



Re-emergence of dengue virus serotype 3 infections in Gabon in 2016–2017, and evidence for the risk of repeated dengue virus infections



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ABSTRACT

Objectives: Dengue outbreaks, mainly caused by dengue virus serotype 2 (DENV-2), occurred in 2007 and in 2010 in Gabon, Central Africa. However, information on DENV infections has been insufficient since 2010. The aim of this study was to investigate the current DENV infection scenario and the risk of repeated infections in Gabon.

Methods: During 2015–2017, serum samples were collected from enrolled febrile participants and were tested for DENV infection using RT-qPCR. DENV-positive samples were analyzed for a history of previous DENV infection(s) using ELISA. The complete DENV genome was sequenced to analyze the phylogeny of Gabonese DENV strains.

Results: DENV-3 was exclusively detected, with a high rate of anti-DENV IgG seropositivity among DENV-3-positive participants. DENV-3 showed higher infection rates in adults and the infection was seasonal with peaks in the rainy seasons. Phylogenetic analysis revealed that Gabonese DENV-3 originated from West African strains and has been circulating continuously in Gabon since at least 2010, when the first DENV-3 case was reported.

Conclusions: These findings indicate stable DENV-3 circulation and the risk of repeated DENV infections in Gabon, highlighting the need for continuous monitoring to control DENV infections.

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Introduction

With a significant increase in incidence in the past decades, dengue has become one of the most prevalent infectious diseases and a major health concern in tropical and subtropical regions of the

world, including more than 100 countries in the Southeast Asia, Africa, North and South America, and Western Pacific regions (Brady et al., 2012). Dengue virus (DENV) is estimated to cause 390 million infections annually, and 3.6 billion people are at risk of DENV infection (Bhatt et al., 2013). The incidence of dengue increased seven-fold between 1990 and 2013, with 8.9 billion dollars of global economic burden for dengue in 2013 (Shepard et al., 2016).

DENV is mainly transmitted by the mosquito vectors *Aedes aegypti* and *Aedes albopictus* and has four serotypes: DENV serotypes 1–4 (DENV-1 to DENV-4) (Simmons et al., 2012). In Africa, DENV infections have been reported in at least 34 countries,

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where DENV-1 and DENV-2 are the main causes of dengue; however, it is quite likely that DENV infection cases have been underreported due to a prevailing clinical picture of undifferentiated fever and the unavailability of appropriate diagnostic tests in some settings (Amarasinghe et al., 2011). Currently, there is a very limited number of reports regarding DENV-3 and DENV-4 infections in Africa; whole-genome information is available for only two African DENV-3 strains (Mozambique, 1985, GenBank accession number FJ882575; Senegal, 2009, GenBank accession number KU509282) and is still unavailable for African DENV-4.

In Gabon, a Central African country, two dengue outbreaks were reported in 2007 and 2010 (Caron et al., 2012). DENV-2 was mainly detected on both occasions and this serotype caused nearly 95% of DENV infections. DENV-1 caused a small number of infections in the 2010 outbreak. In contrast, DENV-3 has only been detected in one individual in Gabon so far (Caron et al., 2013). Since the 2010 outbreak, there has been little information on DENV infections in the country, with no evidence of DENV circulation across rural Gabon, as reported in 2011 (Pourrut et al., 2011). However, in the area around Lambaréné, an anti-DENV immunoglobulin G (IgG) seropositivity rate of 12.3% was found in an analysis of the sera from 30-month-old children that was collected before the 2007 and 2010 outbreaks; this indicated a pre-outbreak circulation of DENV (Gabor et al., 2016). These data on DENV prevalence seem to be contradictory, requiring further surveillance studies to clarify the current scenario of both DENV circulation in Gabon and the anti-DENV IgG seropositivity rate, which depicts repeated DENV infection in febrile patients.

As part of a project towards establishing a surveillance system for DENV infections in Lambaréné, Gabon, we conducted serum sample collection from febrile patients to investigate current DENV infections and the genetic characteristics of DENV strains circulating in Gabon. This showed evidence of endemic DENV-3 circulation that has been maintained since the first DENV-3 detection in 2010 in Gabon. A high rate of anti-DENV IgG seropositivity among DENV-3-positive patients was also found, indicating that the residents around Lambaréné are facing the risk of repeated DENV infections.

Materials and methods

Study population

In this study, the samples were collected at two medical facilities, the Albert Schweitzer Hospital in Lambaréné and the Centre de Recherches Médicales de Lambaréné (CERMEL). The Albert Schweitzer Hospital is one of the main general hospitals in the study area, and CERMEL is the medical institute that has a clinical department. Febrile patients (body temperature $\geq 37.5^\circ\text{C}$) who visited the two medical facilities between January 2015 and June 2017 were recruited for enrolment in a surveillance study on DENV infections. In this study, the age of the participants was restricted to >1 year. Demographic information (age and sex) of the recruited participants was collected.

