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RESEARCH ARTICLE

Long-range movements coupled with heterogeneous incubation period sustain dog rabies at the national scale in Africa

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Abstract

Dog-transmitted rabies is responsible for more than 98% of human cases worldwide, remaining a persistent problem in developing countries. Mass vaccination targets predominantly major cities, often compromising disease control due to re-introductions. Previous work suggested that areas neighboring cities may behave as the source of these re-introductions. To evaluate this hypothesis, we introduce a spatially explicit metapopulation model for rabies diffusion in Central African Republic. Calibrated on epidemiological data for the capital city, Bangui, the model predicts that long-range movements are essential for continuous re-introductions of rabies-exposed dogs across settlements, eased by the large fluctuations of the incubation period. Bangui's neighborhood, instead, would not be enough to self-sustain the epidemic, contrary to previous expectations. Our findings suggest that restricting long-range travels may be very efficient in limiting rabies persistence in a large and fragmented dog population. Our framework can be applied to other geographical contexts where dog rabies is endemic.

Author summary

Rabid dogs constitute the main source of rabies transmission to humans, therefore understanding disease dynamics in dogs is essential to reduce human risk. Mathematical models are invaluable in exploring intervention scenarios where field experiments can be expensive or impractical. We developed a data-driven model to identify the mechanisms for rabies dispersal, accounting for dogs' spatial fragmentation, human-mediated movements, and empirically estimated fluctuations of the incubation period in dogs. Applied to Central African Republic, our model indicates that continuous re-introductions of rabid dogs via human-mediated movements are critical in sustaining the disease in the country. Long-range travel restrictions can be very effective in aiding rabies control. Our approach can be extended to other geographical settings.

Competing interests: The authors have declared that no competing interests exist.

Introduction

Canine rabies is a viral zoonosis responsible for approximately 21,500 (95% C.I. 9,100–58,100) annual human deaths in Africa [1]. Almost all cases result from bites from infected dogs, thus establishing domestic dogs as the primary vector for human rabies in the developing world [2,3]. Despite the existence of effective and efficient preventive measures [4–7], restricted availability [8,9], cost [8–10], cultural opposition [9,11] and lack of policy coordination hinder their full implementation in resource-limited countries [12–14]. As a result, rabies remains an endemic zoonosis throughout Africa, affecting individuals in urban areas as well as in remote or rural locations and continues to represent nowadays a major public health concern [2].

As rabid dogs constitute the main source of transmission to humans, understanding the dynamics of rabies in domestic dog populations is thus essential to identify the key drivers for pathogen maintenance and develop improved control measures to reduce spillover opportunities and, consequently, the number of human fatalities. An important indicator of the transmission potential of a disease is the basic reproductive number, R_0 , measuring the average number of secondary cases generated by a single infectious individual in a fully susceptible population [15,16]. Analyses of outbreak episodes based on surveillance of rabid dog cases reported rather low values of the transmission potential, with R_0 estimates just above the critical value of 1, both in rural and urban areas in Africa [5,17–19]. Despite the low-level endemic stability that would require relatively low degrees of vaccination coverage to eliminate rabies [20], empirical and theoretical evidence suggests that sustained pulsed vaccination campaigns with 70% coverage of the domestic dog population are needed to achieve long-term elimination [17,21,22]. This high coverage is required to prevent the rapid decline of population herd immunity below its critical value between two consecutive campaigns, induced by the large renewal of the naive domestic dog population due to dogs' short life-span and high fecundity rate [19,23].

However, the planning of intervention is made difficult by the geographical dissemination of the disease that may cause rabies reemergence in disease-free areas [19,23–25]. Rabies spatial dispersal is mainly ascribed to human-mediated transport of dogs [24,26–29]. Free-roaming movements of domestic dogs are indeed limited to very short distances of few kilometers per day [30], or less for rabid dogs [17]. Human-mediated movements were previously recognized to be an important vector for spatial dispersal of diseases in wildlife [31–33], domestic [26,27] and farmed animals [34]. For canine rabies, available evidence from household survey data indicates the rather common practice of transporting dogs with frequent movements (daily, monthly, and annual scales) even over large distances (e.g. between different provinces of a country) [29,35,36]. This may compromise vaccination campaigns unable to maintain high coverage over the entire region [24,29]. The key role played by humans in the dispersal of rabies is also evident from the observed relation between rabies virus spatial spread and the geographic location of human settlements, administrative borders, roads and bridges overcoming natural barriers [26,32,37–39].

Transmission of rabies occurs mainly through the bite of an infected animal, with inoculation of viral-loaded saliva in the subcutaneous or muscular tissue of a susceptible host [40,41]. The incubation period is highly variable, with a median close to one month, and a range that can extend from 10 days to more than one year [17]. This is likely related to the process of dissemination of the virus in the infected body, migrating from the site of inoculation to the central nervous system [42,43]. The infectious period is instead rather short, with a median around 3 days and fluctuations typically not exceeding 4 days. While dog movements are extremely restricted during the infective phase because of its duration and of disease

symptoms, infected dogs may move across distant locations while incubating the virus, potentially carrying the disease over large distances, as suggested by phylogeographic data [26,27,35,39].

Several models were introduced to address the role of rabies reintroduction in disease-free areas through importations of rabies-exposed dogs [23,24,29,44–46]. Their findings are however specific to the characteristics of the public health and veterinary health interventions adopted (e.g. frequency of mass vaccination, coverage, movement bans, etc.). An endemic setting where no intervention has been introduced yet offers an ideal scenario to characterize the endemic disease dynamics at the spatial level, being it isolated from the effects resulting from disease control. It may thus provide the opportunity to better understand the underlying key mechanisms to guide the implementation of elimination measures. Central African Republic reported an endemic circulation of canine rabies for at least the last 20 years [18,47–49], with no mass vaccination campaign of domestic dog population implemented during the period. Long-term surveillance data on rabid-dog and exposed-human cases are available for the capital city of Bangui and showed that almost all exposed individuals (93%) are from Bangui or its immediate neighborhood in the suburban areas of Bimbo and Bégoua [47]. Analyses of epidemiological time series and viral sequence data indicated that the epidemiology of rabies virus in Bangui's dog population is shaped by a sequence of successive waves characterized by periods of 53 and 89 months and long intervals (>1 year) of absence of cases [18]. This was associated to local extinction coupled with the introduction of new viral subtypes. Therefore, while canine rabies does not appear to self-sustain in the city of Bangui, its persistence was suggested to be related to frequent introductions of rabies-exposed dogs in the capital city from its neighboring area [18], a transmission pattern similar to those reported also in other African settings [17,19,25].

Here we investigate the factors underlying the observed rabies epidemiology in Bangui through a spatially explicit stochastic metapopulation model parameterized on the geography of Central African Republic and calibrated on 20 years epidemiological data from Bangui. We consider empirical distributions of incubation and infectious periods coupled with mapping of dog population in the country and models of human-mediated movements to study the local vs. long-range spatial transmission, accounting for both urban and rural settlements.

Methods

Metapopulation model structure

We develop a georeferenced discrete stochastic metapopulation epidemic model [50,51] for Central African Republic where domestic dog habitats are represented by model patches and human-mediated movements of dogs are represented by links connecting different patches. The infection dynamics occurs within each patch, and the disease can spread from one patch to another by means of dog movements.

Spatially distributed dog populations in the country are inferred from georeferenced high-resolution human demographic data [52–54], similarly to [45]. We used a publicly available high-resolution dataset of human population distribution in space [55] to map the georeferenced human settlements in the country and the associated population size, population density, and spatial extension. Population data is validated on Bangui population size provided by national census. The resulting human settlements in space correspond to the dog habitats considered in the metapopulation model, i.e. patches. They are classified as *urban* or *rural*, depending on the density of the corresponding human settlements. We follow the OECD (Organisation for Economic Co-operation and Development) definition using a threshold of 1,000 individuals/km² to define a urban settlement [56]. Dog population size in each patch is

computed from the human-to-dog ratio for *urban* ($>1,000$ individuals/km², ratio = 21.20) or *rural* ($<1,000$ individuals/km², ratio = 7.40) settings [53].

Human-mediated movements of dogs from one patch to another are modeled with a geographical distance model consistent with known patterns of human movements [57–60] and compatible with the spatial spread of canine rabies [35]. Similarly to [44], we define the number of dogs transported from one settlement to another to be proportional to $1/d$, where d is the great circle distance between the centroids of the two patches. Movements are divided into three different ranges, similarly to [29]: *short-range* movements ($d < 20$ km) correspond to the average distance within a municipality, *medium-range* movements ($20 \leq d \leq 100$ km) correspond to the average distance traveled to connect two municipalities within the same region, and *long-range* movements ($d > 100$ km) correspond to the remaining ones.

