistical Analysis

Microsoft Excel was used for data entry and data analysis was performed using Statistical Package for the Social Sciences (SPSS) 23.0 software. Chi-square test and Fisher's exact test were used to compare categorical variables. The Chi-square was chosen for determination of

association between a tested variable and a positive dengue result. If a relationship existed between any of the variables, the Chi-square value (p value) would reflect the strength of the association. The Fisher's exact test is used in place of the Chi-square to measure the same association for smaller sample sizes. In cases where the frequency counts are fewer than five in a two by two table, the test statistics (p) used is the Fisher's exact value. A probability value (p) of < 0.05 was considered statistically significant.

3. RESULTS

More than 450 individuals were recruited for this study but only 380 of these gave consent, completed the questionnaire, got evaluated, had venipuncture and were included in the final analysis. Patients included were noted to have come from different ethnic groups of people living in this part of the country. Among the study participants, 38.7% were of mixed ethnicity followed by patients of African descents, 36.6%. The East Indian and Spanish descents were 22.6% and 1.1% respectively. Most of the study participants were females (61.3%) and the median age of all analyzed individuals in the study was 26 years (range, 3 years to 87 years) but the prevalent age group surveyed was between 21 – 30 years (Fig. 2). The median time between onset of illness and collection of specimens was 3 days (range, 1 to 50 days).

As shown in Table 1, the laboratory tests of the blood samples using the ELISA reference for dengue IgM and IgG, initially classified the analysis as 92.5% positive for dengue and 7.5% non-dengue. Of those that tested positive for dengue, females were in the majority (60.5%) and 32.6% of all positive cases were between the ages of 21-30 years old. Based on the clinical history, presentation of fever, body aches and headache, the blood samples and the subjects were further defined or classified as acute cases or phase (74.2%), convalescent cases or phase (18.3%); and based on immune status, as primary 5.4% or secondary, 87.1 %. An acute sample was recorded as one collected within ≤ 7 days post onset of symptoms while those ≥ 7 days post onset of symptoms were recorded as convalescent. Demographics were the first parameters used to determine what would qualify as risk factors in acquiring a dengue infection. Being of particular ethnic group had no bearing or significance on whether the patient tested dengue positive. The majority of the positives (38.4%) were found to be of 'mixed' descent, followed by African descent (37.2%).

There was also no association between living in a particular area and contracting dengue, although most recruits were from the Arima area (Fig. 1 above), and there was a high percentage (47.3%) that tested positive there.

The statistical analysis in this study revealed that retro-orbital pain, respiratory symptoms (cold, cough, runny/stuffy nose) and headache had a significant association with samples that tested positive for dengue ($p < 0.05$), Table 1. More than half (53.3%) of patients surveyed that tested positive for dengue reported experiencing retro-orbital pain; 88.4% of dengue-positive patients experienced headaches while 80.2% experienced respiratory symptoms (Table 1).

Platelet levels of the patients were analyzed and categorized as abnormal ($\leq 150 \times 10^9/L$) and normal ($\geq 150 \times 10^9/L - 450 \times 10^9/L$). As shown in Table 2, the largest numbers of dengue positives were found in the age group 21-30, 27.9% in the normal platelet range and 4.7% in the abnormal platelet range, however, this difference was not statistically significant ($p = 0.172$). The age group 11-20 showed the second highest number of dengue positives with 18.6%. The mean age of those that tested positive was 29 years old, while the mean platelet counts were 130,000 and 293,000 within the abnormal and normal range, respectively. Except for the age groups 21 – 30 that recorded abnormal platelet counts, all the other age groups had no abnormal platelet counts (Table 2).

