EARLY DETECTION OF DENGUE INFECTIONS USING CLUSTER SAMPLING AROUND INDEX CASES

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Abstract. A two-year study using a cluster investigation method was conducted in West Jakarta, Indonesia to demonstrate the detection of dengue cases prior to onset of clinical illness. The clusters consisted of family members and neighbors of 53 hospitalized dengue index cases. Among 785 adult and child volunteers enrolled, 17 (2.2%) post-enrollment dengue (PED) infections were identified. Eight PED cases were asymptomatic and nine were symptomatic. Symptomatic cases included eight with dengue hemorrhagic fever (DHF) (grade II). Among the eight asymptomatic PED cases, viremia was detected in two. Eleven volunteers had acute dengue infections at the time of enrollment. Four of the 11 developed DHF, resulting in a total of five DHF cases detected during the investigation. This study design can serve as a benchmark for future investigations that seek to define early immunologic events following dengue infections that contribute to the development of DHF.

INTRODUCTION

Infection with any of the four serotypes of dengue virus (DEN-1, DEN-2, DEN-3, and DEN-4) can be asymptomatic, result in a mild-to-moderate febrile illness termed dengue fever (DF), or a more severe illness characterized by bleeding and shock called dengue hemorrhagic fever (DHF) or dengue shock syndrome (DSS). After an initial infection with dengue virus, homologous serotype immunity persists for many years; however, individuals remain susceptible to infection with the remaining dengue virus serotypes. In several studies, DHF occurs 15–80 times more frequently in individuals experiencing a second dengue virus infection compared with individuals experiencing their first infection. The biologic events that lead to a higher frequency of DHF in secondary dengue virus infections compared with primary infections are not completely known.

Antibody-dependent enhancement of dengue virus infection leading to increased viral load is a leading theory for severe diseases caused by dengue virus. Increased levels of circulating dengue virus have been documented in DHF cases compared with DF cases. However, the immunologic events that occur in the early stages of infection that lead to severe disease have not been characterized in patients. In the absence of an animal model for DHF, researchers are forced to try to characterize early immunologic events in patients with recently acquired dengue virus infections. A major limiting factor in such an investigation is identifying individuals prior to the onset of their clinical illness so that early events can be characterized. The current study was designed to overcome that problem by identifying individuals very early in the infection. Family members and nearest neighbors of index cases hospitalized with acute dengue virus infections were enrolled and followed longitudinally to determine the incidence rate of dengue virus infection. This proof-of-concept study was conducted to evaluate the hypothesis that monitoring persons clustered around an index case would be an efficient and practical way to obtain blood samples before, during, and after dengue infection.

MATERIALS AND METHODS

Study location and population. West Jakarta (West Java), Indonesia, was selected as the location to pilot the concept of a dengue cluster study for several reasons. The catchment area for cases is in a tropical, urban setting with approximately 2.5 million inhabitants of various socioeconomic strata, mostly of low income. On average, 5–10 persons may live in a single housing unit measuring 10 × 15 meters. Most neighborhoods are densely populated, with dwellings in relatively close proximity to one another (approximately 10 meters apart) and constructed of cement and/or wood. The average house does not have piped water and some communities share toilet facilities. The government supplies potable water in large cans and water for bathing is often stored in a traditional uncovered, tub-like area (bak mandi) located in bathrooms. The bak mandi has long been a potential source of Aedes mosquito larvae, in addition to other open containers used for water storage and various other miscellaneous items (i.e., flower pots, tires) containing water found in close proximity to human dwellings. Aedes aegypti mosquitoes are ubiquitous throughout Indonesia and transmission of all four dengue virus serotypes is known to occur. According to the Indonesian Communicable Disease Control, the three-year incidence of dengue disease in Jakarta was 75.4, 52.4, and 108.5 cases per 100,000 population from 2001 to August 2003, respectively (unpublished data).

Dengue virus laboratory tests. Serum samples were tested for the presence of IgM antibodies to dengue virus using commercial enzyme-linked immunosorbent assay (ELISA) kits (Focus Technologies, Cypress, CA). Hemagglutination inhibition (HI) assays and plaque reduction neutralization tests (PRNTs) were also performed to confirm dengue virus infection and to classify the infection as primary or secondary based on the antibody response. Dengue virus RNA in blood was detected using a reverse transcriptase–polymerase chain reaction (RT-PCR) assay. Blood samples collected from symptomatic volunteers during the acute stage of febrile illness and from suspected asymptptomatically infected volun-
teers were tested for virus by culture in C6/36 cells, and virus was identified using serotype-specific monoclonal antibodies for dengue virus.