Ethics statement

This study was approved by the institutional review boards of CERMEL and Nagasaki University (approval numbers CEI-007 and 170921177, respectively). Written informed consent was obtained from all of the participants or their parents.

Viral RNA extraction and detection by reverse transcription-quantitative polymerase chain reaction (RT-qPCR)

Viral RNA was extracted from 140 μl of each serum sample with a QIAamp Viral RNA Mini Kit (Qiagen, Hilden, Germany) according

to the manufacturer's instructions. RT-qPCR was performed in a 20- μl reaction using a QuantiTect Probe RT-PCR Kit (Qiagen). Each reaction mixture contained 10 μl 2 \times QuantiTect Probe RT-PCR Master Mix, 0.2 μl QuantiTect RT Mix, 0.5 μM of each primer, 0.2 μM TaqMan probe, 2 μl RNA template, and RNase-free water up to 20 μl . RT-qPCR assays were conducted using a LightCycler 480 instrument (Roche, Basel, Switzerland) under the following conditions: 30 min at 50°C ; 15 min at 95°C ; and 45 cycles of 15 s at 95°C , and 60 s at 60°C . The primers and the probe were designed using the sequences reported previously (Santiago et al., 2013). Data from the RT-qPCR assays were analyzed using software included in the LightCycler 480 system. RT-qPCR assays were performed in duplicate and the samples showing cycle threshold (Ct) values under 40 were set as positive.

Enzyme-linked immunosorbent assay (ELISA) of anti-DENV IgG antibodies

Anti-DENV IgG antibodies were detected using the Dengue Virus IgG DxSelect Kit (Focus Diagnostics, Cypress, CA, USA) and the signal intensities were normalized by the calibrator value according to the manufacturer's protocol. The normalized signals that displayed >1.0 were defined as positive. A correlation between the Ct values of RT-qPCR and the normalized signal intensities of ELISA was calculated using GraphPad Prism 7 software (GraphPad Software, San Diego, CA, USA).

Sequencing of the envelope gene for genotyping of detected DENV strains

Amplification of the full-length envelope gene (1479 bp) of DENV-3 strains was performed with the PrimeScript II High Fidelity One Step RT-PCR Kit (Takara Bio, Shiga, Japan) using primers reported previously (Shihada et al., 2017). After agarose gel purification was performed with a QIAquick Gel Extraction Kit (Qiagen), the PCR products were processed using the BigDye Terminator v3.1 Cycle Sequencing Kit (Thermo Fisher Scientific, Waltham, MA, USA) and analyzed using an ABI3500 capillary sequencer (Thermo Fisher Scientific) to obtain sequence data. The genotypes of DENV strains were determined through BLAST analysis of the sequence data (<https://blast.ncbi.nlm.nih.gov/Blast.cgi>).

Whole-genome sequencing

Libraries were prepared from extracted viral RNA samples using a TruSeq RNA Library Preparation Kit v2 (Illumina, San Diego, CA, USA) according to the manufacturer's instructions. After quality and quantity check of each library using an Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA) with a High Sensitivity DNA Kit (Agilent), sequencing was performed using a 300-cycle High Output Kit (Illumina) on a MiniSeq sequencer (Illumina). Mapping of the paired-end reads was performed on CLC Genomics Workbench 11.0.1 software (Qiagen) using the whole-genome sequence of the Senegal 2009 strain as a template. Consensus sequences were extracted and aligned with reference strains on BioEdit 7.0.5.3 software (<http://www.mbio.ncsu.edu/BioEdit/bioedit.html>).

Phylogenetic analysis

To infer the phylogeny of full-length envelope gene sequences of DENV-3 strains, a Bayesian analysis was performed with time-stamped reference sequences that include all complete envelope gene sequences of African DENV-3 strains using BEAST v1.8.0 software (<http://beast.community/>) under the conditions of the

Table 1
Demographic and laboratory data of patients infected with DENV-3.

