Comparison of Three Commercially Available Dengue NS1 Antigen Capture Assays for Acute Diagnosis of Dengue in Brazil

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Abstract

Background: Dengue is associated with explosive urban epidemics and has become a major public health problem in many tropical and subtropical developing countries, including Brazil. The laboratory diagnosis of dengue can be carried out using several approaches; however, sensitive and specific assays useful to diagnose in the early stage of fever are desirable. The flavivirus non-structural protein NS1, a highly conserved and secreted glycoprotein, is a candidate protein for rapid diagnosis of dengue in endemic countries.

Methodology/Principal Findings: We aimed to evaluate the potential use of 3 commercial kits in a panel of 450 serum samples for early diagnosis of dengue in Brazil. The PanBio Early ELISA (PanBio Diagnostics) showed a sensitivity of 72.3% (159/220) and a specificity of 100%, while the sensitivity of the Platelia™ NS1 assay (Biorad Laboratories) was 83.6% (184/220). However, the highest sensitivity (89.6%; 197/220) was obtained by using the NS1 Ag Strip (Biorad Laboratories). A lower sensitivity was observed in DENV-3 cases by all 3 kits. Serum positive by virus isolation were more often positive than cases positive by RT-PCR by all three assays and a higher detection rate was observed during the first four days after the onset of the symptoms. The presence or absence of IgM showed no influence in the confirmation by the pan-E Early ELISA (P = 0.6159). However, a higher confirmation by both Platelia™ NS1 (Biorad) and Dengue NS1 Ag Strip (Biorad) in the absence of IgM was statistically significant (P<0.0001 and P = 0.0008, respectively). Only the Platelia™ NS1 test showed a higher sensitivity in confirming primary infections than secondary ones.

Conclusions/Significance: The results indicate that commercial kits of dengue NS1 antigen are useful for the laboratory diagnosis of acute primary and secondary dengue. It can be used in combination with the MAC-ELISA for case detection and as screening test to complement viral isolation.

Introduction

Dengue is associated with explosive urban epidemics and has become a major public health problem [1]. Annually, the World Health Organization estimates that 50–100 million people are infected with dengue virus (DENV) worldwide with estimated 250,000–500,000 cases of dengue haemorrhagic fever (DHF) and dengue shock syndrome (DSS) with about 25,000 deaths occurring. One or more of four serotypes of DENV (DENV1–4), a mosquito-borne, positive-strand RNA virus in the genus Flavivirus, family Flaviviridae cause the disease in more than 100 endemic countries in tropical areas [2].

The geographical spread of all four DENV serotypes throughout the subtropical regions of the world has led to larger and more severe outbreaks and the accurate and efficient diagnosis of the disease is important for clinical care, surveillance, pathogenesis studies and vaccine research. Furthermore, an efficient diagnosis is an important tool to support Epidemiological Surveillance Programs considering the difficulties in confirming dengue cases based only on the clinical symptoms, especially during inter-epidemic periods.

Dengue is an enveloped virus with a single-stranded, positive sense RNA genome of about 11 kb containing a single open reading frame encoding a single polyprotein co- and post-translationally cleaved into 3 structural (C, prM and E) and 7 nonstructural proteins (NS1, NS2A, NS2B, NS, NS4A, NS4B and NS5) [3].

Dengue is a major public health problem in many tropical and subtropical countries in the world. The accurate and efficient diagnosis of dengue is important for clinical care, surveillance, pathogenesis studies, and vaccine research.

The most used techniques for dengue serodiagnosis are based on the anti-DENV IgM and IgG detection by using MAC-ELISA and IgG-ELISA [4]. However, one of the limitations...
Author Summary

Dengue is one of the most prevalent arthropod-borne viral diseases in tropical regions of the world. Manifestations may vary from asymptomatic to potentially fatal complications. Laboratory diagnosis is essential to diagnose dengue and differentiate it from other diseases. Dengue virus non-structural protein 1 (NS1) may be used as a marker of acute dengue virus infection. Our results, based on the comparison of three NS1 antigen capture assays available, have shown that this approach is reliable for the early diagnosis of dengue infections, especially in the first four days after the onset of the symptoms. A lower sensitivity was observed in DENV-3 cases. Serum positive by virus isolation were more often detected than those positive by RT-PCR by all three assays. Only the Platelia™ NS1 test showed a higher sensitivity in confirming primary infections than secondary ones. In conclusion, NS1 antigen capture commercial kits are useful for diagnosis of acute primary and secondary dengue infections and, in endemic countries where secondary infections are expected to occur, may be used in combination with MAC-ELISA to increase the overall sensitivity of both tests.

Materials and Methods

Ethics statement

The samples belong to a previously-gathered collection from an ongoing Project in the Laboratory approved by the Ethics Committee on Human Research (CEP: 274/05).

Dengue cases and non-dengue cases definitions

Laboratory-positive DENV infection was defined in patients experiencing a febrile illness consistent with dengue according to WHO criteria [20] in which infection was confirmed by DENV isolation [21], detection of DENV RNA by RT-PCR [22], and/or a >4-fold rise in anti-DENV IgG-ELISA titer in paired acute and convalescent sera [24]. Individuals negative for DENV infection by using all the methods described above and health individuals were classified as non-dengue cases.