**Study design.** Each cluster of volunteers was identified by an index case from the pediatric ward at Sumber Waras Hospital in West Jakarta. Physicians at this local community hospital have a well-established relationship with the surrounding community and are experienced in clinically diagnosing dengue virus infections. For the index cases, children 4–14 years of age were recruited. Ward clinicians were asked to select 1–2 cases per week based on clinical manifestations, together with the detection of IgM antibody to dengue virus by ELISA and/or dengue viral RNA by RT-PCR. In most cases, these diagnostic assays were performed within 48 hours of index case identification. Other considerations in selecting index cases included automobile accessibility to their community and the willingness of the family to participate.

Usually within 48 hours of index case identification, family members and nearest neighbors greater than four years old who lived within a 10-meter radius of the index case’s home were invited to participate. Volunteers who were afflicted with or who had a history of severe anemia, bleeding disorder, or any known immunologic disorder were excluded from the study. On enrollment, we measured the otic temperature, and collected blood samples and demographic data from each study volunteer. Volunteers were followed for 14 consecutive days for fever and other clinical manifestations suggestive of acute dengue virus infection. During the monitoring period, we collected additional blood samples from each volunteer every 2–3 days. We also collected blood specimens from each volunteer who developed a fever or dengue-like signs or symptoms and processed them for a diagnosis of dengue. To confirm that dengue virus was the cause of fever, we tested samples within 24–48 hours of collection for dengue viral RNA by RT-PCR and for IgM antibody to dengue virus by ELISA. If a sample collected from a febrile volunteer was positive for IgM antibodies to dengue virus, the enrollment blood sample for that volunteer was tested for IgM antibodies to dengue virus to determine if seroconversion had occurred. These laboratory tests were performed at the Virology Laboratory of the U.S. Naval Medical Research Unit No. 2. Blood samples treated with EDTA were also obtained at the onset of fever and sent to Sumber Waras Hospital for hematocrit and platelet count determinations.

Febrile volunteers with a positive IgM ELISA or RT-PCR result were encouraged to be hospitalized for close monitoring and serial laboratory tests. Volunteers refusing hospitalization were monitored closely as outpatients until they were afebrile for two consecutive days. Convalescent blood samples were obtained two weeks later and again at six months.

Hospitalized volunteers were bled daily until two days after defervescence, two weeks later, and approximately six months after discharge from the hospital. Blood samples were routinely tested for hemoglobin, hematocrit, platelets, white blood cells, protein, and albumin. A tourniquet test was done on admission and daily ultrasound examinations for ascites and pleural effusions were performed until two days after defervescence to detect evidence of plasma leakage. We classified hospitalized volunteers as DF or DHF (World Health Organization, 1997) following evaluation of the clinical data.

We categorized all volunteers clustered around the index cases into one of three groups: non-dengue (ND) infection, dengue infection at enrollment (ED), or post-enrollment dengue (PED) infection. A PED case was defined as a volunteer who developed fever after enrollment and demonstrated seroconversion for IgM antibodies to dengue virus and/or dengue viremia by RT-PCR or virus isolation. Volunteers who developed fever but were negative for IgM antibodies to dengue virus or viremia were later classified as PED infections if there was a four-fold increase in the HI antibody titer to dengue virus between enrollment and the two-week post-enrollment blood samples. For equivocal HI results (less than a four-fold increase in titer), pre-enrollment and two-week post-enrollment PRNT50 titers were compared for a ≥ four-fold increase against one or more dengue virus serotypes. For volunteers who never developed fever, laboratory evaluation of serum samples (dengue serology and virus detection) was still conducted throughout the 14-day monitoring period. Asymptomatic volunteers who were positive for viremia and/or showed a four-fold increase in HI titer between the enrollment and two-week post-enrollment blood samples were also classified as PED cases.

Volunteers that demonstrated evidence of infection (IgM, RT-PCR, or virus isolation), with or without symptoms at the time of enrollment were classified as ED cases. All other volunteers were classified as ND cases.

The U.S. Naval Medical Research Unit No. 2 Institutional Review Board and the Indonesian Ministry of Health Ethical Review Committee reviewed and approved the human use protocol (DoD# 30861) used in the study, in compliance with all U.S. federal regulations governing the protection of human subjects. Informed written consent was obtained from all participants, and if < 15 years of age, from a parent or legal guardian.

**RESULTS**

Over a two-year period from October 2001 to October 2003, 53 confirmed dengue index cases were identified in the pediatrics ward at Sumber Waras Hospital (Table 1 and Figure 1). Infections by all dengue virus serotypes were identified among the index cases. The predominant serotype identified was DEN-1 (10), followed by DEN-2 (8), DEN-3 (5), and DEN-4 (2). The serotype was not determined in 28 index cases. None of the index dengue infections was fatal.

