Peridomestic Infection as a Determining Factor of Dengue Transmission

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Abstract

Background

The study of endemic dengue transmission is essential for proposing alternatives to impact its burden. The traditional paradigm establishes that transmission starts around cases, but there are few studies that determine the risk.

Methods

To assess the association between the peridomestic dengue infection and the exposure to a dengue index case (IC), a cohort was carried out in two Mexican endemic communities. People cohabitating with IC or living within a 50-meter radius (exposed cohort) and subjects of areas with no ICs in a 200-meter radius (unexposed cohort) were included.

Results

Exposure was associated with DENV infection in cohabitants (PRa 3.55; 95%CI 2.37–5.31) or neighbors (PRa 1.82; 95%CI 1.29–2.58). Age, location, toilets with no direct water discharge, families with children younger than 5 and the House Index, were associated with infection. Families with older than 13 were associated with a decreased frequency. After a month since the IC fever onset, the infection incidence was not influenced by exposure to an IC or vector density; it was influenced by the local seasonal behavior of dengue and the age. Additionally, we found asymptomatic infections accounted for 60% and a greater age was a protective factor for the presence of symptoms (RR 0.98; 95%CI 0.97–0.99).
Conclusion

The evidence suggests that dengue endemic transmission in these locations is initially peri-domestic, around an infected subject who may be asymptomatic due to demographic structure and endemicity, and it is influenced by other characteristics of the individual, the neighborhood and the location. Once the transmission chain has been established, dengue spreads in the community probably by the adults who, despite being the group with lower infection frequency, mostly suffer asymptomatic infections and have higher mobility. This scenario complicates the opportunity and the effectiveness of control programs and highlights the need to apply multiple measures for dengue control.

Author Summary

The study of dengue transmission is essential for proposing alternatives to diminish the cases and the cost of dengue treatment and control. The traditional paradigm establishes that transmission chain starts around a case, but there are few studies that determine the risk, therefore, we studied if to live around a dengue case increases the risk to get infected by Dengue virus. We interviewed and took blood samples from people cohabitating with dengue cases and neighbors in two Mexican communities, to compare we interviewed and took blood samples from subjects of areas without dengue cases in these communities. We found that people cohabitating and neighbors had more risk to get infected. Younger and older person, the workers, families with children younger than 5, houses with toilets with no direct water discharge, and areas with more mosquitoes, also had increased infection risk until one month after the fever onset of dengue case. After this month the frequency of dengue infections was only influenced by the seasonal behavior of dengue and the age of the subjects. Also, we found that 60% of infections are asymptomatic and older people have less risk to develop symptoms. This study suggests that dengue transmission in these locations is initially peridomestic, around the houses of infected subject who may be asymptomatic (without symptoms), and it is influenced by other characteristics of the individual, the neighborhood and the community. After this peridomestic transmission, dengue spreads in the community probably by adults who mostly suffer asymptomatic infections and have higher mobility, which complicates the application and affects the results of vector control programs.

Introduction

Dengue is the most important vector-borne viral disease in the world due to its morbidity and economic impact [1]. Because there are no vaccines available to date, the interruption of dengue transmission is based on the vector control. However, despite control programs in endemic countries, the persistence and dispersion of the disease imply a review of the epidemiologic assumptions in which vector control is based on.

It is generally assumed that dengue transmission mainly depends on vector density and although there is an association between dengue incidence and the increase of entomological indices at the regional level, in endemic locations this association is controversial [2,3,4]. In Mexico, the perifocal control, which consists on modifying environments around the home of dengue cases, is the main component of the vector control program and has not proved to
decrease dengue incidence. This may be due to the unsuitable application and vector resistance to pesticides, lack of continuity of control measures or because transmission is not occurring around cases or occurring at a lower rate than expected. The evidence of peridomestic transmission is also controversial; in Thailand, there were more Dengue virus (DENV) infections reported in cohabitants and neighbors of dengue cases compared to neighbors of febrile subjects without dengue, but in Nicaragua, the infection frequency was similar between these groups [5–7].

In addition, perifocal control is based on cases reported to National System of Epidemiologic Surveillance (SINAVE). However, the asymptomatic infections are around to 50% [8–19] and at least 25% of them present viremia [6,11,12,20,21]. Therefore, their contribution to transmission may be considerable for maintaining endemicity and may influence the impact of any vector control measure. Other subject characteristics such as age, intra-locality mobility, some housing characteristics and population density, have also been associated with increased infections [2,22–25].

Identifying factors associated with DENV infection and clarifying dengue transmission pattern in endemic areas would allow to improve prevention and control programs in order to have a greater impact on the dengue burden. In this work we test the hypothesis of dengue transmission occurring mainly in the peridomestic area of Index Cases (ICs) in two Mexican endemic communities; additionally, the relation of some sociodemographic, environmental and vector density variables was determined.

Materials and Methods
Design and study population
We carried out a prospective cohort study in 5 years old (y/o) or older living in Axochiapan and Tepalcingo, Morelos, Mexico. Subjects not susceptible to follow-up were excluded. The study protocol was previously published [26]. The calculated sample size was 1,178 subjects. A summary of the used methods is presented below, mentioning the changes in the protocol.

Recruitment and Follow-up
The exposed cohort included subjects living in the residence of ICs or even in up to 4 neighbor houses within a 50-meter radius. The unexposed cohort included subjects living in areas where no ICs were reported within a 100-meter radius in the 2 months previous to the sampling day; from these areas, up to 5 houses were included in 50-meter radius around the first house where at least one subject accepted to participate. Per each group exposed to an IC (exposed group) an unexposed group was enrolled in the same location during the following three weeks (Tepalcingo, median: 6 day, range: 0–8; Axochiapan, 17.5 days, 2–22 days).

Two visits were carried out, baseline between June and November 2011, and follow-up between August 2011 to March 2012. In each visit, an interview that included the presence of symptoms was performed, and a blood sample was taken in order to measure DENV antibodies, as describe in the protocol [26], but the follow-up visit was made between week 12 and 18 and the active telephonic surveillance was carried out at least once a month. Passive surveillance was also carried out consulting the “Unique Automated System for Epidemiologic Surveillance (SUAVE)”. Variables such as working/studying outside the locality, leaving the locality in the past 15 days and hours of the day staying home in a working day or holiday, and occupation were evaluated. Previous history of DENV exposure (seroprevalence), determined in basal serum by indirect IgG ELISA (Panbio, E-DEN-01G) following manufacturer recommendations, was also considered.
Regarding the vector, each house was geolocated using GPS; a questionnaire was applied and the yard and water containers were inspected at each visit. Home and neighborhood control actions carried out by subjects, the Health Services of Morelos (SSM) or the municipality, and mosquito relative abundance were assessed. At the housing level, larva/pupa home-infections and Container Index were considered. In each group of neighbor houses within a 50-meter radius, the Breteau Index and the House Index, were determined. Environment and weather variables were also measured.

