Zika might not be acting alone: Using an ecological study approach to investigate potential co-acting risk factors for an unusual pattern of microcephaly in Brazil

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Abstract

Zika virus infections can cause a range of neurologic disorders including congenital microcephaly. However, while Zika infections have been notified across all regions in Brazil, there has been an unusual number of congenital microcephaly case notifications concentrated in the Northeast of the country. To address this observation, we investigated epidemiological data (2014–2016) on arbovirus co-distribution, environmental and socio-economic factors for each region in Brazil. Data on arbovirus reported cases and microcephaly were collected from several Brazilian Ministry of Health databases for each Federal unit. These were complemented by environmental management, social economic and Dengue aegypti infestation index data, extracted from multiple databases. Spatial time “ecological” analysis on the number of arboviruses transmitted by Aedes mosquitoes in Brazil show that the distribution of dengue and Zika was widespread in the whole country, with higher incidence in the West-Central region. However, reported chikungunya cases were higher in the Northeast, the region also with the highest number of microcephaly cases registered. Social economic factors (human development index and poverty index) and environmental management (water supply/storage and solid waste management) pointed the Northeast as the less wealthy region. The Northeast is also the region with the highest risk of Aedes aegypti house infestation due to the man-made larval habitats. In summary, the results of our ecological analysis support the hypothesis that the unusual distribution of microcephaly might not be due to Zika infection alone and could be accentuated by poverty and previous or co-infection with other pathogens. Our study reinforces the link between poverty and the risk of disease and the need to understand the effect on pathogenesis of sequential exposure to arboviruses and co-viral infections. Comprehensive large-scale cohort studies are required to corroborate our findings. We recommend that the list of infectious diseases screened, particularly during pregnancy, be regularly updated to include and effectively differentiate all viruses from ongoing outbreaks.
Introduction

Zika, dengue and chikungunya are arboviral diseases transmitted by mosquitoes from the genus *Aedes*. The co-circulation of these viruses in Brazil, sharing the same vectors and causing similar symptoms, represents a major public health challenge. Dengue fever is the most important re-emerging mosquito-borne viral disease worldwide. In Brazil, the incidence of dengue has been frequently high, with 1.49 million cases reported in 2016 [1]. Chikungunya viral infections were first reported in Bahia, Brazil in 2014, in a patient travelling from Angola [2]. Since then, more than 250,000 cases of chikungunya infection have been registered [3]. The first Zika cases reported in Brazil were identified in the Northeast region, in early 2015, in patients presenting symptoms of mild fever, rash, conjunctivitis and arthralgia [4]. Since the first reported autochthonous transmission, more than 200,000 cases have been registered in Brazil [5], mainly during 2015–2016.

The outbreak of Zika infection in Brazil has exposed the high risk this arbovirus imposes during pregnancy to the fetus. Brazil experienced an approximately 20-fold increase in the total number of congenital microcephaly cases from 2014 to 2015, following the confirmation of autochthonous Zika virus transmission [6]. In December 2015, the Brazilian Ministry of Health enhanced congenital microcephaly surveillance by implementing a more sensitive case definition, and added Zika to the list of “TORCHS” pathogens (Toxoplasmosis, Rubella, Cytomegalovirus, Herpes simplex virus, Syphilis) that are screened [7,8]. In February 2016, WHO declared the link between Zika virus and microcephaly to be a Public Health Emergency of International concern. Subsequently, several cases of microcephaly associated with mothers infected with Zika during pregnancy were reported in different Federal Units of Brazil, most of them in the Northeast region [9]. Several studies have shown that the arbovirus can be transmitted vertically during pregnancy and cause congenital problems of the fetus. A recent case-control study performed in Recife, Pernambuco State, revealed that babies with microcephaly were 55 times more likely to have been infected with Zika virus during pregnancy than non-infected [10]. The virus has been detected in the placenta, amniotic fluid and neural tissues of newborns with microcephaly [11–13]. In addition to microcephaly, other birth defects have been reported such as intracranial calcifications and abnormalities of the corpus callosum and the cerebellum, mainly when the exposure to the virus occurs during the first trimester of pregnancy [14].

Before the Brazilian outbreak, no microcephaly-Zika related cases have previously been reported. Zika virus was firstly isolated from a Rhesus monkey in the Zika Forest of Uganda, in 1947 [15]. The first large outbreak of the Zika virus in humans was registered only in 2007 on the Pacific island of Yap, in Micronesia [16]. The same lineage, an Asian type, caused epidemics in the Pacific Islands in 2013–2014 [17–18]. Phylogenetic analyses revealed the Brazilian Zika virus to be closer to those circulating in French Polynesia in 2013, therefore, it might be possible that the Asian-type strains caused unreported microcephaly in the past outbreaks [4,19]. Indeed, a recent retrospective study in French Polynesia has shown that Zika virus infection during the first trimester of pregnancy led to a 1% increase in the risk of congenital microcephaly [20].

Recently, Colombia faced the world’s second largest Zika outbreak, leading to a four-fold increase in the overall microcephaly cases compared to the previous year. However, the relative increase of reported cases (per 10,000 live births) was far fewer than in Brazil, where a nine-fold increase was reported [11]. This observation raises the possibility that additional risk factors might be driving the highest incidence of microcephaly-zika related cases reported so far. It is unclear why a focused cluster of microcephaly cases has occurred in the Northeast region of Brazil. Several theories on the intensification of Zika virus transmission and resulting severe
fetal neural defects have been raised. These have pointed to socio-economic factors such as precarious living conditions and low regional gross domestic product (GDP) [21–25]. Further, sequential exposure to arboviruses and even co-infections could play a role in severe clinical manifestations [19, 26–27].

Here we aim to provide insights into the unusual pattern of microcephaly distribution in Brazil. By analysing the number of cases reported and co-distribution of dengue, chikungunya and Zika virus, their vectors, as well as socioeconomic and environmental data, we sought to investigate whether co-acting risk factors might be contributing to the Zika microcephaly cases in the Brazil Northeast region.

