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Determinants Of Dengue Transmission During Epidemic And Inter-Epidemic Periods In Two Colombian Cities

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DETERMINANTS OF DENGUE TRANSMISSION DURING EPIDEMIC AND INTER-EPIDEMIC PERIODS IN TWO COLOMBIAN CITIES

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Thesis in Candidacy for the Degree of Master of Public Health

Yale School of Public Health

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ABSTRACT

Dengue fever is an urban disease that has a complex epidemiology consisting of alternating periods of intense epidemics and persistence in endemic locations. No studies have compared the patterns of geographic variation and determinants of dengue transmission in neighborhoods during epidemic and non-epidemic periods. Colombia’s individual-based epidemiological surveillance system provides a unique opportunity to study this topic. The goal of this study was to better understand dengue epidemiology in two of the highest dengue fever reporting Colombian cities that vary in climate, Armenia (elevation 1320-1580 m, 21-23°C) and Barranquilla (elevation 5-134 m, 27-30°C). We used a novel ecological approach, Levin’s niche breadth, to define epidemic and inter-epidemic periods in each city. Regression tree models were built with the following outcome variables for each neighborhood: total number of dengue cases reported during the study period and proportion of dengue cases that occur during inter-epidemic periods. The explanatory variables used were elevation, house count (in lieu of population), housing density, and the Colombian socioeconomic class (SEC) indicator. House count was consistently found to be the main determinant of the total number of reported dengue cases in neighborhoods in both cities. The proportion models identified different determinants of persistent dengue virus transmission in the two cities. Lower elevation was the main driver of persistence in Armenia while lower SEC was the main driver in Barranquilla. These findings suggest that although the overall number of dengue cases depend on the impact of population (as represented by house count) on viral introduction, factors that influence the reproductive rate have a larger influence on transmission during inter-epidemic periods. The persistence determinants identified in this study could potentially help vector control programs to identify key areas to focus disease control efforts.
ACKNOWLEDGMENTS

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INTRODUCTION

Objectives
This investigative study was motivated by the need to better understand the dynamics of dengue transmission during epidemic versus inter-epidemic periods in an urban setting. The objectives were two-fold and were met through a study of two cities in Colombia that have variable patterns of endemic dengue transmission—Armenia and Barranquilla. The first objective was to explore the geographic patterns of dengue cases during dengue epidemics and inter-epidemic periods. The second objective was to determine whether different socio-ecological determinants drive dengue transmission in neighborhoods during epidemic compared to inter-epidemic periods.

Statement of General Problem under Investigation
Dengue is a febrile illness caused by a flavivirus and is principally transmitted by the Aedes aegypti mosquito. This is an urban disease that has a complex epidemiological cycle consisting of alternating epidemic and inter-epidemic periods. The pattern of dengue epidemic cycle is highly variable in different locations due to its dependence on natural and built environmental conditions, vectorial capacity, human biology and behavior, viral serotypes, and susceptibility of the population\textsuperscript{1-3}. Dengue was first detected in Colombia in 1971 and its incidence and prevalence has since grown exponentially\textsuperscript{4}. Colombia now experiences about 40-50,000 cases of dengue annually and has one of the highest severe dengue to dengue fever ratio in all of Latin America\textsuperscript{5}. Currently, all four dengue serotypes are circulating in the country and almost all departments (states) experience periodic epidemics\textsuperscript{4}. The National Institute of Health of Colombia (Instituto Nacional de Salud, INS) operates a national dengue surveillance system that collects weekly reports of individual, probable dengue cases for each neighborhood in each city. This study utilized Colombia’s unique dengue surveillance system and georeferencable neighborhood descriptive data to identify the determinants of dengue transmission during epidemic and inter-epidemic periods in two Colombian cities over multiple year time span.
Dengue epidemics occur in multi-year cycles and are characterized by marked increases in the number of dengue cases above expected levels\textsuperscript{2,6}. When epidemics occur in the city, there are high levels of viral introductions occurring in the neighborhoods. Inter-epidemic periods, on the other hand, are characterized by persistent transmission of dengue at low or undetectable levels and are influenced by seasonal cycles\textsuperscript{1,2,7–11}. Persistent transmission is initially started by viral introduction, but its continuation depends on dengue’s effective reproductive number \((R)\). While the highly visible epidemics often overshadow the inter-epidemic periods, recent studies have elucidated the importance of dengue persistence through inter-epidemic periods. First, separate studies in Australia, Puerto Rico, Brazil, Argentina, Venezuela, and Cuba concurred that the spatial patterns of dengue persistence is stable over time, thereby resulting in clustering of dengue in a few key areas during inter-epidemic periods\textsuperscript{1,12–15}. Secondly, these key areas are also responsible for producing a significant portion of the total dengue cases in a city\textsuperscript{1,12–15}. Third, persistent pockets can also act as virus reservoirs and cause outbreaks in other locations\textsuperscript{1,16,17}. Given the evidence, it is logical to conclude that neighborhoods that have continuous transmission even during inter-epidemic periods must have the ideal conditions for the proliferation of the dengue virus. These neighborhoods are likely to be the first places where new viral introductions take hold and subsequently spread to neighboring communities.