Infection dynamics and spatial spread

We model rabies virus transmission through mixing between individual hosts within the patches, and spatial dissemination through the explicit movements of discrete infected dogs between patches. Viral infection dynamics in dogs is described with a susceptible-exposed-infectious-removed (SEIR) compartmental model [15,16], as previously done for dog rabies [23,24,29,44–46,61], where hosts can be in one of the following states: susceptible (S), i.e. healthy individuals who may acquire the infection; exposed (E) hosts who have contracted the infection but do not shed the virus during an average incubation period ε^{-1} ; infectious (I) hosts who can transmit the virus for an average infectious period μ^{-1} following incubation; hosts removed (R) from the population as rabies is fatal once clinical symptoms appear [40]. More details are provided in the [S1 Text](#). Accounting for the behavioral changes induced by the disease [17], rabid dogs are assumed to be restricted to the patch in which they are located at symptoms onset, whereas exposed dogs can migrate to different locations while incubating the virus according to the defined mobility model. The metapopulation model is parameterized with the empirical distributions for the duration of the incubation and infectious periods obtained for rabies virus in African domestic dog population [17]. To assess the role of the large fluctuations reported for the incubation period, we also consider another version of the compartmental model where disease progression occurs with constant rates, i.e. with exponentially distributed disease stages with the same average duration of incubation and infection, as done in [5,23,44,45]. We explore a range of values of the basic reproductive number ($1.01 \leq R_0 \leq 1.16$) consistent with previous estimates [5,17–19]. Vital dynamics is considered in the model to account for dogs' high fecundity rate leading to a renewal of susceptibles and their relatively short average life span. Birth rate is found to vary drastically across different geographical areas [17,23,36,52,62–65], therefore we considered it as an unknown parameter of the model to be explored. Average life expectancy is parameterized with available estimates for domestic dogs in Africa (see [S1 Text](#)).

Numerical simulations and analyses

We perform numerical simulations of rabies virus transmission in the modelled dog population. Simulations are discrete and stochastic to account for the discrete nature of hosts and for stochastic extinction events that may be favored by small host population sizes. Time is considered to be discrete with a daily timescale. Since the disease is endemic in the country [47], simulations are seeded in each patch with a proportion of exposed and infective dogs computed from surveillance data on rabid-dog cases in the capital city of Bangui [18], considering a detection rate of 20% as in [18]. Other initial conditions are explored in the [S1 Text](#).

We considered different scenarios of transmission and birth rate. For each model parameterization and under each hypothesis considered, we run 10^3 stochastic simulations for a long

enough period (>300 years) to evaluate the occurrence of an endemic condition for rabies virus diffusion.

Simulations provide at each time step the number of domestic dogs in each compartment in each patch, and of those moving from one patch to another. We compute the persistence probability of rabies virus in the domestic dog population as the fraction of stochastic simulations reaching the endemic condition (i.e. pathogen survival in the dog population). Endemic prevalence per patch is computed as the average prevalence in time, excluding the initial transient period (~10 years), over each simulation run where rabies virus continued circulating in the dog population.

To estimate the reproductive number R_0 and the birth rate, we fitted the metapopulation model to the epidemiological situation reported for Bangui. We used a Monte Carlo procedure to evaluate the likelihood, similar to what previously done in [66,67]. More details are reported in the [S1 Text](#).

To estimate the degree of urbanization of rabies epidemic in Central African Republic compared to its dispersion in rural regions, we plot the national-level epidemic concentration curve (ECC) [68] by ranking patches according to increasing density ϕ of rabid dogs ($\phi_1 \leq \phi_2 \leq \dots \leq \phi_N$) and computing the indicator $c_j = \frac{100}{\sum_i \phi_i} \sum_{i \geq j} \phi_i$.

The metapopulation framework is implemented in C++, and technical details for simulations are reported in the [S1 Text](#). A sensitivity analysis is performed to assess the impact of numerical choices.

Role of spatial fragmentation and human-mediated mobility

We numerically assess the impact of the landscape of human settlements in Central African Republic and of human-mediated movements on the persistence of canine rabies in the country by comparing our findings with synthetic scenarios altering the spatial structure of the dog population and/or its mobility, on the full parameter space explored. The scenarios considered are the following:

- Only Bangui: Bangui is considered as an isolated patch, the rest of the metapopulation model is ignored;
- Only Bangui neighborhood: transmission is considered exclusively in Bangui's patch and its immediate neighborhood (<20km), assuming these patches are isolated from the other settlements;
- Only urban patches: the metapopulation model is constituted exclusively of urban patches; rural patches and links departing from them are removed;
- Only rural patches: the metapopulation model is constituted exclusively of rural patches; urban patches and links departing from them are removed.
- Only short travels: all movements except short travels are restricted;
- Only medium travels: as above for medium travels;
- Only long travels: as above for long travels;
- Only short + medium travels: long travels are restricted;
- Only short + long travels: medium travels are restricted;
- Only medium + long travels: short travels are restricted;

Additional scenarios (e.g. the whole country without the capital Bangui) are considered in the [S1 Text](#).

Results

Domestic dogs' spatial demographics and human-mediated movements

Dog population in Central African Republic is estimated to be of approximately 80,000 animals distributed in a rather dispersed spatial pattern comprising 137 settlements divided in 58 (42%) urban and 79 (58%) rural patches ([Fig 1A](#)). Only 10% of the total dog population lives in rural patches, whereas almost 50% is located in Bangui alone. Other top populated urban areas are Berbérati and Carnot in the western region of the country, accounting for 4% and 2.5% of the total dog population, and Bambari (3%) in the center-east.

Settlements are mainly gathered in the center and western regions of the country, with very few urban areas in the eastern region where national parks and natural reserves are located. Most rural settlements (>70%) have an estimated population size of less than 100 dogs, whereas the number of dogs in urban patches is rather heterogeneous, ranging from few tens up to the largest size constituted by the population in Bangui (almost 40,000 dogs, [Fig 1B](#)). Low populated urban patches are mostly geographically located in the proximity of the main cities. While dog population sizes change considerably between rural and urban settlements,

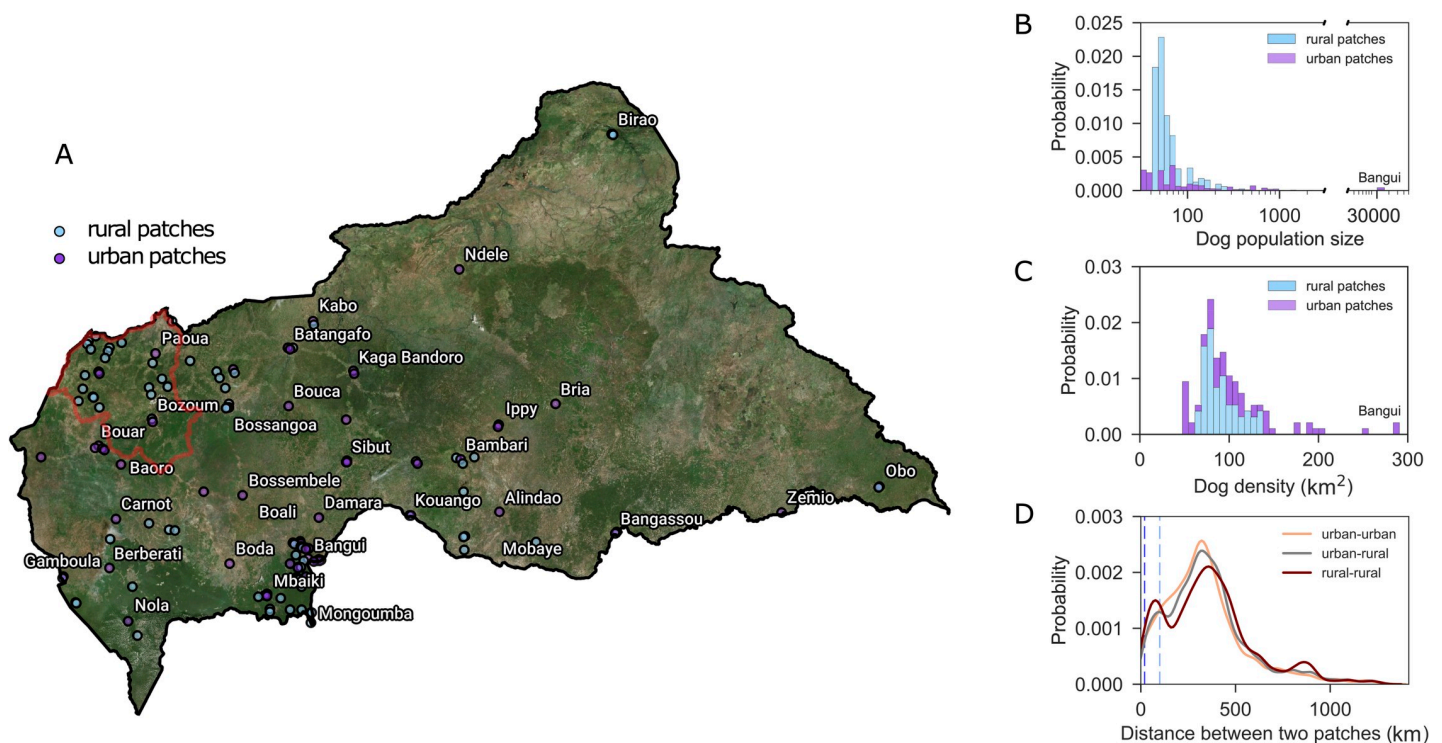


Fig 1. Modeled domestic dog population in Central African Republic. **A)** Geographical distribution of the predicted dog subpopulations divided in rural (light blue) and urban (violet) patches. The red shaped area indicates the Ouham-Pendé prefecture. **B)** Probability distribution of the estimated dog population size per patch in rural (light blue) and urban (violet) patches. **C)** Probability distribution of the estimated dog population density per km² per patch in rural (light blue) and urban (violet) patches. **D)** Probability distribution of the distance between any two patches in the metapopulation network, whether connecting two urban patches (orange), two rural patches (dark red), a urban with a rural patch (black). The two dashed vertical lines indicate the separation between short, medium, and long-range movements. The satellite map was generated using data sourced from OpenStreetMap (OpenStreetMap contributors) and created through QGIS software (QGIS Development Team (2019). QGIS Geographic Information System. Open Source Geospatial Foundation Project. <http://qgis.osgeo.org>).

