Burden of Chikungunya and its seasonal trend in south Karnataka- A study in a tertiary care centre

Trupti B. Naik1, Sathish J. V.2*, Jayashree S.3

1Assistant Professor, 2Associate Professor, 3Tutor, Dept. of Microbiology, Chamarajanagar Institute of Medical Sciences, Chamarajanagar, Karnataka, India

*Corresponding Author: Sathish J. V
Email: javagalsathish37@gmail.com

Received: 2nd August, 2018 Accepted: 17th September, 2018

Abstract

Introduction: Mosquito borne arboviral infections have become major public health problem at present due to many emerging and re – emerging infections and Chikungunya is one of them. Although Chikungunya causes severe clinical manifestations like arthritis, it is not routinely tested at the health care facilities and therefore goes undiagnosed and as such its prevalence is likely to be underestimated. Climate is one of the important factor influencing the occurrence and distribution of disease.

Objectives: a) To estimate seroprevalence of Chikungunya virus infection among clinically suspected cases. b) To determine the seasonal variation of Chikungunya infection.

Materials and Methods: A cross sectional study was conducted at a tertiary care hospital using secondary data maintained in the microbiology laboratory registers, for clinically suspected patients of Chikungunya infection who reported to the hospital from January- December 2017. Prevalence was estimated by analyzing data for the results of IgM antibody by ELISA tests and any variations in disease reporting by age, gender and season were assessed. Statistical analysis was done using WHO Epi info software version 3.5.4.

Results: A total of 1308 serum samples were analyzed during the study period. Out of which 123 (9.4%) samples were found positive for Chikungunya infection by IgM ELISA. The proportion of Chikungunya cases was higher in monsoon season with maximum rate of positivity in the month of July 48 (39.02%). Age group of 16-30 years was the most commonly affected i.e. (35.77%).

Conclusion: The present study confirms that Chikungunya is mainly a disease of rainy season and also identifies certain vulnerable groups for effective planning of interventions.

Keywords: Chikungunya, Immunoglobulin M. Seasonal variation, Seroprevalence.

Introduction

Arthropod-borne viral infections cause major disease burden in tropical and subtropical countries worldwide and Chikungunya is one among them. Chikungunya virus (Arbovirus), a member of the genus Alpha virus and family Togaviridae is mainly transmitted by the bite of mosquito Aedes aegypti and less frequently by Aedes albopictus. Climate is an important factor contributing to change in the disease epidemiology. Studies recently reported in Southeast Asia, found that Chikungunya outbreaks were negatively correlated (P< 0.05) with drought conditions, but positively correlated with warmer-than-normal temperatures and rainfall.

India is severely affected with over 1.4 million infections as per recent estimates and the disease is known to cause epidemics when the proportion of immune naive population increases. By the latest report in 2010, Chikungunya has spread to more than 18 states/union territories within the country and the outbreak is still continuing. This rapid spread of Chikungunya demonstrated the devastating magnitude of infection and its ability of transmission in temperate regions. Resurgence has been attributed to various factors including urbanization, globalization, increase in the mosquito population, loss of herd immunity and the mutation A226V in the E1 gene causing a significant increase in CHIKV infectivity for Aedes albopictus.

Chikungunya causes fever and nonspecific clinical manifestations like arthralgia, chills, headache, nausea, vomiting, low back pain, and rash lasting for a period of 1-7 days similar to malaria, dengue and other bacterial infections, but it is not routinely tested at the health care facilities and therefore goes undiagnosed and as such its prevalence is likely to be underestimated. The probable diagnosis of Chikungunya can be made on the basis of clinical manifestations, where as confirmatory diagnosis can be made only by laboratory tests. Currently, used tests include RT-PCR for confirming the presence of CHIKV, while sensitive IgM antibody ELISA is used to distinguish from dengue. Hence the present study was undertaken with the following objectives 1) To estimate seroprevalence of Chikungunya infection among clinically suspected cases 2) To determine the seasonal variation of Chikungunya infection.

Materials and Methods

The present cross-sectional study was conducted at a tertiary care hospital of Chamarajanagar Institute of Medical Sciences, Chamarajanagar using secondary
data maintained in the microbiology laboratory registers, of clinically suspected chikungunya cases reported to various inpatient and out-patient departments of the hospital for a period of one year from January–December 2017, after obtaining prior permission from the head of the Institute. Prevalence was estimated by analyzing data for the results of IgM antibody by ELISA tests and any variations in disease reporting by gender, age and season were assessed.