Table 1. Characteristic features of individuals surveyed for dengue virus infection in the north central regional health authority, Trinidad and Tobago, 2016 – 2017

Characteristics		Negative (%)	Positive (%)	p-value
Demographics	Male	8 (28.6)	139 (39.5)	0.702
	Female	20 (71.4)	213 (60.5)	0.702
	African descent	12 (42.9)	131 (37.2)	1.000
	East Indian descent	4 (14.3)	82 (23.3)	1.000
	Mixed	12 (42.9)	135 (38.4)	1.000
	Spanish	0	4 (1.2)	1.000
Symptoms	Rash	4 (14.3)	41 (11.6)	1.000
	Headache	16 (57.1)	311 (88.4)	0.054*
	Retro-orbital pain	0	188 (53.5)	0.012*
	Body pain	20 (71.4)	274 (77.9)	0.654
	Joint pain	4 (14.3)	176 (50)	0.115
	Diarrhoea	8 (28.6)	119 (33.7)	1.000
	Cough, cold, runny nose	8 (28.6)	282 (80.2)	0.007*
	Gum/nose bleeds	0	33 (9.3)	1.000
	Previous infections	None	28 (100)	254 (72.1)
Dengue		0	8 (2.3)	1.000
Chikungunya		0	29 (8.1)	1.000
Zika		0	4 (1.2)	1.000
Co-morbidities	Hypertension	0	17 (4.7)	1.000
	Diabetes	0	8 (2.3)	1.000
	Diabetes + HTN	0	4 (1.2)	1.000
	Asthma	0	37 (10.5)	1.000
	Other – Arthritis, PCOS, etc.	4 (14.3)	29 (8.1)	0.479
	None	24 (85.7)	254 (72.1)	0.670
Mosquito Conditions	Many mosquitoes in area	24 (85.7)	237 (67.4)	0.428
	Nets/screens at home	0	61 (17.4)	0.593
	Blocked drains around house	0	70 (19.8)	0.342
	Get bitten often	20 (71.4)	193 (54.7)	0.459
	No mosquito problems	4 (14.3)	111 (31.4)	0.670

* $p < 0.05$ is considered statistically significant. P-values were determined using Chi-square tests. Data are presented as n (%) or median (interquartile range); HTN – hypertension, PCOS – polycystic ovary syndrome

Table 2. Age distribution of participants for dengue who were ELISA positive categorized by platelet counts

Age groups	Negative ELISA		Positive ELISA	
	Abnormal (%)	Normal (%)	Abnormal (%)	Normal (%)
1 – 10	0 (.0)	16 (4.3)	0 (.0)	45 (11.8)
11 – 20	0 (.0)	8 (2.1)	0 (.0)	65 (17.2)
21 – 30	4 (1.0)	0 (.0)	16 (14.3)	98 (25.8)
31 – 40	0 (.0)	0 (.0)	0 (.0)	38 (9.7)
41 – 50	0 (.0)	0 (.0)	0 (.0)	33 (8.6)
51 – 60	0 (.0)	0 (.0)	0 (.0)	41 (10.8)
61 – 70	0 (.0)	0 (.0)	0 (.0)	4 (1.1)
71 – 80	0 (.0)	0 (.0)	0 (.0)	8 (2.2)
81+	0 (.0)	0 (.0)	0 (.0)	4 (1.1)
TOTAL	4 (1.0)	24 (6.4)	16 (4.3)	336 (88.3)

The Platelet counts were considered as abnormal ($\leq 150 \times 10^9/L$) and normal ($\geq 150 \times 10^9/L - 450 \times 10^9/L$)