While the dengue vaccine is still under development, dengue prevention and control is heavily based on combating the mosquito vector. However, the outcomes of vector control programs have not been very effective or long lasting\textsuperscript{1,15,17,18}. Studies in Singapore determined that the cause of dengue reemergence was due to the adaptation of a case-reactive approach to vector control in combination with other factors including increased global travel and urbanization\textsuperscript{17}. Case-reactive vector control programs cannot identify subclinical infections or viral persistence, thereby allowing dengue to reemerge when herd immunity wanes\textsuperscript{17}. In addition, conditions that favor proliferation of mosquitos such as uncontrolled urban development and sprawl, reintroduction of vectors, and climate change may also hinder vector
control programs\textsuperscript{17,19,20}. Since most vector control programs have limited resources, strategic allocation of these resources to combat \emph{A. aegypti} may lead to better and more sustainable outcomes\textsuperscript{1}.

Given the studies on inter-epidemic periods described above, factors that drive persistent viral transmission need to be better understood. Vector control programs should focus its resources to stop dengue transmission at its weakest during the inter-epidemic periods in these key neighborhoods\textsuperscript{7}. Furthermore, identification of persistent areas can help direct immunization campaigns once dengue vaccines are approved for general use\textsuperscript{7}. Lastly, disease persistence in a neighborhood lowers the overall quality of life of its residents and it is important for public health interventions to focus on these areas to decrease health disparity across a city.

**Relevant Studies**

To our knowledge, no studies have compared geographic variation and determinants of dengue transmission during epidemic and inter-epidemic periods. In order to fulfill this knowledge gap, we need to first address several challenges in studying this topic. The first challenge is in establishing a definition to differentiate between epidemic and inter-epidemic periods. The transition between the two periods is a gradual process and not readily apparent. In addition, definitions used in past studies are based on total cases within a certain time period, which was useful for areas being studied but lacked the capability to be used in comparative investigations between cities because case thresholds are relative to each location’s dengue burdens\textsuperscript{1,3,8}. For example, the estimated overall dengue prevalence (in cases per 10,000 inhabitants) is 339 in Brazil, 116.5 in Colombia, but only 36.7 in Peru, therefore a cases-based definition can represent different epidemic extents in each of these places\textsuperscript{21}.

A study on dengue persistence by Barrera et al. showed that there is a statistical significant correlation between the number of cases and the total area that report cases—meaning that as the number of dengue cases increase over time, the geographical areas that experience dengue transmission also expands\textsuperscript{13}. Since epidemics are initiated by introductions, this observation suggests that epidemics (city
level events) are perpetuated by multiple local introductions (neighborhood level events). This phenomenon underlines the importance of studying epidemic cycles from the perspectives of both the city and local areas such as neighborhoods. Therefore, in order to encompass conditions of both the city and neighborhoods, we used a novel ecological approach based on the correlation of geographic dengue distribution with number of cases to define epidemic and inter-epidemic periods.

After defining epidemic and inter-epidemic periods, the second challenge lies in finding outcome measurements for regression analysis that can distinguish between transmissions occurring during these two periods. If the determinants of these two periods are different, then disease control would benefit greatly from this knowledge and could adapt their programs accordingly. The main ecological difference between epidemic and inter-epidemic periods is that inter-epidemic periods experience lower rates of viral introductions among neighborhoods. Modeling studies have shown that it is important to distinguish between introduced cases and local transmissions, in other words, transmission among groups and transmission within groups, respectively. Furthermore, strictly local transmission that do not have new introductions are maintained based on the $R$ of the system, which can be influenced by social, environmental, herd immunity, and vectorial capacity factors.

The term persistence had also been interpreted inconsistently in relation to inter-epidemic periods in literature. For example, Barrera et al. stratified neighborhoods in a city into three-levels of persistence, where persistence was defined as having a certain number of consecutive months reporting dengue. Barrera et al.’s definition of persistence captured dengue transmission only with respect to conditions within the neighborhood. Our study is the first to specifically focus on dengue persistence in neighborhoods during a time when a majority of neighborhoods in the city does not have dengue (inter-epidemic periods). This second definition of persistence is more selective and depends on conditions in both the neighborhoods and the remainder of the city. Past studies have found evidence that supports differential transmission dynamics when examining the drivers of local and global dengue transmission patterns. Preliminary studies at our study sites found that these two definitions result in
identification of different key neighborhoods, thereby also suggesting that different mechanisms may underline transmission at the local level than transmission at the city level. Therefore, we hypothesized that the determinants of total dengue cases in each neighborhood would be different from the number of cases that occur during inter-epidemic periods (standardized by the total cases in each respective neighborhoods) in those neighborhoods.

Research on dengue epidemiology is often limited by the lack of vector or seroprevalence data. Vector data is not available for every neighborhood in a city simply because it is a time and resource-consuming process. In addition, both modeling and seroprevalence evidence found that fluctuations in vector abundance are not associated with fluctuations in epidemic size and often moderated by other factors such as socioeconomic status (SEC)\textsuperscript{3,24}. Seroprevalence data would be useful to determine the source of infection (introductory versus autochthonous), but it is expensive to obtain and impractical for city-wide studies. In terms of operability of the model, however, surveillance data is superior to seroprevalence data because surveillance data is likely to reflect the temporal dynamics of epidemic and persistent cycles, unlike cross-sectional seroprevalence data.