Sample ID	Age (years)	Sex	Ct value ^a	IgG signal ^b	Collection Date
01	19	M	24.19	+++	9 May 2016
02	33	F	20.24	+	17 May 2016
03	28	M	18.55	+++	2 Jun 2016
04	46	F	30.33	+++	17 Jun 2016
05	18	F	34.84	+++	21 Jun 2016
06	43	F	34.55	+++	05 Jul 2016
07	36	F	23.95	+++	14 Jul 2016
08	4	M	33.91	++	15 Jul 2016
09	29	F	18.32	–	22 Jul 2016
10	6	F	30.08	+	08 Aug 2016
11	26	M	34.54	+++	22 Aug 2016
12	26	F	35.11	–	28 Sep 2016
13	21	M	36.79	+++	08 Oct 2016
14	44	M	36.81	+++	10 Nov 2016
15	29	M	33.99	+++	05 Dec 2016
16	24	F	33.00	+++	10 May 2017
17	57	M	27.78	+++	15 May 2017

DENV-3, dengue virus serotype 3; M, male; F, female.

^a Cycle threshold (Ct) value of virus detection by RT-qPCR.^b Signal intensities of ELISA tests for anti-DENV IgG are presented as normalized signal intensities: –, signal <1.0 (negative); +, signal 1.0–3.0; ++, signal 3.1–5.0; +++, signal >5.0.

SRD08 model, the relaxed lognormal clock, and 80 million generations. For countries outside Africa, reference strains were widely selected from each continent and from various time points of the collection date. The effective sample size (ESS) values were verified to show more than 1000. For complete genome sequences of DENV-3 strains, all whole-genome information of reference strains was obtained from each continent and from various time points of the collection date, as described above. Consensus sequences of DENV-3 strains detected in this study and reference sequences were aligned and analyzed using a maximum-likelihood method with a general time reversible model and gamma distribution of substitution rate. A total of 1000 bootstrap replicates were generated in this analysis. Phylogenetic trees were visualized and modified using FigTree v1.4.2 software (<http://tree.bio.ed.ac.uk/software/figtree>).

Statistical analysis

The statistical data analysis was performed using GraphPad Prism 7 software (GraphPad Software, San Diego, CA, USA). The Chi-square test or Fisher's exact test was used to determine the significance of differences in the comparisons of general

categorical variables. Results were considered to be statistically significant when the *p*-value was less than 0.05.

Sequence data availability

The sequences of the DENV-3 strains obtained in this study have been deposited in GenBank with accession numbers of **LC379193–LC379202**.

Results

Study population and DENV detection

During 2015–2017, 1007 serum samples were collected from febrile patients (body temperature >37.5 °C) who were enrolled in this study. The age range was 1–82 years and the mean age was 10.83 ± 14.4 years (± standard deviation). The samples were obtained from 186 adult patients (≥18 years of age) and 805 pediatric patients (<18 years of age); no age information was available for 16 samples. Samples were screened for DENV (serotypes 1–4) by RT-qPCR, as described previously (Santiago et al., 2013). Overall, 17 samples were found to be positive for DENV-3 (Table 1), including 15 samples from adult patients (positivity rate in adults, 8.06%) and two from pediatric patients (positivity rate in children, 0.25%) (Table 1). Comparison of the mean age for all samples showed significant differences between the DENV-3-positive and negative populations (Table 2). However, there were no statistically significant differences in DENV-3 positivity rates in each of the adult or pediatric population, and between the female and male populations (Table 3).

As serum samples were collected continuously from January 2015 to June 2017 (Figure 1A), the seasonality of DENV-3 infections was analyzed. Considering that most of the DENV-3-positive samples were derived from adult patients, the seasonal change in the DENV-3 positivity rate was analyzed in the adult population. The result showed peaks of DENV-3 infections in May and September–October, during the rainy season (Figure 1B).

Detection of anti-DENV IgG antibodies in DENV-3-positive patients

To screen DENV-positive patients for potential previous DENV infection(s), an ELISA test was performed against anti-DENV IgG antibodies. The ELISA results showed that most of the DENV-positive patients identified by RT-qPCR were also positive for anti-DENV IgG, with a positivity rate of 88.24% (Table 1). To investigate the suppression of viral replication by IgG production in the febrile phase of patients who had experienced a previous DENV infection (s), correlation was analyzed between Ct values of RT-qPCR and the

Table 2
Comparison of age between the patients who were positive for DENV-3 and those who were negative for DENV-3.

	Sex	Negative		Positive		<i>p</i> -Value
		Age (years) ^a	(95% CI)	Age (years)	(95% CI)	
All patients	Mix	10.38	(9.50–11.26)	28.76	(21.68–35.85)	<0.001
	F	10.80	(9.48–12.13)	29.00	(19.45–38.55)	<0.001
	M	9.95	(8.77–11.14)	28.50	(15.07–41.93)	<0.001
Adults (≥18 years)	Mix	36.42	(34.08–38.75)	31.93	(25.73–38.14)	0.27
	F	36.91	(33.57–40.25)	31.88	(23.89–39.86)	0.38
	M	36.02	(32.68–39.37)	32.00	(19.36–44.64)	0.50
Pediatrics (<18 years)	Mix	4.84	(4.59–5.09)	5.00	(–7.71–17.71)	0.95
	F	4.77	(4.42–5.12)	6.00	N/A ^b	N/A
	M	4.90	(4.54–5.26)	4.00	N/A	N/A

DENV-3, dengue virus serotype 3; F, female; M, male; 95% CI, 95% confidence interval; N/A, not analyzed.