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their density per squared km are predicted to be rather similar, except for few urban patches with an estimated density of >150 dogs/km² (Fig 1C).

Given the country's geography and population distribution, estimated human-mediated transport of dogs range from few kilometers for neighboring settlements to more than 1,000 km connecting far away locations (Fig 1D). The most probable distance traveled is predicted to be approximately 300 km for long-range movements connecting two urban patches or urban and rural patches, and it is slightly bigger (~350 km) for connections between two rural settlements. The latter type of movements displays in addition two marked peaks. The first is estimated at around 100 km and corresponds to a high spatial concentration of close rural habitats such as around the capital city or in the Ouham-Pendé prefecture in the northwest of the country (Fig 1A). The second is found for distances of approximately 850 km, typically connecting rural areas located in the northwest with the most populated areas in the center of the country (Fig 2A).

Multiple scales of connectivity are predicted by the mobility model for human-mediated movements of dogs (Fig 2). While long-range links are estimated to ensure the overall connectivity of the metapopulation, medium-range transports characterize mobility in the prefectures of Bangui and Ouham-Pendé in the north-west. Short-range transports move the largest

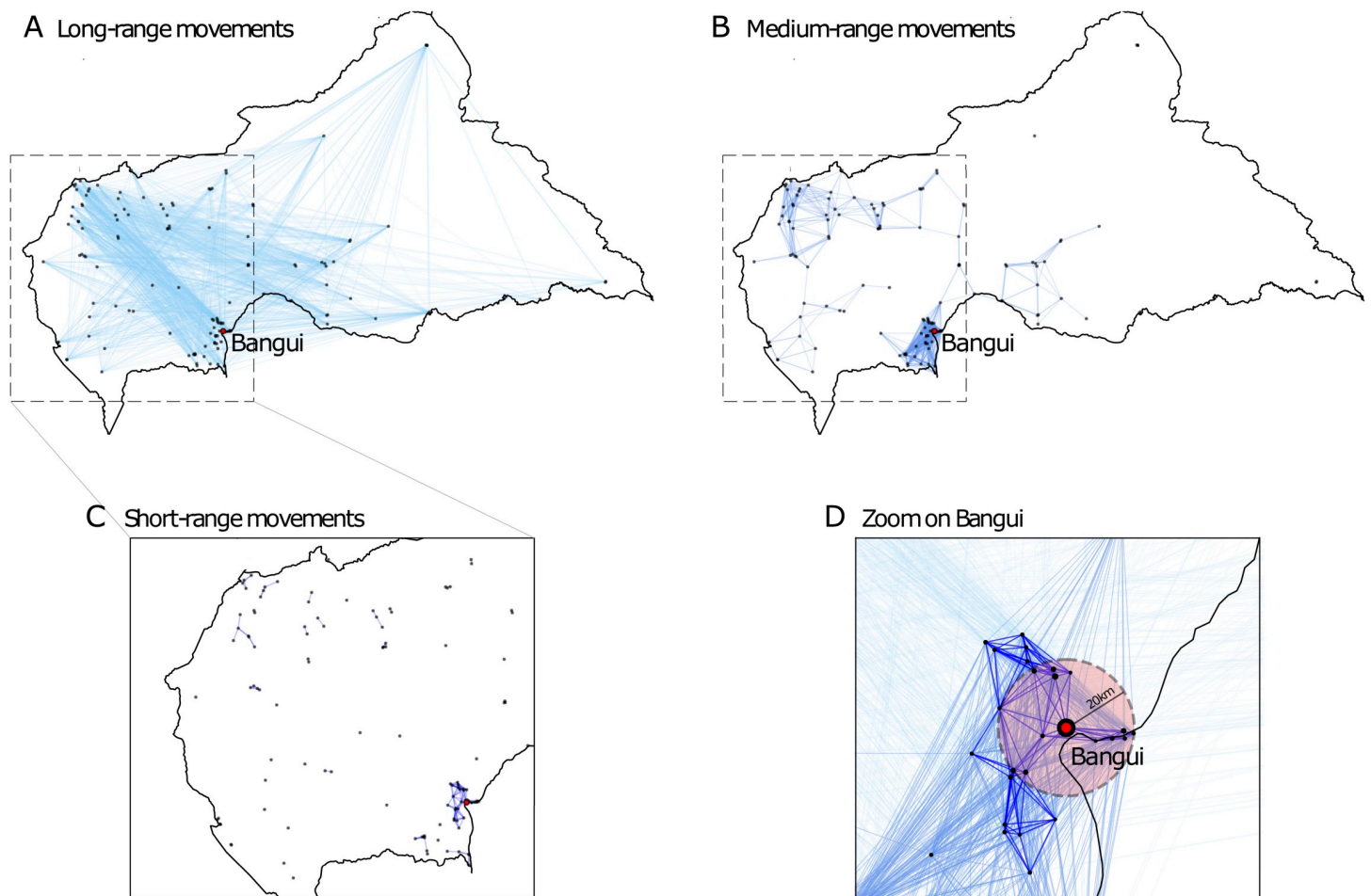


Fig 2. Georeferenced multiscale network of estimated domestic dog mobility. Dots represent patches and links represent the estimated human-mediated transports of dogs on long A), medium B), and short scales C). D) Bangui and its surrounding region. The red circle defines Bangui's neighborhood. The Central African Republic borders were generated using data from "Global Administrative Areas (2019)". University of California, Berkely. Available online: <http://www.gadm.org> [21/07/2019].

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Table 1. Modeled dog movements in Central African Republic. For each movement range we report the predicted number of links (and percentage of the total), and the predicted daily moving rate (dog/day).

Movement range	Estimated number of links (%)	Estimated % population moving daily
short (0–20 km)	183 (2%)	0.035 (95% CI: 0.003, 0.2)
medium (20–100 km)	951 (10%)	0.0046 (95% CI: 0.0001, 0.0146)
long (20 km–CAR borders)	8,182 (88%)	0.00064 (95% CI: 0.00002, 0.0025)

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number of dogs (estimated average of 0.035% of the dog population per day, Table 1) and provide connectivity to the local communities (40% of them are predicted to be located around Bangui).

Rabies virus persistence

Rabies virus is predicted to remain endemic with high probability ($\geq 80\%$) in Central African Republic for $R_0 \geq 1.01$ and a large range of birth rate values (Fig 3A); moreover, except for very low values of the basic reproductive number ($R_0 \leq 1.02$), persistence probability remains rather constant for increasing R_0 . Birth rates smaller than ≈ 1 dog/year lead to considerably rare maintenance of the virus, and a rapid transition toward high probability is predicted for increasing birth rates (Fig 3B). These findings are obtained parameterizing the infection dynamics with empirically estimated distributions of incubation and infectious periods. If instead we assume exponentially distributed disease stages, rabies virus would not be maintained in the domestic dog population, unless for higher values of the basic reproductive number ($R_0 \geq 1.12$, Fig 3C).

We fitted parameters to the epidemiological surveillance data from Bangui estimating R_0 equal to 1.03 (95% CI: 1.02–1.04) and birth rate equal to 1.59 (95% CI: 1.19–1.99). The monthly number of rabid dogs predicted in the city of Bangui ranges from zero to a few tens (Fig 4A), consistently with previous observations [18]. Wavelet analysis [69] was used to identify the periodicity of the numerical trajectories in Bangui and compare them with the estimates obtained from surveillance data in the city [18]. Numerically predicted epidemic waves in Bangui are characterized by a full spectrum of periodicities with the highest probability reached for the empirically estimated period of 89 months (Fig 4B). The distribution of the

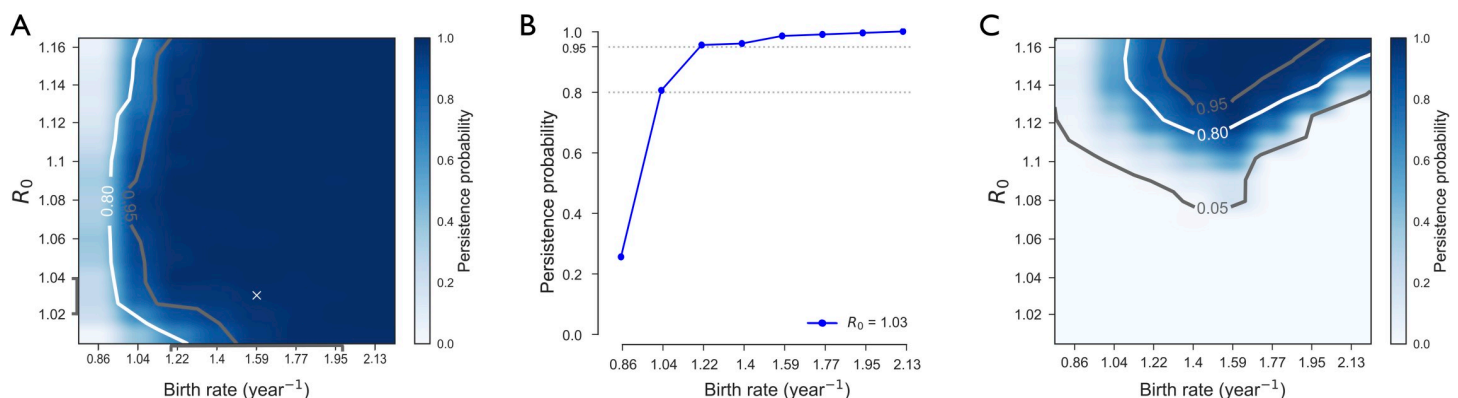


Fig 3. Dog rabies persistence probability in Central African Republic. A) Predicted persistence probability of rabies in the domestic dog population of Central African Republic as a function of the basic reproductive number R_0 and of the annual birth rate. Results are obtained using empirically distributed incubation and infectious periods. The symbol 'x' corresponds to the maximum likelihood estimate. B) Predicted persistence probability as a function of the annual birth rate for the maximum likelihood estimate $R_0 = 1.03$. The two dotted lines indicate 80% and 95% persistence probability. C) As in A) assuming that incubation and infectious periods are exponentially distributed with the same average duration of incubation and infection of the empirical distributions.