Approximately 5 ml of blood samples were collected from all the suspects of Chikungunya as a part of the routine laboratory work and the sera were separated by centrifugation of the whole blood sample and tested for the Chikungunya IgM antibody by using IgM antibody capture ELISA kit developed by National Institute of Virology (Arbovirus Diagnostic, NIV, Pune, India). Manufacturer’s instructions were followed strictly while performing the ELISA. Values were calculated and results were interpreted as per manufacturer’s guidelines.

**Principle of IgM Capture ELISA for CHIK:** Anti-human IgM are coated on to the solid surface (wells) and if IgM antibodies are present in the patient’s sera, they are captured by it. If IgM and CHIK antigen, which is added in the next step, are homologous, the antigen binds to captured IgM. During the washing step unbound antigen is removed. Biotinylated anti-CHIK monoclonal antibody (CHIK-B) is added in the subsequent steps followed by avidin-histidine rich protein (HRP). Then substrate is added and development of color is monitored. 1N H2SO4 is used to stop the reaction. The optical density (OD) is monitored at 450 nm. OD values are directly proportional to the amount of CHIK virus specific IgM antibodies present in the sample. Kit controls and in-house positive and negative controls were used to validate the test.

**Statistical Analysis:** Data was analyzed using WHO Epi info software version 3.5.4.

**Results**

A total of 1308 serum samples from suspected cases of Chikungunya were tested for presence of specific IgM antibodies by ELISA method and of which 123 (9.4%) samples were found positive for Chikungunya. Majority i.e. 44 (35.77%) of the positive cases were in the age group of 16-30 years and the proportion of positive cases was slightly higher in males with male to female ratio of 1:0.75. A seasonal peak of Chikungunya infection was seen in the months of July 48 (39.02%) followed by June i.e. 31 (25.2%).

**Fig. 1:** Illustrates seroprevalence of Chikungunya infection. Among 1308 clinically suspected cases, 123 (9.4%) were found to be positive for Chikungunya infection by IgM ELISA in the present study.

**Table 1 and Fig. 2:** Of the 123 total positive cases of Chikungunya, though the number of positive cases 44 (35.77) was higher in the age group of 16-30 years, the rate of positivity was higher in the age group of 31-60 years (23.02%). The study found no significant association between the prevalence of Chikungunya and the age group of patients (P > 0.05).

**Table 2:** Gender wise distribution of samples of Chikungunya suspects and positives. Out of 1308 samples tested, 661 (50.53%) were females and the rest males. Out of the 123 total positive cases of Chikungunya, both the proportion of cases i.e. 70 (56.91%) and the rate of seropositivity i.e. 10.81% was slightly higher in males as compared to females. However, there was no significant association between the prevalence of Chikungunya and gender of patients (P > 0.05).

**Fig. 3 and Table 3:** Seasonal trend showed that suspected cases of Chikungunya infection were highest in the month of June and July followed by August and slowly tapered by September with maximum positive cases in the month of July 48 (39.02%) followed by June i.e. 31 (25.2%). Serologically confirmed Chikungunya cases were highest i.e. 108 (87.8%) during the monsoon period compared to other seasons of the year and it was found to be statistically significant (P < 0.05).

**Table 1: Age wise distribution of samples of Chikungunya suspects and positives**

<table>
<thead>
<tr>
<th>Age in years</th>
<th>Suspects</th>
<th>Positives</th>
<th>Rate of positivity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No. (%)</td>
<td>No. (%)</td>
<td>%</td>
</tr>
<tr>
<td>&lt; 15</td>
<td>284 (21.71)</td>
<td>25 (20.32)</td>
<td>8.8</td>
</tr>
<tr>
<td>16-30</td>
<td>531 (40.59)</td>
<td>44 (35.77)</td>
<td>8.28</td>
</tr>
<tr>
<td>31-45</td>
<td>304 (23.24)</td>
<td>35 (28.45)</td>
<td>11.51</td>
</tr>
<tr>
<td>46-60</td>
<td>139 (10.62)</td>
<td>16 (13)</td>
<td>11.51</td>
</tr>
<tr>
<td>&gt;60</td>
<td>50 (3.82)</td>
<td>3 (2.43)</td>
<td>6</td>
</tr>
<tr>
<td>Total</td>
<td>1308 (100)</td>
<td>123 (100)</td>
<td>9.4</td>
</tr>
</tbody>
</table>

\[ \chi^2 = 3.89 \; P = 0.42 \]
Table 2: Gender wise distribution of Chikungunya suspects and positives