From these 53 dengue index cases, 53 cluster investigations were performed where family members and nearest neigh-

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**Table 1**

Demographics of volunteers and corresponding dengue infections identified in West Jakarta, Indonesia*  

<table>
<thead>
<tr>
<th>Number of index cases</th>
<th>Community volunteers</th>
<th>ED cases</th>
<th>PED cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>53</td>
<td>785</td>
<td>175</td>
</tr>
<tr>
<td>Males:females</td>
<td>23:30</td>
<td>317:468</td>
<td>84:91</td>
</tr>
<tr>
<td>Adults</td>
<td>NA</td>
<td>453</td>
<td>87</td>
</tr>
<tr>
<td>Children (4–14 years old)</td>
<td>53</td>
<td>332</td>
<td>88</td>
</tr>
<tr>
<td>Mean age (years)</td>
<td>6.6</td>
<td>22.57</td>
<td>19.2</td>
</tr>
<tr>
<td>Family members</td>
<td>NA</td>
<td>164</td>
<td>32</td>
</tr>
<tr>
<td>Nearest neighbors</td>
<td>NA</td>
<td>621</td>
<td>143</td>
</tr>
</tbody>
</table>

*ED = enrollment dengue cases; PED = post-enrollment dengue cases; NA = not applicable.
bors surrounding the residence of each index case were monitored for dengue virus infection. A total of 785 volunteers were recruited from the 53 community clusters, of which 541 completed the entire 14-day monitoring period. The 244 volunteers who did not complete the entire monitoring period were included in the analysis since at least one blood specimen was available for evaluation from each volunteer. Table 1 shows the demographics of the volunteers in the study.

Among the 785 cluster volunteers, 175 (22.3%) were classified as ED cases. Table 1 also shows the demographics of ED cases. Eleven ED cases were found to be viremic for dengue at enrollment by the RT-PCR. The DEN-1 serotype was identified in three cases and DEN-2 in eight cases. Confirmatory virus isolation was accomplished in 5 of the 11 cases. The remaining 164 ED cases had evidence of recent dengue virus infection, as indicated by IgM antibodies to dengue virus at enrollment and were classified as convalescent infections (Figure 1).

Nine of the 11 viremic ED cases had fever at enrollment. The nine febrile volunteers were hospitalized for close monitoring within 24–48 hours of enrollment after dengue serologic and/or RT-PCR results were known. Four of the febrile hospitalized ED cases were subsequently diagnosed with DHF (three grade I and one grade II). All four DHF cases were caused by a DEN-2 virus. Analysis of HI and PRNT titers suggested that two of the DHF cases were secondary infections and two were primary infections. The volunteers with primary infections were 10 and 8 years old, whereas those with the secondary infections were 20 and 21 years old.

Seventeen (2.2%) volunteers were diagnosed as PED cases during the 14-day monitoring period (Table 2), and most (15 of 17, 88%) were among nearest neighbors (Table 1). Of the 17 PED cases, 10 were positive by the RT-PCR (1 DEN-1, 7 DEN-2, 1 DEN-3, and 1 DEN-4), and the serotype was confirmed by virus isolation in 5 cases. The other seven PED cases became positive for IgM antibodies to dengue virus during monitoring, and dengue infections were also evident by seroconversion (≥ 4-fold increase in antibody titer) by HI or PRNT. Four of the 17 PED cases were primary infections based on HI and PRNT results, and all others were secondary infections.

Nine of the 17 PED cases experienced symptomatic illnesses. Eight of the nine were DF and one was diagnosed as DHF grade II. The HI and PRNT patterns comparing pre-illness and convalescent blood samples suggested a secondary infection in all but three cases (Table 2).

The one PED DHF case was hospitalized five days after enrollment with a fever of 38.8°C and a positive tourniquet test result, along with a history of epistaxis prior to admission that continued for two days while hospitalized. Small pleural

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**Figure 1.** Map of index cases and corresponding dengue infections identified in respective clusters in West Jakarta, Indonesia. Each circle represents an index case. Each square represents an acute dengue case identified at enrollment by detection of virus (ED acute infection) or during a two-week period of monitoring by detection of virus or by seroconversion (post-enrollment dengue infection). The numbers inside the circles represent convalescent dengue infections detected by the presence of IgM antibody only at enrollment (ED convalescent infection). The study hospital (Sumber Waras Hospital) is indicated.
Characteristics of 17 post-enrollment dengue (PED) infection cases identified