**Material and methods**

**Mapping the incidence of dengue, chikungunya and Zika virus**

To better understand the co-distribution of dengue, chikungunya and Zika virus in Brazil we collated data on reported cases (per 100,000 inhabitants) for all three pathogens from 2014 to 2016. Data including the total number of cases per year and per state for each virus were obtained from epidemiological Bulletins from the Brazilian Ministry of Health database, (until Volume 48, representing cumulative data until week 52 of 2016) [28], confirmed by information available at each Brazil Federal Unit’s Secretary of Health. Bulletins from the Brazilian Ministry of Health database are technical-scientific publications edited by the Department of Health Surveillance, are circulated with monthly and weekly frequency, and are used for reporting the monitoring activities and investigation of specific seasonal diseases. Specifically, they report the total number of cases notified by each State Secretary. The data obtained for Zika in 2015 consist of suspected and/or confirmed cases, as most of them were only confirmed in 2016. Confirmation is based on RT-PCR and serology methodologies. PubMed and Web of Science databases were searched for studies that reported outbreaks of dengue (family Flaviviridae), Zika and chikungunya (family Togaviridae) from 2010, including cases of co-infections. The search terms used were: “Outbreak”, “Zika”, “Chikungunya”, “Dengue”, “Brazil”, and “Co-infections”. Maps of arbovirus incidence (per 100,000) per Brazilian Federal Unit were constructed using the package `tmap` in R software [29].

**Distribution of the notified cases of microcephaly**

Microcephaly is a congenital malformation where babies are born with a skull size smaller than expected when compared to those of the same sex and age [30]. Specifically, microcephaly is defined as a head circumference that is two standard deviations (SDs) below the mean for the appropriate age and sex, or gestational age if measured at birth. The geographic regions where the microcephaly cases were registered refer to the mother’s place of residence. Data on microcephaly from 2010–2016 were extracted from the Brazilian Ministry of Health website (http://portalsauade.saude.gov.br/) and from the epidemiological Bulletins (until volume 47, week 52 of 2016) available at the Brazilian Ministry of Health database [31]. Additional information was extracted from the System of Strategic Management Support (SAGE) [32], the “Registro de Eventos em Saúde Pública” (RESP-Microcefaalia) [33]. It includes the epidemiological information (2015–2016) regarding microcephaly and/or Central Nervous System changes, provided under the “Protocol on Surveillance and Response to Occurrence of Microcephaly and/or Central Nervous System” [34].

Data on the cumulative incidence of microcephaly from 2015 to 2016 are based on cases that fulfilled the previous definition of cephalic perimeter (33 cm), in addition to the new definitions adopted in the Surveillance Protocol 2015, that defined the 32 cm for boys and 31.5 cm for girls, born with 37 or more weeks of gestation. Notified cases in fetuses, abortions,
stillbirths or newborns were confirmed positive for microcephaly by Zika or other undetermined infectious diseases when: a) typical alterations indicative of congenital infection were found, such as cerebral calcifications, ventricular and posterior fossa alterations among other signs observed by an imaging method, or/and b) when confirmed by laboratory-based Zika virus identification. Data are presented per state.

Socioeconomic data, environmental management

Demographic, socioeconomic, and environmental management data were obtained from the national census performed by the Brazilian Institute of Geography and Statistics (IBGE) [35]. Data for 2016 have been estimated from the last national census (2010). The demographic and socioeconomic variables used, include: (i) Human development index (scale from 0 to 1, where lower values indicate lower development), which considers education (average years of studies), longevity (population life expectancy) and gross domestic product per capita; (ii) Poverty index (scale from 0 to 100, where lower values indicate greater poverty), which considers health (nutrition and child mortality), education (school attendance) and living standards (sanitation, water, electricity, cooking fuel, assets, house conditions). Environmental management data considers: Garbage accumulation index (number of houses with accumulated and uncollected garbage) and rainwater storage (number of houses with storage of rainwater in containers). Data are presented per year and per state.

Aedes aegypti surveillance

Aedes aegypti infestation levels in Brazil in 2016, was obtained from the Aedes aegypti Infestation Index Rapid Survey (LIRAA), performed every year by the Federal Government. This house infestation index measures the percentage of searched buildings with the presence of larvae of A. aegypti. The average house infestation index (HII) of each municipality is calculated and classified in different risk levels (low-risk or satisfactory, HII < 0.9; mid-risk or alert, 1 < HII < 3.9; high risk of A. aegypti infestation, HII > 4) [36].

Statistical analysis

All the data were merged within the R statistical software. We performed an ecological analysis where pairwise relationships between microcephaly, virus infection, social economic factors, arbovirus and Aedes aegypti infestation index were assessed using Spearman’s and Pearson’s correlations and regression models.

Results

Distribution of microcephaly and arbovirus infections in Brazil

During the period from 2010–2014, Brazil registered an average of 156 cases of microcephaly per year. Surprisingly, by the end of 2015, the number of cases was 20 times higher (Fig 1A). Pernambuco was the first state reporting the unusual number of microcephaly cases in newborns, with 10 times more notifications than the average for the whole country during the preceding 5 years (Fig 1B) [37]. The number of reported cases kept increasing, with 10,867 notifications reported from November 2015 until the end of 2016, from which 7,023 were registered only in the Northeast region (Fig 1B). Ongoing laboratory investigations confirmed 2,366 positive cases for congenital microcephaly suspected of Zika virus infection or other infectious agents. Of these positive cases, molecular or serological laboratory investigations confirmed Zika infection in 697 microcephaly cases. The top 10 Brazil Federal Units with the
The highest number of infection-related microcephaly cases includes all the 9 states from the Northeast region and one state from the Northern Brazil (Figs 1 and 2, S1 Fig).

It was also in the Northeast region that the first cases of Zika infection were reported in May 2015 [4]. However, a study based on the genome of Zika virus strains from Brazil and ecological and epidemiological data, revealed that the Zika virus was already present in the Northeast by February 2014 [38]. By the end of 2015 there was a higher number of suspected Zika cases in the States of Bahia, Mato Grosso, Rio Grande do Norte and Mato Grosso do Sul (Fig 3A). One year later, autochthonous cases of Zika virus infection had been confirmed in all the 27 Brazil Federal Units and more than 200,000 cases have already been registered [5] (Fig 3A).

In 2016, the analysis of the reported cases according to geographical regions shows that the Center-West region had the highest incidence rate (222/100,000) followed by Northeast (134.4/100,000) (Fig 3B).