Definition of recent DENV infection

A recent infection by DENV, symptomatic or asymptomatic, was considered whenever IgM or IgG capture ELISA tests resulted positive in any blood sample (Panbio Cat No. E-DENO1M y E-DENO2G). The tests were carried out and interpreted according to manufacturer recommendations. Furthermore, infections were considered recent when suspected dengue case were confirmed by the SSM [26].

Recent infection was subclassified as 1. Pre-enrollment infection when IgM or IgG capture ELISA was positive in the baseline sample or when a subject was confirmed dengue by the SSM. A subject was considered with no pre-enrollment infection when the two capture tests in the baseline sample were negative. We decided to include positive IgG capture subjects as recent infections, because the people who live in these towns show high seroprevalence (>80% in 25 and older), and it has been observed that in secondary infections negative IgM subjects could still develop an IgG response [27, 28]. 2. Post-enrollment Infection when IgM and IgG capture tests were negative in the baseline sample and any test was positive in the following-up sample or the case was confirmed by the SSM (S1 Table). A subject was considered as having no post-enrollment infection when the two capture tests were negative in the baseline and follow-up samples. Otherwise, a symptomatic infection was considered when the subject presented fever in the two months before the baseline evaluation in pre-enrollment infection or when the subject presented at least a fever once at some point during follow-up in post-enrollment infections.

Statistical Analysis

Initially, house locations from Global Positioning System were verified using satellite photography; ICs reported in the SINAVE [29] and not evaluated in the cohort, were located on a map. To verify the house groups and the peridomestic exposure in the area, buffer areas of 200 m, 100 m, and 50 m in diameter were traced, with the center at the IC house, using ArcGIS 10 software. Considering the 200 m buffer, ICs symptoms starting date and unexposed group recruitment date, 5 unexposed groups had to be reclassified as exposed in Axochiapan (Fig 1). The Fig 1 was developed in ArcGIS 10 Software of ESRI, the ArcGIS Software include access to Map Image Services web property of ESRI. The map Image Services includes NASA Blue Marble: Next Generation 500m resolution imagery at small scales (above 1:1,000,000), i-cubed 15m eSAT imagery at medium-to-large scales (down to 1:70,000) for the world, and USGS 15m Landsat imagery for Antarctica. The map also includes i-cubed Nationwide Prime 1m or better resolution imagery for the contiguous United States, Getmapping 1m imagery for Great Britain, and GeoEye IKONOS 1m resolution imagery for Hawaii, parts of Alaska, and several hundred metropolitan areas around the world. I-cubed Nationwide Prime is a seamless, color mosaic of various commercial and government imagery sources, including Aerials Express 0.3 to 0.6m resolution imagery for metropolitan areas and the best available United States Department of Agriculture (USDA) National Agriculture Imagery Program (NAIP) imagery and enhanced versions of United States Geological Survey (USGS) Digital Ortho Quarter Quad
Comparisons were made for both individual and housing models, using the Chi-squared or Mann-Whitney tests. The proportions (pre-recruitment infection prevalence and post-recruitment infection incidence) were analyzed using binomial regression. When there were convergence problems the adjusted measures of association were obtained using Breslow-Cox regression with a constant in the time variable and the robust option [30–32]. Counting of pre-recruitment infections was modeled using Poisson regression. According to the analysis level, an adjustment of standard errors was made considering the 359 houses as cluster units for individual analysis; and, the 91 clusters (houses in 50 meters around IC) for comparison of housing. In such models, only variables based on subject matter knowledge and \( p < 0.20 \) value were evaluated by manual backward elimination, keeping those variables with a \( p < 0.05 \) or those that modified the estimator of IC exposure by more than 10%. Subsequently, eliminated variables during the backward elimination process were evaluated in the resulting model, one by one. Additionally, we tested vector indexes, month of recruitment and family composition. After, this resulting model was evaluated with a multilevel regression, with two and three levels, with random and mixed effects. We use de Akaike’s Information Criterion (AIC) to select the better model. The analysis was made with Stata SE 12.

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Results

Description of ICs

46 ICs were visited (31 in Axochiapan); 45% were men; there was only one case younger than 5; 28.3% were reported as hemorrhagic dengue fever. 95.7% were visited between 11 and 33 days after the fever onset. In 39.1% of the houses, inhabitants had carried out some kind of measure against the vector within the last month, contrasting with those carried out by the SSM (91.3%) or by the local government (69.6%); 15.2% of the houses were infested with Aedes larvae/pupae. The serotype from IC was DENV-1.

Cohort Characteristics

46 exposed groups (15 in Tepalcingo) from the ICs and 44 unexposed groups (15 in Tepalcingo) were selected. 392 houses were visited, 1,893 subjects lived in them, in addition to the ICs. 231 of them were younger than 5; however, 2 of them were included in the cohort because they were pre-enrollment infection diagnosed by the SSM. Out of the 1,662 subjects older than 5, 1,196 were interviewed (71.8%) (Fig 2). No significant differences were present among participants and non-participants (468, 28.2%) regarding location (p = 0.26) and age (p = 0.31), but there were differences for gender (68.7% non-participating men vs. 41.7% of participants, p<0.001). The losses were not differential between exposed and unexposed (5.2% vs 5%, p = 0.854).

After the reclassification of 5 groups, 58.4% of the houses were exposed and 65.6% were located in Axochiapan. The exposition was differential between localities (p = 0.001). In exposed houses a greater frequency of adulticide application due to focal fumigation around ICs was observed. House Index and Breteau Index showed a significant difference (Table 1).

59.5% (712/1,196) of the subjects were exposed. Among the exposed and the unexposed, a significant difference was observed in the health insurance (p = 0.001), the location where the subject lives (p <0.001), fever reports within the last two months and nutritional status (p <0.001) (Table 2). Seroprevalence was 43.5% in subjects 5 to 14 y/o and higher than 92% in older than 35; both globally and per decade, seroprevalence was similar between exposed and unexposed subjects.