Clinical samples

The serum samples (days 1st to 9th after the onset of the symptoms) analyzed in this study by the pan-E Early ELISA (PanBio Diagnostics, Brisbane, Australia—first generation), Platelia™ (Biorad Laboratories, Marnes-La-Coquette, France) and NS1 Ag Strip (Biorad Laboratories, Marnes-La-Coquette, France) belong to a previously-gathered serum collection of the Laboratory of Flavivirus at Oswaldo Cruz Institute, FIOCRUZ, Brazil, from epidemics occurred from 1996 to 2008. A panel of 450 sera (220 dengue positive sera and 230 non-dengue sera) was divided into eleven Groups as follows: Groups A to C, sera from patients infected with DENV-1 (n = 50), DENV-2 (n = 50), and DENV-3 (n = 50), respectively; Group D, sera from patients with dengue infection serologically confirmed by MAC-ELISA with negative virus isolation and RT-PCR (n = 62); Group E, sera from healthy individuals (n = 30); Group F, sera from individuals negative for dengue (n = 86); Group G, sera from yellow fever positive individuals (n = 20); Group H, sera from individuals vaccinated for yellow fever and negative for anti-DENV antibodies (n = 44); Group I, sera from measles patients (n = 16) and Group J, sera from rubella patients (n = 34).

Dengue virus isolation

Virus isolation was performed by inoculation into C6/36 Aedes albopictus cell line [21] and isolates were identified by indirect fluorescent antibody test (IFAT) using serotype-specific monoclonal antibodies [25].

Reverse transcriptase (RT)–PCR

RT—PCR for detecting and typing DENV was performed as described previously [22]. Briefly, consensus primers were used to anneal to any of the four DENV types and amplify a 511-bp product in a reverse transcriptase-polymerase reaction. A cDNA copy of a portion of the viral genome was produced in a reverse transcriptase reaction. After a second round of amplification (nested PCR) with type-specific primers, DNA products of unique sizes for each DENV serotype were generated and analyzed by gel electrophoresis.

Immunoglobulin M (IgM) antibody capture ELISA (MAC-ELISA)

The in-house MAC-ELISA was carried out for dengue cases confirmation as described previously [23].

Immunoglobulin G (IgG) antibody detection ELISA (IgG—ELISA)

The IgG—ELISA previously described by Miagostovich [24] was performed for the characterization of dengue immune response as primary or secondary infections in dengue cases previously confirmed by virus isolation, RT—PCR and/or MAC-ELISA. Briefly, 96-well plates were coated with hyper immune ascitic fluid (a mixture of anti-DENV-1 to 4), followed by the addition of a mixture of the four DENV antigens. Serum diluted 1:40 was added to the first well and four-fold dilutions were carried out up to the eighth well. After incubation, anti-human IgG conjugated to horseradish peroxidase was added. Acute phase serum samples (<6 days after onset of symptoms) with IgG-ELISA titers of 1:160 or greater are considered to be secondary infections.
Likewise, samples with titers >1:10, 240 on days 6–9, or >1:40, 960 on days 10–15 after onset are secondary responses.

NS1 antigen capture assays

Pan-E Dengue Early ELISA (PanBio Diagnostics, Brisbane, Australia). This first generation test is based on a one-step sandwich format microplate enzyme immunoassay to detect DENV NS1 antigen in human sera. The test uses horseradish peroxidase conjugated anti-NS1 monoclonal antibody with preservative. If NS1 antigen is present in the sample, an immune-complex MAb-NS1-MAb/peroxidase will be formed. The acute serum specimens were allowed to thaw to laboratory ambient temperature (21–22°C). Briefly, 100 µl diluted test samples and controls were pipetted into their respective microwells and incubated for 60 min at 37°C. After a six-time washing steps, 100 µl HRP conjugate anti-NS1 MAb were pipetted into each well and plate was incubated for 60 min at 37°C. After a six-times washing step, 100 µl of substrate was added into each well and incubated for 10 min at room temperature in the dark. The presence of immune-complex was demonstrated by a color development and the enzymatic reaction was stopped by adding 100 µl 1M H3PO4. The optical density (OD) reading was taken with a spectrophotometer at a wavelength of 450–620 nm and the amount of NS1 antigen present in an individual serum sample was determined by comparing the OD of the sample to the OD of the cut-off control serum. Results were calculated as “Panbio units” with results <0.0, 0.0–11.0, and ≥11.0 defined as negative, inconclusive, and positive, respectively. Inconclusive samples were re-tested to confirm the result.