Although it is now established that Zika virus is the cause of severe fetal complications in pregnancy, including microcephaly, the distribution of reported Zika cases and infection-related microcephaly do not overlap (Fig 3A). In fact, there is no correlation between the number of Zika reported cases per 100,000 inhabitants in 2015 or 2016 and the distribution of infection-related microcephaly in Brazil (Parts A and D of S2 Fig) (Correlation R^2 ≤ 0.25; P ≥ 0.212). During the same period Brazil experienced simultaneous transmission of dengue and chikungunya. During January-September 2016, Brazil recorded 200,465 Zika cases, 236,287 chikungunya cases and 1,438,624 dengue cases [39]. It is possible that co-infection increases the severity of symptoms, as previously shown [40]. In Brazil, few studies have reported the co-circulation and/or co-infection of dengue, Zika, and chikungunya [19, 27, 41]. Interestingly, the number of chikungunya cases reported from 2014 to 2015 (Fig 3A) is predominant in the Northeast region. By 2016 the incidence of chikungunya virus increased in
the whole country but it remained mostly concentrated in the Northeast (407.7/100,000) followed by the North (44/100,000) region (Fig 3B). All of the 9 Federal Units from the Northeast region recorded chikungunya cases in excess of 80 per 100,000 habitants. In addition, Amapa and Tocantins states in the North, and Rio de Janeiro in the Southeast, also had a very high number of reported cases (Fig 3A). There is a significant overlap in the distribution of infection-related microcephaly cases and of chikungunya infection (Part B and E of S2 Fig) ($R^2 = 0.57; P < 0.002$). There have been reports that associated chikungunya infection with an increase in neurologic manifestations, including Guillain-Barre syndrome and encephalitis, encephalopathy and microcephaly [42–48]. An investigation in the state of Salvador reported the intense chikungunya transmission between June and November 2015 and the co-occurrence with the Guillain-Barre syndrome outbreak in the city [49]. It has also been suggested that the local health authorities have underestimated chikungunya transmission, as attention was focused on the Zika and microcephaly outbreaks [50].

Dengue fever has a wider distribution in Brazil. Between 2014 and 2015, the incidence of dengue increased in almost all geographic regions, keeping similar rates during 2016 (Fig 3A). In the last three consecutive years, the highest number of reported dengue cases has been observed in the Center-West and Southeast regions (Fig 3B). There is no correlation between infection-related microcephaly cases (2015–2016) and the distribution of suspected dengue fever cases (2014–2016) ($R^2 \leq 0.22; P \geq 0.272$) (Part C and F of S2 Fig). This is consistent with other studies of the dengue virus infection in pregnancy, where although there is evidence of
vertical transmission and increased risk of preterm birth and low birthweight [51], no microcephaly or other congenital brain abnormalities have been reported [52].

**Infection-related microcephaly, socio-economic and environmental management factors**

To evaluate the potential effect of socioeconomic factors on the incidence of Zika-suspected microcephaly, as suggested by others [21–23, 53], we considered the Human development index (HDI) and the poverty index calculated in 2010. The HDI values range from 0.6 to 0.8 across the country. The Northeast and North regions concentrate the Federal Units with the lowest HDI (= 0.6) (Fig 4A). The highest poverty index is observed in the Northeast region (43.5 to 59.5) (Fig 4B), the epicentre of microcephaly notified cases. There is a strong correlation between the distribution of infection-related microcephaly cases and poverty index (Part A of S3 Fig) ($R^2 = 0.68; p<0.0001$).

The Northeast region also had the highest numbers of houses that accumulate garbage and store rainwater (Fig 4C and 4D) that could lead to an increase in vector proliferation and therefore Zika transmission. These results are consistent with a recently reported association of an increase in Zika and chikungunya infections with garbage destination, type of sanitary installation and pipe-borne water [54]. These environmental management indicators may assist an understanding of the population’s behaviours, which may be responsible for increasing the chances of *Aedes* proliferation through underlying human-driven increased mosquito breeding. It also indicates the lack of sanitation and water distribution.
In the Americas, the domestic behavior of *Ae. aegypti* makes it the most important vector for chikungunya, Zika and dengue virus in urban and suburban areas [55, 56]. A surveillance program of *Ae. aegypti* density (house infestation index) was introduced in Brazil aiming to assist the implementation of measures for vector population control. The 2016 *Ae. aegypti* house infestation index classified most of the regions in the Northeast region (Ceará, Rio Grande do Norte, Paraíba, Pernambuco, Alagoas, Sergipe, Bahia) or in the North (Para) (Fig 4E), as being at high risk of *Ae. aegypti* infestation. In fact, data on the entomological index corroborate the findings on the environmental management in the Northeast region (Fig 4C and 4D). We did observe a correlation across the federal Units of the risk of *A. aegypti* infestation and the incidence of chikungunya, but not dengue or Zika (S4 Fig) (dengue: $R^2 = 0.01, P = 0.962$; chikungunya: $R^2 = 0.44, P = 0.020$; zika: $R^2 = 0.25, P = 0.202$). It is important to consider that arboviral epidemics are most likely determined by multiple factors including environmental
conditions, host status and herd immunity and virus genetic mutations that might cause changes in virulence, or vector competence [57].

Discussion

Several studies have confirmed the association of Zika infection during pregnancy and congenital microcephaly [58]. Since February 2017, 31 countries or territories have reported microcephaly cases potentially associated with Zika infection [59] However, the overwhelming level of cases in the Brazilian Northeast region has not been reported elsewhere.

During the Zika virus outbreak in Brazil there was simultaneous transmission of dengue and chikungunya. These viral infections are transmitted by the same Aedes spp. mosquitoes and have close clinical manifestations. In Brazil, the prediction values for the distribution and prevalence of *Ae. Aegypti*, found mainly in urban environment [55], are higher than for *Ae. albopictus*, a predominantly peri-urban and rural vector [60–63]. A study of the geographic distribution of spatially unique occurrence data of Aedes mosquitoes for the Americas shows that *A. aegypti* abundance is 1.5 times higher than for *Ae. albopictus* in Brazil, and with particularly high concentrations in Northern Brazil [62]. A survey performed in 2014 revealed that *Ae. Albopictus* is currently present in at least 59% of the Brazilian municipalities, particularly in the Southeast region [63].

Recent data have shown that in Brazilian urban areas there is an increase number of Zika and chikungunya cases [54], which coincides with the *Ae. aegypti* strong affinity to urbanised regions [64]. Conversely, relatively lower numbers of cases occur in forest or agricultural areas [54]. Densely populated areas with poor water infrastructure may provide more breeding opportunities for *Ae. Aegypti* [65]. In the Northeast of Brazil, over 75% of the breeding sites are due to precarious water storage [54]. Dengue infections have also been shown to be higher in urban areas in Thailand [66].

Although dengue, chikungunya and Zika viruses share the same vector, there was not a perfect overlap of their spatial and temporal distributions in Brazil. Several factors could lead to the observed distribution, such as herd immunity, vector competence for the different viruses, the proportion of asymptomatic cases and under notification of cases. The Center-West region has the highest reported cases of dengue and Zika virus infections, whereas chikungunya reported infection is much higher in the Northeast region compared to the rest of the country. Chikungunya transmission efficiency seems to be lineage specific. Studies on *Ae. aegypti* and *Ae. albopictus* from Florida have shown differences in vector competence depending on the chikungunya strain. *Ae. albopictus* were more susceptible to infection with the La Réunion strain than sympatric *Ae. Aegypti* [67]. A more recent study showed that the chikungunya Asian strain is better transmitted by *Ae. aegypti* species than by *Ae. albopictus*, whereas the Indian Ocean strain leads to higher body infection and transmission in *Ae. albopictus* mosquitoes [68]. In addition, virus mutations can interfere with the transmission rate, for instance, an alanine-valine substitution at position 226 of the E1 envelope glycoprotein (E1-A226V) can improve transmission by *Ae. albopictus* [69, 70] Comparative genome studies on the chikungunya virus circulating in different Brazil regions could provide insights on the higher number of chikungunya cases found in the Northeastern Brazil.