^a Age is presented as mean years.^b DENV-3 positive numbers in pediatric patients were too small to analyze statistically.

Table 3

Comparison of sex between the patients who were positive for DENV-3 and those who were negative for DENV-3.

	Sex	Negative ^a	Positive	OR	p-Value
All patients	F	475	9	1.19	0.72
	M	502	8	–	–
Adults (≥18 years)	F	84	8	1.07	0.90
	M	90	7	–	–
Pediatrics (<18 years)	F	385	1	1.09	>0.99
	M	418	1	–	–

DENV-3, dengue virus serotype 3; OR, odds ratio; F, female; M, male.

^a Thirteen DENV-3 negative patients were excluded from the analysis due to the lack of sex information.

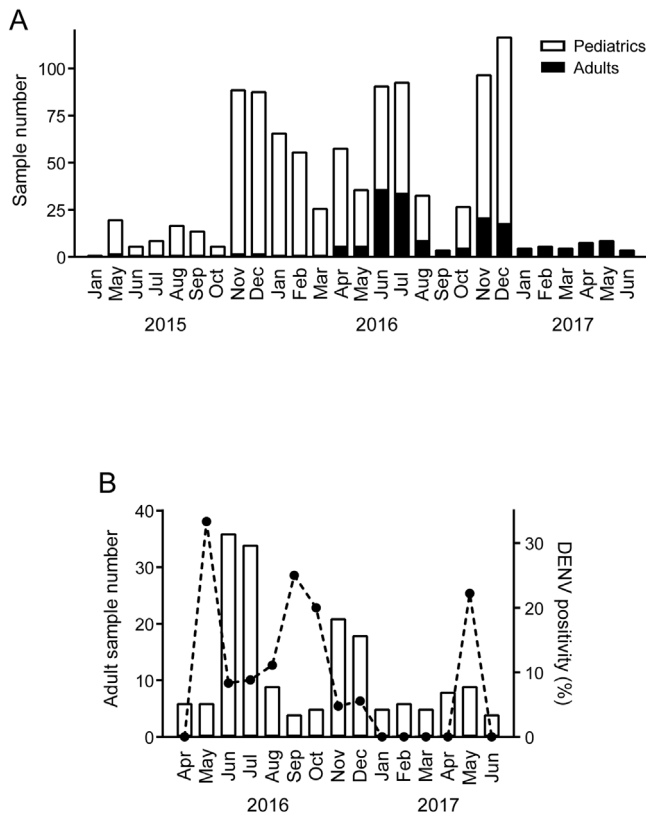


Figure 1. Seasonal data of sample collection and DENV positivity. (A) Sample numbers collected from febrile patients during the study. White and black columns show pediatric and adult sample numbers, respectively. (B) DENV positivity values in adults from April 2016 to June 2017. Columns and the dotted line depict the adult sample number and DENV positivity, respectively.

signal intensities of ELISA tests. Although there were two ELISA-negative samples, Ct values were significantly correlated with ELISA signal intensities, indicative of effective blockade of viral replication by rapid IgG production (Figure 2).

Phylogenetic analysis of detected DENV-3 strains

To investigate the genetic characteristics of DENV-3 strains detected in this study, sequences of the full-length envelope glycoprotein gene were obtained from 10 samples (eight samples from adults and two from pediatric participants, with GenBank accession numbers [LC379193–LC379202](#)), using a method described previously (Shihada et al., 2017). Phylogenetic analysis

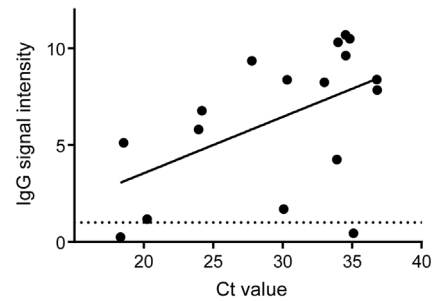


Figure 2. Correlation between the cycle threshold (Ct) values of RT-qPCR and signal intensities of anti-DENV IgG as analyzed by ELISA. Pearson's correlation coefficient and significance were calculated using GraphPad Prism 7 software ($n = 17$, $p < 0.05$, $r = 0.499$). The dotted line depicts the threshold to determine ELISA positivity (signal intensity = 1.0).