Platelia™ Dengue NS1 Ag-ELISA (Biorad Laboratories, Marnes-La-Coquette, France). The test is based on a one-step sandwich format microplate enzyme immunoassay to detect DENV NS1 antigen in human serum or plasma. The assay uses murine monoclonal antibody for capture and revelation. If NS1 antigen is present in the sample, an immune-complex MAb-NS1-MAb/peroxidase will be formed. Briefly, the acute serum specimens were allowed to thaw to laboratory ambient temperature (21–22°C). Sample diluent (50µl), respective samples and controls (50 µl each) and 100 µl of diluted conjugate were incubated for 90 min at 37°C within the respective microwells coated with purified mouse anti-NS1 monospecific antibodies. After a six-times washing step, 160µl of substrate was added into each well and incubated for 30 min at room temperature in the dark. The presence of immune-complex was demonstrated by a color development and the enzymatic reaction was stopped by adding 100µl 1N H2SO4. The optical density (OD) reading was taken with a spectrophotometer at a wavelength of 450–620 nm and the amount of NS1 antigen present in an individual serum sample was determined by comparing the OD of the sample to the OD of the cut-off control serum.

Dengue NS1 Ag Strip (Biorad Laboratories, Marnes-La-Coquette, France). Dengue NS1 Ag Strip is an immuno-chromatographic test (ICT) for the rapid detection of NS1 antigen. Briefly, one drop of migration buffer was added to 50 µL serum in a specimen tube and a strip was placed in the tube. The strip has two lines: a control line (C) (‘biotin – gold colloidal particles coated with streptavidin’ complex) and a test line (T) (‘monoclonal anti-NS1 antibodies (mAb) – NS1 Ag – gold colloidal particles coated with anti-NS1 mAb’ complex). The appearance of the T and C lines after a migration time of 15 minutes (min) indicates a positive result. The appearance of the C line alone indicates a negative result. If the C line is not present, the test is considered invalid and is repeated. It is recommended that strips giving ambiguous (faint color at the T line) or negative results are put back in the tube after the initial reading and left for a further 15 min for re-evaluation. We evaluated all samples at 15 min (ICT 15 min) and then at 30 min (ICT 30 min).

Data and statistical analysis

The sensitivities, specificities, efficiency, negative and positive predicted values were calculated as follows:

Sensitivity: a/a+c X 100%
Specificity: d/d+b X 100%
Efficiency: a+d+a+b+c+d X 100%
Negative Predicted Value: d/d+c X100%
Positive Predicted Value: a/a+b X 100%; where: a = number of true positive, b = number of false positive, c = number of false negative and d = number of true negative.

The derived data was tabulated in appropriate worksheets using the Microsoft Excel and evaluated by chi-square test using the Epi Info 6 (Center for Disease Control and Prevention, Atlanta) for any statistical significant association.

Results

Overall sensitivities of the NS1 antigen capture tests

A panel of 450 (n=220 dengue cases and n=230 non-dengue cases) was used to evaluate three NS1 antigen capture tests commercially available. The overall sensitivities were 72.3% (159/220) for the pan-E Early ELISA (PanBio) test, 83.6% (194/220) for the Platelia™ NS1 (Biorad) kit, and 89.6% (197/220) for the NS1 Ag Strip kit (BioRad), Table 1. The differences observed in the sensitivities between the three kits analyzed were statistically significant (P= 0.0009).

NS1 sensitivities in relation to viral serotype

The pan-E Early ELISA (PanBio) showed a higher sensitivity in confirming DENV-2 infections (Group B; 92.0%) than confirming DENV-1 and DENV-3 infections. The Platelia™ NS1 kit (Biorad) was more sensitive in the detection of DENV-1 cases (Group A; 90.0%) than in the detection of DENV-2 and DENV-3 infections. The Dengue NS1 Ag Strip kit (BioRad) showed the same sensitivity in confirming DENV-1 and DENV-2 infections (98.0%), DENV-3 infections were detected less often by all the tests. The overall sensitivities were 72.3% (159/220) for the Platelia™ NS1 kit (Biorad), respectively, based on the analysis of sera of healthy individuals (Group E) and individuals negative for dengue (Group F), Table 1. No cross-reactivity was observed with sera from yellow fever infected patients (Group G); however both Biorad kits showed cross-reactivity with one yellow fever vaccinee (Group H). None of the measles sera (Group I) were recognized by the NS1 tests. One rubella positive case (Group J) showed cross-reactivity with one yellow fever vaccinee (Group H).

NS1 specificities and cross-reactivity in patients with other confirmed diagnoses and yellow fever vaccination

Specificities were 100%, 98.7% and 99.1% for the PanBio kit, for the Platelia™ NS1 kit (Biorad) and for the NS1 Ag Strip kit (BioRad), respectively, based on the analysis of sera of healthy individuals (Group E) and individuals negative for dengue (Group F), Table 1. No cross-reactivity was observed with sera from yellow fever infected patients (Group G); however both Biorad kits showed cross-reactivity with one yellow fever vaccinee (Group H). None of the measles sera (Group I) were recognized by the NS1 tests. One rubella positive case (Group J) showed cross-reactivity with both Platelia™ NS1 kit (Biorad) and for the NS1 Ag Strip kit (BioRad) kit. The overall evaluations according to the different Groups analyzed are shown in Table 1.

NS1 sensitivities according to the different acute diagnosis methods

A higher sensitivity (71.5%, 94.8% and 98.7%) was observed in cases positive by virus isolation only than in cases previously
positive by RT-PCR (62.3%, 82.3% and 82.3%) for the pan-E Early ELISA, Platelia™ NS1 and Dengue NS1 Ag Strip, respectively (Table 2).