Alternative mechanisms by which dengue virus persists during inter-epidemic periods include vertical transmission of virus to new mosquito generations and virus replication in alternative vectors\textsuperscript{25,26}. Vertical transmission alone is the least probable cause of virus persistence since mosquito life cycles are short and the virus will be diluted quickly\textsuperscript{25}. In addition, the strength of association of vertical transmission is most likely very small when other factors are present. In Colombia, virus replication in alternative vectors is also not likely since \textit{A. aegypti} has been the sole \textit{Aedes} mosquito found in vector surveillance in the two study cities. Therefore, these mechanisms were not examined in this study.

This study focused on drivers of dengue transmission at the neighborhood level for practical, scientific, and political reasons. Neighborhood level analysis is practical because dengue surveillance and prevention programs are often conducted at the neighborhood level\textsuperscript{7,27}. Neighborhoods are good units of measurements scientifically because characteristics such as SEC are generally homogenous within
neighborhoods and movement patterns generally revolve around neighborhoods\textsuperscript{28,29}. Finally, study results based on neighborhood-level analysis can be directly applied to policies targeting community improvement and health projects. Studies have found that policies and programs targeting health disparities can be especially effective if it engages the community at the neighborhood level\textsuperscript{30}.

**METHODS**

**Locations under Study**

**Armenia, Quindio Department, Colombia**

Armenia is located on the Cordillera Central within the mountainous regions of Western Colombia. The city has a large altitude range that varies from 1320 – 1580 m and has an annual average ambient temperature ranges between 20 - 20.3°C\textsuperscript{31,32}. According to the 2005 census, the city has an estimated population of 272,574 people\textsuperscript{33}.

**Barranquilla, Atlántico Department, Colombia**

Barranquilla is located on the Caribbean coast of Colombia. The city is located at approximately sea level has an altitude range between 5 - 134 m and has average ambient temperature range between 26.8 - 28.3°C\textsuperscript{31,32}. The population of the city is approximately 1,112,889 people\textsuperscript{33}.

**Sources of Data**

**Epidemiological data**

Dengue surveillance data consisting of weekly, individual-based probable cases were obtained from each city. The data was originally collected as part of the national dengue surveillance system of the Colombian National Institute of Health. Probable dengue cases from 2001 - 2011 were obtained for Armenia. Laboratory confirmed dengue cases from 2004 - 2006 and probably dengue cases from 2007 -
2011 were obtained for Barranquilla. The surveillance data reported diagnosis date, doctor notification date, patient’s age, and patient’s neighborhood of residence. Prior to analysis, the weekly surveillance data were organized into three-week periods based on doctor-visit-dates to account for the time lag between initial infection and doctor visit date. In instances where the doctor-visit-date was not available, dengue-case-notification-date was used in lieu of doctor-visit-date. Based on the 2010-2011 Armenia dengue surveillance reports, the average time lag between doctor-visit date and notification date was 0.53 days for 99.9% of the data. Neighborhoods were included in the analysis if they reported greater than 5 cases during the study period in order to account for neighborhood identification errors that may have occurred during surveillance reporting.

**Spatial data**

Geographically projected shapefiles of the cities were obtained from municipal governments (projection-Transverse Mercator, datum-Bogota). The shapefiles contained digitalized polygons of neighborhoods and houses in each city.

**Other sources of data**

Orthorectified digital elevation data (90 m resolution, datum-WGS_1984) for the study areas were downloaded from The CGIAR Consortium for Spatial Information in raster format (http://srtm.csi.cgiar.org/). The elevation data was projected to the same coordinates of the city shapefiles using the Project Raster Tool of the Data Management Tools in ArcMap Software v.10.0 (ESRI, Redlands, CA). Mean elevation of each neighborhood was calculated using the Zonal Statistics function in ArcMap10. SEC ratings of the neighborhoods and house count of the neighborhoods were obtained from municipal governments. The SEC ratings were based on a scale of 1-6 (from less to more affluent) and measures socioeconomic status relative to each municipality (not comparable between cities). House counts were used as a proxy for population because population data was not available. Interpretation of this proxy is limited because house count does not take into account possible differences in household
size. The estimated area in hectare of each neighborhood was calculated from the neighborhood shapefiles using Zonal Statistics function in ArcMap10. Housing density of each neighborhood in houses per hectare was calculated using its corresponding house count and area.

**Analysis**

**Defining Epidemic and Inter-epidemic Periods**

We observed a statistically significant correlation between the total number of dengue cases reported in each 3-week period and the number of neighborhoods reporting five or more cases of dengue in the same 3-week period in both cities (Armenia, Pearson Correlation Coefficient 0.91605, p<0.0001; Barranquilla, Pearson Correlation Coefficient 0.94476, p<0.0001). When the total dengue cases increased, the area that experience dengue cases expanded and vice versa. This geographic variation between epidemic and inter-epidemic periods was the rational for using an ecological approach, specifically Levin’s standardized niche breadth index \((B_n)\), to define and standardize dengue epidemic and inter-epidemic periods.