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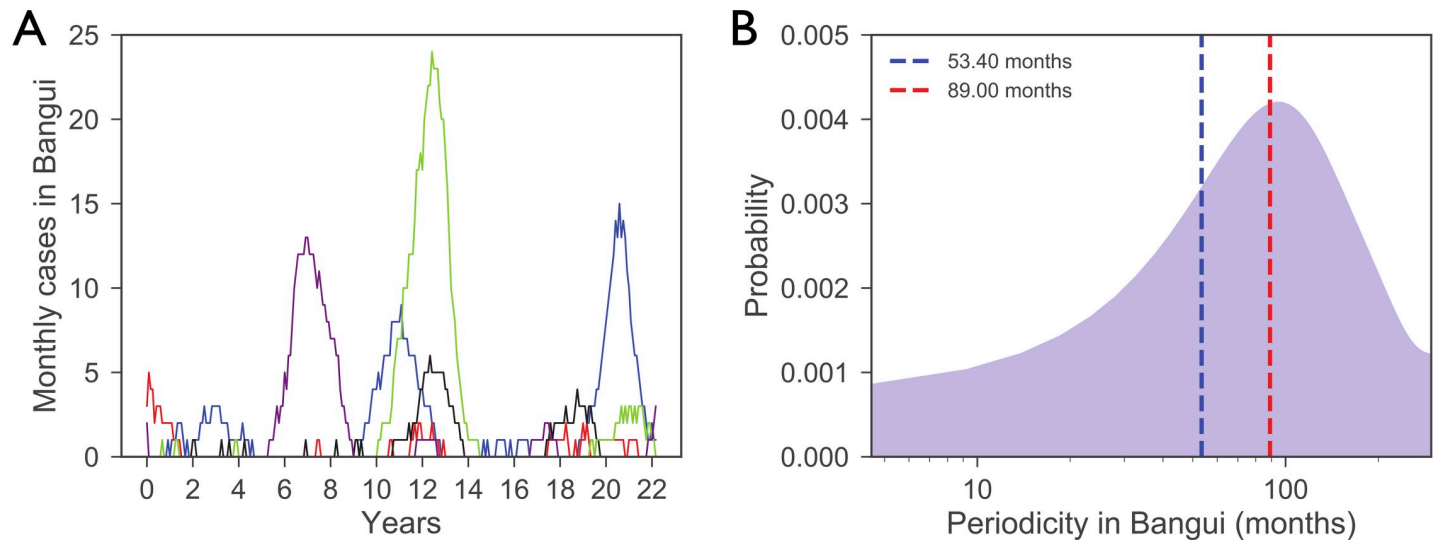


Fig 4. Numerical trajectories for Bangui. A) Simulated number of monthly cases in Bangui. Colors correspond to five different numerical runs. B) Probability distribution of the periods of the simulated epidemic cycles in Bangui compared to the two empirical oscillation periods of 53.4 (blue) and 89.0 months (red) reported in [18]. The distribution is obtained through wavelet analysis. Its peak is at 92 months. The x axis is in log scale.

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number of cases periodicity is obtained through wavelet analysis. Its peak is at 92 months. Additional numerical results are provided in the [S1 Text](#).

Spatial heterogeneity of rabies epidemic

The map of [Fig 5A](#) provides a spatial visualization of the simulated endemic situation for canine rabies in the country. Most of the prefectures are predicted to have an average prevalence in the 5–20% range, with the exception of the most eastern prefecture of Haut-Mbomou at lower prevalence, and the prefectures of Bangui, Ombella-M’Poko and Lobaye at higher prevalence. These last prefectures also contain the settlement with the highest proportion of rabid dogs, in both rural and urban patches. Prevalence is predicted to vary depending on the settlement type (Kolmogorov-Smirnov test, $p = 0.0001$, [Fig 5B](#)) and correlates with the density of infected dogs in each patch (Pearson correlation $r = 0.7, p < 10^{-6}$, [Fig 5C](#)).

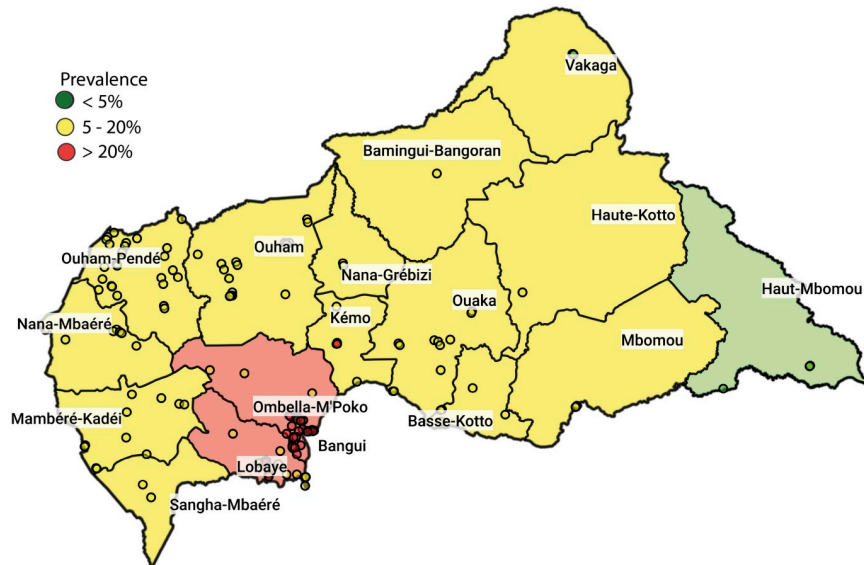
[Fig 5D](#) shows the lack of marked degree of urbanization of rabies epidemic in the country, with both urban and rural patches continuously contributing to both prevalence and density of infections.

Role of spatial fragmentation and human-mediated mobility

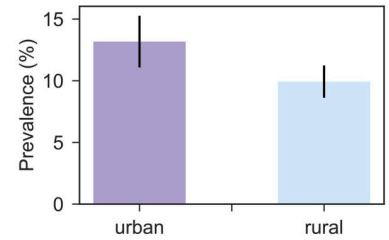
Transmission in Bangui only, or in the network of Bangui and its neighborhood would not self-sustain ([Fig 6](#), panels a and b, and [Fig 2](#) in [S1 Text](#)). Focusing on the role of urban vs. rural areas, our model predicts that rabies would not be able to circulate in a set of connected rural patches only (panels a and d), whereas its persistence probability would increase if dog population were structured in urban patches only ([Fig 6C](#)). Removing Bangui leads to results similar to urban patches only scenario ([Fig 6](#) in [S1 Text](#)).

Restrictions of human-mediated movements is predicted to dramatically impact viral persistence at the country level. Considering one single range of movement—whether short, or medium, or long—would practically prevent the virus to persist ([Fig 6E](#), [6F](#) and [6G](#)), even if the mobility network still ensures connectivity between the country’s settlements (see [Table 2](#) in [S1 Text](#)). Coupling two scales of movements in the model would re-establish similar

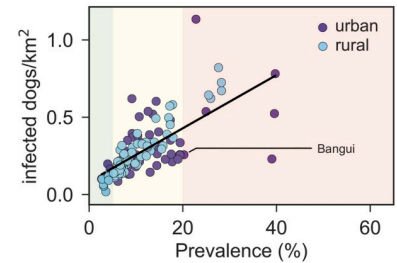
A



B



C



D

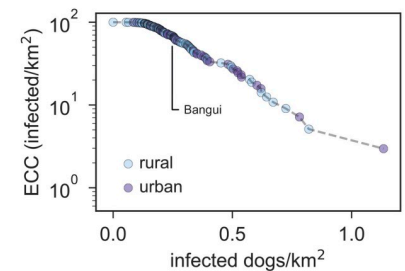


Fig 5. Dog rabies endemic prevalence in Central African Republic. A) Geographical representation of the predicted dog rabies endemic prevalence by administrative prefectures and by patch. Color codes range from low prevalence (<5%, green), to moderate prevalence (between 5% and 20%, yellow), to high prevalence (>20%, red). B) Predicted endemic prevalence by patch type. C) Scatter plot of the average density of infected dogs per km² vs. the endemic patch prevalence. D) Epidemic concentration curve (ECC) for rabies prevalence in the domestic dog population in Central African Republic. ECC is used to address the relative contribution of high density and low density areas to endemic infection [68]. The Central African Republic borders and administrative areas were generated using data from "Global Administrative Areas (2019)". University of California, Berkeley. Available online: <http://www.gadm.org> [21/07/2019].