**NS1 sensitivities in the presence and absence of IgM**

The detection rate by the pan-E Early ELISA, Platelia™ NS1 and Dengue NS1 Ag Strip in the presence of IgM was 69.4%, 64.5% and 77.4%, respectively (Table 2). In this study, the presence or absence of IgM did not influence detection by the pan-E Early ELISA ($P = 0.6159$). However, a higher detection rate by both Platelia™ NS1 (Biorad) and Dengue NS1 Ag Strip (Biorad) in the absence of IgM was statistically significant ($P < 0.0001$ and $P = 0.0008$, respectively).

**Sensitivities of the NS1 tests by the number of days after the onset of the symptoms**

The sensitivities of all NS1 tests were evaluated according to the number of days of illness. A higher detection rate by the three tests analyzed was during the first four days after the onset of the symptoms (Day 3, in Figure 1, considering day 0 as the first day of fever). The sensitivity of Platelia™ NS1 (Biorad) decreased to 75% of detection after that and maintained the same rate until day 6 of illness. However, after the 4th day, the NS1 Ag Strip (Biorad) showed 89.0% of sensitivity up to the 7th of symptoms. From day five to the 7th, the pan-E Early ELISA (Panbio) confirms about 60.0% of the cases (Figure 1). Although dengue NS1 antigen detections up to the 9th day are observed, here we plotted cases only up to the 7th day due to the low number of samples representing 8th and 9th days in our population.

We also aimed to compare the cases confirmation by the dengue NS1 antigen capture to the confirmation by other methodologies used in this study according to the number of the days of illness. In this comparison, we considered a NS1 positive case, as a case positive in any of the three tests used. Figure 2 shows NS1 confirmation around 90% of the cases up to the 7th day of illness, as previously shown. RT-PCR and virus isolation detections rate

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**Table 1.** Sensitivity and specificity of 3 commercially available kits for NS1 antigen capture in dengue sera and controls.

| Groups* | NS1 antigen capture kit | No. of sera with indicated result/total no. testedd |
|---------|-------------------------|-----------------------------------------------|
|         | pan-E DENGUE EARLY ELISA | Platelia™ Dengue NS1 Ag-ELISA | Dengue NS1 Ag STRIP |
| A (DENV1 cases; \( n = 50 \)) | 13/50 (26.0) | 37/50 (74.0) | 1/50 (2.0) | 49/50 (98.0) | 1/50 (2.0) | 49/50 (98.0) |
| B (DENV2 cases; \( n = 50 \)) | 9/50 (18.0) | 41/50 (82.0) | 5/50 (10.0) | 45/50 (90.0) | 1/50 (2.0) | 49/50 (98.0) |
| C (DENV3 cases; \( n = 58 \)) | 20/58 (34.5) | 38/58 (65.5) | 8/58 (13.8) | 50/58 (86.2) | 7/58 (12.0) | 51/58 (88.0) |
| D (IgM positive cases; \( n = 62 \)) | 19/62 (31.0) | 43/62 (69.0) | 22/62 (35.5) | 40/62 (64.5) | 14/62 (22.6) | 48/62 (77.4) |
| Total of Groups A–D | 61/220 (27.7) | 159/220 (72.3) | 36/220 (16.4) | 184/220 (83.6) | 23/220 (10.5) | 197/220 (89.5) |
| E (healthy individuals; \( n = 30 \)) | 30/30 (100) | 0/30 | 30/30 (100) | 0/30 | 30/30 (100) | 0/30 |
| F (individuals negative for dengue; \( n = 86 \)) | 86/86 (100) | 0/86 | 85/86 (98.8) | 1/86 | 86/86 (100) | 0/86 |
| G (yellow fever positive cases; \( n = 20 \)) | 20/20 (100) | 0/20 | 20/20 (100) | 0/20 | 20/20 (100) | 0/20 |
| H (individuals vaccinated for yellow fever; \( n = 44 \)) | 44/44 (100) | 0/44 | 43/44 (97.7) | 1/44 | 43/44 (97.7) | 1/44 |
| I (measles cases; \( n = 16 \)) | 16/16 (100) | 0/16 | 16/16 (100) | 0/16 | 16/16 (100) | 0/16 |
| J (rubella cases; \( n = 34 \)) | 34/34 (100) | 0/34 | 33/34 (97.0) | 1/34 | 33/34 (97.0) | 1/34 |
| Total of Groups E–J | 230/230 (100) | 0/230 | 227/230 (97.8) | 3/230 | 228/230 (99.1) | 2/230 |

*Subjects in groups A to D had confirmed DENV infection; subjects in groups E to J had no DENV infection.

dValues in parentheses are percentages.