Niche breadth measures the extent of utilization of a single type of resource by a species within an environment at a given time\(^{34}\). The concept of niche breadth was applied to the dengue system as follows: dengue was considered the species, the city was the species environment, each neighborhood in the city was a unique resource, and each dengue case reported in a neighborhood was interpreted as a single utilization of the resource\(^{34}\). In summary, the extent of resource utilization as measured by the niche breadth corresponded to the geographical distribution of dengue cases within a city. The niche breadth was then standardized by the total number of neighborhoods in each city. The standardization step in the formula ensured that \(B_n\) from different surveillance data sources could be integrated. It also allowed comparative analysis between cities that have different underlying transmission dynamics.
$B_n$ was calculated for each 3-week period using Equation 1, thereby obtaining a measure of geographic spread of dengue for each 3-week time period. $B_n$ ranges from 0 to 1. The time period was considered an epidemic period if $B_n$ was close to 1.0, representing dengue was equally distributed amongst all neighborhoods. The time period was considered as an inter-epidemic period if $B_n$ was close to 0.0, representing dengue concentrated in a small number of neighborhoods. The specific cut-offs were determined based on the histogram of each niche breadth over time, which showed a clear binary transition between low niche periods (the majority, skewed right) and a high niche periods (long tail). Histogram bin widths were selected using an optimization method of Shimazaki and Shinomoto which minimized the mean integrated squared error of variance within each bin. The relationship among total number of dengue cases, total number of neighborhood that report 5 or more dengue cases, and $B_n$ was graphically represented in Figure 1 and Figure 2.

**Equation 1.** Levin’s Standardized Niche Calculation

$$B_n = \frac{1}{R \sum p_i}$$

$B_n$ = Standardized Niche Breadth Index

R = total number of neighborhoods (resource states) in a city in the study period

$p_i$ = proportion of dengue cases in $i^{th}$ neighborhood in each three-week period out of total number of dengue cases observed in each three-week period
Figure 1: Epidemic curve of Armenia (2001-2011) overlaid on the number of neighborhoods that report 5 or more cases of dengue. Right axis describes the Levin's Standardized Niche Breadth Index. (Surveillance data was not available for the three week periods 78-87)

Figure 2: Epidemic curve of Barranquilla (2004-2011) overlaid on the number of neighborhoods that report 5 or more cases of dengue. Right axis describes the Levin's Standardized Niche Breadth Index.
Determinants of Dengue Transmission during Epidemic and Inter-epidemic Periods

Regression tree analyses were used to model epidemic and inter-epidemic periods in order to determine the drivers of dengue transmission during these periods. Regression trees are robust models that utilize a hierarchical approach to handle complex interactions, non-linear relationships, over-fitting, and missing variables—problems commonly encountered when fitting statistical linear models to dengue systems\textsuperscript{36}. The tree is built by splitting a single response variable (either categorical or continuous) into branches of homogenous groups based on a single explanatory variable using the method of least squares\textsuperscript{36}. This is a fully automated process that explores all splits and branch orders possible within a given set of explanatory variables. The final tree is identified through a pruning process to obtain the tree with the largest number of branches while having the smallest cross-validation error. Colinearity of explanatory variables is presented through surrogate branches with the level of congruence indicated for each surrogate. The surrogate splits answers the question “which other splits would classify the same outcomes in the same way\textsuperscript{37}.” The regression tree results are presented visually and are easy to interpret—each branch is labeled with the explanatory variable value dictating the split, outcome variables that satisfy the split criteria are grouped to the left, and each leaf is labeled with the mean value of the outcome variable and the number of observations in the group\textsuperscript{9,36}. The length of the vertical branches is directly proportion of the total sum of squares the division accounted for\textsuperscript{36}.

Regression tree models were built using the Recursive Partitioning and Regression Tree (rpart) package of the statistical software R (http://www.r-project.org/). Two models representing transmission during epidemic or inter-epidemic periods were explored for each city. Their respective outcome variables were 1) the total number of dengue cases reported by the neighborhood reflecting transmission during epidemic periods and 2) the proportion of cases of that occurred during the inter-epidemic periods in each neighborhood reflecting persistent transmission. The explanatory variables used to predict the responses were elevation, SEC, house count, and housing density.
The first outcome, the total number of dengue cases reported by the neighborhood, was chosen to represent transmission during epidemic periods because it was the most obvious indicator of the total disease burden in a neighborhood. Since transmission cannot indefinitely persist in a neighborhood, the continuation of cases necessarily requires an introduction event. Increase in cases observed in a neighborhood is associated with increase in the rate of introduction into the neighborhood. Therefore, the disease burden in a neighborhood is dependent on the rate of introduction, which in turn is proportional to the level of the epidemic in the city. Inter-epidemic periods are characterized by transmission that occurs locally with limited outside introductions. However, the cases identified through surveillance do not differentiate between locally acquired cases and introductory cases. Therefore, the second outcome was chosen to reflect inter-epidemic periods because it captures the timing of cases. Specifically, this outcome captures the autochthonous cases that occur only through persistent mechanisms and not through introduction.

RESULTS

Epidemiology Trends

A total of 11777 cases in 269 neighborhoods in Armenia were reported to the national dengue surveillance system from 2001-2011. Dengue cases were included in data analysis if they reported the patient’s neighborhood of residence (5.04% of total cases did not report neighborhoods). In order to minimize reporting error, a neighborhood was included in data analysis if it could be located in the city’s GIS shapefile and had at least 50 houses. Niche breadth analysis was conducted with a total of 9622 cases reported in 181 neighborhoods over 186 three-week periods in 2001 - 2011 in Armenia.

Two sets of surveillance data were obtained in Barranquilla. From 2004 to 2006, 2227 laboratory confirmed cases occurred in 150 neighborhoods in Barranquilla and from 2007 to 2011, 6461 probably cases occurred in 294 neighborhoods in Barranquilla. 0.4% of total cases during 2004-2006 and 11.5% of total cases during 2007-2011 did not report neighborhoods and were excluded from analysis. The
difference between epidemic and inter-epidemic periods is much more pronounced in Armenia than in Barranquilla.