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persistence probability profiles as in the full model if long-range movements are present (Fig 6I and 6J). If dogs were not allowed to move on such long-range connections, the probability for a considerably high persistence would decrease substantially (Fig 6H).

Discussion

Controlling and eliminating rabies in domestic dogs through the implementation of strict control measures including dog mass vaccination campaigns is now considered as the most cost-effective measure to prevent rabies in humans [6,13,17,24]. Thousands of people however continue to die every year from rabies disease transmitted by infected dogs, and rabies remains endemic throughout the African continent. In most African countries rabies surveillance is possible only in capital cities, where external re-introductions of the disease are suggested to sustain persistence [18,19,70]. On the other hand, very little is generally known about how the transmission in rural and periurban areas can affect the epidemic in the cities. Understanding the key drivers for the endemic circulation of the virus on a large geographical area is thus essential to design improved interventions for a sustainable elimination of dog rabies. Focusing on Central African Republic where the virus has been endemic in the domestic dog population for more than 20 years [18,47], we introduce a spatially explicit model to simulate rabies

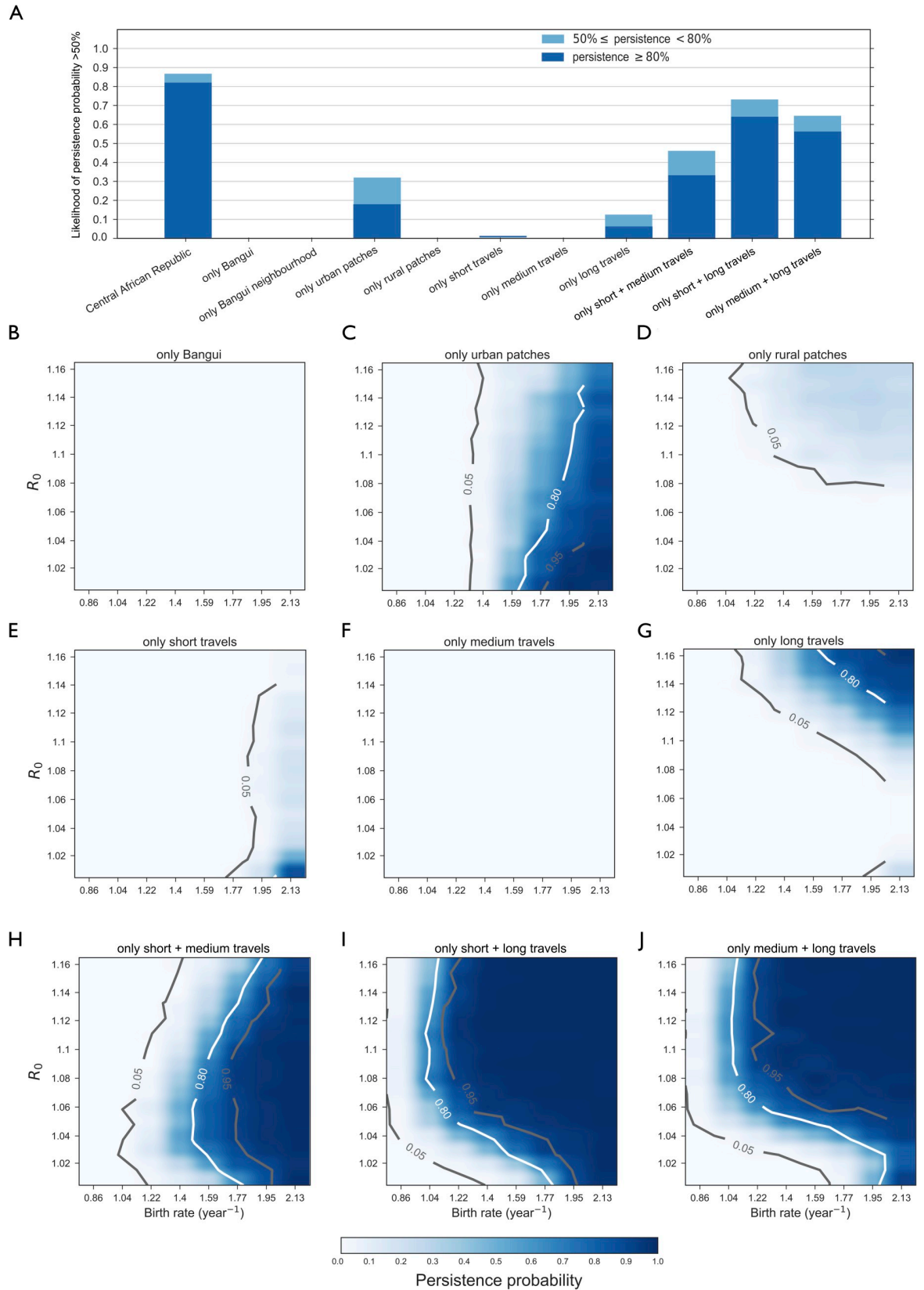


Fig 6. Role of spatial fragmentation and mobility. A) Likelihood to have persistence probability between 50% and 80%, or larger than 80% (light blue/dark blue stacked histogram) in the full parameter space (R_0 , birth rate) investigated and for different experimental scenarios (panels B–J). B), C), D), E), F), G), H), I), J) Predicted persistence probability of rabies in the domestic dog population as a function of the basic reproductive number R_0 and of the annual dogs birth rate for the scenarios considering: only Bangui population B); only urban patches and their connections, discarding rural patches C); only rural patches and their connections, discarding urban patches D); all patches but connected only through short travels E), medium travels F), or long travels G); all patches but connected only through short and medium travels H), short and long travels I), or medium and long travels J).

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transmission within both villages and urban areas and the spatial diffusion of the epidemic throughout the country led by human-mediated movements. Through numerical simulations, we can investigate the role of the various components acting on the infection dynamics at the population level.

First, we find that rabies would go extinct not only in the capital city alone, but also considering continuous re-introductions from the neighborhood of the city. This is in sharp contrast with previous expectations of the pattern of viral circulation proposed for Bangui [18] and other capital cities, such as e.g. N'Djamena in Chad [5,19], which suggested that importations might occur from the peri-urban areas adjacent to the city [18,19]. Our model predicts that longer range movements, connecting far-away settlements, are essential for rabies persistence. They allow occasional reintroductions of the disease sustaining its spatial spread.

Further, our work clearly shows that persistence of rabies in Central African Republic is favored by the non-trivial interplay of long-range movements of infected dogs with unusually long incubation periods. Thus, the virus can slowly diffuse in the country and the resulting continuous seeding of asynchronous rabies waves in distant settlements can efficiently counteract the local viral fade out while susceptibles are swiftly renewed elsewhere. This effect was already found in theoretical studies and in prior work addressing other diseases [71–75]. It was only recently considered for canine rabies persistence [44,45]. Moreover, this effect may be relevant for rabies in other carnivore species behaving in a similar way. It is indeed now evident that rabies virus has long evolved mechanisms to avoid extinction and successfully maintain itself in different reservoir host species during several thousand years, often in fragmented and poorly connected populations [76,77]. The intensification and the change of scale of human-mediated dissemination starting in the 15th century only plaid a role in the rapid global spread and maintenance of rabies observed in the 18th and 19th century in the Americas, Africa, Asia and Europe [76]. The prolonged incubation periods occurring in nature in dogs could then be an intrinsic factor of rabies virus pathogenesis that have constantly been selected and maintained during rabies virus genetic evolution as it played an instrumental role to ensure viral dispersion to new dog populations allowing the renewal of naïve domestic dog populations.

Our model also allows to address the respective role of urban and rural areas and shows that both are important for disease persistence, but extinction is more likely in a network of connected rural patches (as if, for example, urban areas undergo massive vaccination). This seems to be unrelated to the density of dogs—as it is rather similar in urban and rural patches with few exceptions, leading to a predicted spatial epidemic not showing a marked degree of urbanization—and it is more likely associated with the overall larger dog population size localized in urban patches.

Finally, numerically obtained periodic patterns of infected dogs in time are compatible with observations in Bangui [18] and in other African countries [17,25]. While we find a continuum spectrum of possible period durations that were not captured by surveillance data, this is likely due to the size effects of the limited observations (20 years) compared to the numerical extension of our simulations (>300 years). Analysis of longer historical data, when available, can further validate our predictions.

Our modeling findings have significant implications for the control of canine rabies in Central African Republic. The predictions indicate that sustainable elimination of the disease will indeed require a substantial effort targeting a much larger geographical area than focusing exclusively on the capital city and its surrounding peri-urban settlements. To be effective, mass vaccination campaigns would require targeting the entire set of urban areas in the country (42% of the settlements), which may result to be unfeasible. Banning movements on the long range (associated to the smallest fluxes of dogs moved) may be very effective in supporting sustained and prolonged vaccination campaigns in the country.

Our estimates indicate a low-level endemic stability of the disease consistent with previous values obtained in Bangui and in other African countries, and allowing rather steady dog population sizes [17–19]. With no mass vaccination in Central African Republic, higher transmission scenarios and the fatality rate of the disease would indeed not be compatible with the observed rather stable host population [24,78,79].