The initial epidemic of the Zika virus was also reported in the Northeast region, spreading eventually to other states. Few studies in Brazil have reported the co-circulation and/or co-infection of dengue, Zika, and chikungunya in the Northeast region [19, 41]. In Bahia, a small study using metagenomic next-generation sequencing approach revealed that 13.3% of patients with confirmed Zika virus infection were actually co-infected with chikungunya virus [27]. Other studies worldwide have also reported co-infections with various combinations of chikungunya/dengue and Zika in endemic and epidemic regions [71–72]. It has also been
shown that *A. aegypti* mosquitoes can co-transmit all combinations of these viruses simultaneously without affecting vector competence [73]. The effect of co-infections in the development of infection and disease outcome is poorly defined, particularly due to the limited clinical information, misdiagnosis and lack of laboratory testing. However, some studies have indicated that previous arboviral infections or co-infections may represent a risk factor for severe clinical manifestations [26]. Pre-existing anti-dengue immunity can promote substantial enhancement of Zika virus infection *in vitro* as well as increased morbidity and mortality in mice [40]. Also, dengue-specific antibodies enhance the infection of a primary Brazilian Zika isolate in K562 cell line [74]. Even though our data did not show an association of infection-related microcephaly cases and the distribution of dengue fever, it is important to note that dengue is endemic in the Northeast region, with 573.3 reported cases per 100,000 inhabitants in 2016 [39]. Dengue transmission is permanent in all regions where Zika and chikungunya infection were reported [75], and there is a high probability of previous or co-infection of dengue with either virus. In relation to the observed outbreaks of chikungunya and Zika, these infections partially overlap by time and space [76], as described in more detail for regions in the Northeast including Pernambuco, Recife [77, 78] and Bahia [49], indicating that previous or co-infection of these virus is possible. It is important to note that the overlap in space-time distribution of Zika, dengue, and chikungunya cases is challenging to report, particularly due to the number of asymptomatic cases and the possible presence of non-specific clinical manifestations which are difficult to diagnose.

The pathogenic effect of Zika and chikungunya virus co-infection, or of the three viruses, has not been studied. We observed a correlation between the distribution of chikungunya infection and the congenital microcephaly in the Northeast region, which may indicate that previous arboviral infections or co-infection with chikungunya could increase Zika severity. Others have linked the outbreak of Guillain-Barre syndrome, firstly related to the Zika virus outbreak in 2015 in the city of Salvador, with the concurrent intense chikungunya transmission [50]. It is important to note that chikungunya infection can result in neurologic manifestations such as encephalitis [79], encephalopathy [43,80], peripheral neuropathy (including Guillain-Barre syndrome) [46,81–82]. In addition, chikungunya virus infection may result in complications during pregnancy either to the mother or to the newborn [83–84]. Complications for the mother, such as chronic inflammatory rheumatism, as well as neurocognitive impairment in infants and microcephaly have been reported in congenital chikungunya transmission [47,49,85–86].

The Northeast region of Brazil has the highest poverty index, with the lowest social economic factors and poorest environmental management, which can be responsible for increasing *Aedes* proliferation. Poverty can drive malnutrition and general poor health that might affect host immunity and the response and clinical progression of an infection.

Altogether, these findings corroborate previous study that suggests that the Zika virus may not be the only factor responsible for the high frequency of congenital microcephaly observed in the Northeast region [55]. Here, using ecological data analysis, we highlight the possible co-circulation of the three arboviruses, together with the socio-economic and environmental factors specific for that region. To understand the complete pathogenesis and severity of Zika infection in the presence of other viruses and other possible co-factors, it is necessary to conduct comprehensive cohort studies involving large patient groups, with detailed socio-economic and environmental factors and effective differential diagnosis for these viruses.

**Limitations of the study**

It is important to consider that the findings in this study are subject to at least six limitations. Firstly, the cases of infection-related microcephaly reported in the databases include congenital
microcephaly caused by Zika virus or other infectious agents. Therefore, it is not known if the distribution of microcephaly cases is caused by laboratory confirmed Zika virus infection. Secondly, cases of co-infections (dengue, Zika, chikungunya) were not investigated or not reported. Third, the ascertainment of birth defects generally does not capture infants or fetuses whose birth defects are not apparent prenatally or at delivery, but rather are identified several months after birth. Fourth, there was no report on *Ae. albopictus* distribution by the LIRAa survey. Fifth, the molecular characterization and phylogenetic analysis of the current circulating strains of Zika, dengue and chikungunya viruses in Brazil is important, but was not possible. Sixth, these data are analyzed at a State-level, which are large regions, and differences between smaller areas (municipalities) were not investigated. Future studies of the genomic epidemiology of these viruses can assist with improving an understanding of the biology, disease phenotypes and transmission, and support the design of diagnostic and vaccine strategies to control the next epidemics.

Overall, we analysed a set of robustly collected and curated data on epidemiological, environmental and socio economic factors of all regions in Brazil. Our results support the hypothesis that the high rates of microcephaly in the Northeast state in Brazil might not be due to Zika infection alone. Our study reinforces the need for comprehensive large-scale cohort studies, as well as public-health measures and guidance to better inform the population under the higher risk of infection-related microcephaly in Northeast of Brazil.

**Conclusions**

The unusual distribution of microcephaly-Zika associated cases in Brazil is likely to be caused by a combination of epidemiological, environmental and socio economic factors. Our work highlights the overlap between the distribution of chikungunya infection with the co-incidence of infection-related microcephaly in the Northeast region. It emphasizes the link between poverty and the risk of disease, and the impact that poor environments can have on human health and the spread of infections. To understand the impact of co-infection on disease outcomes, we recommend that the list of infectious diseases screened during epidemics, particularly for pregnant woman, is updated to include and effective differentiate arboviruses from ongoing outbreaks or epidemics.