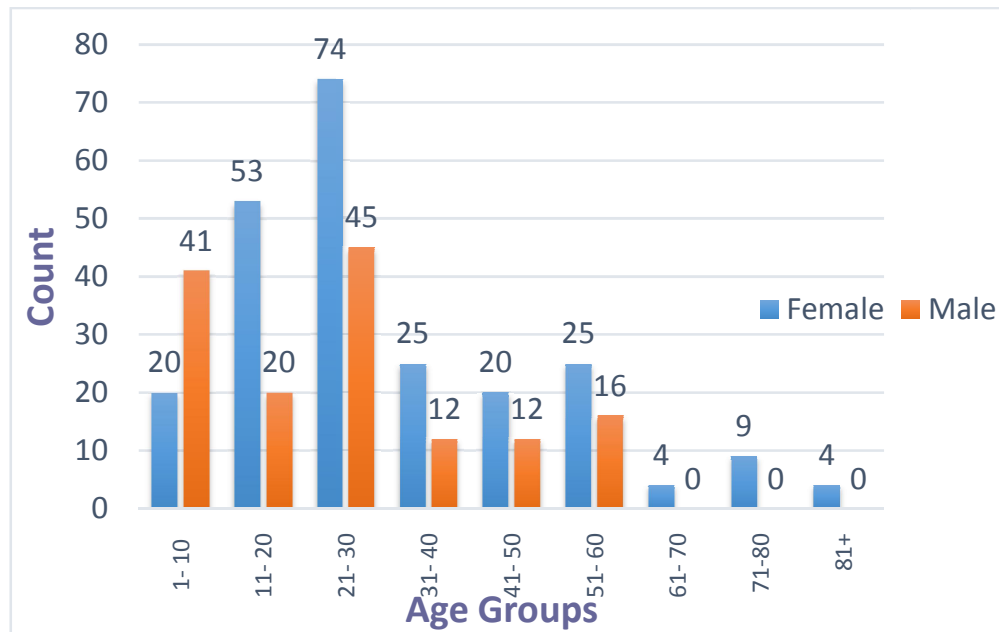


Fig. 2. Age and gender distribution of participants surveyed for dengue virus infections in Trinidad and Tobago

4. DISCUSSION

The objective of this study was to use serological analysis to determine the frequency of dengue virus infection and make the association between epidemiological risk factors that may exist among the patients suspected of the infection in a cross section of individuals in Trinidad and Tobago. Results from studies such as this can assist physicians in making a definitive diagnosis of dengue in our locality since many cases go unnoticed or recorded as acute viral illness (AVIs). While accurate laboratory diagnosis can be beneficial in confirming the disease, it will also

provide key data on the epidemiology and health burden of dengue, which is very useful for accurate public health surveillance [15]. Detection of seropositive cases of dengue in this region of study still suggest that vector control operations that have previously been carried out in this region failed to achieve the desired target of reducing mosquito densities in the eight counties to below the disease transmission threshold as previously reported by Chadee et al. [13] These authors had reported two decades ago that the Trinidad vector control program relied on the chemical approach with the application of insecticides in artificial containers;

[13] and this has continued to date. Perhaps more intensive and aggressive efforts may turn to health education.

Females were noted to be the majority (60.5%) of the dengue cases in our study which is different from what has been reported in other countries [16]. Adults were more affected in our analysis with ages 21-30 having 32.6% of all positive cases. This again was not in agreement with Anker and Arima that reported more of their positive cases occurring more in those over 15 years in the countries they studied [16]. Anker and Arima attributed the dominance of the males and the age group to cultural and economic reasons. Female was more perhaps because more took part in the study despite the fact that participants were systematically selected randomly. Economic differences could not have influenced our results as reported by others [16] since medical care is free in our country and so all are afforded the opportunity to seek medical care.

In this study, a similar number of individuals reported their ethnicity to be either of African descent or mixed race; and many of these tested positive for dengue virus infections. This was not in agreement with what was reported by Rojas et al in Colombia that Afro-Colombians population had a significantly lower risk of getting dengue and its complications, compared with the non-Afro-Colombians population [17]. Trinidad and Tobago is a cosmopolitan society with several ethnic groups, although the African and Indian descents dominate in number; but dengue virus infection could not be selective because all the different groups live together. Also, the majority of the participants surveyed gave their location to be Arima area which was also noted to be a significant factor in this study. The high number of positive results in each of these categories appears to only reflect the majority within the sampled population.

Symptoms were statistically analyzed to determine their associations with a dengue virus infection although dengue infections may initially be asymptomatic in 50 – 90% of individuals [18]. The significant ones include retro-orbital pain (eye pain), headaches and respiratory symptoms which are similar to a previous report [19]. Eye pain is particularly common in dengue infection along with headaches but the degree to which they are experienced are not quantifiable and so they remain non-specific. Most patients who tested positive for dengue antibodies also

complained of body pains; but this was not found to be significant. Reporting of having a previous infection of either dengue, chikungunya or zika, also did not show any differences for those who tested positive. Among the several patients that had already suffered from a dengue infection, none of them showed signs or symptoms that were more severe than those who said they never were infected with dengue. As dengue is one of the most under reported tropical diseases [6], it is very possible that patients who claimed to have never had dengue may be unaware of the past diagnoses seeing that symptoms are non-specific and home remedies are administered by patients themselves until symptoms subside. This way, there is and can be no accurate monitoring of the actual disease or possible burden of infection.