inferred that the detected DENV-3 strains belonged to the African cluster of genotype III (Figure 3). The Gabonese DENV-3 strain detected in 2010 (GenBank accession number JX080299), the only DENV-3 strain that has been reported so far in Gabon, was genetically the closest to the strains detected in this study, with a sequence homology that ranged between 99.47% and 99.79%. In spite of the existence of some other African strains detected after 2010 (e.g. Djibouti 2012, KT187294; West Africa–Germany 2013, KJ922394), these post-2010 African strains were all located outside of the Gabonese cluster, indicating that DENV-3 was stably maintained in Gabon (Figure 3).

It was attempted to sequence the complete genome of the DENV-3 strains detected in this study by next-generation sequencing. Complete genome sequences of DENV-3 strains were obtained from five samples collected from adult patients (GenBank accession numbers [LC379193–LC379197](#)). Phylogenetic analysis was performed with whole-genome sequences including two strains that constitute all African DENV-3 complete sequences available (Mozambique, 1985 and Senegal, 2009). The result indicated that the Senegal 2009 strain is genetically close to the DENV-3 strains sequenced in this study (Figure 4). Interestingly, a few Chinese strains detected in 2009 were also inferred as the genetically closest strains, with no relationships to other Chinese strains detected in different years. The amino acid sequence homology between the DENV-3 strains detected in this study and the Senegal 2009 strain was 99.00–99.16%. The amino acid substitutions tended to be located in non-structural proteins, especially in the NS5 protein (Table 4). These substitutions, however, have not yet been reported to correlate with the replication and pathogenicity of DENV.

Discussion

There were two dengue outbreaks in Gabon in 2007 and 2010, which were caused by DENV-2 in the vast majority of cases and DENV-1 in a small number of patients (Caron et al., 2013). In this study, DENV-3 was exclusively detected in febrile patients during the years 2016 and 2017, possibly indicative of serotype shift on mainly circulating DENV in Gabon, as shown previously in a report of longitudinal surveillance (Nisalak et al., 2003). As recognized generally (WHO, 2019), seasonal changes in DENV-3 positivity rate with peaks in the rainy seasons were also observed, probably reflecting an increase in mosquito vector activity.

A meta-analysis study suggested the possibility of DENV-3 infections being significantly related to severe cases (Soo et al., 2016), raising the alert for the careful continuous monitoring of the dissemination of DENV-3 infections. So far, limited cases of