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**Table 2.** Sensitivities of 3 NS1 antigen capture assays in patients with clinical diagnosis of acute dengue confirmed by RT-PCR and/or virus isolation and by MAC-ELISA.

| Dengue case confirmation | NS1 antigen capture kits | No. of sera with indicated result/total no. testedd |
|--------------------------|--------------------------|-----------------------------------------------|
|                          | pan-E DENGUE EARLY ELISA | Platelia™ Dengue NS1 Ag-ELISA | Dengue NS1 Ag STRIP |
| RT-PCR only \( (n = 45) \) | 28/45 (62.3) | 37/45 (82.3) | 37/45 (82.3) |
| Virus isolation only \( (n = 77) \) | 55/77 (71.5) | 73/77 (94.8) | 76/77 (98.7) |
| RT-PCR and virus isolation \( (n = 36) \) | 33/36 (91.7) | 34/36 (94.5) | 36/36 (100) |
| MAC-ELISA only | 43/62 (69.4) | 40/62 (64.5) | 48/62 (77.4) |
| Total | 159/220 (72.3) | 184/220 (83.6) | 197/220 (89.6) |

Values in parentheses are percentages.

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we were around 80% in the first three days of illness, decreasing after that. However, on the other hand IgM detection rates increase only after the 4th day of illness.

NS1 sensitivity in primary and secondary infections

The serologic response could be characterized by IgG-ELISA in 54 samples, where a second specimen was available. There were 40 primary and 14 secondary infections. No differences were observed by the pan-E Early ELISA (Panbio) \((P=0.96)\) and by the NS1 Ag Strip (Biorad) \((P=0.76)\) in confirming primary and secondary infections (Table 3). However, the Platelia™ NS1 test showed a higher sensitivity in confirming primary infections than secondary ones \((P=0.01)\).

NS1 tests efficiency and predicted values

In our study, the pan-E Early ELISA test (Panbio) was less efficient in detecting acute dengue infections (86.1%) when compared to the Platelia™ NS1 test (91.3%) and the NS1 Ag Strip (95.0%). Positive predictive values were 98.3%, 99.5% and 100% for the Platelia™ NS1 (Biorad), NS1 Ag Strip (Biorad) and pan-E Early ELISA tests (Panbio), respectively. However, the pan-E Early ELISA (Panbio) showed the lowest negative predictive value (78.3%), followed by the Platelia™ NS1 test (Biorad) with 86.3% and the NS1 Ag Strip (Biorad) with 91.1% (Table 4).

Discussion

The techniques of dengue serologic diagnosis which have been widely used are based on the detection of IgM antibodies by MAC-ELISA and IgG by IgG-ELISA. However, one of the limitations of these techniques is the inability to detect antibodies to DENV in the acute phase of disease [5,26]. It takes 3 to 5 days for IgM antibodies and anti-DENV 10–14 days for IgG anti-DENV to become detectable. Moreover, primary and secondary infections have different profiles of production of these antibodies [27].

According to previous studies the presence of NS1 in human sera can be confirmed between days 0 to 9 [28,29,30] and with a peak at days 6 to 10 [31]. Currently, commercial kits such as the Dengue EARLY ELISA (Panbio Diagnostics, Brisbane, Australia), Platelia™ Dengue NS1 Ag-ELISA and Dengue NS1 Ag STRIP (BioRad Laboratories Marnes La Coquette, France) are available for early diagnosis of dengue based on NS1 antigen capture and several studies have been conducted in many laboratories [12,15,18,19,29,31,32,33,34,35,36].

In this study, we had the opportunity to evaluate and compare three NS1 antigen capture kits available with a panel of samples \((n=450)\) from cases occurred since the introduction of dengue in Rio de Janeiro, Brazil in 1986 to 2008. The NS1 Ag Strip test (Biorad) was the most sensitive in confirming dengue cases,
followed by Platelia™ NS1 (BioRad). The least sensitive was the pan-E Early ELISA (PanBio) with 72.3% of sensitivity. However, in this study PanBio kit was the most specific (100%) while both kits from BioRad showed 98.7% and 99.1% of specificity, respectively. A recent evaluation in Malaysia showed that the NS1 Ag Strip had 90.4% of sensitivity and 99.5% of specificity [17]. Studies performed in Vietnam [18] and French Guyana [29] showed sensitivities of 82% and 88%, respectively for the Platelia™ NS1 test. However, sensitivities varying from 63.2% to 93.3% have also been reported for this kit [12,15]. Even though different DENV genotypes may circulate in the Americas and Asia, NS1 kits evaluations in countries from those areas show the ability of those tests to detect DENV in infected patients. Our observations are consistent with previous studies in which the pan-E Early ELISA had lower sensitivities [13,14,16,37]. However, to increase diagnostic performance, Panbio has recently released an improved second generation for their NS1 capture kit with changes in key reagents and procedure [38].