Recent DENV Infections in the Cohort

253 recent infections were identified, 99 subjects (39.1%) reported fever sometime during the two months prior to recruitment (pre-enrollment infections) or during follow-up (post-enrollment infections). Out of these, 33 did not seek medical attention (33.3% underreported); and out of those 66 subjects who consulted a doctor, 45 were not reported to the SINAVE (68.2% undernotified). Importantly, no one of the 17 dengue infections that consulted to the private sector were reported to the SINAVE.

60.9% of infections were asymptomatic. A relation between asymptomatic infection and age was observed; younger presented a lower percentage of asymptomatic infections (p<0.001). The median age for asymptomatic infections was 27 y/o (Interquartile Range—IQR 16–54) and for symptomatic infections was 16 y/o (IQR 10–25) (Mann-Whitney, p<0.0001). A greater age was a protective factor for the presence of symptoms (Risk Ratio—RR 0.98; 95%CI 0.97–0.99). Even though recent infections were detected in the all age groups, most were present in
those younger than 30 (Figs 3A and 4A). Recent infection was 27% (182/675) in exposed subjects and 15.4% (71/460) in unexposed subjects (RR 1.73; 95%CI 1.37–2.19). 213 (84.2%) were pre-enrollment and 40 (15.8%) post-enrollment infections.

Pre-enrollment DENV Infections

Present in all age groups, but a greater infection frequency was observed in younger than 30 and older than 64 (Fig 3A and 3B). Out of 392 visited houses, subjects were evaluated in 388 of them, 62.1% (241) did not present any infection, 25.8% (100) presented at least one pre-enrollment infection, 8.8% (34) presented two, 2.3% (9) presented three, 0.5% (2) presented four and 0.5% (2) presented up to five infections. The Prevalence Ratio to IC exposure was 1.96 (95%CI 1.5–2.56). An association was also observed with locality (Axochiapan), age, and the presence of younger than 5 in the houses, but weather variables were not associated. In multivariate analyses, consistency among the used methods for found relationships was observed (Table 3,
Table 1. Description of Houses from the Cohort.

| Characteristic                        | Total (n = 392) | Unexposed (n = 163) | Exposed (n = 229) | p     |
|---------------------------------------|----------------|---------------------|-------------------|-------|
| **Cohabitants**<br>Median (range)     | 5 (1–21)       | 5 (1–15)            | 5 (1–21)          | 0.834 |
| **Children under 5 years of age**<br>n (%) | 163 (41.6)     | 76 (46.6)           | 87 (38)           | 0.087 |
| **Location**<br>Tepalcingo            | 135 (34.4)     | 71 (43.6)           | 64 (28)           | 0.001 |
| Axochiapan                            | 257 (65.6)     | 92 (56.4)           | 165 (72.1)        | 0.111 |
| **Toilet**<br>Direct discharge        | 206 (52.6)     | 80 (49.1)           | 126 (55)          |       |
| Manual discharge                      | 174 (44.4)     | 78 (47.9)           | 96 (41.9)         |       |
| Toilet with no water intake           | 4 (1.0)        | 0                   | 4 (1.8)           |       |
| No toilet available                   | 7 (1.8)        | 5 (3.1)             | 2 (0.9)           |       |
| **Piped water availability**<br>Public network within the house | 300 (76.5) | 130 (79.8) | 170 (74.2) | 0.576 |
| Public network outside the house but within housing grounds | 10 (2.6) | 3 (1.8) | 7 (3.1) | |
| Water from a public faucet or hydrant | 3 (0.8) | 0 | 3 (1.3) | |
| Water from another house              | 3 (0.8)        | 1 (0.6)             | 2 (0.9)           |       |
| Water from a water tank truck         | 4 (1.0)        | 1 (0.6)             | 3 (1.3)           |       |
| Water from a well                     | 72 (18.4)      | 28 (17.2)           | 44 (19.2)         |       |
| **Socioeconomic level**<br>n (%)      |                |                     |                   | 0.211 |
| Low                                   | 131 (33.5)     | 55 (33.7)           | 76 (33.3)         |       |
| Medium                                | 130 (33.3)     | 61 (37.4)           | 69 (30.3)         |       |
| High                                  | 130 (33.3)     | 47 (28.8)           | 83 (36.4)         |       |
| **Mosquito nets in windows**          |                |                     |                   | 0.522 |
| None                                  | 215 (54.9)     | 86 (52.8)           | 129 (56.3)        |       |
| All of them                           | 93 (23.7)      | 38 (23.3)           | 55 (24)           |       |
| Some of them                          | 83 (21.2)      | 38 (23.3)           | 45 (19.7)         |       |
| **Measures against the vector within the last month**<br>Carried out by the subjects | 198 (50.5) | 83 (50.9) | 115 (50.2) | 0.697 |
| House fumigation                      | 114 (29.1)     | 45 (27.6)           | 69 (30.1)         | 0.409 |
| Breeding elimination                  | 62 (15.8)      | 26 (16)             | 36 (15.7)         | 0.489 |
| Other measures<sup>b</sup>            | 81 (20.7)      | 34 (20.9)           | 47 (20.5)         | 0.489 |
| **Carried out by the SSMs**           | 277 (70.7)     | 89 (54.6)           | 188 (82.1)        | <0.001|
| Larvicides                            | 93 (33.6)      | 66 (74.2)           | 27 (14.4)         | <0.001|
| Adulticides                           | 95 (34.3)      | 14 (15.7)           | 81 (43.1)         |       |
| Larvicides and adulticides            | 89 (32.1)      | 9 (10.1)            | 80 (42.6)         |       |
| **Carried out by the Municipality**   | 239 (61)       | 96 (58.9)           | 143 (62.5)        | 0.283 |
| Larvicides                            | 8 (3.4)        | 3 (3.1)             | 5 (3.5)           | 0.117 |
| Adulticides                           | 198 (82.9)     | 73 (76)             | 125 (87.4)        |       |
| Larvicides and adulticides            | 16 (6.7)       | 9 (9.4)             | 7 (4.9)           |       |
| Dejunking                              | 13 (5.4)       | 9 (9.4)             | 4 (2.8)           |       |
| Other combinations of actions         | 4 (1.7)        | 2 (2.1)             | 2 (1.4)           |       |
| **Infested Houses (larvae/pupae)**    | 88 (22.5)      | 44 (27)             | 44 (19.2)         | 0.069 |
| Containers Index<sup>c</sup> Median (IQR) | 0 (0–0%) | 0 (0–5.3%) | 0 (0–0%) | 0.111 |
| **House Index**<sup>d</sup>           | 20 (0–40)      | 20 (0–40)           | 20 (0–33.3)       | 0.0001|
| **Breteau Index**<sup>d</sup>         | 20 (0–50)      | 20 (20–60)          | 20 (0–40)         | 0.0001|

<sup>a</sup>An exposed house did not specify (0.4%) the type of toilet.