Co-morbidities such as hypertension, diabetes mellitus and asthma are among the non-communicable illnesses that are most prevalent in Trinidad and Tobago [20]. If left unmanaged they can lead to high morbidity and mortality rates. Whether or not either of these had any effects on the prevalence of dengue infection was also investigated. Most of those that were found positive for dengue infection showed no significant associations with having any medical conditions (asthma, diabetes, hypertension), being on any particular medications or having received any vaccines in the last two months prior to being enrolled. However, a study in Asia, attempted to show the association of diabetes mellitus with DHF. The study found that female, Chinese, age group 30-49 years with pre-existing diabetes mellitus or diabetes with hypertension were risk factors of developing DHF during an epidemic while dengue serotype 2 was predominant [21]. As stated above, neither of these characteristics were found to show any significant differences in our current study despite age group (21-30 years), gender (more females than males) or ethnicity (more of mixed ethnic group descents) gave more numbers; and also the fact that 25.5% of the sampled population in this study suffered from comorbidities.

In our locality where we do not have problem of distinguishing dengue from malaria that produces low platelet counts [22], hence platelet counts have been one of the most important factors in tracking the progress of dengue infection. Monitoring platelet levels, however, should not be the sole criteria to presume dengue infection as many patients in this study tested dengue

positive without abnormal platelet counts that are indicative of plasma leakage. In a study by Lovera et al., they investigated platelet count as a risk factor for shock. Using a cut-off of $< 100 \times 10^9/L$ they found that children who did not develop shock exhibited similar percentage level of thrombocytopenia compared to patients who eventually developed it (47% vs 49%). The results were similar when the comparison included patients only with platelet counts $< 50,000/mL$ (28% vs 25.6 %). [23] In this present study, the mean platelet count for positive samples in patients 1- 10 years of age was $295 \times 10^9/L$. Those with abnormal counts were only found in the 21 – 30 year-old age group and 80% of them tested positive for dengue virus. This adds up to 4.3% of those who tested positive but was not of any significance. None of the patients had platelet levels that were $< 50 \times 10^9/L$. Lam et al reported and the WHO guideline states that, daily platelet counts can be used to predict the development of DSS. [24,25]. Also in an extensive review, Leal de Azeredo et al. concluded that thrombocytopenia, coagulopathy, and vasculopathy are hematological abnormalities related to platelet and endothelial dysfunction generally observed in severe dengue [26]. We do not have proven explanations why majority of the patients who were suspected of dengue in our study had normal plate counts, but we can only speculate that their platelets were normal because they may have recovered.

The Pan American Health Organization (PAHO) has already issued a release of the number of reported cases of dengue and severe dengue in the Americas by country for epidemiological week 39 (updated October 13, 2017). After week 32 in Trinidad and Tobago the number of probable reported cases were 206, none of which were laboratory confirmed [10]. This is as a result of non-availability of the laboratory facilities because of lack of economic resources. It is however very critical that identification, isolation of the virus or confirmation of the dengue diagnosis be made so that dengue can successfully be managed and differentiated from other viral infections. It is also of utmost importance that all probable cases not only be reported but confirmed, especially if headway is to be made on curbing infection and development/implementation of a vaccine. The first dengue vaccine – the Sanofi CYD-TDV vaccine, has now been licensed by several endemic countries for use in 9-45 years and 9-60-year-olds. The vaccine was unusual in that the recommended target population for

vaccination was not only defined by age but also by transmission setting as defined by seroprevalence. The WHO has stated their position on the newly developed vaccine (CYD-TDV) saying that countries should consider the introduction of the dengue vaccine only in geographic settings where epidemiological data indicate a high burden of disease [27,28]. The vaccine, also known as Dengvaxia, is a live attenuated (recombinant) tetravalent vaccine that was created to be administered by 3 doses of 0.5ml given at 6-month intervals. We cannot indicate high burden of disease if the epidemiological data being collected is recorded incorrectly or disregarded. Hence, all assumptions for diagnoses need to be confirmed by the most accurate methods.