Our study is affected by some limitations. First, model inputs of host population size and mobility are inferred from human data and mobility models, similarly to [44,45]. Lack of data characterizing hosts' spatial distribution and mobility is a central issue for epidemic models, especially in African countries, where also data on human movements are typically scarce [80]. For this reason, well-established models of human mobility, such as e.g. the gravity model [81], are largely used in spatially explicit epidemic models for human diseases. Here, we use the same approach to model human-mediated movements of dogs from one settlement to another. Additional refinements to the gravity model can be made in future work (e.g. a refined dependence on the distance between settlements), if surveillance data from multiple locations (and not only from the capital city) become available. Although a direct comparison is limited by the differences in contexts, we find that our parsimonious choice of the gravity model provide projections for canine population moving daily on short, medium, and long-range movements that are close to the upper estimates resulting from a 2013 survey on human-mediated dog movements in the Philippines [29]. While lack of data remains a limitation, our modeling approach offers the theoretical framework where to evaluate the role that spatial fragmentation of human settlements in a particular geographical area may have on rabies persistence, based on expected human mobility flows. As such, it can be readily applied to other countries and different landscapes. This may additionally provide numerical evidence for context-specific elements favoring rabies dispersal (e.g. geography, natural barriers, etc.). Second, we did not consider rabies virus transmission from wildlife to domestic dogs, as dogs are recognized to be the main reservoir for rabies in sub-saharan Africa [32,63,64] and dog-to-dog transmission is estimated to be approximately eight times as common as transmission between dogs and other carnivores [30,63,82]. Moreover, domestic dogs live mostly in human settlements where interactions with wildlife are sporadic. Indeed epidemiological cycles of the virus in non-flying wildlife mammals are geographically limited in Africa and, to our knowledge, not present in Central African Republic [76]. Third, like other studies [83], our model did not consider the role of density transmission of RABV and assume a homogeneously mixing dog population within each patch. Therefore, it neither accounts for the fine scale heterogeneities in dog densities nor for the network structure of dog-to-dog contacts within the patches [84]. The study of the role of contact patterns between dogs is still at its infancy, and additional empirical data are needed to infer synthetic models that can be applicable in space. Fourth, we did not consider possible importation events across the country borders. This may occur for instance in the north along the border with Chad, separating densely populated areas in both countries. Future work may further extend our framework to a larger region including several countries where rabies disease is endemic in the domestic dog population. Increasing evidence calls indeed for prolonged and sustained efforts that are concerted across different

key stakeholders and countries in the region [4,25,85,86]. By parameterizing the model to national contexts, including the spatial fragmentation of the host population and implemented interventions, and fitting it to available surveillance data, it would be possible to numerically investigate the impact of different degrees of cross-border mobility on rabies persistence, and provide novel fundamental understanding to guide the successful elimination of rabies in the region.

Supporting information

S1 Text. The file contains: mathematical formulation of the models, additional numerical results and input values for the models.

(DOCX)

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References

1. Hampson K, Coudeville L, Lembo T, Sambo M, Kieffer A, Atllan M, et al. Estimating the Global Burden of Endemic Canine Rabies. Carvalho MS, editor. *PLoS Negl Trop Dis*. 2015; 9: e0003709. <https://doi.org/10.1371/journal.pntd.0003709> PMID: 25881058
2. World Health Organization. WHO Expert Consultation on Rabies: Second Report. 2013.
3. Wandeler AI, Matter HC, Kappeler A, Budde A. The ecology of dogs and canine rabies: a selective review. *Rev Sci Tech*. 1993; 12(1): 51–71.
4. Bögel K, Meslin FX. Economics of human and canine rabies elimination: guidelines for programme orientation. *Bull World Health Organ*. 1990; 68(3): 281–291.
5. Zinsstag J, Durr S, Penny MA, Mindekem R, Roth F, Gonzalez SM, et al. Transmission dynamics and economics of rabies control in dogs and humans in an African city. *Proc Natl Acad Sci*. 2009; 106: 14996–15001. <https://doi.org/10.1073/pnas.0904740106> PMID: 19706492
6. Lembo T, Hampson K, Kaare MT, Ernest E, Knobel D, Kazwala RR, et al. The Feasibility of Canine Rabies Elimination in Africa: Dispelling Doubts with Data. Rupprecht CE, editor. *PLoS Negl Trop Dis*. 2010; 4: e626. <https://doi.org/10.1371/journal.pntd.0000626> PMID: 20186330
7. Fitzpatrick MC, Hampson K, Cleaveland S, Meyers LA, Townsend JP, Galvani AP. Potential for Rabies Control through Dog Vaccination in Wildlife-Abundant Communities of Tanzania. Reithinger R, editor. *PLoS Negl Trop Dis*. 2012; 6: e1796. <https://doi.org/10.1371/journal.pntd.0001796> PMID: 22928056

8. Jibat T, Hogeveen H, Mourits MCM. Review on Dog Rabies Vaccination Coverage in Africa: A Question of Dog Accessibility or Cost Recovery? Rupprecht CE, editor. *PLoS Negl Trop Dis*. 2015; 9: e0003447. <https://doi.org/10.1371/journal.pntd.0003447> PMID: 25646774
9. Davlin SL, VonVille HM. Canine rabies vaccination and domestic dog population characteristics in the developing world: A systematic review. *Vaccine*. 2012; 30: 3492–3502. <https://doi.org/10.1016/j.vaccine.2012.03.069> PMID: 22480924
10. Mulipukwa CP, Mudenda B, Mbewe AR. Insights and efforts to control rabies in Zambia: Evaluation of determinants and barriers to dog vaccination in Nyimba district. Rupprecht CE, editor. *PLoS Negl Trop Dis*. 2017; 11: e0005946. <https://doi.org/10.1371/journal.pntd.0005946> PMID: 28991898
11. Cleaveland S, Kaare M, Knobel D, Laurenson MK. Canine vaccination—Providing broader benefits for disease control. *Vet Microbiol*. 2006; 117: 43–50. <https://doi.org/10.1016/j.vetmic.2006.04.009> PMID: 16701966
12. Dodet B, Tejiokem MC, Aguemou A-R, Bourhy H. Human rabies deaths in Africa: breaking the cycle of indifference. *Int Health*. 2015; 7: 4–6. <https://doi.org/10.1093/inthealth/ihu071> PMID: 25303941
13. Abela-Ridder B, Knopf L, Martin S, Taylor L, Torres G, De Balogh K. 2016: the beginning of the end of rabies? *Lancet Glob Health*. 2016; 4: e780–e781. [https://doi.org/10.1016/S2214-109X\(16\)30245-5](https://doi.org/10.1016/S2214-109X(16)30245-5)
14. Nel LH. Discrepancies in Data Reporting for Rabies, Africa. *Emerg Infect Dis*. 2013; 19: 529–533. <https://doi.org/10.3201/eid1904.120185> PMID: 23628197
15. Anderson RM, May RM. *Infectious Diseases of Humans: Dynamics and Control*. OUP Oxford; 1992.
16. Keeling MJ, Rohani P. *Modeling Infectious Diseases in Humans and Animals*. Princeton University Press; 2011.
17. Hampson K, Dushoff J, Cleaveland S, Haydon DT, Kaare M, Packer C, et al. Transmission Dynamics and Prospects for the Elimination of Canine Rabies. Rupprecht CE, editor. *PLoS Biol*. 2009; 7: e1000053. <https://doi.org/10.1371/journal.pbio.1000053> PMID: 19278295
18. Bourhy H, Nakouné E, Hall M, Nouvellet P, Lepelletier A, Talbi C, et al. Revealing the Micro-scale Signature of Endemic Zoonotic Disease Transmission in an African Urban Setting. Parrish C, editor. *PLOS Pathog*. 2016; 12: e1005525. <https://doi.org/10.1371/journal.ppat.1005525> PMID: 27058957
19. Zinsstag J, Lechenne M, Laager M, Mindekem R, Naïssengar S, Oussiguéré A, et al. Vaccination of dogs in an African city interrupts rabies transmission and reduces human exposure. *Sci Transl Med*. 2017; 9: eaaf6984. <https://doi.org/10.1126/scitranslmed.aaf6984> PMID: 29263230
20. Lee J-H, Lee J-B, Kim J-S, Bae C-S, Lee W-C, Lee M-J. Review of canine rabies prevalence under two different vaccination programmes in Korea. *Vet Rec*. 2001; 148: 511–512. <https://doi.org/10.1136/vr.148.16.511> PMID: 11345995
21. Coleman PG, Dye C. Immunization coverage required to prevent outbreaks of dog rabies. *Vaccine*. 1996; 14: 185–186. [https://doi.org/10.1016/0264-410X\(95\)00197-9](https://doi.org/10.1016/0264-410X(95)00197-9)
22. Kitala PM, McDERMOTT JJ, Coleman PG, Dye C. Comparison of vaccination strategies for the control of dog rabies in Machakos District, Kenya. *Epidemiol Infect*. 2002; 129: <https://doi.org/10.1017/S0950268802006957> PMID: 12211590
23. Bilinski AM, Fitzpatrick MC, Rupprecht CE, Paltiel AD, Galvani AP. Optimal frequency of rabies vaccination campaigns in Sub-Saharan Africa. *Proc R Soc B Biol Sci*. 2016; 283: 20161211. <https://doi.org/10.1098/rspb.2016.1211> PMID: 27852799
24. Townsend SE, Sumantra IP, Pudjiatmoko, Bagus GN, Brum E, Cleaveland S, et al. Designing Programs for Eliminating Canine Rabies from Islands: Bali, Indonesia as a Case Study. Rupprecht CE, editor. *PLoS Negl Trop Dis*. 2013; 7: e2372. <https://doi.org/10.1371/journal.pntd.0002372> PMID: 23991233
25. Hampson K, Dushoff J, Bingham J, Bruckner G, Ali YH, Dobson A. Synchronous cycles of domestic dog rabies in sub-Saharan Africa and the impact of control efforts. *Proc Natl Acad Sci*. 2007; 104: 7717–7722. <https://doi.org/10.1073/pnas.0609122104> PMID: 17452645
26. Talbi C, Holmes EC, de Benedictis P, Faye O, Nakoune E, Gamatie D, et al. Evolutionary history and dynamics of dog rabies virus in western and central Africa. *J Gen Virol*. 2009; 90: 783–791. <https://doi.org/10.1099/vir.0.007765-0>
27. Bourhy H, Reynes J-M, Dunham EJ, Dacheux L, Larrous F, Huong VTQ, et al. The origin and phylogeography of dog rabies virus. *J Gen Virol*. 2008; 89: 2673–2681. <https://doi.org/10.1099/vir.0.2008/003913-0> PMID: 18931062
28. Lemey P, Rambaut A, Welch JJ, Suchard MA. Phylogeography Takes a Relaxed Random Walk in Continuous Space and Time. *Mol Biol Evol*. 2010; 27: 1877–1885. <https://doi.org/10.1093/molbev/msq067> PMID: 20203288

