

Initiation of Infection

Urbanization and geographic expansion of zoonotic arboviral diseases: mechanisms and potential strategies for prevention

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Arthropod-borne viruses (arboviruses) mainly infect people via direct spillover from enzootic cycles. However, dengue, chikungunya, and yellow fever viruses have repeatedly initiated urban transmission cycles involving human amplification and peridomestic mosquito vectors to cause major epidemics. Here, I review these urban emergences and potential strategies for their prevention and control.

Arthropod-borne viruses (arboviruses) are transmitted by mosquitoes and other biting arthropods. They occur nearly worldwide and all were, until recently, zoonotic in nature (circulating enzootically among wild animals but sometimes infecting people). Arboviruses have undoubtedly caused human disease in people for at least millennia, but a few have increased in importance during recent decades because of human population expansion and activities that have increased exposure to infection. Some have simply expanded the geographic range of their enzootic cycles, often due to human trade or travel, to infect more people via direct spillover (transmission from animals via enzootic or bridge vectors to humans, who typically do not develop sufficient viremia to extend the transmission chain and are thus dead-end hosts). An example is West Nile virus (WNV; *Flaviviridae: Flavivirus*), which circulates in an avian-mosquito cycle and has caused small epidemics for many decades, because of spillover, in the Old World. However, in 1999 WNV was introduced into New York, where it amplified and spread rapidly to cause the largest epidemic of arboviral encephalitis ever documented in the USA, followed by a major resurgence in 2012 [1]. Several other arboviruses may have comparable abilities to invade new geographic regions; the evidence that little or no adaptive evolution was needed for the WNV epidemic to spread across the Americas [2] suggests that such events could occur with other deadly flaviviruses that exploit multiple vectors and hosts [3].

Several of the most important arboviral pathogens are those that not only infect people via enzootic spillover but also use humans as amplification hosts. The most important is the flavivirus dengue virus (DENV), which infects more than 50 million people annually nearly throughout the tropics and subtropics, with secondary infections

carrying a high risk of hemorrhagic disease that can be fatal [4]. DENV, which includes four serotypes with limited antigenic cross-reactivity, evolved thousands of years ago in Southeast Asia, where divergence of the four serotypes probably occurred in arboreal nonhuman primate (NHP)–mosquito cycles (Figure 1). Later, DENV-2 was introduced into an African enzootic cycle (probably via spillback or infection of arboreal vectors or NHPs by human strains) and all four serotypes ultimately emerged into independent endemic or epidemic cycles involving humans and the anthropophilic mosquito vectors *Aedes aegypti* and *Aedes albopictus* [5]. Extensive urbanization, the nearly global tropical invasion by *A. aegypti*, and increased commerce and travel led to an explosion of urban dengue fever after World War II [4,6]. Evidence suggests that there was little or no adaptation when enzootic DENV-2 first emerged in the urban cycle, and that human DENV infections with enzootic strains in West Africa and Southeast Asia can be highly pathogenic, like urban infections [5].

Another flavivirus with a long history of emergence into a human–*A. aegypti* cycle is yellow fever virus (YFV), which is highly virulent (case fatality rate of 20–50%) [7]. YFV originated in Africa (Figure 1) in a mosquito–NHP arboreal transmission cycle nearly indistinguishable from that of DENV [8]. Like DENV, YFV caused periodic epidemics in port cities of the Americas, including temperate regions, during the 17th to early 20th centuries, as well as in many parts of Africa. Urban YFV epidemics involving *A. aegypti* transmission have recently been infrequently reported, especially in Latin America. However, unlike DENV, YFV developed enzootic transmission cycles in the Neotropics, probably via spillback, during the early years of the slave trade from West Africa. Also unlike DENV, YFV has no documented history of spread into Asia, and its urban cycle has never become permanent (Figure 1). An effective YFV vaccine has been available since the 1930s. Unfortunately, incomplete coverage in regions at risk of enzootic spillover and urban epidemics allows hundreds to thousands of human infections, many of which are fatal, to continue annually in Africa and South America, with many more undoubtedly unreported [7].

The third major arboviral disease with a history of urbanization is chikungunya virus (CHIKV; *Togaviridae: Alphavirus*), which originated in Africa (Figure 1) and, like DENV and YFV, was probably transported by sailing ships