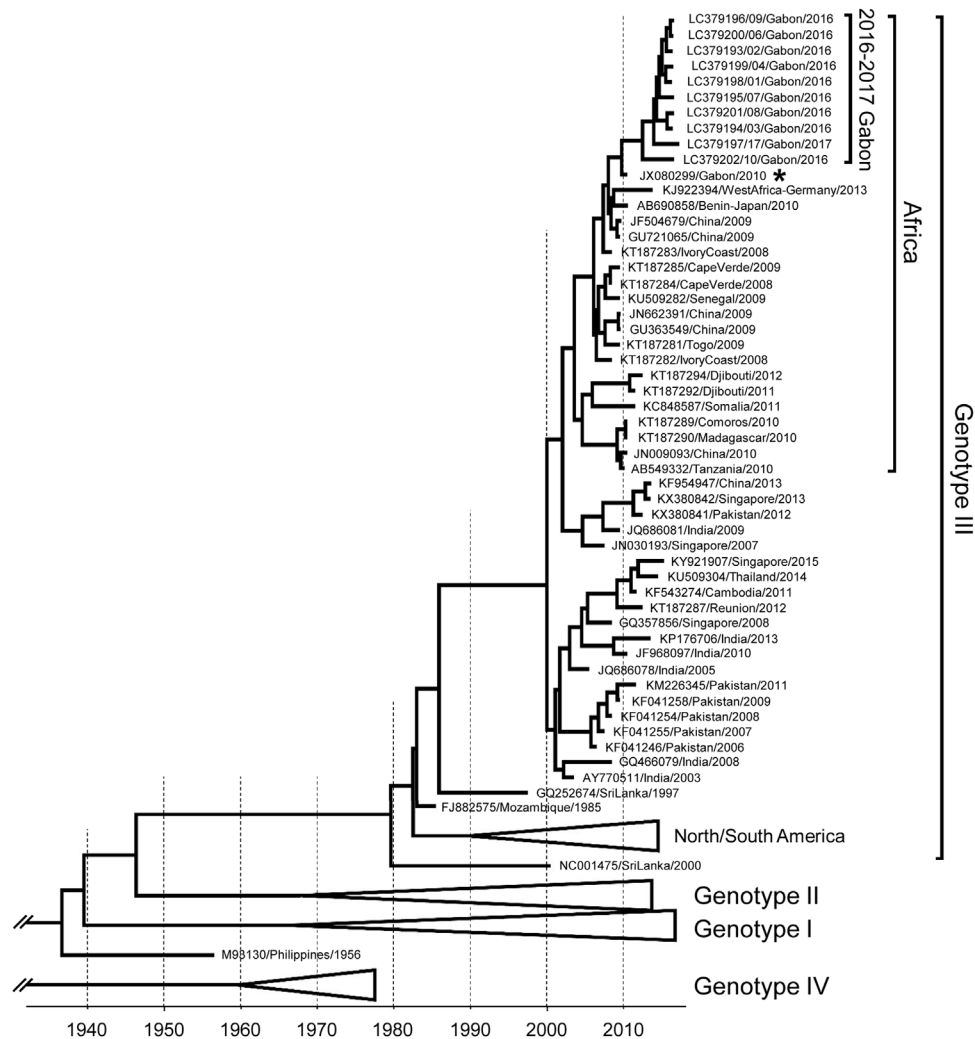


Figure 3. Phylogenetic analysis of the whole envelope glycoprotein gene sequence of DENV-3. A time-scaled Bayesian maximum clade credibility tree was inferred using all African strains with full-length envelope sequences available. For better visualization of sequence positions, several clusters were collapsed and shown as triangles. Virus lineages are shown on the right. The asterisk indicates the Gabonese strain detected in 2010. An entire picture of the phylogeny, including 95% CI bars at each node, is provided in the Supplementary material (Figure S1).

DENV-3 infection have been reported in Central Africa: Cameroon in 2006 (Domingo et al., 2011) and Gabon in 2010 (Caron et al., 2013). Both of the two Central African cases showed DENV-3 detection from only one individual; however, it is quite likely that DENV-3 infection cases have been underreported due to the unavailability of appropriate diagnostic tests in Africa (Amarasinghe et al., 2011). The latest report in Central Africa showed that 7.02% (8/114) of febrile patients in Douala, Cameroon were infected with DENV in 2017 (Yousseu et al., 2018), suggesting a similar infection rate of DENV in adults in Gabon (8.06%, 15/186 in this study) and also, presumably, in neighboring Central African countries.

Unlike Southeast Asian countries, pediatric samples showed lower detection rates than adults in the present study, requiring future virological characterization of Gabonese DENV-3 for the possibility of preferential infection in adults and the identification of the mosquito vectors around the study area. The surveillance of symptomatic patients can provide information on circulating DENV strains, but may possibly miss approximately 65–67% of DENV-infected asymptomatic patients (Fritzell et al., 2018; Salje et al., 2018). A combination of molecular genetic data with epidemiological seroprevalence

would be informative for understanding the whole context of DENV infections.

In Africa, DENV-3 has mainly been reported in West and East Africa, and in islands of the southeastern region (Virus Pathogen Resource, <https://www.viprbrc.org/>). African DENV-3 was first detected in Mozambique in 1985 (Gubler et al., 1986) and has been detected independently in East Africa since 1993 (e.g. Djibouti, Somalia, Tanzania, Madagascar) and in West Africa since 2006 (e.g. Senegal, Côte d'Ivoire, Togo, Benin). Phylogenetic studies showed that these occurrences of African DENV-3 infections were caused by viruses that originated from the Indian sub-continent (Messer et al., 2003). The time-scaled phylogenetic analysis in the present study revealed that the West African strains were likely transmitted into Gabon via Côte d'Ivoire, Togo, and Benin around 2008, before the first detection of DENV-3 in Gabon in 2010 (Figure 5). The phylogenetic tree also clarified that Gabonese DENV-3 strains were derived from a lineage distinct from the one that caused outbreaks in Cape Verde and Senegal in 2009.