Tables 1 and 2 describes the distribution of cases and neighborhoods in epidemic and inter-epidemic periods in Armenia and Barranquilla, respectively. Although the epidemic curves (Figures 1 and 2) were drastically different between the cities, the epidemic trends and geographic patterns were consistent. In both cities, the epidemic period contained the majority of the cases. In addition, the epidemic periods had larger number of neighborhoods report dengue than during the inter-epidemic periods.

<table>
<thead>
<tr>
<th></th>
<th>Number of 3-week Periods (% total time)</th>
<th>Dengue Cases (% total)</th>
<th>Average Number of Cases Per 3-week Period</th>
<th>Average Number of Neighborhoods Per 3-week Period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>186 (100)</td>
<td>9622 (100)</td>
<td></td>
<td>33</td>
</tr>
<tr>
<td>Epidemic</td>
<td>59 (30)</td>
<td>7191 (75)</td>
<td>123</td>
<td>51</td>
</tr>
<tr>
<td>Inter-epidemic</td>
<td>118 (63)</td>
<td>2431 (25)</td>
<td>21</td>
<td>14</td>
</tr>
<tr>
<td>Unknown (no data)</td>
<td>9 (7)</td>
<td>0</td>
<td></td>
<td>-</td>
</tr>
</tbody>
</table>

Table 1. Distribution of cases and neighborhoods in Armenia during epidemic and inter-epidemic periods.

<table>
<thead>
<tr>
<th></th>
<th>Number of 3-week periods (% total time)</th>
<th>Dengue cases (% total)</th>
<th>Average Number of Cases per 3-week Period</th>
<th>Average number of Neighborhoods per 3-week Period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>134 (100)</td>
<td>7658 (100)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Epidemic</td>
<td>22 (16)</td>
<td>3417 (45)</td>
<td>155</td>
<td>71</td>
</tr>
<tr>
<td>Inter-epidemic</td>
<td>108 (81)</td>
<td>4241 (55)</td>
<td>39</td>
<td>25</td>
</tr>
<tr>
<td>Unknown (no data)</td>
<td>4 (3)</td>
<td>0</td>
<td></td>
<td>-</td>
</tr>
</tbody>
</table>

Table 2. Distribution of cases and neighborhoods in Barranquilla during epidemic and inter-epidemic periods.

**Regression Tree Results**

Outcome 1 – Total number of cases in each neighborhood

The regression tree models built using the total number of cases in each neighborhood showed that house count was the main determinant of the number of cases in both cities and was responsible for
multiple branches (Figures 3 and 4). In both cities, house was responsible for 5 splits and there is a positive correlation between the number of houses and the total number of dengue cases in each neighborhood. Interestingly, in Armenia, SEC becomes an important predictor in neighborhoods that have medium number of houses. SEC was also a major surrogate split explanatory variable at the second node in Armenia that would have resulted in 65% agreement. SEC and house count is shown in Armenia to be colinearly related. The highest colinearity values in Barranquilla were between house and elevation at the first node (73% agreement) and between housing density and house at the third node (69% agreement). Since house was used as a proxy for population, we conclude that population was the main driver of epidemic periods.

Figure 3. Regression tree predicting the total number of dengue cases in each neighborhood in Armenia based on the neighborhood’s elevation, SEC, house, and housing density. Threshold criteria are labeled at the top of each branch and the values at the leaves indicate the mean expected number of dengue cases and the number of neighborhoods (n) grouped into that branch.
Outcome 2 – Proportion of cases that occur during inter-epidemic periods in each neighborhood

The regression trees that examined the proportion of cases in inter-epidemic periods identified very different drivers of persistence in the two cities (Figure 5 and 6). In Armenia, elevation was the main driver of persistent dengue transmission during inter-epidemic periods while SEC appeared to be the main driver in Barranquilla. In Armenia, there was an inverse relationship between elevation and proportion of cases that occur during inter-epidemic periods. In Barranquilla, there was also an inverse relationship between SEC and proportion of cases that occur during inter-epidemic periods. The number of splits was also significantly fewer than the trees from outcome 1. In Armenia, SEC was a surrogate split for elevation at both nodes (77% agreement at node one and 97% agreement at node two). In Barranquilla, elevation was a surrogate split for SEC (73% agreement). This indicates that there is strong colinearity between SEC and elevation for both cities.
DISCUSSION

The regression tree analysis results showed that determinants of epidemic transmission differ from that of inter-epidemic transmission but the mechanisms behind the different drivers of transmission during the two periods warrant further discussion.

Population was an important determinant in both cities for the epidemic models. The number of cases was directly reflective of the extent of epidemics and is correlated with the geographic expansion of
dengue in a city. This showed that epidemics occur due to multiple introductions occurring between neighborhoods. High population levels would increase the probability that introductions would occur through increased human movement and increased number of susceptibles. A recent $R_0$ modeling study showed that even if $R_0$ was sufficiently high to favor local transmission, epidemics (propagation of disease across a city) would not occur without a large enough population and high levels of movement. The sheer number of branches driven by housing suggested that the number of cases was sensitive to population levels.