29. Ferguson EA, Hampson K, Cleaveland S, Consunji R, Deray R, Friar J, et al. Heterogeneity in the spread and control of infectious disease: consequences for the elimination of canine rabies. *Sci Rep*. 2015; 5: 18232. <https://doi.org/10.1038/srep18232> PMID: 26667267
30. Woodroffe R, Donnelly CA. Risk of contact between endangered African wild dogs *Lycaon pictus* and domestic dogs: opportunities for pathogen transmission: Contact between domestic and wild dogs. *J Appl Ecol*. 2011; 48: 1345–1354. <https://doi.org/10.1111/j.1365-2664.2011.02059.x>
31. Nettles VF, Shaddock JH, Sikes RK, Reyes CR. Rabies in translocated raccoons. *Am J Public Health*. 1979; 69: 601–602. <https://doi.org/10.2105/AJPH.69.6.601> PMID: 443502
32. Smith DL, Lucey B, Waller LA, Childs JE, Real LA. Predicting the spatial dynamics of rabies epidemics on heterogeneous landscapes. *Proc Natl Acad Sci*. 2002; 99: 3668–3672. <https://doi.org/10.1073/pnas.042400799> PMID: 11904426
33. Wilson ML, Bretsky PM, Cooper GH, Egbertson SH, Van Kruiningen HJ, Cartter ML. Emergence of raccoon rabies in Connecticut, 1991–1994: spatial and temporal characteristics of animal infection and human contact. *Am J Trop Med Hyg*. 1997; 57: 457–463.
34. Bajardi P, Barrat A, Savini L, Colizza V. Optimizing surveillance for livestock disease spreading through animal movements. *J R Soc Interface*. 2012; 9: 2814–2825. <https://doi.org/10.1098/rsif.2012.0289> PMID: 22728387
35. Talbi C, Lemey P, Suchard MA, Abdelatif E, Elharrak M, Jalal N, et al. Phylodynamics and Human-Mediated Dispersal of a Zoonotic Virus. *Emerman M*, editor. *PLoS Pathog*. 2010; 6: e1001166. <https://doi.org/10.1371/journal.ppat.1001166> PMID: 21060816
36. Gsell AS, Knobel DL, Kazwala RR, Vounatsou P, Zinsstag J. Domestic dog demographic structure and dynamics relevant to rabies control planning in urban areas in Africa: the case of Iringa, Tanzania. *BMC Vet Res*. 2012; 8: 236. <https://doi.org/10.1186/1746-6148-8-236> PMID: 23217194
37. Kulonen K, Smreczak M, Kissi B, Tordo N, Holmes EC, Audry L, et al. Ecology and evolution of rabies virus in Europe. *J Gen Virol*. 1999; 80: 2545–2557. <https://doi.org/10.1099/0022-1317-80-10-2545> PMID: 10573146
38. Tenzin, Sharma B, Dhand NK, Timsina N, Ward MP. Reemergence of Rabies in Chhukha District, Bhutan, 2008. *Emerg Infect Dis*. 2010; 16: 1925–1930. <https://doi.org/10.3201/eid1612.100958> PMID: 21122223
39. Dellicour S, Rose R, Faria NR, Vieira LFP, Bourhy H, Gilbert M, et al. Using Viral Gene Sequences to Compare and Explain the Heterogeneous Spatial Dynamics of Virus Epidemics. *Mol Biol Evol*. 2017 [cited 28 Jul 2017]. <https://doi.org/10.1093/molbev/msx176> PMID: 28651357
40. Hemachudha T, Laothamatas J, Rupprecht CE. Human rabies: a disease of complex neuropathogenic mechanisms and diagnostic challenges. *Lancet Neurol*. 2002; 1: 101–109. [https://doi.org/10.1016/S1474-4422\(02\)00041-8](https://doi.org/10.1016/S1474-4422(02)00041-8)
41. Rupprecht CE, Hanlon CA, Hemachudha T. Rabies re-examined. *Lancet Infect Dis*. 2002; 2: 327–343. [https://doi.org/10.1016/S1473-3099\(02\)00287-6](https://doi.org/10.1016/S1473-3099(02)00287-6)
42. Dean DJ, Evans WM, McClure RC. Pathogenesis of rabies. *Bull World Health Organ*. 1963; 29: 803–811.
43. Hemachudha T. Human rabies: clinical aspects, pathogenesis, and potential therapy. *Curr Top Microbiol Immunol*. 1994; 187: 121–143.
44. Beyer HL, Hampson K, Lembo T, Cleaveland S, Kaare M, Haydon DT. The implications of metapopulation dynamics on the design of vaccination campaigns. *Vaccine*. 2012; 30: 1014–1022. <https://doi.org/10.1016/j.vaccine.2011.12.052> PMID: 22198516
45. Beyer HL, Hampson K, Lembo T, Cleaveland S, Kaare M, Haydon DT. Metapopulation dynamics of rabies and the efficacy of vaccination. *Proc R Soc B Biol Sci*. 2011; 278: 2182–2190. <https://doi.org/10.1098/rspb.2010.2312> PMID: 21159675
46. Dürr S, Ward MP. Development of a Novel Rabies Simulation Model for Application in a Non-endemic Environment. Zinsstag J, editor. *PLoS Negl Trop Dis*. 2015; 9: e0003876. <https://doi.org/10.1371/journal.pntd.0003876> PMID: 26114762
47. Tricou V, Bouscaillou J, Kamba Mebourou E, Koyanongo FD, Nakouné E, Kazanji M. Surveillance of Canine Rabies in the Central African Republic: Impact on Human Health and Molecular Epidemiology. Zinsstag J, editor. *PLoS Negl Trop Dis*. 2016; 10: e0004433. <https://doi.org/10.1371/journal.pntd.0004433> PMID: 26859829
48. Nakouné E, Digol M, Konamna X, Selekon B, Le Faou A. New introduction and spread of rabies among dog population in Bangui. *Acta Trop*. 2012; 123: 107–110. <https://doi.org/10.1016/j.actatropica.2012.04.005> PMID: 22569561