All NS1 tests were more sensitive in confirming cases positive by virus isolation than in cases positive by RT-PCR. Dussart [29] confirmed 94.1% of cases positive by virus isolation and 85% of the cases RT-PCR positive using the Platelia™ NS1 test. Recently, McBride [16] showed that the NS1 antigen capture was positive in 87% of the cases positive by RT-PCR. In our study, the Dengue NS1 Ag Strip confirmed 98.7% of the cases positive by virus isolation and 92.3% of RT-PCR positive cases, results similarly observed by Zainah [17]. In the presence of IgM antibodies, the Dengue NS1 Ag Strip confirmed more cases (77.4%) than the pan-E Early ELISA (69.4%) and the Platelia™ NS1 (64.5%). The presence or absence of IgM did not influence in the cases confirmation by the pan-E Early ELISA (P=0.6159). However, the higher confirmation by both Platelia™ NS1 and the Dengue NS1 Ag Strip in the absence of IgM were statistically significant. Sekaran [32] showed that the NS1 detection rates decrease as IgM levels rise, in agreement with our results.

The pan-E Early ELISA (PanBio) showed a higher sensitivity in confirming DENV-2 infections and the Platelia™ NS1 kit (BioRad) in DENV-1 infections. However, the Dengue NS1 Ag Strip kit (BioRad) showed the same sensitivity in confirming DENV-1 and DENV-2 infections. DENV-3 infections were the least confirmed by all three kits. The apparent inability in confirming infection by this serotype has been shown previously [33]. Furthermore, differences in the inter-serotype sensitivities have been reported for all three kits. McBride [16] recently showed lower sensitivities by the pan-E Early ELISA (PanBio) in DENV-2 and DENV-4 infections. The latter was also found in previous studies performed by Bessoff [37] and Dussart [14] and most recently in a study performed in Venezuela [36]. Due to the absence of DENV-4 circulating in Brazil, we were not able to access the assays sensitivities in cases infected by this serotype. Both Biorad kits (Platelia™ NS1 and Dengue NS1 Ag Strip) showed a lower sensitivity in DENV-2 infections from Vietnam [18] and Venezuela [36].

A higher detection rate by the three tests was found during the first four days after the onset of the symptoms. Although dengue NS1 antigen detections up to the 9th day are described, here we analyzed cases only up to the 7th day due to the low number of samples representing 8th and 9th days in our population. The lack of later samples in this study did not allow us to determine when NS1 detection would decrease. However, previous studies found NS1 antigen in 82% to 83% of patients with dengue from day 1 to 9th after the onset of fever [11,30].

The Platelia™ NS1 test showed a higher sensitivity in confirming primary infections than secondary ones, as previously observed [12,15,17,18,32,34]. False negative results by NS1 antigen capture in secondary infections may also be due to the immune-complexes formation by the anti-DENV IgG sequestration [39]. Efforts to dissociate immune complexes by acid treatment can enhance the assays sensitivities, as previously shown [15]. However, in our study no attempts were made to dissociate

### Table 3. Sensitivities of NS1 antigen capture assays in patients with primary and secondary dengue infections (n=54).

| Patients immune response | NS1 antigen capture kits | pan-E DENGUE EARLY ELISA P value | Platelia™ Dengue NS1 Ag-ELISA P value | Dengue NS1 Ag STRIP P value |
|--------------------------|--------------------------|---------------------------------|--------------------------------------|---------------------------|
| Primary infection (n=40)  | 26/40 (65.0)             | 0.96                            | 38/40 (95.0)                        | 0.01                      |
| Secondary infection (n=14)| 9/14 (64.3)              | 10/14 (71.4)                    | 13/14 (93.0)                        |                           |

*Values in parentheses are percentages.

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### Table 4. Sensitivity, specificity, efficiency and predictive values of 3 commercial dengue NS1 capture assays.

| NS1 antigen capture tests                          | pan-E Early ELISA (PanBio) (%) | Platelia NS1 (BioRad) (%) | NS1 Ag Strip (BioRad) (%) |
|---------------------------------------------------|-------------------------------|--------------------------|--------------------------|
| Sensitivity                                       | 72                            | 84                       | 90                       |
| Specificity                                       | 100                           | 99                       | 99                       |
| Efficiency                                        | 86                            | 91                       | 95                       |
| Positive Predicted Value                          | 100                           | 98                       | 99                       |
| Negative Predicted Value                          | 78                            | 86                       | 91                       |

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those complexes. To further analyze the sensitivity of those tests in confirming secondary cases, a larger number of cases should be tested.

Among the kits evaluated, the Dengue NS1 Ag Strip (BioRad) was the most efficient in confirming dengue infections by capturing NS1 antigen from infected patients. Moreover, it was more convenient to be used, as the results can be obtained in 15 minutes, easy to perform and its performance does not involve the use of special laboratory equipment.

Previous studies have demonstrated a diagnostic strategy combining NS1 Ag detection in acute-phase sera and DENV IgM detection in early-convalescent-phase sera, providing a sensitivity of about 90% for dengue diagnosis [29,34].

In conclusion, this evaluation has shown that NS1 antigen capture assays are indeed an alternative tool for the early diagnosis of dengue infections, may be used as a screening test prior virus isolation and used in combination with IgM capture can increase the rate of cases confirmation, especially in endemic areas where secondary infections are expected to occur due to the cocirculation of the different DENV serotypes, such as seen in Brazil. This evaluation was performed for research purposes only and authors have no financial interest. The pan-E Early ELISA from Panbio and the Dengue NS1 Ag Strip from BioRad were kindly provided for evaluation.