The use of proportion as an outcome in the inter-epidemic models allowed us to tease apart introduction cases and persistent autochthonous cases. House was not present in the final proportion models, thereby indicating that the effect of population was diminished during autochthonous transmissions and factors that influence $R$ become more important. For example, elevation was the main driver of dengue persistence in Armenia, where there was a high elevation gradient within the city. Elevation affects an area’s ambient temperature, humidity, and other climate characteristic, which in turn affects $R_0$ factors associated with vectorial capacity. This showed that the transmission system at high altitudes may be sensitive to temperature changes.

In contrast, elevation was not a significant branch in the proportion model in Barranquilla. This was probably due to the overall low elevation, low elevation gradient, and high temperature conditions throughout Barranquilla. This observation agreed with a past study that showed where temperature and precipitation are already high, increase in either have little effect on transmission rates. One study on two neighboring cities with identical climate conditions along the US-Mexico border found different seroprevalence levels due to differences in SEC. The US city have higher mosquito infestation levels but its residents have very limited exposure to mosquitoes due to continuous air-condition use and therefore the city has essentially no dengue transmission. Given similar climate conditions amongst the neighborhoods, SEC emerged as a main driver of persistence in Barranquilla.
LIMITATIONS

The scope of this study was limited by the explanatory variables available for the study areas. First, we were not able to obtain temperature and climate data at the neighborhood level. Elevation was used as a proxy. Second, population data at the neighborhood level was not available in the two cities. We used house count as a proxy for population, but house count did not account for variations in household size. In addition, housing density did not account for non-urbanized areas in a neighborhood nor variations in household size. Furthermore, there were relatively high levels of colinearity amongst the variables used in the regression trees (60-90%). However, colinearity was expected for a vector-borne disease study and the regression tree approach was selected as the most appropriate method to account for colinearity.

Finally, spatial autocorrelation was not explored in the study and should be analyzed in the future. Spatial autocorrelation could capture the effects of human movement amongst neighborhoods. Preliminary analysis of spatial autocorrelation of residuals from the epidemic regression tree models using Global Moran’s I in Arc GIS showed that both cities have statistically significant autocorrelation. Local Moran’s I hot spot analysis identified various hot spots in the city at 2000 m, 3000 m, and 4000 m radius. Spatial autocorrelation of inter-epidemic periods was not significant. Further analysis of spatial distribution of the various response variables used in regression tree could supplement the regression tree findings.

CONCLUSIONS

Our analysis in Colombia found that there was a geographic expansion of cases during epidemic periods and a geographic recession of cases during inter-epidemic periods. In addition, we found that the determinants of dengue transmission during epidemic periods differed from that of inter-epidemic periods.
Epidemic transmission was driven by population while persistent transmission during inter-epidemic periods was driven by factors unique to each city. Lower elevation was the main driver of persistence in Armenia while lower SEC was the main driver in Barranquilla.

This study used a novel application of Levin’s Standardized Niche Breadth Index to define dengue outbreak and persistent periods. This study also found that regression tree modeling gave insight into factors that cause certain neighborhoods to experience persistent dengue transmission even when there is low dengue introduction between neighborhoods and low overall dengue transmission at the city-level is low. Elevation was the main drivers of persistence in Armenia while SEC was the main driver in Barranquilla. Results from the models could help the identification of key neighborhoods that experience persistent dengue transmission and the mechanism that drive the persistence. Dengue prevention programs can benefit greatly by focusing their resources on addressing these mechanisms within key neighborhoods during the low-transmission periods. Shifting dengue prevention and vector control strategies towards a persistence focused direction could be effective for both short-term control of dengue outbreaks and long term sustained halt of transmission.
REFERENCES


32. IDEAM Instituto de Hidrología, Meteorología y Estudios Ambientales. at <http://institucional.ideam.gov.co/jsp/index.jsf>


37. No Title.

APPENDIX

rPart protocol

Recommended introduction reading – An Introduction to Recursive Partitioning Using the RPART Routines. Elizabeth J. Atkinson & Terry M. Therneau. Mayo Foundation

Blue=code
Green=output
***Words preceded by stars are author’s notes

Data saved in .csv format

Annotated output

R version 2.13.1 (2011-07-08)
Copyright (C) 2011 The R Foundation for Statistical Computing
ISBN 3-900051-07-0
Platform: i386-pc-mingw32/i386 (32-bit)

R is free software and comes with ABSOLUTELY NO WARRANTY.
You are welcome to redistribute it under certain conditions.
Type 'license()' or 'licence()' for distribution details.

Natural language support but running in an English locale

R is a collaborative project with many contributors.
Type 'contributors()' for more information and
'citation()' on how to cite R or R packages in publications.

Type 'demo()' for some demos, 'help()' for on-line help, or
'help.start()' for an HTML browser interface to help.
Type 'q()' to quit R.

***code for loading the data and printing the data

> library (rpart)
> Arm <-read.csv ("C:/.../Arm_reg_Mar31_thesis.csv", header=T)
> print (Arm)
barrioID            GISname
  1      79        BARRIO ALAMOS
  2      29        BARRIO ALCAZAR
  3     128     CONDOMINIO LA ALDEA
  4      30    ALFONSO LOPEZ
  5     170      BARRIO ARCO IRIS

***remaining data omitted

***code for building the tree
> prop11_100 <- rpart (prop11_above100 ~ house + SES + elevation + hdens, data=Arm, method="anova", minsplit=15)

***code for plotting the tree
> plot (prop11_100)

***code for labeling the tree
> text (prop11_100, use.n=T)

*** “anova” method is used for continuous outcome variables, “class” method can be used for categorical variables.
“minsplit” command specifies the minimum number of observations needed to further split the outcomes.
“use.n=T” command labels the number of observations in each leaf.