<table>
<thead>
<tr>
<th>Gender</th>
<th>Suspects</th>
<th>Positive</th>
<th>Rate of positivity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No. (%)</td>
<td>No. (%)</td>
<td>(%)</td>
</tr>
<tr>
<td>Male</td>
<td>647 (49.46)</td>
<td>70 (56.91)</td>
<td>10.81</td>
</tr>
<tr>
<td>Female</td>
<td>661 (50.53)</td>
<td>53 (43.08)</td>
<td>8.01</td>
</tr>
<tr>
<td>Total</td>
<td>1308 (100)</td>
<td>123 (100)</td>
<td>9.4</td>
</tr>
</tbody>
</table>

χ² = 3.01 P = 0.082

Table 3: Season wise distribution of Chikungunya suspects and positives.

<table>
<thead>
<tr>
<th>Months</th>
<th>Suspects</th>
<th>Positives</th>
<th>Rate of positivity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No. (%)</td>
<td>No. (%)</td>
<td>(%)</td>
</tr>
<tr>
<td>Pre Monsoon</td>
<td>90 (6.88)</td>
<td>4 (3.25)</td>
<td>4.44</td>
</tr>
<tr>
<td>Monsoon</td>
<td>1173 (89.67)</td>
<td>108 (87.8)</td>
<td>9.2</td>
</tr>
<tr>
<td>Post Monsoon</td>
<td>45 (3.44)</td>
<td>11 (8.94)</td>
<td>23.91</td>
</tr>
<tr>
<td>Total</td>
<td>1308 (100)</td>
<td>123 (100)</td>
<td>9.4</td>
</tr>
</tbody>
</table>

χ² = 14.60 P = 0.0006

Fig. 1: Seroprevalence of Chikungunya infection

Fig. 2: Age wise distribution of Chikungunya suspects and positives
chikungunya virus is reported with other studies. Higher humidity
National Institute of Allergy
reness and undertake
at Navi Mumbai
ada
Ahmedabad
5
Ahmedabad
28.45
19
Trupti B. Naik et al. Burden of Chikungunya and its seasonal trend in south Karnataka…. Indian Journal of Microbiology...

Susceptible with few studies reporting both genders to be equally susceptible.17,24,26

Seasonal trend of chikungunya infection was analyzed by the month wise data of samples. In India the first CHIKV outbreak in 1963 was observed during July to December, coinciding with the monsoon and post monsoon seasons. However, in the present study, CHIKV was detected more during the monsoon season with positive cases being raised gradually from June with peak in the month of July and declined in the month of October. This finding correlated with findings of Tomar A. et al. at Navi Mumbai,1 Bharti N et al. at Tamil Nadu27 Nandi J. et al. at Delhi,28 Ray P. et al. in a multicentric study19 Divya P. et al. at Ballar18 Manimunda SP et al. at Dakshin Kannada29 and Mohanty I et al. at Southern Odissa.7 This may be due to increased transmission of disease at the start of the rainy season with increased breeding of both Aedes aegypti and Aedes albopictus. Higher humidity lengthens their life span and increased temperatures shortens the extrinsic incubation period. As mosquitoes are sensitive to changes in temperature and available moisture, they decrease in number in dry and cool seasons so cases declined in rest of the year. Certain studies showed highest number of cases during July to December (Ahmedabad)22 September-December (Lucknow)12 November (Varanasi)13 sept- nove (Mumbai)14 sep-dec (Ahmedabad)15 sep-jan (Ahmedabad).16 These seasonal peaks emphasize the need to create public awareness and undertake necessary vector control measures including improvements in sanitation and hygiene during pre monsoon period.

Conclusion
The seroprevalence of Chikungunya in our study was 9.4% which reiterates the fact that Chikungunya is a major health concern in our setting. The present study also confirms that Chikungunya is mainly a disease of
monsoon and identifies certain vulnerable groups for effective planning of interventions.

References

How to cite this article: Naik T. B, Sathish J. V, Jayashree S. Burden of Chikungunya and its seasonal trend in south Karnataka- A study in a tertiary care centre. Indian J Microbiol Res. 2018;5(4):492-496.