**Supporting information**

**S1 Fig. A map of Brazil highlighting regional divisions and Federal Units.** (JPG)

**S2 Fig. Association between infection-related microcephaly and incidence of arboviruses.** Pearson’s correlation test and linear regression was used to investigate the association between infection-related microcephaly and incidence of: Zika in 2015 (A) and 2016 (D); chikungunya in 2015 (B) and 2016 (E); dengue in 2015 (C) and 2016 (F); all per region of Brazil, and results were considered significant for $P<0.05$. (JPG)

**S3 Fig. The correlation between infection-related microcephaly and poverty index.** Microcephaly versus poverty index. Results were considered significant when $P<0.05$. (JPG)

**S4 Fig. The correlation between *Aedes aegypti* infestation risk (LIRAa) and arboviruses incidence in the year 2016.** The number of municipalities at high risk of *A. aegypti* infestation, per Brazil Federal Unit, versus incidence of dengue (A), chikungunya (B) and Zika (C). Linear
correlation was considered significant when $P < 0.05$.

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**References**

1. World Health Organisation. http://www.who.int/mediacentre/factsheets/fs117/en/.

2. Teixeira MG, Andrade AM, Costa Mda C, Castro JN, Oliveira FL, Goes CS et al. East/Central/South African genotype chikungunya virus, Brazil, 2014. Emerg Infect Dis. 2015; 21(5):906–7. https://doi.org/10.3201/eid2105.141727 PMID: 25898939

3. Boletim Epidemiológico, Ministério da Saúde, Brasil. 2017; 48:6. Monitoramento dos casos de febre de chikungunya até a Semana Epidemiológica 52, 2016. Available from http://portalarquivos.saude.gov.br/images/pdf/2017/fevereiro/05/2017_002-Dengue%20SE52_corrigido.pdf.

4. Zanluca C, Melo VC, Mosimann AL, Santos GI, Santos CN, Luz K. First report of autochthonous transmission of Zika virus in Brazil. Mem Inst Oswaldo Cruz. 2015; 110(4):569 –72. https://doi.org/10.1590/0074-02760150192 PMID: 26061233

5. Boletim Epidemiológico, Ministério da Saúde, Brasil. 2017; 48:9. Monitoramento dos casos de febre pelo vírus Zika até a Semana Epidemiológica 52, 2016. Available from http://portalarquivos.saude.gov.br/images/pdf/2017/fevereiro/05/2017_002-Dengue%20SE52_corrigido.pdf.

6. Possas C, Brasil P, Marzochi MC, Tanuri A, Martins RN, Marques ET et al. Zika puzzle in Brazil: peculiar conditions of viral introduction and dissemination—A Review. Mem Inst Oswaldo Cruz. 2017; 112(5):319–327. https://doi.org/10.1590/0074-02760160510 PMID: 28443985

7. Victora CG, Schuler-Faccini L, Matijasevich A, Ribeiro E, Pessoa A, Barros FC. Microcephaly in Brazil: how to interpret reported numbers? Lancet. 2016; 387(10019):621–4. https://doi.org/10.1016/S0140-6736(16)00273-7 PMID: 26864961

8. Souza WV, Araújo TV, Albuquerque Mde F, Braga MC, Ximenes RA, Miranda-Filho Dde B et al. Microcephaly in Pernambuco State, Brazil: epidemiological characteristics and evaluation of the diagnostic accuracy of cutoff points for reporting suspected cases. Cad Saude Publica. 2016; 32(4):e00017216. https://doi.org/10.1590/0102-311X00017216 PMID: 27143306

9. de Fatima Vasco Araoao M, van der Linden V, Brainer-Lima AM, Coeli RR, Rocha MA, Sobral da silav P et al. Clinical features and neuroimaging (CT and MRI) findings in presumed Zika virus related congenital infection and microcephaly: retrospective case series study. BMJ. 2016; 353:i1901. https://doi.org/10.1136/bmj.i1901 PMID: 27075009

10. de Araújo TV, Rodrigues LC, de Alencar Ximenes RA, de Barros Miranda-Filho D, Montarroyos UR, de Melo AP et al. Association between Zika virus infection and microcephaly in Brazil, January to May, 2016: preliminary report of a case-control study. Lancet Infect Dis. 2016; (12):1356–1363. https://doi.org/10.1016/S1473-3099(16)00318-8 PMID: 27641777
11. Cuevas EL, Tong VT, Rozo N, Valencia D, Pacheco O, Gilboa SM, et al. Preliminary Report of Microcephaly Potentially Associated with Zika Virus Infection During Pregnancy—Colombia, January-November 2016. MMWR Morb Mortal Wkly Rep. 2016; 65(49):1409–1413. https://doi.org/10.15585/mmwr.mm6549e1 PMID: 27977645

12. Benjamin I, Fernández G, Figueira JV, Parpacén L, Urbina MT, Medina R et al. Zika virus detected in amniotic fluid and umbilical cord blood in an in vitro fertilization-conceived pregnancy in Venezuela. 2017; 107(6):1319–1322. https://doi.org/10.1016/fernstiert.2017.02.112 PMID: 28390691

13. Mikar J, Korva M, Tui N, Popovic M, Poljsak-Prijatelj M, Mraz J et al. Zika Virus Associated with Microcephaly. N Engl J Med. 2016; 374(10):951–8. https://doi.org/10.1056/NEJMoai1600651 PMID: 26862926

14. Honein MA, Dawson AL, Petersen EE, Jones AM, Lee EH, Yazdy MM, et al. Birth Defects Among Fetuses and Infants of US Women With Evidence of Possible Zika Virus Infection During Pregnancy. JAMA. 2017; 317(1):59–68. https://doi.org/10.1001/jama.2016.19006 PMID: 27960197

15. Dick GW, Kitchen SF, Haddow AJ. Zika virus. I. Isolations and serological specificity. Trans R Soc Trop Med Hyg. 1952; 46(5):509–20. PMID: 12995440

16. Duffy MR, Chen TH, Hancock WT, Powers AM, Kool JL, Lanciotti RS et al. Zika virus outbreak on Yap Island, Federated States of Micronesia. N Engl J Med. 2009; 360(24):2536–43. https://doi.org/10.1056/NEJMo805715 PMID: 19516034

17. Bulletin épidémiologique hebdomadaire international, BHll n° 424 (2014) Available from: http://www.irvs.sante.fr/fr/Publications-ets-outils/Bulletin-ebedomadaireinternational/Tous-les-numeros/2013/Bulletin-ebedomadaire-international-du-30-octobre-au-5-novembre-2013-N-424.