***This is the full tree. The criteria for the split is at the top of each branch, the mean expected outcome value is at the bottom of each leaf, and the number of observations in each leaf is shown under the
outcome values. Branches to the left are those that obey the criteria of the split, branches to the right are those that do not.

***code for printing the summary statistical results of the tree
> summary (prop11_100)

Call:
rpart(formula = prop11_above100 ~ house + SES + elevation + hdens, 
data = Arm, method = "anova", minsplit = 15)
n=140 (37 observations deleted due to missingness)

CP nsplit rel error xerror xstd
1 0.22795022 0 1.0000000 1.0296171 0.1234651
2 0.07485321 1 0.7720498 0.9972420 0.1395282
3 0.03185386 2 0.6971966 0.9881304 0.1332446
4 0.03144762 3 0.6653427 0.9744629 0.1434450
5 0.02983496 4 0.6338951 0.9875267 0.1460658
6 0.02571645 5 0.6040601 1.0021328 0.1464100
7 0.01857509 6 0.5783437 1.0159742 0.1559440
8 0.01571209 7 0.5597686 1.0399030 0.1554440
9 0.01381519 8 0.5440565 1.0387945 0.1554124
10 0.01291739 9 0.5302413 1.0387263 0.1559440
11 0.01000000 10 0.5173239 1.0453323 0.1567865

Node number 1: 140 observations,  complexity param=0.2279502
mean=0.3694357, MSE=0.02278631
left son=2 (46 obs) right son=3 (94 obs)
Primary splits:
  elevation < 1464.04 to the right, improve=0.22188280, (1 missing)
  house < 113 to the left, improve=0.05388324, (0 missing)
  SES < 2.991489 to the right, improve=0.05049566, (0 missing)
  hdens < 3710 to the left, improve=0.04728225, (0 missing)
Surrogate splits:
  SES < 3.017793 to the right, agree=0.770, adj=0.304, (1 split)
  hdens < 3655.97 to the left, agree=0.698, adj=0.087, (0 split)

***Node number 1 is shown on the tree (a). “left son=2” indicates the total number of observations in subsequent branches to the left and the number indicates the node number of the first subsequent branch. “Primary splits” shows the possible variables to be used in the first node of the tree. “improve” shows how much the tree would improve if the specific variable is used. Improvement is obtained by n times the change in impurity index. The relative size of improvement is important. The variable that gives the best improvement is chosen as the split.
“Surrogate splits” lists possible variables to be used instead of the one chosen. “agree” shows the level of agreement in the results if that variable was used. Ex: 77% of the results in a tree with SES as primary split would agree with using Elevation as primary split.

Node number 2: 46 observations,  complexity param=0.03185386
mean=0.2664109, MSE=0.01162481
left son=4 (41 obs) right son=5 (5 obs)
Primary splits:
- elevation < 1516.875 to the left, improve=0.19002920, (0 missing)
- house < 250.5 to the left, improve=0.16268700, (0 missing)
- SES < 4.861742 to the left, improve=0.13185070, (0 missing)
- hdens < 9020.833 to the right, improve=0.09123028, (0 missing)
Surrogate splits:
- SES < 4.992772 to the left, agree=0.935, adj=0.4, (0 split)

Node number 3: 94 observations, complexity param=0.07485321
mean=0.4198521, MSE=0.02051236
left son=6 (75 obs) right son=7 (19 obs)
Primary splits:
- elevation < 1364.175 to the right, improve=0.12865650, (1 missing)
- house < 127 to the left, improve=0.08165134, (0 missing)
- SES < 2.982206 to the right, improve=0.03583874, (0 missing)
- hdens < 3710 to the left, improve=0.02924879, (0 missing)
Surrogate splits:
- house < 650 to the left, agree=0.828, adj=0.158, (1 split)

Node number 4: 41 observations, complexity param=0.02983496
mean=0.2499976, MSE=0.008965589
left son=8 (19 obs) right son=9 (22 obs)
Primary splits:
- house < 298.5 to the left, improve=0.25891950, (0 missing)
- SES < 1.268889 to the right, improve=0.12161650, (0 missing)
- elevation < 1480.28 to the left, improve=0.10215740, (0 missing)
- hdens < 9020.833 to the right, improve=0.08392677, (0 missing)
Surrogate splits:
- SES < 2.585672 to the left, agree=0.732, adj=0.421, (0 split)
- elevation < 1475.49 to the left, agree=0.683, adj=0.316, (0 split)
- hdens < 2938.034 to the left, agree=0.659, adj=0.263, (0 split)

Node number 5: 5 observations
mean=0.401, MSE=0.0131071

Node number 6: 75 observations, complexity param=0.03144762
mean=0.394484, MSE=0.01545287
left son=12 (10 obs) right son=13 (65 obs)
Primary splits:
- house < 127 to the left, improve=0.08656038, (0 missing)
- SES < 2.976296 to the right, improve=0.04736354, (0 missing)
- hdens < 3710 to the left, improve=0.03553696, (0 missing)
- elevation < 1401.9 to the right, improve=0.03415841, (1 missing)