In recent years, dengue caused by DENV-3 infection has mainly been reported in Sub-Saharan Africa in sentinel cases involving febrile returning travelers (Shihada et al., 2017). Interestingly, BLAST analysis of the complete DENV-3 genome sequenced in this study

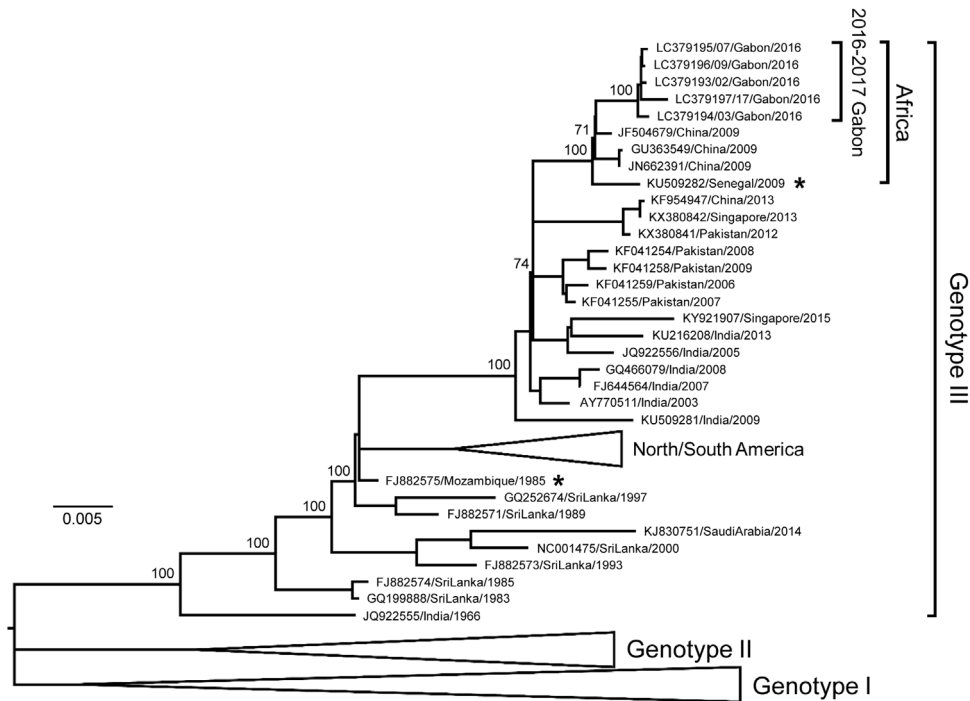


Figure 4. Phylogenetic analysis of the complete genome sequence of DENV-3. A maximum-likelihood tree was inferred using all African strains available. Bootstrap values of $\geq 70\%$ are shown at nodes of the tree. For better visualization of sequence positions, several clusters were collapsed and shown as triangles. Virus lineages are shown on the right. The asterisk indicates the African strain. The scale bar indicates nucleotide substitutions per site. An entire picture of the phylogeny is provided in the Supplementary material (Figure S2).

Table 4
Amino acid substitutions in the whole polyprotein sequences of the Senegal 2009 strain and the strains sequenced in this study.

Genes	Amino acid position ^a	Reference	Gabon ^b					
			Senegal 2009	#02 2016	#03 2016	#07 2016	#09 2016	#17 2017
C	93	S	*	*	*	*	*	G
PrM	171	T	*	*	A	A	*	*
PrM	254	L	*	*	*	*	*	I
E	508	T	I	I	I	I	I	I
E	546	T	*	*	A	*	*	*
E	724	A	*	*	V	*	*	*
NS1	813	R	K	K	K	K	K	K
NS1	955	R	K	*	*	*	*	*
NS1	1101	D	G	G	G	G	G	G
NS2A	1144	L	F	F	F	F	F	F
NS3	1682	V	I	I	I	I	I	I
NS3	1731	V	I	I	I	I	I	I
NS3	1943	N	*	*	*	*	*	H
NS4A	2155	L	*	*	*	*	*	S
NS4B	2353	A	V	V	V	V	V	V
NS5	2520	K	R	R	R	R	R	R
NS5	2764	D	*	*	*	*	*	N
NS5	2784	V	D	D	D	D	D	D
NS5	2791	K	T	T	T	T	T	T
NS5	2796	R	G	G	G	G	G	G
NS5	2913	A	*	*	*	*	*	V
NS5	2978	P	L	L	L	L	L	L
NS5	3166	D	N	N	N	N	N	N
NS5	3240	E	*	D	*	*	*	*
NS5	3354	S	P	P	P	P	P	P

C, capsid; M, membrane; E, envelope.

^a Number depicts the amino acid position in the polyprotein of the Senegal 2009 strain.

^b Asterisk shows identical amino acid to the Senegal 2009 strain.

identified Chinese DENV-3 strains isolated in 2009 in Zhejiang and Guangzhou as highly homologous strains (GenBank accession numbers JF504679, GU363549, and JN662391). However, these African-like Chinese strains were supposedly not maintained in China, because other recent Chinese strains isolated after 2012 were composed of Asian circulating strains in genotype II (Figures 3 and 4). These results may be indicative of international transmission and a sporadic spread of infections due to returning Chinese travelers or workers importing African DENV-3 strains. Similar to China, around 2009, a variety of European countries and Japan reported imported cases of DENV-3 strains belonging to genotype III from West Africa, possibly reflecting a large epidemic of DENV-3 in West Africa at that time (e.g. *Moi et al., 2010; Ninove et al., 2009*).