18. Kochakarn T, Kotanan N, Kümpornsin K, Loesbanluechai D, Auewarakul P et al. Comparative genome analysis between Southeast Asian and South American Zika viruses. Asian Pac J Trop Dis. 2016; 6(11):1048–1054. https://doi.org/10.1016/j.ajptd.2016.10.002 PMID: 27890363

19. Cardoso CW, Paploski IA, Kikuti M, Rodrigues MS, Campos GS et al. Outbreak of Exantheatic Illness Associated with Zika, Chikungunya, and Dengue Viruses, Salvador, Brazil. Emerg Infect Dis. 2015; 21(12):2274–6. https://doi.org/10.3201/eid2112.151167 PMID: 26584464

20. Cauchemez S, Besnard M, Bompard P, Dub T, Eronde-Guignot D et al. Association between Zika virus and microcephaly in French Polynesia, 2013–15: a retrospective study. Lancet. 2016; 387(10033):2125–2132. https://doi.org/10.1016/S0140-6736(16)00651-6 PMID: 26993883

21. Souza WV, Albuquerque MFPM, Bezerra LCA, Mendes ADCG, Lyra TM et al. Microcephaly epidemic related to the Zika virus and living conditions in Recife, Northeast Brazil. BMC Public Health. 2018; 18(1):130. https://doi.org/10.1186/s12889-018-5039-z PMID: 29329574

22. Gardner LM, Bota A, Gangavarapu K, Kraemer MUG, Grubaugh ND. Inferring the risk factors behind the geographical spread and transmission of Zika in the Americas. PLoS Negl Trop Dis. 2018; 12(1):e0006194. https://doi.org/10.1371/journal.pntd.0006194 PMID: 29346387

23. Human Rights Watch, Neglected and Unprotected: The Impact of the Zika Outbreak on Women and Girls in Northeastern Brazil, 12 July 2017, available at: http://www.refworld.org/docid/59671dcb4.html [accessed 16 February 2018].

24. Ali S, Guglielmini O, Harber S, Harrison A, Houle L, Ivory J, et al. Environmental and Social Change Drive the Explosive Emergence of Zika virus in the Americas. PLoS Negl Trop Dis. 2017; 11(2):e0005135. https://doi.org/10.1371/journal.pntd.0005135 PMID: 28182667

25. Pelizzo G, Calcetta V, Fusillo M, Nakib G, Ierullo AM, Alfei A, et al. Malnutrition in pregnancy following bariatric surgery: three clinical cases of fetal neural defects. Nutr J. 2014; 13:59. https://doi.org/10.1186/1475-2891-13-59 PMID: 24929556

26. Paul LM, Carlin ER, Jenkins MM, Tan AL, Barcellona CM, Nicholson CO, et al. Dengue virus antibodies enhance Zika virus infection. Clin Transl Immunology. 2016; 5(12):e117.

27. Sardi SI, Somasekar S, Naccache SN, Bandeira AC, Tauro LB, Campos GS et al. Coinfections of Zika and Chikungunya Viruses in Bahia, Brazil, Identified by Metagenomic Next-Generation Sequencing. J Clin Microbiol. 2016; 54(9):2348–53. https://doi.org/10.1128/JCM.00877-16 PMID: 27413190

28. Portal da Saúde. Ministério da Saúde. Brasil. Available from: http://portalsaude.gov.br/boletins-epidemiologicos (under assunto: Zika

29. https://www.r-project.org/

30. World Health Organisation (WHO) 2017. Microcephaly definition. Available from: http://www.who.int/mediacentre/factsheets/microcephaly/en/

31. Portal da Saúde. Ministério da Saúde. Brasil. Available from: http://portalsaude.gov.br/boletins-epidemiologicos (under assunto: Microcefalia e/ou alterações do Sistema Nervoso Central (SNC) associadas à infecção por vírus Zika)
32. Saia de Apoio a Gestão Estratégica (SAGE). Ministério da Saúde. Brasil. Available from: http://sage.saude.gov.br/paineis/microcefalia).

33. Registro de Eventos em Saúde Pública. (RESP-Microcefalia). Available from: http://www.resp.saude.gov.br/microcefalia.

34. Portal da Saúde. Ministério da Saúde. Brasil. Microcefalia. Available from: http://portalsaude.saude.gov.br/index.php/cidadao/principal/agencia-saude/22553-brasil-adota-recomendacao-da-oms-e-reduz-medida-para-microcefalia.

35. Instituto Brasileiro de Geografia e Estatística. Available from: http://www.ibge.gov.br/home/

36. Levantamento Rápido de Índices para Aedes aegypti—LIRAa—para Vigilância Entomológica do Aedes aegypti no Brasil. Available from: http://portalarquivos.saude.gov.br/images/pdf/2016/novembro/24/Tabela—LIRAaNacional-2016.pdf.

37. Vargas A, Saad E, Dimach GS, Santos RH, Sivini MA, Albuquerque LC, et al. Characteristics of the first cases of microcephaly possibly related to Zika virus reported in the Metropolitan Region of Recife, Pernambuco State, Brazil. Epidemiol Serv Saude. 2016; 25(4):691–700. https://doi.org/10.5123/S1679-49742016000400003 PMID: 27869982

38. Faria NR, Quick J, Claro IM, Theze J, de Jesus JG, Giovannetti M, et al. Establishment and cryptic transmission of Zika virus in Brazil and the Americas. Nature. 2017; 546(7634):406–410. https://doi.org/10.1038/nature22401 PMID: 28538727

39. Boletim Epidemiológico, Ministério da Saúde, Brasil. 2017; 48:3. Monitoramento dos casos de dengue até a Semana Epidemiológica 52, 2016. Available from http://portalarquivos.saude.gov.br/images/pdf/2017/fevereiro/05/2017_002-Dengue%20SE52_corrigido.pdf.

40. Bardina SV, Bunduc P, Tripathi S, Duehr J, Frere JJ, Brown JA, et al. Enhancement of Zika virus pathogenesis by preexisting antiflavivirus immunity. Science. 2017; 356(6334):175–180. https://doi.org/10.1126/science.aal4365 PMID: 28360135

41. Pessoa R, Patriota JV, Lourdes de Souza Md, Felix AC, Mamede N, Sanabani SS. Investigation into an outbreak of Dengue-like illness in Pernambuco, Brazil, Revealed a Cocirculation of Zika, Chikungunya, and Dengue Virus Type 1. Medicine (Baltimore). 2016; 95(12):e3201.