<sup>b</sup>Fishes, fumes, mosquito coils, private application of abate, repellent, bed nets.

<sup>c</sup>Containers Index = (# of infested containers/ # of inspected containers with water)x100.

<sup>d</sup>Obtained for each group of neighbor houses within a 50-meter radius. House Index = ([# of positive houses / # inspected houses in the group]x100).

Breteau Index = ( [# of positive containers / # of inspected houses in the group])x100.

IQR: Interquartile Range.
An increased risk was observed with proximity to an IC, locality and younger than 30 and older than 64. The occupation (workers) and the insurance (government health coverage), that were associated in the Breslow-Cox Regression, lost the association in the multilevel analysis (p = 0.059 and p = 0.053, respectively), but were remained in the model because the AIC of this model was lower (Table 3).

Regarding family structure, the presence of younger than 5 was associated with a higher risk of infection, both in the individual and the house model. On the other hand, in the individual

### Table 2. Description of Participating from the Cohort.

| Characteristic                        | Total (n = 1,196) | Unexposed (n = 484) | Exposed (n = 712) | p     |
|---------------------------------------|-------------------|---------------------|------------------|-------|
| **Age**                               |                   |                     |                  |       |
| Median (range)                        | 30 (3–86)         | 31 (5–85)           | 29 (3–86)        | 0.213 |
| **Sex**                               |                   |                     |                  |       |
| Male n (%)                            | 499 (41.7)        | 197 (40.7)          | 302 (42.4)       | 0.555 |
| **Occupation**                        |                   |                     |                  | 0.094 |
| Student                               | 296 (24.8)        | 111 (22.9)          | 185 (26)         |       |
| Student and Employee                  | 37 (3.1)          | 10 (2.1)            | 27 (3.8)         |       |
| Employee                              | 224 (18.7)        | 84 (17.4)           | 140 (19.7)       |       |
| Independent                           | 211 (17.6)        | 84 (17.4)           | 127 (17.8)       |       |
| Housewife                             | 345 (28.9)        | 156 (32.2)          | 189 (26.5)       |       |
| Other                                 | 83 (6.9)          | 39 (8.1)            | 44 (6.2)         |       |
| **Education in older than 6**         | n = 1152          | n = 472             | n = 680          | 0.084 |
| Illiterate                            | 123 (10.7)        | 62 (13.1)           | 61 (9)           |       |
| Reads and Writes                      | 309 (26.8)        | 122 (25.9)          | 187 (27.5)       |       |
| Elementary or Middle School           | 550 (47.7)        | 227 (48.1)          | 323 (47.5)       |       |
| High School or more                   | 170 (14.8)        | 61 (12.9)           | 109 (16)         |       |
| **Insurance**                         |                   |                     |                  | 0.001 |
| Seguro Popular                        | 860 (71.9)        | 378 (78.1)          | 482 (67.7)       |       |
| Uninsured                             | 185 (15.5)        | 66 (13.6)           | 119 (16.7)       |       |
| IMSS: Instituto Mexicano del Seguro Social | 71 (5.9)   | 17 (3.5)            | 54 (7.6)         |       |
| ISSSTE: Instituto de Seguridad y Servicios Sociales de los Trabajadores del Estado. | 68 (5.7) | 20 (4.1) | 48 (6.7) |
| **Location**                          | <0.001            |                     |                  |       |
| Tepalcino                             | 392 (32.8)        | 216 (44.6)          | 176 (24.7)       |       |
| Axochiapan                            | 804 (67.2)        | 268 (55.4)          | 536 (75.3)       |       |
| **Has always lived in Morelos**       | 960 (80.3)        | 395 (81.6)          | 565 (79.4)       | 0.336 |
| **Fever in the last 2 months**        | 201 (16.8)        | 52 (10.7)           | 149 (20.9)       | <0.001|
| Consultation due to fever             | 110 (54.7)        | 27 (51.9)           | 83 (55.7)        | 0.637 |
| Private consultation                   | 29 (26.6)         | 9 (33.3)            | 20 (24.1)        | 0.344 |
| **Nutritional State**                 | 0.026             |                     |                  |       |
| Malnutrition                          | 26 (2.4)          | 11 (2.4)            | 15 (2.3)         |       |
| Normal                                | 431 (39)          | 186 (41.3)          | 245 (37.4)       |       |
| Overweight                            | 336 (30.4)        | 114 (25.3)          | 222 (33.8)       |       |
| Obesity                               | 313 (28.3)        | 139 (30.9)          | 174 (26.5)       |       |
| **Socioeconomic Level**               | 0.007             |                     |                  |       |
| Low                                   | 399 (33.5)        | 157 (32.4)          | 242 (34.1)       |       |
| Medium                                | 402 (33.7)        | 187 (38.6)          | 215 (30.3)       |       |
| High                                  | 392 (32.9)        | 140 (28.9)          | 252 (35.5)       |       |
| **Losses**                            | 61 (5.1)          | 24 (5)              | 37 (5.2)         | 0.854 |
| **Seroprevalence** (positive baseline indirect IgG) | 887/1162 (76.3)  | 362/479 (75.6)      | 525/683 (76.9)   | 0.858 |

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model, the number of cohabitants between 13 and 29 y/o was associated with protection, but in the house model, protection was associated with the 30 to 64 y/o group. Also in individual model, having a toilet with no direct discharge was associated with a higher infection risk. Both in individual and house models, a higher risk was related to vector density, as evaluated with Breteau Index and House Index. The final model included the House Index, as both indexes presented a similar explanation and this one was easier to measure. We observed that with every 10 percent increase in House Index, pre-enrollment infection increased by 10% (Table 3, S2 Table).
Post-enrollment DENV Infections