49. Cori A, Nouvellet P, Garske T, Bourhy H, Nakouné E, Jombart T. A graph-based evidence synthesis approach to detecting outbreak clusters: An application to dog rabies. Pascual M, editor. *PLOS Comput Biol*. 2018; 14: e1006554. <https://doi.org/10.1371/journal.pcbi.1006554> PMID: 30557340
50. Lloyd AL, May RM. Spatial Heterogeneity in Epidemic Models. *J Theor Biol*. 1996; 179: 1–11. <https://doi.org/10.1006/jtbi.1996.0042> PMID: 8733427
51. Grenfell B. (Meta)population dynamics of infectious diseases. *Trends Ecol Evol*. 1997; 12: 395–399. [https://doi.org/10.1016/S0169-5347\(97\)01174-9](https://doi.org/10.1016/S0169-5347(97)01174-9)
52. Butler JRA, Bingham J. Demography and dog-human relationships of the dog population in Zimbabwean communal lands. *Vet Rec*. 2000; 147: 442–446. <https://doi.org/10.1136/vr.147.16.442> PMID: 11079440
53. Knobel DL, Cleaveland S, Coleman PG, Fèvre EM, Meltzer MI, Miranda MEG, et al. Re-evaluating the burden of rabies in Africa and Asia. *Bull World Health Organ*. 2005; 83: 360–368.
54. Knobel DL, Laurenson MK, Kazwala RR, Boden LA, Cleaveland S. A cross-sectional study of factors associated with dog ownership in Tanzania. *BMC Vet Res*. 2008; 4: 5. <https://doi.org/10.1186/1746-6148-4-5> PMID: 18230137
55. Linard C, Gilbert M, Snow RW, Noor AM, Tatem AJ. Population Distribution, Settlement Patterns and Accessibility across Africa in 2010. Schumann GJ-P, editor. *PLoS ONE*. 2012; 7: e31743. <https://doi.org/10.1371/journal.pone.0031743> PMID: 22363717
56. <https://www.oecd.org/cfe/regional-policy/Definition-of-Functional-Urban-Areas-for-the-OECD-metropolitan-database.pdf>.
57. Balcan D, Colizza V, Gonçalves B, Hu H, Ramasco JJ, Vespignani A. Multiscale mobility networks and the spatial spreading of infectious diseases. *Proc Natl Acad Sci*. 2009; 106: 21484–21489. <https://doi.org/10.1073/pnas.0906910106> PMID: 20018697
58. Erlander Sven, Stewart Neil F. *The Gravity Model in Transportation Analysis: Theory and Extensions*. 1990.
59. Ortúzar J, de D, Willumsen LG. *Modelling Transport*. (Wiley, Chichester, United Kingdom); 2001.
60. Xia Y, Bjørnstad ON, Grenfell BT. Measles Metapopulation Dynamics: A Gravity Model for Epidemiological Coupling and Dynamics. *Am Nat*. 2004; 164: 267–281. <https://doi.org/10.1086/422341> PMID: 15278849
61. Zinsstag J, Durr S, Penny MA, Mindekem R, Roth F, Gonzalez SM, et al. Transmission dynamics and economics of rabies control in dogs and humans in an African city. *Proc Natl Acad Sci*. 2009; 106: 14996–15001. <https://doi.org/10.1073/pnas.0904740106> PMID: 19706492
62. Eng TR, Fishbein DB, Talamante HE, Hall DB, Chavez GF, Dobbins JG, et al. Urban epizootic of rabies in Mexico: epidemiology and impact of animal bite injuries. *Bull World Health Organ*. 1993; 71: 615.
63. Lembo T, Hampson K, Haydon DT, Craft M, Dobson A, Dushoff J, et al. Exploring reservoir dynamics: a case study of rabies in the Serengeti ecosystem. *J Appl Ecol*. 2008; 45: 1246–1257. <https://doi.org/10.1111/j.1365-2664.2008.01468.x> PMID: 22427710
64. Kitala P, McDermott J, Kyule M, Gathuma J, Perry B, Wandeler A. Dog ecology and demography information to support the planning of rabies control in Machakos District, Kenya. *Acta Trop*. 2001; 78: 217–230. [https://doi.org/10.1016/S0001-706X\(01\)00082-1](https://doi.org/10.1016/S0001-706X(01)00082-1)
65. New JCJ, Kelch WJ, Hutchison JM, Salman MD, King M, Scarlett JM, et al. Birth and death rate estimates of cats and dogs in U.S. households and related factors. *J Appl Anim Welf Sci JAAWS*. 2004 [cited 20 Jun 2016]. Available: <http://agris.fao.org/agris-search/search.do?recordID=US201301007213>
66. Balcan D, Hu H, Gonçalves B, Bajardi P, Poletto C, Ramasco JJ, et al. Seasonal transmission potential and activity peaks of the new influenza A (H1N1): a Monte Carlo likelihood analysis based on human mobility. *BMC Med*. 2009; 7: 45.
67. Poletto C, Pelat C, Levy-Bruhl D, Yazdanpanah Y, Boelle P, Colizza V. Assessment of the Middle East respiratory syndrome coronavirus (MERS-CoV) epidemic in the Middle East and risk of international spread using a novel maximum likelihood analysis approach. *Eurosurveillance*. 2014; 19: 3.
68. Coburn BJ, Okano JT, Blower S. Using geospatial mapping to design HIV elimination strategies for sub-Saharan Africa. *Sci Transl Med*. 2017; 9: eaag0019.
69. Torrence C, Compo GP. *A Practical Guide to Wavelet Analysis*. *Bull Am Meteorol Soc*. 1998; 79: 61–78. [https://doi.org/10.1175/1520-0477\(1998\)079<0061:APGTWA>2.0.CO;2](https://doi.org/10.1175/1520-0477(1998)079<0061:APGTWA>2.0.CO;2)
70. Brunner K, Hampson K, Horton DL, Biek R. Integrating the landscape epidemiology and genetics of RNA viruses: rabies in domestic dogs as a model. *Parasitology*. 2012; 139: 1899–1913. <https://doi.org/10.1017/S003118201200090X> PMID: 22814380
71. Conlan AJK, Grenfell BT. Seasonality and the persistence and invasion of measles. *Proc R Soc B Biol Sci*. 2007; 274: 1133–1141. <https://doi.org/10.1098/rspb.2006.0030> PMID: 17327206

72. Wearing HJ, Rohani P, Keeling MJ. Appropriate Models for the Management of Infectious Diseases. Ellner SP, editor. *PLoS Med.* 2005; 2: e174. <https://doi.org/10.1371/journal.pmed.0020174> PMID: [16013892](https://pubmed.ncbi.nlm.nih.gov/16013892/)
73. Lloyd AL. Realistic Distributions of Infectious Periods in Epidemic Models: Changing Patterns of Persistence and Dynamics. *Theor Popul Biol.* 2001; 60: 59–71. <https://doi.org/10.1006/tpbi.2001.1525> PMID: [11589638](https://pubmed.ncbi.nlm.nih.gov/11589638/)
74. Feng Z, Xu D, Zhao H. Epidemiological Models with Non-Exponentially Distributed Disease Stages and Applications to Disease Control. *Bull Math Biol.* 2007; 69: 1511–1536. <https://doi.org/10.1007/s11538-006-9174-9> PMID: [17237913](https://pubmed.ncbi.nlm.nih.gov/17237913/)
75. Keeling MJ, Grenfell BT. Understanding the persistence of measles: reconciling theory, simulation and observation. *Proc R Soc B Biol Sci.* 2002; 269: 335–343. <https://doi.org/10.1098/rspb.2001.1898> PMID: [11886620](https://pubmed.ncbi.nlm.nih.gov/11886620/)
76. Troupin C, Dacheux L, Tanguy M, Sabeta C, Blanc H, Bouchier C, et al. Large-Scale Phylogenomic Analysis Reveals the Complex Evolutionary History of Rabies Virus in Multiple Carnivore Hosts. Parrish C, editor. *PLOS Pathog.* 2016; 12: e1006041. <https://doi.org/10.1371/journal.ppat.1006041> PMID: [27977811](https://pubmed.ncbi.nlm.nih.gov/27977811/)
77. Fisher CR, Streicker DG, Schnell MJ. The spread and evolution of rabies virus: conquering new frontiers. *Nat Rev Microbiol.* 2018; 16: 241–255. <https://doi.org/10.1038/nrmicro.2018.11> PMID: [29479072](https://pubmed.ncbi.nlm.nih.gov/29479072/)
78. Windiyarningsih C, Wilde H, Meslin FX, Suroso T, Widarso HS. The rabies epidemic on Flores Island, Indonesia (1998–2003). *J Med Assoc Thai Chotmaihet Thangphaet.* 2004; 87 11: 1389–93.
79. Randall DA, Williams SD, Kuzmin IV, Rupprecht CE, Tallents LA, Tefera Z, et al. Rabies in Endangered Ethiopian Wolves. *Emerg Infect Dis.* 2004; 10: 2214–2217. <https://doi.org/10.3201/eid1012.040080> PMID: [15663865](https://pubmed.ncbi.nlm.nih.gov/15663865/)
80. Wesolowski A, Buckee CO, Bengtsson L, Wetter E, Lu X, Tatem AJ. Commentary: Containing the Ebola Outbreak—the Potential and Challenge of Mobile Network Data. *PLoS Curr.* 2014 [cited 13 Apr 2019]. <https://doi.org/10.1371/currents.outbreaks.0177e7fc52217b8b634376e2f3efc5e> PMID: [25642369](https://pubmed.ncbi.nlm.nih.gov/25642369/)
81. Garcia AJ, Pindolia DK, Lopiano KK, Tatem AJ. Modeling internal migration flows in sub-Saharan Africa using census microdata. *Migr Stud.* 2015; 3: 89–110. <https://doi.org/10.1093/migration/mnu036>
82. Hanlon CA, Niezgodna M, Rupprecht CE. Rabies in Terrestrial Animals. Rabies. Elsevier; 2007. pp. 201–258. <https://doi.org/10.1016/B978-012369366-2/50007-5>
83. Laager M, Léchenne M, Naissengar K, Mindekem R, Oussiguere A, Zinsstag J, et al. A metapopulation model of dog rabies transmission in N'Djamena, Chad. *J Theor Biol.* 2019; 462: 408–417. <https://doi.org/10.1016/j.jtbi.2018.11.027> PMID: [30500602](https://pubmed.ncbi.nlm.nih.gov/30500602/)
84. Laager M, Mbilo C, Madaye EA, Naminou A, Léchenne M, Tschopp A, et al. The importance of dog population contact network structures in rabies transmission. Rupprecht CE, editor. *PLoS Negl Trop Dis.* 2018; 12: e0006680. <https://doi.org/10.1371/journal.pntd.0006680> PMID: [30067733](https://pubmed.ncbi.nlm.nih.gov/30067733/)
85. Cleaveland S, Hampson K. Rabies elimination research: juxtaposing optimism, pragmatism and realism. *Proc R Soc B Biol Sci.* 2017; 284: 20171880. <https://doi.org/10.1098/rspb.2017.1880> PMID: [29263285](https://pubmed.ncbi.nlm.nih.gov/29263285/)
86. Cleaveland S, Lankester F, Townsend S, Lembo T, Hampson K. Rabies control and elimination: a test case for One Health. *Vet Rec.* 2014; 175: 188–193. <https://doi.org/10.1136/vr.g4996> PMID: [25172649](https://pubmed.ncbi.nlm.nih.gov/25172649/)