Supporting Information

Alternative Language Abstract S1 Translation of the abstract into Portuguese by Flavia Barreto dos Santos

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Author Contributions

Conceived and designed the experiments: RMRN HGS FBdS. Performed the experiments: MdRQL. Analyzed the data: MdRQL. Wrote the paper: MdRQL, FBdS.

References

1. Kyle JL, Harris E (2000) Global spread and persistence of dengue. Annu Rev Microbiol 62: 71–92.
2. WHO (2002) Dengue and Dengue haemorrhagic fever. Fact sheet No. 117, Geneva.
3. Chambers TJ, Hahn G, Gallay P, Rice CM (1990) Flavivirus genome organization, expression and replication. Annu Rev Microbiol 44: 649–669.
4. Huang JL, Huang JH, Shyu RH, Teng GW, Lin YL, et al. (2001) High-level expression of recombinant dengue viral NS-1 protein and its potential use as a diagnostic antigen. J Med Virol 63: 553–569.
5. Schilling S, Lindenbach BD, Van Au L, Schilling H (2004) Laboratory diagnosis of primary and secondary dengue infection. J Clin Virol 31(3): 179–184.
6. Mason PW (1989) Maturation of Japanese encephalitis virus glycoproteins produced by infected mammalian and mosquito cells. Virology 169: 334–364.
7. Falconar AK (1997) The dengue virus nonstructural-1 protein (NS1) generates antibodies to common epitopes on human blood clotting, integrin/adhesin proteins and binds to human endothelial cells: potential implications in haemorrhagic fever pathogenesis. Arch Virol 142(5): 897–916.
8. Westaway EG, Mackenzie JM, Kenney MT, Jones AR, Khasnovych AA (1997) Ultrastructure of Kunjin virus-infected cells: colocalization of NS1 and NS3 with double-stranded RNA, and of NS2B with NS3, in virus-induced membrane structures. J Virol 71: 6650–6661.
9. Flamand M, Megret F, Mathieu M, Lepault J, Rey FA, et al. (1999) Dengue virus type 1 nonsstructural glycoprotein NS1 is secreted from mammalian cells as a soluble hexamer in a glycosylation-dependent fashion. J virology 73(7): 4610–4619.
10. Young PR, Hilditch PA, Fisher R, Conboy M, Jarman RG, et al. (2002) Diagnostic antigen strip for the rapid diagnosis of patients with dengue infection. Diagn Microbiol Infect Dis 46: 39–44.
11. Alcon S, Talarmín A, Debruyne M, Falconar A, Deuchel V, et al. (2002) Enzyme-linked immunosorbent assay specific to Dengue virus type 1 nonstructural protein NS1 reveals circulation of the antigen in the blood during the acute phase of disease in patients experiencing primary or secondary infections. J Clin Microbiol 40: 576–81.
12. Kumarasamy V, Wahab AH, Chua SK, Hassan Z, Chem YK, et al. (2007) Dengue serotyping based on envelope and membrane and nonstructural protein NS1 sero-type-specific capture immunoglobulin M enzyme-linked immunosorbent assay assays. J Clin Microbiol 45(6): 2049–94.
13. Shimizu S, Lai SK, Lee NH, Iwanaga H, Nakayama M, et al. (2008) Evaluation of an IgG enzyme-linked immunosorbent assay for dengue diagnosis. J Clin Virol 41: 138–143.
14. Lanciotti RS, Calisher CH, Gubler DJ, Chang GJ, Vorndam AV (1992) Rapid detection and typing of dengue viruses from clinical samples by using reverse transcription-polymerase chain reaction. J Clin Microbiol 30: 545–551.
15. Nogueira RM, Miagostovich MP, Lampè E, Souza RW, Zague SMO, et al. (1993) Dengue epidemic in the stage of Rio de Janeiro, Brazil, 1990-1: cocirculation of Dengue 1 and dengue 2 serotypes. Epidemiol Infect 111: 163–170.
16. Miandad M, Gunther A, Lautsch S, et al. (2009) Evaluation of dengue nonstructural protein 1 serotyping based on envelope and membrane and nonstructural protein NS1 serotype-specific capture immunoglobulin M enzyme-linked immunosorbent assays. J Clin Microbiol 47(4): 1224–1230.
17. Chaiyaratana W, Chuanumsmit P, Pongthanapisith V, Tangnararatuchat K, Lertwongrath S, et al. (2009) Evaluation of dengue nonstructural protein 1 antigen strip for the rapid diagnosis of patients with dengue infection. Diagn Microbiol Infect Dis 69: 7–9.
18. WHO (1997) Dengue haemorrhagic fever: diagnosis, treatment and control. World Health Organization, Geneva, Switzerland. http://www.who.int/csr/resources/publications/dengue/Denguepublication/en/index.html.
19. Igarka A (1978) Isolation of a Singh’s Aedes aegypti cell clone sensitive to Dengue and Chikungunya viruses. J Gen Virol 40: 531–544.
20. WHO (1997) Dengue haemorrhagic fever: diagnosis, treatment and control. World Health Organization, Geneva, Switzerland. http://www.who.int/csr/resources/publications/dengue/Denguepublication/en/index.html.
21. Igarashi A (1978) Isolation of a Singh’s Aedes aegypti cell clone sensitive to Dengue and Chikungunya viruses. J Gen Virol 40: 531–544.
22. Lanciotti RS, Calisher CH, Gubler DJ, Chang GJ, Vorndam AV (1992) Rapid detection and typing of dengue viruses from clinical samples by using reverse transcription-polymerase chain reaction. J Clin Microbiol 30: 545–551.
23. Nogueira RM, Miagostovich MP, Lampè E, Souza RW, Zague SMO, et al. (1993) Dengue epidemic in the stage of Rio de Janeiro, Brazil, 1990-1: cocirculation of Dengue 1 and dengue 2 serotypes. Epidemiol Infect 111: 163–170.
24. Miandad M, Gunther A, Lautsch S, et al. (2009) Evaluation of dengue nonstructural protein 1 serotyping based on envelope and membrane and nonstructural protein NS1 serotype-specific capture immunoglobulin M enzyme-linked immunosorbent assays. J Clin Microbiol 47(4): 1224–1230.
25. Lanciotti RS, Calisher CH, Gubler DJ, Chang GJ, Vorndam AV (1992) Rapid detection and typing of dengue viruses from clinical samples by using reverse transcription-polymerase chain reaction. J Clin Microbiol 30: 545–551.
26. PAHO (1997) Resurgimiento del dengue en las Américas. Boletín Epidemiológico 10: 1–6.
27. Shu PY, Chen LK, Chang SF, Su CL, Chen LJ, et al. (2004) Dengue virus serotyping based on envelope and membrane and nonstructural protein NS1 serotype-specific capture immunoglobulin M enzyme-linked immunosorbent assays. J Clin Microbiol 42(6): 2409–94.
28. Falconer AK (1997) The dengue virus nonstructural-1 protein (NS1) generates antibodies to common epitopes on human blood clotting, integrin/adhesin proteins and binds to human endothelial cells: potential implications in haemorrhagic fever pathogenesis. J Virol 142(5): 897–916.
29. Dussart P, Laureau B, Lagautu G, Louis P, Nunes MR, et al. (2006) Evaluation of an enzyme immunoassay for detection of dengue virus NS1 antigen in human serum. Clin Vaccine Immunol 13: 1105–1109.
30. Shu PY, Chen LK, Chang SF, Su CL, Chen LJ, et al. (2002) Potential application of nonstructural protein NS1 sero-type-specific immunoglobulin G enzyme-linked immunosorbent assay in the seroepidemiologic study of dengue virus infection: correlation of results with those of the plaque reduction neutralization test. J Clin Microbiol 40(5): 1840–1844.
31. Xu H, Bi B, Pan YY, Qiu LW, Wang YD, et al. (2006) Serotype 1-specific monoclonal antibody-based antigen capture immunoassay for detection of circulating nonstructural protein NS1: Implications for early diagnosis and serotyping of dengue virus infections. J Clin Microbiol 44: 2872–2878.
32. Selenar SD, Ew CL, Kantesh BM, Appana R, Subramaniam G (2007) Evaluation of a dengue NS1 capture ELISA assay for the rapid detection of dengue virus infections. J Infect Developing Countries 1: 182–188.
33. Blacksell SD, Bell D, Kelley J, Mannen Jr MP, Gibbons RV, et al. (2007) Prospective study to determine accuracy of rapid serological assays for diagnosis of acute dengue virus infection in Laos. Clin Vaccine Immunol 14: 1458–64.