5. CONCLUSION

Despite the limitations of this study that include the small sample size and lack of use of molecular tests, viral isolation or virus detection using indirect immunofluorescence for confirmation of dengue virus, the study still detected positive cases of dengue virus infections in the country. Except for nonspecific symptoms observed among patients suspected of dengue fever, there were no other significant factors that were exclusive in identifying dengue infection among the subjects studied. Platelet monitoring may not be the only parameter to use in determining deteriorating dengue patients. Vector eradication activities in the country may not have been fully effective after all and so attention may also focus on other areas such as the education program.

CONSENT

Informed consent was also obtained from each of the patients, along with assent from children that were included in the study. Patients under the age of 18 were considered as children.

ETHICAL APPROVAL

Ethics approval for this study was obtained from the Campus Ethics Committee of the University of the West Indies St. Augustine Campus and the North Central Regional Health Authority (NCRHA) Ethics Committees. The study was carried out in accordance with the ethical standards laid down in the 1964 declaration of Helsinki.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Velayudhan R. A WHO report on global strategy for dengue prevention and control 2012 – 2020; WHO/HTM/NTD/VEM/2012.5.; 2018.
Available:<http://www.who.int/en/>
2. Wesolowski A, Qureshi T, Boni MF, Sundsøy PR, Johansson MA, Rasheed BS, et al. Impact of human mobility on dengue epidemics. *Proceedings of the National Academy of Sciences* 2015; 112(38):11887-11892.
DOI: 10.1073/pnas.1504964112
3. Tuyet-Hanh TT, Cam NN, Thanh Huong LT, Khanh Long T, Mai Kien T, Kim Hanh DT, Huu Quyen N, et al. Climate variability and dengue hemorrhagic fever in Hanoi, Vietnam, during 2008 to 2015. *Asia Pacific Journal of Public Health*; 2018.
Available:<https://doi.org/10.1177/1010539518790143>
4. Guzman A, Isturiz RE. Update on the global spread of Dengue. *Int J Antimicrobial Agents*. 2010;(Suppl 1):S40-S42.
5. Bhatt S, Gething PW, Brady OJ, Messina JP, Farlow AW, Moyes CL, et al. The global distribution and burden of dengue. *Nature*. 2013;496:504-7.
6. Gubler DJ. Dengue and dengue hemorrhagic fever. *Lin Microbiol Rev*. 1998;11:480-496.
7. Mustafa MS, Rasotgi V, Jain S, Gupta V. Discovery of fifth serotype of dengue virus (DENV-5): A new public health dilemma in dengue control. *Med J Armed Forces India* 2015;71(1):67-70.
DOI: 10.1016/j.mjafi.2014.09.011
8. Vaughn DW, Green S, Kalayanaroj S, Innis BL, Nimmannitya S, Suntayakorn S, et al. Dengue viremia titer, antibody response pattern, and virus serotype correlate with disease severity. *J Infect Dis* 2000;181(1):2-9.
DOI: 10.1086/315215
9. Thomas SJ, Endy TP, Rothman AL, Barrett AD. Flaviviruses (Dengue, Yellow Fever, Japanese Encephalitis, West Nile Encephalitis, St. Louis Encephalitis, Tick-Borne Encephalitis, Kyasanur Forest Disease, Alkhurma Hemorrhagic Fever, Zika). In Mandell, Douglas, and Bennett's *Principles and Practice of Infectious Diseases*, 8th edition. Edited by: Bennett JE, Dolin R, Blasser MJ. Philadelphia, PA 19103-2899: Elsevier Saunders. 2015; (Chapter 155):1881–1906.
10. PAHO. Number of Reported Cases of Dengue and Severe Dengue (SD) in the Americas, by Country. Figures for 2016 (to week noted by each country). *Epidemiological Week / EW* 52; 2017.
(Accessed online June 10, 2018)
Available:www.paho.org/hq/dmdocuments/2016/2016-cha-dengue-cases-jan-26-ew-52.pdf
11. Chadee DD, Shivnauth B, Rawlins SC, Chen AA. Climate, mosquito indices and the epidemiology of dengue fever in Trinidad (2002–2004). *Annals of Tropical Medicine & Parasitology*. 2007;101(1):69-77.
DOI: 10.1179/136485907X157059
12. Campbell CA, George A, Salas RA, Williams SA, Doon R, Chadee DD. Seroprevalence of dengue in Trinidad using rapid test kits: A cord blood survey. *Acta Trop*. 2007;101(2):153-158.
DOI: 10.1016/j.actatropica.2006.11.009
13. Chadee DD, Williams FL, Kitron UD. Impact of vector control on a dengue fever outbreak in Trinidad, West Indies, in 1998. *Trop Med Int Health*. 2005;10(8):748-754.
DOI: 10.1111/j.1365-3156.2005.01449
14. WHO. *Dengue: Guidelines for Diagnosis, Treatment, Prevention and Control: New Edition 2009*; World Health Organization. (Accessed online June 10, 2018)
Available:www.who.int/rpc/guidelines/9789241547871/en
15. Parkash O, Shueb RH. Diagnosis of dengue infection using conventional and biosensor based techniques. *Viruses*. 2015;7(10):5410-5427.
DOI: 10.3390/v7102877
16. Anker M, Arima Y. Male-female differences in the number of reported incident dengue fever cases in six Asian countries. *Western Pacific Surveillance and Response Journal*. 2011;2(2):17-23.
DOI: 10.5365/wpsar.2011.2.1.002
17. Rojas PJH, Alzate A, Martínez Romero HJ, Concha-Eastman AI. Afro-Colombian ethnicity, a paradoxical protective factor