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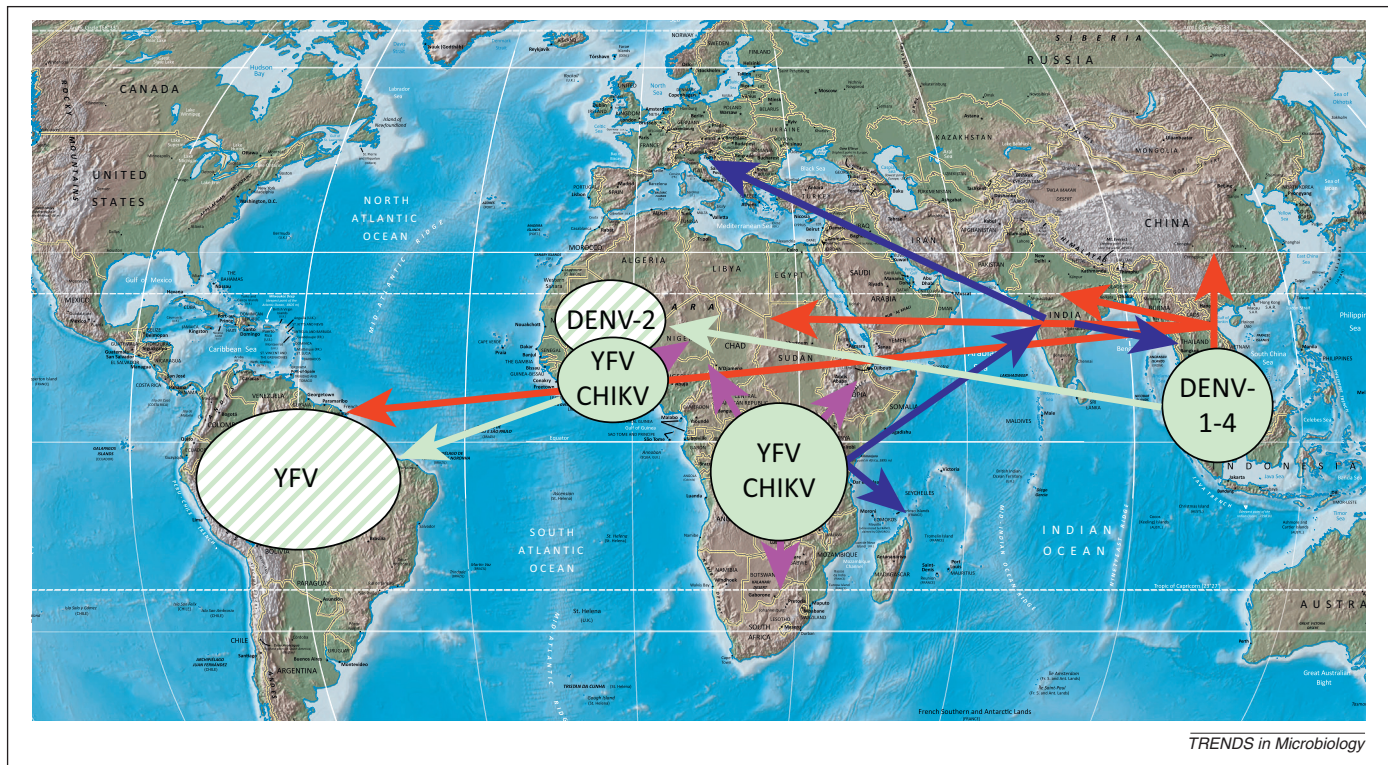


Figure 1. Map showing approximate enzootic origins of dengue virus (DENV), chikungunya virus (CHIKV), and yellow fever virus (YFV) (green circles), introductions (green arrows) that established additional documented sites of enzootic transmission (striped green circles), and general patterns (not comprehensive) of the urbanized spread of DENV (red), CHIKV (blue), or YFV (magenta) leading to extant endemic or epidemic strains.

to port cities during the 18th and 19th centuries [3]. However, CHIKV has spread from eastern Africa into Asia at least twice to establish permanent urban transmission. One emergence occurred during or before the early 1950s. The most recent emergence began in coastal Kenya in 2004, before spreading into islands of the Indian Ocean and into Asia to cause severe, often chronic arthralgia in millions of people (Figure 1). The recent epidemics were facilitated by *A. albopictus*-adaptive mutations that led to more efficient interhuman transmission, even in temperate regions of Europe where this mosquito survives cold winters.

History suggests that enzootic arboviruses that use NHPs as hosts probably represent the greatest risk for urbanization. This group includes the arthralgic alphavirus Mayaro (MAYV), which circulates enzootically in South America among NHPs, transmitted by arboreal mosquitoes in the genera *Haemagogus* and *Sabethes*. Another threat is the flavivirus Zika, which circulates widely in Africa and Asia, probably using NHPs as enzootic hosts, and which recently caused an epidemic of febrile illness, apparently involving human amplification, on Yap Island in the Pacific [9]. Although its enzootic cycle does not appear to involve NHPs, Oropouche virus (OROV; *Bunyaviridae: Bunyavirus*), an enzootic in South America, has caused major epidemics of febrile illness involving human amplification and urban transmission by the midge *Culicoides paraensis* [10].

Five strategies for preventing or mitigating arbovirus urbanization are outlined below.

- (i) Intervention in enzootic cycles. In theory, the prevention of arbovirus urbanization could be mitigated at

several stages of the emergence process (Figure 2). Reduction or elimination of enzootic circulation via vector control or prevention of reservoir and amplification host infection would be highly challenging considering the widespread and often remote locations of sylvatic foci, the environmental concerns of large-scale insecticide use, and the extreme difficulty of vaccinating wild animals. Modulation of mosquito transmission by the introduction of endosymbiotic *Wolbachia* spp. bacteria, which suppress the replication of some arboviruses, holds promise for reducing urban transmission of DENV, YFV, and CHIKV [11]. However, it is unknown if this approach could be applied to the more diverse enzootic vectors that are poorly understood.

- (ii) Prevention of enzootic spillover. If enzootic transmission of arboviruses within their sylvatic NHP cycles offers a poor target for interruption, then prevention of introductions into the urban cycle represents the next step potentially amenable to intervention (Figure 2). DENV, YFV, and CHIKV apparently emerge from enzootic cycles by infecting people who live near forested habitats and who then initiate the human-mosquito cycle, and ultimately transport it to urban centers. Thus, reductions in the exposure of these human populations to enzootic vectors, or vaccination of such persons, could reduce the risk of further urbanization. Some measures successfully implemented to reduce infection with malaria parasites and other vector-borne pathogens, such as bed nets, may offer some protection against enzootic arbovirus spillover infections. Nevertheless, anthropogenic

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