During follow-up, new infections occurred in almost all age groups (Fig 4A and 4B). Of 359 houses followed, 21 (8.7%) presented at least one infection, 3 (0.8%) presented two, and 1 (0.3%) presented three infections during follow-up. Living in the peridomestic area of an IC did not increase the risk of new infections (RR 0.97; 95%CI 0.53–1.78). In the individual multilevel model, an increased risk infection was observed in younger than 30 (Adjusted Risk Ratio—RRa 2.42; 95%CI 1.16–5.04) compared to the 30 to 64 y/o group; although older than 64 exhibited similar tendency, it was not significative (RRa 2.89; 95%CI 0.99–8.47, p = 0.052). Furthermore, the June-August enrollment period was associated with post-enrollment ingestion.
infections (Individual: RRa 3.82; 95%CI 2.05–7.12 and House: 2.56; 95%CI 1.32–4.98) compared to those enrolled between September–November 2012. Likewise, infection risk increased 17% at the house per each additional family member between 5 and 29 y/o (RRa 1.17; 95%CI 1.01–1.36). When the dependent variable was analyzed as a rate using the Poisson model, the use of adulticides in the baseline evaluation was associated with a lower infection rate in houses

**Table 3. Multilevel Pre-enrollment Infection Models (n = 1,172 subjects; 213 infections).**

| Variable                                              | n     | Infected n (%) | PR     | Model 1 (with individual-level variables) PRa (95% CI) | Model 2 (with contextual-level variables) PRa (95% CI) |
|-------------------------------------------------------|-------|----------------|--------|------------------------------------------------------|------------------------------------------------------|
| Main exposition                                       |       |                |        |                                                      |                                                      |
| Exposure to an Index Case (IC)                        |       |                |        |                                                      |                                                      |
| Unexposed                                             | 479   | 53 (11.1)      | 1      |                                                      |                                                      |
| Lives within 50m of an IC                             | 534   | 107 (20)       | 1.81 (1.33–2.46) | 1.82 (1.29–2.58) |
| Cohabits with an IC                                   | 159   | 53 (33.3)      | 3.01 (2.15–4.21) | 3.55 (2.37–5.31) |
| Individual variables                                  |       |                |        |                                                      |                                                      |
| Age                                                   |       |                |        |                                                      |                                                      |
| <30 years old                                         | 573   | 138 (24.1)     | 2.42 (1.79–3.27) | 2.8 (1.94–4.05) | 2.94 (2–4.34) |
| 30–64 years old                                       | 492   | 49 (10)        | 1      |                                                      |                                                      |
| >64 years old                                         | 107   | 26 (24.3)      | 2.44 (1.59–3.74) | 2.5 (1.53–4.08) | 2.68 (1.64–4.40) |
| Occupation                                             |       |                |        |                                                      |                                                      |
| Student                                               | 320   | 66 (20.6)      | 1      |                                                      |                                                      |
| Non-studying workers                                  | 426   | 74 (17.4)      | 0.84 (0.62–0.14) | 1.39 (0.95–2.02) | 1.45 (0.99–2.14) |
| Housewives and othersa                                 | 426   | 73 (17.1)      | 0.83 (0.62–1.12) | 1.22 (0.84–1.79) | 1.33 (0.90–1.96) |
| Insuranceb                                            |       |                |        |                                                      |                                                      |
| Uninsured                                             | 183   | 45 (24.6)      | 1      |                                                      |                                                      |
| Seguro Popular                                        | 840   | 139 (16.5)     | 0.67 (0.50–0.90) | 0.75 (0.53–1.06) | 0.71 (0.50–1) |
| Other                                                 | 145   | 29 (20)        | 0.81 (0.54–1.23) | 0.97 (0.6–1.57) | 1.01 (0.62–1.65) |
| Contextual variables                                  |       |                |        |                                                      |                                                      |
| Location                                              |       |                |        |                                                      |                                                      |
| Tepalcino                                             | 386   | 49 (12.7)      | 1      |                                                      |                                                      |
| Axochiapan                                            | 786   | 164 (20.9)     | 1.64 (1.22–2.21) | 1.65 (1.18–2.31) |
| Toilet                                                |       |                |        |                                                      |                                                      |
| Direct discharge                                      | 573   | 97 (16.9)      | 1      |                                                      |                                                      |
| Other                                                 | 599   | 116 (19.4)     | 1.14 (0.90–1.46) | 1.38 (1.04–1.83) |
| Younger than 5 at homec                                |       |                |        |                                                      |                                                      |
| No                                                    | 638   | 100 (15.7)     | 1      |                                                      |                                                      |
| Yes                                                   | 534   | 113 (21.2)     | 1.35 (1.06–1.72) | 1.34 (1–1.8) |
| No. of cohabitants between 13 and 29 years oldd        | 2 (1–2)| 1.04 (0.96–1.13) | 0.88 (0.79–0.98) |
| House Index (10 points %)d                            | 20 (0–60) | 1.04 (0.98–1.10) | 1.1 (1.03–1.17) |
| Log-likelihood value                                  |       |                |        |                                                      |                                                      |
| AIC                                                   |       |                |        |                                                      |                                                      |
|                                                      | 1125.13 | 1083.13       |      |                                                      |                                                      |

*aOthers including unemployed, retirees, handicapped persons.

*bInsured status not specified in 4 uninfected subjects.

*cFamily members were categorized: <5 y/o, between 5 and 12, between 13 and 29, between 30 and 64, and older than 64.

*dMedian (Interquartile Range).

PR: Prevalence Ratio. PRa: Adjusted Prevalence Ratio.

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Incidence Rate Ratio 0.50; 95%CI 0.26–0.96); however, this relationship loses statistical significance when adjusting for enrollment month.

Discussion

This is the first study that has estimated the magnitude of the association between a dengue case detected by SINAVE and the occurrence of other peridomestic infections with a follow-up longer than 21 days. The frequency of recent infections around an IC (27%) was similar in Indonesia (24.5%) and in a multicenter study (25.6%), but greater than in Thailand (12.4% and 16%), which may have occurred due to a longer follow-up and the inclusion of older than 15, and in Vietnam (11.9%) because in this study IgM was the only criteria used to define recent infection [5,7,10,21,33]. Our data contrasts with the low infection frequency in Nicaragua (2.4%), where there is probably a lower force of infection [6].