34. Chuansumrit A, Chaiyaratana W, Pongthanapisith V, Tangnararatchakit K, Lertwongrath S, et al. (2008) The use of dengue nonstructural protein 1 antigen for the early diagnosis during the febrile stage in patients with dengue infection. Pediatr Infect Dis J 27: 43–48.

35. Phuong HL, Thai KT, Nga TT, Giao PT, Hung le Q, et al. (2009) Detection of dengue nonstructural 1 (NS1) protein in Vietnamese patients with fever. Diagn Microbiol Infect Dis 63: 372–8.

36. Ramirez AH, Moros Z, Comach G, Zambrano J, Bravo L, et al. (2009) Evaluation of dengue NS1 antigen detection tests with acute sera from patients infected with dengue virus in Venezuela. Diagn Microbiol Infect Dis.

37. Bessoff K, Delorey M, Sun W, Hunsperger E (2008) Comparison of two commercially available dengue virus (DENV) NS1 capture enzyme-linked immunosorbent assays using a single clinical sample for diagnosis of acute DENV infection. Clin Vaccine Immunol 15: 1513–8.

38. Panbio. Available http://www.panbiocience.com/story/improved-panbio-dengue-early-elsa-available-now. Accessed 2010 Apr 13.

39. Koraka P, Burghoorn-Maas CP, Falconar A, Setiati TE, Djamiatun K, et al. (2003) Detection of immune-complex-dissociated nonstructural-1 antigen in patients with acute dengue virus infections. J Clin Microbiol 41(9): 4154–9.