Epidemiological studies have suggested that severe symptoms might occur during secondary DENV infections with a different serotype (*Guzman and Harris, 2015*). Recently, two clinical studies demonstrated that a specific range of pre-existing DENV antibody titers were associated with the highest risk of severe symptoms, supporting the theory of antibody-dependent enhancement (ADE) of DENV infections (*Katzelnick et al., 2017; Salje et al., 2018*). The present study revealed a high anti-DENV IgG detection rate in DENV-3-positive patients with the risk of repeated DENV infections in Gabon, which is related to a report of anti-DENV seropositivity before the 2007 outbreak in Gabon (*Gabor et al., 2016*). Although a higher IgG titer correlated with a lower viral titer in this study, which might show an effective suppression of viral replication (Figure 2), the ELISA results showed that physicians need to pay attention to the occurrence of severe dengue in the country.

In conclusion, the present DENV surveillance conducted in febrile populations revealed the latest situation of DENV circulation in

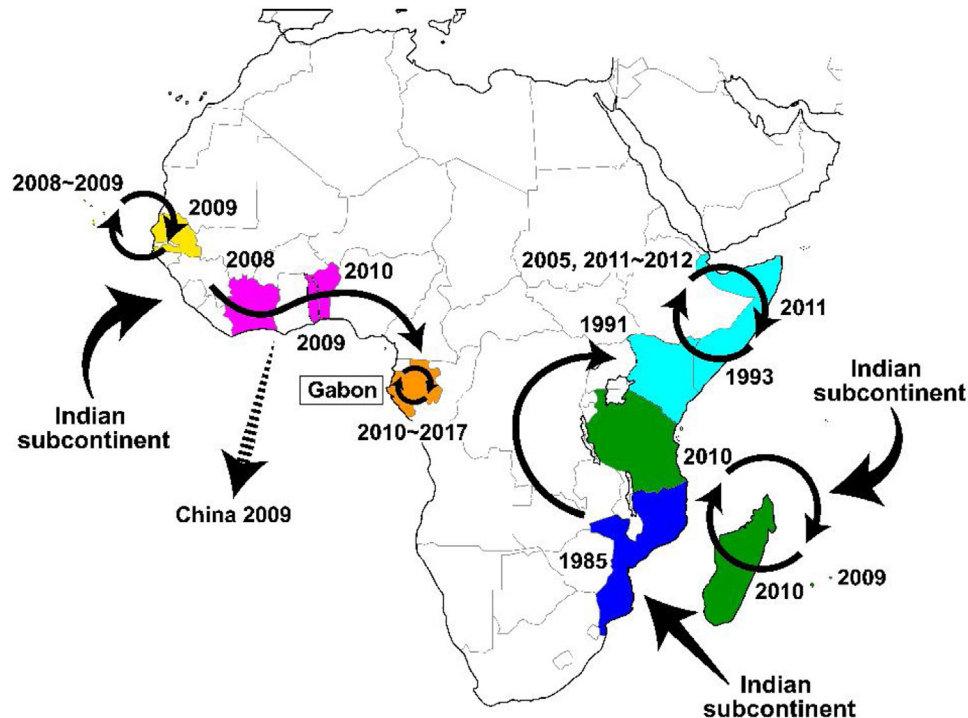


Figure 5. Geographical patterns of DENV-3 transmission in Africa. The transmission pattern is based on the Bayesian tree shown in Figure 3, studies reporting phylogeny of African DENV-3 strains (Messer et al., 2003), and information of outbreaks (Brady et al., 2012; Franco et al., 2010). Each color depicts the lineage of DENV-3 strains that belongs to a single cluster. Years show the time point of DENV-3 detection in the country.

Gabon, with a tendency to infection in adults rather than the pediatric population, and with peaks during the rainy seasons. The dissemination pattern of DENV-3 from West Africa to Gabon was also clarified by the phylogenetic tree. These findings will be informative in suggesting countermeasures for the spread of DENV in Gabon. However, it will be necessary to conduct continuous DENV surveillance to understand the current situation in order to help in the immediate alert and control of DENV infections.

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Ethical approval

This study was approved by the Institutional Review Boards of CERMEL and Nagasaki University (approval numbers CEI-007 and 170921177, respectively). Written informed consent was obtained from all participants or their parents.

Conflict of interest

All authors have no conflicts of interest to declare.

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Appendix B. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.ijid.2019.12.002>.

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