- against Dengue. *Colomb Med (Cali)*. 2016; 47(3):133-41.
18. Kyle JL, Harris E. Global spread and persistence of dengue. *Anual Rev Microbiol*. 2008;62:71-92.
19. Gregory J, Santiago LM, Arguello DF, Hunsperger E, Tomashek KM. Clinical and laboratory features that differentiate dengue from other febrile illness in an endemic area – Puerto Rico, 2007 – 2008. *Am J Trop Med Hyg*. 2010;82:922-929.
20. WHO. Non communicable Diseases (NCD) Country Profiles: Trinidad and Tobago; 2014.
(Accessed online June 10, 2018)
Available:www.who.int/nmh/publications/ncd-profiles-2014/en
21. Pang J, Salim A, Lee VJ, Hibberd ML, Chia KS, Leo YS, Lye DC. Diabetes with hypertension as risk factors for adult dengue hemorrhagic fever in a predominantly dengue serotype 2 epidemic: A case control study. *PloS Negl Trop Dis*. 2012;6(5):e1641.
DOI: 10.1371/journal.pntd.0001641
22. Chadwick D, Arch B, Wilder-Smith A, Panton N. Distinguishing dengue fever from other infections on the basis of simple clinical and laboratory features: Application of logistic regression analysis. *J Clin Virology*. 2006;35:147-153.
23. Lovera D, Martinez de Cuellar C, Araya S, Amarilla S, Gonzalez N, Aguiar C, Arbo A. Clinical characteristics and risk factors of dengue shock syndrome in children. *Pediatr Infect Dis J*. 2016;35(12):1294-1299.
DOI: 10.1097/INF.0000000000001308
24. Lam PK, Ngoc TV, Thu Thuy TT, Hong Van NT, Nhu Thuy TT, Hoai Tam DT, et al. The value of daily platelet counts for predicting dengue shock syndrome: Results from a prospective observational study of 2301 Vietnamese children with dengue. *PLoS Negl Trop Dis*. 2017;11(4): e0005498.
Available:<https://doi.org/10.1371/journal.pntd.0005498>
25. WHO. Dengue haemorrhagic fever: Diagnosis, treatment, prevention and control. World Health Organization, Geneva, Switzerland; 1997.
(Accessed online June 10, 2018)
Available:<http://www.who.int/csr/resources/publications/dengue/Denguepublication/en/>
26. Leal de Azeredo E, Monteiro RQ, Pinto LM-O. Thrombocytopenia in dengue: Interrelationship between virus and the imbalance between coagulation and fibrinolysis and inflammatory mediators. *Mediators of Inflammation*; 2015.
(Article ID 313842, 16, 2015)
Available:<https://doi.org/10.1155/2015/313842>
27. WHO. Dengue vaccine: WHO position paper *Weekly Epidemiological Report*. 2016;30(91):349-364.
28. Imai N, Ferguson NM. Targeting vaccinations for the licensed dengue vaccine: Considerations for serosurvey design. *PLoS ONE*. 2018;13(6):e0199450.
Available:<https://doi.org/10.1371/journal.pone.0199450>

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