Node number 7: 19 observations, complexity param=0.01291739
mean=0.5199895, MSE=0.02791621
left son=14 (7 obs) right son=15 (12 obs)
Primary splits:
house < 334 to the left, improve=0.07769026, (0 missing)
SES < 2.040568 to the left, improve=0.05998758, (0 missing)
elevation < 1348.24 to the right, improve=0.04116785, (0 missing)
hdens < 6884.091 to the right, improve=0.03967865, (0 missing)

Surrogate splits:
elevation < 1343.335 to the right, agree=0.684, adj=0.143, (0 split)

Node number 8: 19 observations, complexity param=0.01857509
mean=0.1981526, MSE=0.008765521
left son=16 (11 obs) right son=17 (8 obs)
Primary splits:
  SES < 1.990132 to the right, improve=0.35579630, (0 missing)
hdens < 2948.611 to the right, improve=0.10123190, (0 missing)
elevation < 1480.315 to the left, improve=0.06701270, (0 missing)
house < 207.5 to the right, improve=0.06661416, (0 missing)
Surrogate splits:
  hdens < 2600 to the right, agree=0.737, adj=0.375, (0 split)
  house < 135.5 to the right, agree=0.632, adj=0.125, (0 split)
elevation < 1472.365 to the left, agree=0.632, adj=0.125, (0 split)

Node number 9: 22 observations
mean=0.2947727, MSE=0.004812193

Node number 12: 10 observations
mean=0.30124, MSE=0.03366514

Node number 13: 65 observations, complexity param=0.02571645
mean=0.4088292, MSE=0.01110759
left son=26 (9 obs) right son=27 (56 obs)
Primary splits:
  hdens < 9540 to the right, improve=0.11362650, (0 missing)
elevation < 1404.75 to the right, improve=0.09773067, (1 missing)
  house < 145.5 to the right, improve=0.08072609, (0 missing)
  SES < 2.002366 to the right, improve=0.03731130, (0 missing)

Node number 14: 7 observations
mean=0.4590143, MSE=0.04640722

Node number 15: 12 observations
mean=0.5555583, MSE=0.01369583

Node number 16: 11 observations
mean=0.1505273, MSE=0.007524257

Node number 17: 8 observations
mean=0.2636375, MSE=0.003065252

Node number 26: 9 observations
mean=0.3202111, MSE=0.008249723
Node number 27: 56 observations, complexity param=0.01571209
mean=0.4230714, MSE=0.01010194
left son=54 (40 obs) right son=55 (16 obs)
Primary splits:
elevation < 1405.5 to the right, improve=0.10079410, (1 missing)
house < 145.5 to the right, improve=0.07771955, (0 missing)
SES < 1.001055 to the right, improve=0.05851398, (0 missing)
hdens < 7263.333 to the right, improve=0.03776983, (0 missing)
Surrogate splits:
house < 140 to the right, agree=0.727, adj=0.062, (1 split)
SES < 1.001055 to the right, agree=0.727, adj=0.062, (0 split)

Node number 54: 40 observations
mean=0.40415, MSE=0.01026688

Node number 55: 16 observations, complexity param=0.01381519
mean=0.470375, MSE=0.006556902
left son=110 (11 obs) right son=111 (5 obs)
Primary splits:
elevation < 1394.3 to the left, improve=0.42008780, (0 missing)
SES < 2.450207 to the left, improve=0.15027910, (0 missing)
house < 245 to the left, improve=0.08025302, (0 missing)
hdens < 4552.5 to the left, improve=0.01319235, (0 missing)
Surrogate splits:
SES < 2.911518 to the left, agree=0.75, adj=0.2, (0 split)

Node number 110: 11 observations
mean=0.4349909, MSE=0.003495164

Node number 111: 5 observations
mean=0.54822, MSE=0.004478406

***code for printing the cp values of the tree. Cp values are also included in summary output
> printcp (prop11_100)

Regression tree:
rpart(formula = prop11_above100 ~ house + SES + elevation + hdens,
data = Arm, method = "anova", minsplit = 15)

Variables actually used in tree construction:
[1] elevation hdens house SES

Root node error: 3.1901/140 = 0.022786

n=140 (37 observations deleted due to missingness)

<table>
<thead>
<tr>
<th>CP</th>
<th>nsplit</th>
<th>rel error</th>
<th>xerror</th>
<th>xstd</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.227950</td>
<td>0.000000</td>
<td>1.02962</td>
<td>0.12347</td>
</tr>
</tbody>
</table>

— 32 —
2 0.074853 1 0.77205 0.99724 0.13953
3 0.031854 2 0.69720 0.89813 0.13324
4 0.031448 3 0.66534 0.97446 0.14345
5 0.029835 4 0.63390 0.98753 0.14607
6 0.025716 5 0.60406 1.00213 0.14641
7 0.018575 6 0.57834 1.01597 0.15162
8 0.015712 7 0.55977 1.03990 0.15537
9 0.013815 8 0.54406 1.03879 0.15541
10 0.012917 9 0.53024 1.03873 0.15594
11 0.010000 10 0.51732 1.04533 0.15679

*** “xerror” (b) lists the cross validation error resulting from the splits. We used 0-SE method to prune the tree.

0-SE method: the cp value of the nsplit that results in the smallest xerror is used as threshold for pruning (see next code). In this example, smallest xerror is 0.89813, corresponding cp value is 0.031854

*** code for pruning the tree

> p11_100 <- prune (prop11_100, cp=0.031854)

*** code for plotting the pruned tree
> plot (p11_100)

*** code for labeling the pruned tree
> text (p11_100, use.n=T)

*** final pruned tree