42. Tandale BV, Sathe PS, Arankalle VA, Wadia RS, Kulkarni R, Shah SV, et al. Systemic involvements and fatalities during Chikungunya epidemic in India, 2006. J Clin Virol. 2009; 46(2):145–9. https://doi.org/10.1016/j.jcv.2009.06.027 PMID: 19239775

43. Bank AM, Batra A, Colorado RA, Lyons JL. Myeloradiculopathy associated with chikungunya virus infection. J Neurovirol. 2016; 22(1):125–8. https://doi.org/10.1007/s13365-015-0372-9 PMID: 26306687

44. Chusri S, Siripaitoon P, Hirunpat S, Silpaopajakul K. Case reports of neuro-Chikungunya in southern Thailand. Am J Trop Med Hyg. 2011; 85(2):386–9. https://doi.org/10.4269/ajtmh.2011.10-0725 PMID: 21813863

45. Oehler E, Fournier E, Leparc-Goffart I, Larre P, Cubizolle S, Sookhareea C et al. Increase in cases of Guillain-Barré syndrome during a Chikungunyaoutbreak, French Polynesia, 2014 to 2015. Euro Surveill. 2015; 20(48):30079. https://doi.org/10.2807/1560-7917.ES.2015.20.48.30079 PMID: 26690898

46. Lebrun G, Chadda K, Reboux AH, Martinet O, Guazère BA. Guillain-Barré syndrome after chikungunya infection. Emerg Infect Dis. 2009; 15(3):495–6. https://doi.org/10.3201/eid1503.071482 PMID: 19239775

47. Gérardin P, Sampañiz S, Ramful D, Boumahni B, Bintner M, Alessandri JL, et al. Neurocognitive outcome of children exposed to perinatal mother-to-child Chikungunya virus infection: the CHIMERE cohort study on Reunion Island. PLoS Negl Trop Dis. 2014; 8(7):e2996. https://doi.org/10.1371/journal.pntd.0002996 PMID: 25033077

48. Fritel X, Rollot O, Gerardin P, Gauzere BA, Bideault J, Lagarde L et al. Chikungunya virus infection during pregnancy, Reunion, France, 2006. Emerg Infect Dis. 2010; 16(3):495–9. https://doi.org/10.3201/eid1603.091403 PMID: 20204216

49. Malta JM, Vargas A, Leite PL, Percio J, Coelho GE, Ferraro AH, et al. Guillain-Barré syndrome and other neurological manifestations possibly related to Zika virus infection in municipalities from Bahia, Brazil, 2015. Epidemiol Serv Saude. 2017; 26(1):9–18. https://doi.org/10.5123/S1679-49742017000100002 PMID: 28226004

50. Cardoso CW, Kikutli M, Prates AP, Paploski IA, Tauro LB, Silva MM et al. Unrecognized Emergence of Chikungunya Virus during a Zika Virus Outbreak in Salvador, Brazil. PLoS Negl Trop Dis. 2017; 11(1):e0005334. https://doi.org/10.1371/journal.pntd.0005334 PMID: 28114414

51. Friedman EE, Dallah F, Harville EW, Myers L, Buekens P, Breart G et al. Symptomatic Dengue infection during pregnancy and infant outcomes: a retrospective cohort study. PLoS Negl Trop Dis. 2014; 8(10):e3226. https://doi.org/10.1371/journal.pntd.0003226 PMID: 25299383
52. Phongsamart W, Yoksan S, Vanaprapa N, Chokephaibulkit K. Dengue virus infection in late pregnancy and transmission to the infants. Pediatr Infect Dis J. 2008; 27(6):500–4. https://doi.org/10.1097/INF.0b013e318167917a PMID: 18434933

53. Butler D. Brazil asks whether Zika acts alone to cause birth defects. Nature. 2016; 535(7613):475–6. https://doi.org/10.1038/nature.2016.20309 PMID: 27466104

54. Aguiar BS, Lorenz C, Virginio F, Suesdek L, Chiaravalloti-Neto F. Potential Risks of Zika and Chikungunya Outbreaks in Brazil: a Modelling Study. International Journal of Infectious Diseases https://doi.org/10.1016/j.ijid.2018.02.007 [Epub ahead of print]

55. Powell JR, Tabachnick WJ. History of domestication and spread of Aedes aegypti—a review. Mem Inst Oswaldo Cruz. 2013; 108 Suppl 1:11–7.

56. Carvalho MS, Honorio NA, Garcia LMT, Carvalho LCS. Aedes aegypti control in urban areas: A systemic approach to a complex dynamic. PLoS Negl Trop Dis. 2017; 11(7):e0005632. https://doi.org/10.1371/journal.pntd.0005632 PMID: 28749942

57. Liang G, Gao X, Gould EA. Factors responsible for the emergence of arboviruses; strategies, challenges and limitations for their control. Emerg Microbes Infect. 2015; 4(3):e18. https://doi.org/10.1038/emi.2015.18 PMID: 26038768

58. Krauer F, Riesen M, Reivez L, Oladapo OT, Martinez-Vega R, Porgo TV et al. Zika Virus Infection as a Cause of Congenital Brain Abnormalities and Guillain-Barré Syndrome: Systematic Review. PLoS Med. 2017; 14(1):e1002203. https://doi.org/10.1371/journal.pmed.1002203 PMID: 28045901

59. WHO reports 2017. Zika virus and complications: 2016 Public Health Emergency of International Concern. Available from: http://www.who.int/emergencies/zika-virus/en/.

60. Delatte H, Dehecq JS, Thiria J, Domerg C, Paupy C, Fontenille D. Geographic distribution and developmental stages of Aedes albopictus (Diptera: Culicidae) during a Chikungunya epidemic event. Vector Borne Zoonotic Dis. 2008; 8(1):25–34. https://doi.org/10.1089/vbz.2007.0649 PMID: 18171104

61. Santos J, Meneses BM. An integrated approach for the assessment of the Aedes aegypti and Aedes albopictus global spatial distribution, and determination of the zones susceptible to the development of Zika virus. Acta Trop. 2017; 168:80–90. https://doi.org/10.1016/j.actatropica.2017.01.015 PMID: 28111332

62. Kraemer MU, Sinka ME, Duda KA, Mylne AQ, Shearer FM, Barker CM, et al. The global distribution of arbovirus vectors Aedes aegypti and Ae. albopictus. Elife. 2015; 4:e08347. https://doi.org/10.7554/elif.e08347 PMID: 26126267

63. Carvalho RG, Lourenço-de-Oliveira R, Braga IA. Updating the geographical distribution and frequency of Aedes albopictus in Brazil with remarks regarding its range in the Americas. Mem Inst Oswaldo Cruz. 2014; 109(6):787–96. https://doi.org/10.1590/0074-0276140304 PMID: 25317707

64. Rochlin I, Ninivaggi DV, Hutchinson ML, Farajollahi A. Climate change and range expansion of the Asian tiger mosquito (Aedes albopictus) in Northeastern USA: implications for public health practitioners. PLoS One. 2013; 8(4):e60874. https://doi.org/10.1371/journal.pone.0060874 PMID: 23665282