<table>
<thead>
<tr>
<th>Study no.</th>
<th>Age (years)/sex</th>
<th>DEN serotype</th>
<th>Clinical classification*</th>
<th>Serologic response</th>
</tr>
</thead>
<tbody>
<tr>
<td>CL 0102</td>
<td>9/F</td>
<td>4†</td>
<td>DF</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 0118</td>
<td>18/M</td>
<td>Asymptomatic</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 0417</td>
<td>37/F</td>
<td>Asymptomatic</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 0710</td>
<td>14/F</td>
<td>Asymptomatic</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 1002</td>
<td>7/M</td>
<td>2</td>
<td>DF</td>
<td>Primary</td>
</tr>
<tr>
<td>CL 1211</td>
<td>5/M</td>
<td>DF</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 1514</td>
<td>30/F</td>
<td>2</td>
<td>DF</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 1806</td>
<td>10/F</td>
<td>2†</td>
<td>DHF grade II</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 2202</td>
<td>8/M</td>
<td>2†</td>
<td>DF</td>
<td>Primary</td>
</tr>
<tr>
<td>CL 2204</td>
<td>40/F</td>
<td>2†</td>
<td>Asymptomatic</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 2408</td>
<td>47/F</td>
<td>1</td>
<td>Asymptomatic</td>
<td></td>
</tr>
<tr>
<td>CL 3213</td>
<td>24/M</td>
<td>2†</td>
<td>DF</td>
<td>Primary</td>
</tr>
<tr>
<td>CL 3716</td>
<td>9/M</td>
<td>Asymptomatic</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 3911</td>
<td>32/F</td>
<td>2</td>
<td>DF</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 4202</td>
<td>11/M</td>
<td>3</td>
<td>DF</td>
<td>Secondary</td>
</tr>
<tr>
<td>CL 5115</td>
<td>36/M</td>
<td>Asymptomatic</td>
<td>Secondary</td>
<td></td>
</tr>
<tr>
<td>CL 5304</td>
<td>7/F</td>
<td>Asymptomatic</td>
<td>Primary</td>
<td></td>
</tr>
</tbody>
</table>

* DF = dengue fever; DHF = dengue hemorrhagic fever.
† Positive by both dengue virus reverse transcriptase-polymerase chain reaction and virus isolation in C3/06 cells.

Our cluster investigation method was designed as an alternative approach to the commonly used prospective cohort study method for investigating the natural history of dengue virus infection. Unlike the cohort study, the cluster investigation method allows the collection of clinical information and biologic samples during the early pre-infection stage. This allows for more precise characterization of the early immune pathologic events that contribute to either mild or severe clinical manifestations of dengue virus infection.

Among 785 volunteers in this investigation, we observed 17 PED (new) dengue infections that were both symptomatic and asymptomatic, including one case of DHF. The calculated incidence rate of dengue infection was 567 cases per 1,000 person-years of follow-up, based on the one observed DHF case, the calculated DHF incidence rate was 33 DHF cases per 1,000 person-years of follow-up, resulting in a DHF to DF incidence rate ratio of 1:18. This ratio closely approximates other epidemiologic observations, showing that DHF occurs in roughly 5% of dengue virus infections in areas where all four serotypes of dengue virus are endemic.

Virologic analysis of dengue cases confirmed that all four serotypes circulated in West Jakarta. The DEN-1 serotype was the predominant virus identified among the index cases, but DEN-2 caused most of the cluster cases, including all five serotypes.
Dengue/Dengue Haemorrhagic Fever

The natural history of dengue virus infection, such silent viremia would seem plausible, but with the design of typical longitudinal cohort studies, infected volunteers are only identified upon the development of symptoms. Asymptomatic infections are only detected by seroconversion of blood samples collected at pre-determined time points. Using a cluster design approach, we were able to document eight asymptomatic dengue infections, two of which demonstrated viremia. To our knowledge, this is the first documentation of asymptomatic viremia in naturally acquired dengue infections.

The one obvious disadvantage of the cluster method is that volunteers are required to participate in an intensive period of surveillance where, despite being healthy at the time, they are asked to donate blood samples every 2–3 days for a period of two weeks. We had little difficulty recruiting the requisite 10–15 people per cluster and often had to refuse additional enrollment of willing volunteers. With augmented resources and experience, we plan to expand our study design to new areas (East Jakarta and another major city in West Java, Indonesia) and to increase the number of index cases (6–8 per week) and persons recruited per cluster (up to 20). The major challenge to our cluster study design is maintaining adequate support (finances and trained personnel) for the intensive community monitoring. We believe that the yield of the cluster investigation method is equal to or substantially greater than that of other prospective study methods used to examine the epidemiology of naturally acquired dengue virus infection. As this method is applied to other countries with varying cultural sensitivities and expectations, differing levels of success may be encountered.

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REFERENCES