In the current study, we demonstrate that the risk to get infected living around 50 meters of a IC (vicinity), drops to the half comparing with the risk of a cohabitant of a IC, therefore it is highly possible that the risk disappear after a equivalent distance away. Since the estimated flying range of the vector is about 500 meters, and the highest risk is inside the first 50 meters, therefore, the vector is for all that matters static (Table 3). Likewise, in Thailand, a gradient of infection frequency was observed depending on the distance to the IC houses and a higher frequency of DENV infected mosquitoes in IC houses. This findings in small-scale space-time groups support the hypothesis of dengue transmission being predominantly perifocal and that it is likely dengue spreads in a location by the movement of infected humans more than by the movement of infected mosquitoes [7,25,34].

The frequency of recent infections in the unexposed group (15.4%) was greater than the reported in Thailand (0% and 1.1%), Nicaragua (2.5%) and Vietnam (5.1%); this may be explained by the larger follow-up time, the inclusion of older than 15 [5–7,33] or possibly by unexposed group contamination due to suspicious dengue cases that were not diagnosed in Axochiapan (42.5%) and in Tepalcingo (18.7%) or to underreported/undernotified cases (78.8%), especially subjects consulting private clinics, who were not reported to the SINAVE.

Furthermore, age was independently associated with pre-enrollment infection (Fig 3B). Interestingly, the infection rate in older than 64 was similar to younger than 30; this may be because they remain longer where there is infected vector. In line with this interpretation, we found an marginal association with occupation, since, compared to students, workers had a greater infection risk, similar to housewives/other, even though trend were non-significant because of a lack of power. This could occur, because health promotion school campaigns are carried out in these locations; this could make schools places with lower vector density than houses, work places or other destinations. Also, we have the hypothesis that the students, which are younger with higher DENV infection frequency, get infected in their own houses.

Independently living in Axochiapan was also associated with pre-enrollment infection in agreement with the incidence detected by the SINAVE in 2011. Regarding house characteristics, we find a higher risk in those with toilets with no direct discharge, which probably reflects the way water is used and this is related with an increase of potential breedings. We also found an association with the age structure of the family; families had highest risk if they had children younger than 5. We can speculate they need care providers that must spend more time in houses where the transmission is occurring, or these children have a greater frequency of primary infections, which would present higher viremias and consequently they would transmit the infection more efficiently to nearby vectors.

On the other hand, we found asymptomatic infections accounted for 60% of the total, similar to the magnitude determined in Brazil and Thailand [5,13,16,18]. The frequency of
asymptomatic infections increases with age which is in contrast with the observation in Nicaragua where symptomatic subjects were on average 1.2 years older than asymptomatic, but the cohorts have demographic differences [35]. Factors related to asymptomatic infection include the time between infections, infection type (post-secondary, primary or secondary) and dengue incidence during the past year, and we found evidence of some of those in our study [35–37]. A significant proportion of asymptomatic subjects shows a detectable viremia [6,8–21]. Furthermore, it was found that at least 75% of hospitalized and outpatient cases exceed the viremia threshold required to infect 50% of *Aedes aegypti*, despite outpatients presenting lower viremia [38]. Overall, these findings suggest that asymptomatic subjects could participate in DENV transmission. Overall, these findings suggest that asymptomatic subjects could participate in DENV transmission. If asymptomatic subjects are relevant in the transmission the consequences of these on surveillance and control programs will be enormous, hence further prospective studies must be conducted to clarify this point.

We observed that risk to IC exposure disappears over time because post-enrollment infection was not associated with this exposure, similar to observed in Vietnam [33]. Also, vector density was not associated, probably because once transmission starts, it is maintained by a basal vector level. However, post-enrollment infections were associated with the age and dengue local seasonal behavior (enrollment period), and family composition (subjects between 5 and 29 y/o).

We observed that peridomestic transmission is the main determinant of endemic incidence and accordingly we found high frequency of perifocal antivectorial activities and low entomological indexes in the exposed groups, however there was a high frequency of pre-enrollment infections (Table 1), which may be interpreted as effectiveness on vector indexes but with low impact on perifocal transmission. This may be explained by delayed of control measures, because when a case is detected by the SINAVE an accumulation of asymptomatics and unreported/undernotified symptomatics may have previously occurred, which may start and maintain dengue transmission, or the infection is occurring almost simultaneously in the ICs, cohabitants and neighbors. Although, there are a report from Cuba in which the risk to be infected is associated with high House Index and Breteau Index at the vicinity of ICs [4], in general, there is no evidence that perifocal control impacts dengue transmission, in part because the sensitivity of the SINAVE, but also because the participation of asymptomatic subjects [39]. However, considering that had association between pre-enrollment infections and vector density, we propose to strengthen vector control activities that are carried out throughout the year and to conduct specific studies to evaluate the impact of these activities and perifocal activities on local DENV transmission.

The main limitation of this study is that the start of the transmission chain could not be established because pre-enrollment infections may have occurred one or two months before IC. This was because subjects were enrolled to one a month after the IC fever onset, serologic tests can remain positive during this time and most infections were asymptomatic; even then, we could establish a peridomestic risk pattern that will allow to conduct more specific studies to determine how the transmission chain begins.

Taking together all the data, we propose the hypothesis that in these communities, dengue endemic transmission is initially peridomestic for about 3 months, since we detected high recent infection frequency in exposed groups (Table 3); after 3 months the risk became equal in both exposed groups and unexposed groups (RR 0.97; 95%CI 0.53–1.78). As a consequence, the dengue peak during the year is determined by this period of peridomestic transmission. Also, the decrease of infection incidence related to the increase of distance from IC house occurs because the vector is essentially static (Table 3). Consequently, the spread of infection within a community will mainly depend on human mobility (Fig 5). In this sense, we propose
subjects between 30 and 64 years old, despite they probably have lower force of infection, being asymptomatic and economically active, move to daily destinations where they remain long enough to be bitten by nearby vectors, transmitting DENV to other close subjects which go to their homes and start a new peridomestic transmission cluster. On the other hand, young and elderly subjects, even though they probably develop greater force of infection, would have a lower participation in the local DENV dissemination because they have limited mobility, since younger individuals are typically symptomatics while older ones have lower mobility. In this scenario, the neighbors who live inside 50 meter-radio, although they have a lower risk of infection, are comparable in the dispersion of infection with subjects that live in the same house of the ICs, since they are much more than cohabitants. Therefore all the persons inside 50 meter-radio contribute in the dengue dispersion in these endemic communities.