65. Tsuzuki A1, Vu TD, Higa Y, Nguyen TY, Takagi M. High potential risk of dengue transmission during the hot-dry season in Nha Trang City, Vietnam. Acta Trop. 2009 Sep; 111(3):325–9. https://doi.org/10.1016/j.actatropica.2009.05.010 PMID: 19467217

66. Nakhapakorn K1, Tripathi NK. An information value based analysis of physical and climatic factors affecting dengue fever and dengue haemorrhagic fever incidence. Int J Health Geogr. 2005 Jun 8; 4:13. https://doi.org/10.1186/1476-072X-4-13 PMID: 15943863

67. Pesko K, Westbrook CJ, Mores CN, Lounibos LP, Reiskind MH. Effects of infectious virus dose and bloodmeal delivery method on susceptibility of Aedes aegypti and Aedes albopictus to chikungunya virus. J Med Entomol. 2009: 46(2):395–9 PMID: 19351094

68. Alto BW, Wiggins K, Eastmond B, Velez D, Lounibos LP, Lord CC. Transmission risk of two chikungunya lineages by invasive mosquito vectors from Florida and the Dominican Republic. PLoS Negl Trop Dis. 2017; 11(7):e0005724. https://doi.org/10.1371/journal.pntd.0005724 PMID: 28749964

69. Vega-Ru a A, Zouache K, Caro V, Diancourt L, Delaunay P, Grandadam M, et al. 2013. High efficiency of temperate Aedes albopictus to transmit chikungunya and dengue viruses in the Southeast of France. PLoS One 8:e59716. https://doi.org/10.1371/journal.pone.0059716 PMID: 23527259

70. Arias-Goeta C, Mousson L, Rougeon F, Failloux AB. 2013. Dissemination and transmission of the E1-226V variant of chikungunya virus in Aedes albopictus are controlled at the midgut barrier level. PLoS One 8:e57548. https://doi.org/10.1371/journal.pone.0057548 PMID: 23437397

71. Cherubuddi K, Iovine NM, Shah K, White SK, Paisie T, Salemi M, et al. Zika and Chikungunya virus co-infection in a traveller returning from Colombia, 2016: virus isolation and genetic analysis. JMM Case Rep. 2016; 3(6):e005072. https://doi.org/10.1099/jmmcr.0.005072 PMID: 28348794
72. Waggoner JJ, Gresh L, Vargas MJ, Ballesteros G, Tellez Y, Soda KJ et al. Viremia and Clinical Presentation in Nicaraguan Patients Infected With Zika Virus, Chikungunya Virus, and Dengue Virus. Clin Infect Dis. 2016; 63(12):1584–1590. https://doi.org/10.1093/cid/ciw589 PMID: 27578819

73. Rücker C, Weeger-Lucarelli J, García-Luna SM, Young MC, Byas AD, Marrieta RA, et al. Impact of simultaneous exposure to arboviruses on infection and transmission by Aedes aegypti mosquitoes. Nat Commun. 2017; 8:15412. https://doi.org/10.1038/ncomms15412 PMID: 28524874

74. Castanha PMS, Nascimento EJM, Braga C, Cordeiro MT, de Carvalho OV, de Mendonça LR, et al. Dengue Virus-Specific Antibodies Enhance Brazilian Zika Virus Infection. J Infect Dis. 2017; 215(5):781–785. https://doi.org/10.1093/infdis/jiw638 PMID: 28039355

75. Lowe R, Barcellos C, Brasil P, Cruz OG, Honório NA, Kuper H, et al. The Zika Virus Epidemic in Brazil: From Discovery to Future Implications. Int J Environ Res Public Health. 2018; 15(1). pii: E96.

76. WHO- Pan America report 2016- www.paho.org

77. Magalhaes T, Braga C, Cordeiro MT, Oliveira ALS, Castanha PMS, Maciel APR et al. Zika virus displacement by a chikungunya outbreak in Recife, Brazil. PLoS Negl Trop Dis. 2017; 11(11):e0006055. https://doi.org/10.1371/journal.pntd.0006055 PMID: 29108009

78. Chusri S, Sripaioon P, Hirunpat S, Silpapojakul K. Case reports of neuro-Chikungunya in southern Thailand. Am J Trop Med Hyg. 2011; 85(2):386–9. https://doi.org/10.4269/ajtmh.2011.10-0725 PMID: 21813863

80. Tandale BV, Sathe PS, Arankalle VA, Wadia RS, Kulkarni R, Shah SV et al. Systemic involvements and fatalities during Chikungunya epidemic in India, 2006. J Clin Virol. 2009; 46(2):145–9. https://doi.org/10.1016/j.jcv.2009.06.027 PMID: 19640780

81. Agarwal A, Vibha D, Srivastava AK, Shukla G, Prasad K. Guillain-Barre syndrome complicating chikungunya virus infection. J Neurovirol; 23(3):504–507. https://doi.org/10.1007/s13365-017-0516-1 PMID: 28194661

82. Oehler E, Fournier E, Leparc-Goffart I, Larre P, Cubizolle S, Sookharea C et al. Increase in cases of Guillain-Barre syndrome during a Chikungunya outbreak, French Polynesia, 2014 to 2015. Euro Surveill. 2015; 20(48):30079. https://doi.org/10.2807/1560-7917.ES.2015.20.48.30079 PMID: 26690898

83. Robillard PY, Boumahni B, Gérardin P, Michault A, Fourmaintraux A, Schuffenecker I et al. Vertical maternal fetal transmission of the chikungunya virus. Ten cases among 84 pregnant women. Presse Med. 2006; 35(5 Pt 1):785–8.

84. Ramful D, Carbonnier M, Pasquet M, Boughani B, Ghazouani J, Noormahomed T et al. Mother-to-child transmission of Chikungunya virus infection. Pediatr Infect Dis J. 2007; 26(9):811–5. https://doi.org/10.1097/INF.0b013e3181004372 PMID: 17721376

85. Villamil-Gómez W, Alba-Silvairita L, Menco-Ramos A, Gonzalez-Vergha A, Molinares-Palacios T, Barrios-Corales M, et al 2015. Congenital Chikungunya Virus Infection in Sincelejo, Colombia: A Case-Series. J Trop Pediatr; 61(5):386–92. https://doi.org/10.1093/tropmed/frm051 PMID: 26246086

86. Lyra PP, Campos GS, Bandeira ID, Sardi SI, Costa LF, Santos FR, et al. Congenital Chikungunya Virus Infection after an Outbreak in Salvador, Bahia, Brazil. AJP Rep. 2016; 6(3):e299–300. https://doi.org/10.1055/s-0036-1587323 PMID: 27555980