This hypothesis provides an explanation to the relative ineffectiveness of perifocal measures; the opportunity of these measures is limited by the asymptomatic/symptomatic infections, underreported/undernotified cases and the spread of infection by asymptomatic or by a
combination of those factors. Finally, the absence of case notifications from private care centers presents an opportunity for health services to improve the Mexican dengue surveillance.

**Supporting Information**

S1 Checklist. STROBE Checklist.

S1 Table. Laboratory diagnostic of DENV recent infected subjects from the cohort study.

S2 Table. Multilevel Pre-enrollment Infection House Model (n = 388 houses, 91 groups).

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**Author Contributions**

Conceived and designed the experiments: RAMV FADQ JRC. Performed the experiments: RAMV JRC SRP RSL. Analyzed the data: RAMV RDL FADQ JVH RSL SRP PKM JRC. Contributed reagents/materials/analysis tools: RSL SRP. Wrote the paper: RAMV RDL FADQ JVH RSL SRP PKM JRC.

**References**

1. Gubler DJ. The economic burden of dengue. Am J Trop Med Hyg. 2012; 86:743–44. doi:10.4269/ajtmh.2012.12-0157 PMID: 22556068

2. Thammapalo S, Chongsuphatwong V, Geater A, Dueravee M. Environmental factors and incidence of dengue fever and dengue haemorrhagic fever in an urban area, Southern Thailand. Epidemiol Infect. 2008; 136:135–43. PMID: 17359563

3. Honório NA, Nogueira RM, Codeço CT, Carvalho MS, Cruz OG, Magalhães Mde A, et al. Spatial evaluation and modeling of Dengue seroprevalence and vector density in Rio de Janeiro, Brazil. PLoS Negl Trop Dis. 2009; 3:e545. doi:10.1371/journal.pntd.0000545 PMID: 19901983

4. Sanchez L, Vanlerberghe V, Alfonso L, Marquetti Mdel C, Guzman MG, Bisset J, et al. Aedes aegypti larval indices and risk for dengue epidemics. Emerg Infect Dis. 2006; 12:800–6. PMID: 16704841

5. Mammen MP, Pimgate C, Koenraadt CJ, Rothman AL, Aldstadt J, Nisalak A, et al. Spatial and temporal clustering of dengue virus transmission in Thai villages. PLoS Med. 2008; 5:e205. doi:10.1371/journal.pmed.0050205 PMID: 18986209

6. Reyes M, Mercado JC, Standish K, Matute JC, Ortega O, Moraga B, et al. Index cluster study of dengue virus infection in Nicaragua. Am J Trop Med Hyg. 2010; 83:683–89. doi: 10.4269/ajtmh.2010.10-0023 PMID: 20810839

7. Yoon IK, Getis A, Aldstadt J, Rothman AL, Tannitsupawong D, Koenraadt CJ, et al. Fine scale spatio-temporal clustering of dengue virus transmission in children and Aedes aegypti in rural Thai villages. PLoS Negl Trop Dis. 2012; 6:e1730. doi:10.1371/journal.pntd.0001730 PMID: 22816001

8. Yew YW, Ye T, Ang LW, Ng LC, Yap G, James L, et al. Seroepidemiology of dengue virus infection among adults in Singapore. Ann Acad Med Singapore. 2009; 38:667–75. (pdf) PMID: 19736569

9. Rodríguez Rodríguez D, Garza Rodríguez M, Chavarria AM, Ramos-Jiménez, Rivera MA, Taméz RC, et al. Dengue virus antibodies in blood donors from an endemic area. Transfus Med. 2009; 19:125–31. doi: 10.1111/j.1365-3148.2009.00922.x PMID: 19566669

10. Dussart P, Baril L, Petit L, Beniguel L, Quang LC, Ly S, et al. Clinical and virological study of dengue cases and the members of their households: the multinational DENFRAME Project. PLoS Negl Trop Dis. 2012; 6:e1482. doi:10.1371/journal.pntd.0001482 PMID: 22232098
11. Xu G, Dong H, Shi N, Liu S, Zhou A, Cheng Z, et al. An outbreak of dengue virus serotype 1 infection in Cixi, Ningbo, People’s Republic of China, 2004, associated with a traveler from Thailand and high density of Aedes albopictus. Am J Trop Med Hyg. 2007; 76:1182–88. PMID: 17556633
12. Méndez F, Barreto M, Arias JF, Rengifo G, Muñoz J, Burbano ME, et al. Human and mosquito infections by dengue viruses during and after epidemics in a dengue-endemic region of Colombia. Am J Trop Med Hyg. 2006; 74:678–83. PMID: 16607005
13. Vanwambekke SO, van Benthem BH, Khantikul N, Burghoom-Maas C, Panart K, Oskam L, et al. Multi-level analyses of spatial and temporal determinants for dengue infection. Int J Health Geogr. 2006; 5:5. PMID: 16420702
14. Porter KR, Beckett CG, Kosasih H, Tan RI, Alisjahbana B, Rudiman PI, et al. Epidemiology of dengue and dengue hemorrhagic fever in a cohort of adults living in Bandung, West Java, Indonesia. Am J Trop Med Hyg. 2005; 72:60–6. PMID: 15728868
15. Rodrigues EM, Dal-Fabbro AL, Salomao R, Ferreira IB, Rocco IM, Fonseca BA. Epidemiologia da infecção pela dengue em Ribeirão Preto, SP, Brasil. Rev Saude Publica. 2002; 36:160–65. PMID: 12045796
16. Endy TP, Chunsuttiwat S, Nisalak A, Libraty DH, Green S, Rothman, et al. Epidemiology of inapparent and symptomatic acute dengue virus infection: a prospective study of primary school children in Kamphaeng Phet, Thailand. Am J Epidemiol. 2002; 156:40–51. PMID: 12076887
17. Vásconcelos PF, Lima JW, da Rosa AP, Timbo MJ, da Rosa ES, Lima HR, et al. Epidemia de dengue em Fortaleza, Ceará: Inquérito soro-epidemiológico aleatório. Rev Saude Publica. 1998; 32:447–54. PMID: 10030061
18. Da Cunha RV, Dias M, Nogueira RM, Chagas N, Miagostovich MP, Schatzmayr HG. Secondary dengue infection in schoolchildren in a dengue endemic area in the state of Rio de Janeiro, Brazil. Rev Inst Med Trop Sao Paulo. 1995; 37:517–21. PMID: 8731265
19. Vásconcelos PF, Travassos da Rosa ES, Travassos da Rosa JF, de Freitas RB, Dégallier N, Rodrigues SG, et al. Epidemia de febre clássica de dengue causada pelo sorotipo 2 em Araguaina, Tocantins, Brasil. Rev Inst Med Trop Sao Paulo. 1993; 35:141–48.
20. Chevillon C, Failloux A-B. Questions on viral population biology to complete dengue puzzle. TRENDS in Microbiol. 2003: 11:415–21.
21. Beckett CG, Kosasih H, Faisal I, Nurhayati, Tan R, Widjaja S, et al. Early detection of dengue infections using cluster sampling around index cases. Am J Trop Med Hyg. 2005; 72:777–82. PMID: 15967759
22. Siqueira JB, Martelli CM, Maciel IJ, Oliveira RM, Ribeiro MG, Amorim FP, et al. Household survey of dengue infection in central Brazil: spatial point pattern analysis and risk factors assessment. Am J Trop Med Hyg. 2004; 71:646–51. PMID: 15569799
23. Braga C, Luna CF, Martelli CM, Souza WV, Cordeiro MT, Alexander N, et al. Seroprevalence and risk factors for dengue infection in socioeconomically distinct areas of Recife, Brazil. Acta Trop. 2010; 113:234–40. doi: 10.1016/j.actatropica.2009.10.021 PMID: 19896921
24. Stoddard ST, Morrison AC, Vazquez-Prokopec GM, Paz Soldan V, Kochel TJ, Kitron U, et al. The role of human movement in the transmission of vector-borne pathogens. PLoS Negl Trop Dis. 2009; 3: e481. doi: 10.1371/journal.pntd.0000481 PMID: 19621090
25. Adams B, Kapan DD. Man bites mosquito: understanding the contribution of human movement to vector-borne disease dynamics. PLoS One 2009; 4:e6763. doi: 10.1371/journal.pone.0006763 PMID: 19707544
26. Martínez-Vega RA, Danis-Lozano R, Velasco-Hernández J, Díaz-Quijano FA, González-Fernández M, Santos R, Román S, et al. A prospective cohort study to evaluate peridomestic infection as a determinant of dengue transmission: protocol. BMC Public Health. 2012; 12:262. doi: 10.1186/1471-2458-12-262 PMID: 22471857
27. Amaya-Larios IY, Martínez-Vega RA, Mayer SV, Galeana-Hernández M, Comas-García A, Sepúlveda-Salinas KJ, et al. Seroprevalence of neutralizing antibodies against dengue virus in two localities in the state of Morelos, Mexico. Am J Trop Med Hyg. 2014; 91:1057–65. doi: 10.4269/ajtmh.14-0145 PMID: 25294613
28. Sa-Ngasang A, Anantapreecha S, A-Nuegoonpipat A, Chanama S, Wibulwattanakij S, Pattanakul K, et al. Specific IgM and IgG responses in primary and secondary dengue virus infections determined by enzyme-linked immunosorbent assay. Epidemiol Infect. 2006; 134:820–5. PMID: 16371180
29. Sistema Nacional de Vigilancia Epidemiológica [Internet]. México: Dirección General de Epidemiología. http://www.sinave.gob.mx
30. Lee J, Chia KS. Estimation of prevalence rate ratios for cross sectional data: an example in occupational epidemiology. Br J Ind Med. 1993; 50:861–64. PMID: 8398881
31. Nijem K, Kristensen P, Al-Khatib A, Bjertness E. Application of different statistical methods to estimate relative risk for self-reported health complaints among shoe factory workers exposed to organic solvents and plastic compounds. Nor Epidemiol. 2005; 15:111–16.

32. Diaz-Quijano FA. A simple method for estimating relative risk using logistic regression. BMC Med Res Methodol. 2012; 12:14. doi: 10.1186/1471-2288-12-14 PMID: 22335836

33. Anders KL, Ng Le H, Thuy NT, et al. Households as foci for dengue transmission in highly urban Vietnam. PLoS Negl Trop Dis. 2015; 9:e0003528. doi: 10.1371/journal.pntd.0003528 PMID: 25680106

34. Smith DL, Perkins TA, Reiner RC Jr, Barker CM, Niu T, Chaves LF, et al. Recasting the theory of mosquito-borne pathogen transmission dynamics and control. Trans R Soc Trop Med Hyg. 2014; 108:185–97. doi: 10.1093/trstmh/tru026 PMID: 24591453

35. Montoya M, Gresh L, Mercado JC, Williams KL, Vargas MJ, Gutierrez G, et al. Symptomatic versus inapparent outcome in repeat dengue virus infections is influenced by the time interval between infections and study year. PLoS Negl Trop Dis. 2013; 7:e2357. doi: 10.1371/journal.pntd.0002357 PMID: 23951377

36. Olkowski S, Forshey BM, Morrison AC, Rocha C, Vilcarromero S, Halsey ES, et al. Reduced risk of disease during postsecondary dengue virus infections. J Infect Dis. 2013; 208:1026–33. doi: 10.1093/infdis/jit273 PMID: 23776195

37. Grange L, Simon-Loriere E, Sakuntabhai A, Gresh L, Paul R, Harris E. Epidemiological risk factors associated with high global frequency of inapparent dengue virus infections. Front Immunol. 2014; 5:280. doi: 10.3389/fimmu.2014.00280 PMID: 24966859

38. Nguyet MN, Duong TH, Trung VT, Nguyen TH, Tran CN, Long VT, et al. Host and viral features of human dengue cases shape the population of infected and infectious Aedes aegypti mosquitoes. Proc Natl Acad Sci U S A. 2013; 110:9072–77. doi: 10.1073/pnas.1303395110 PMID: 23674683

39. Barmak DH, Dorso CO, Otero M, Solari HG. Modelling interventions during a dengue outbreak. Epidemiol Infect. 2014; 142:545–61. doi: 10.1017/S0950268813001301 PMID: 23800514