




Socioeconomic risk markers of arthropod-borne virus (arbovirus) infections: a systematic literature review and meta-analysis

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To cite: Power GM, Vaughan AM, Qiao L, *et al.* Socioeconomic risk markers of arthropod-borne virus (arbovirus) infections: a systematic literature review and meta-analysis. *BMJ Global Health* 2022;**7**:e007735. doi:10.1136/bmjgh-2021-007735

Handling editor Seema Biswas

► Additional supplemental material is published online only. To view, please visit the journal online (<http://dx.doi.org/10.1136/bmjgh-2021-007735>).

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Received 18 October 2021

Accepted 2 March 2022



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ABSTRACT

Introduction Arthropod-borne viruses (arboviruses) are of notable public health importance worldwide, owing to their potential to cause explosive outbreaks and induce debilitating and potentially life-threatening disease manifestations. This systematic review and meta-analysis aims to assess the relationship between markers of socioeconomic position (SEP) and infection due to arboviruses with mosquito vectors.

Methods We conducted a systematic search on PubMed, Embase, and LILACS databases to identify studies published between 1980 and 2020 that measured the association of SEP markers with arbovirus infection. We included observational studies without geographic location or age restrictions. We excluded studies from grey literature, reviews and ecological studies. Study findings were extracted and summarised, and pooled estimates were obtained using random-effects meta-analyses.

Results We identified 36 observational studies using data pertaining to 106 524 study participants in 23 geographic locations that empirically examined the relationship between socioeconomic factors and infections caused by seven arboviruses (dengue, chikungunya, Japanese encephalitis, Rift Valley fever, Sindbis, West Nile and Zika viruses). While results were varied, descriptive synthesis pointed to a higher risk of arbovirus infection associated with markers of lower SEP, including lower education, income poverty, low healthcare coverage, poor housing materials, interrupted water supply, marital status (married, divorced or widowed), non-white ethnicities and migration status. Pooled crude estimates indicated an increased risk of arboviral infection associated with lower education (risk ratio, RR 1.5 95% CI 1.3 to 1.9); $I^2=83.1\%$, interruption of water supply (RR 1.2, 95% CI 1.1 to 1.3; $I^2=0.0\%$) and having been married (RR 1.5 95% CI 1.1 to 2.1; $I^2=85.2\%$).

Conclusion Evidence from this systematic review suggests that lower SEP increases the risk of acquiring arboviral infection; however, there was large heterogeneity across studies. Further studies are required to delineate the relationship between specific individual, household and community-level SEP indicators and arbovirus infection risks to help inform targeted public health interventions.

PROSPERO registration number CRD42019158572.

WHAT IS ALREADY KNOWN?

- ⇒ Arboviruses with mosquito vectors are of notable global public health importance owing to their potential to cause explosive outbreaks and induce debilitating and potentially life-threatening disease manifestations.
- ⇒ In regions with established arboviral circulation, factors indicative of socioeconomic position, such as increased population density, inadequate water management and poor housing conditions, may exacerbate vector proliferation and elevate infection risks.

WHAT ARE THE NEW FINDINGS?

- ⇒ Descriptive synthesis pointed to a higher risk of arboviral infection associated with markers of lower socioeconomic position, including lower education, income poverty, low healthcare coverage, poor housing materials, interruptions of water supply, marital status (married, divorced or widowed) and non-white ethnicity.
- ⇒ Pooled crude estimates from meta-analyses indicated an increased risk of arboviral infection associated with having lower education, interruption of water supply and having ever been married.

WHAT DO THE NEW FINDINGS IMPLY?

- ⇒ This review underscores the importance of evaluating the arbovirus-related impacts of social protection policies that aim to reduce the consequences of poverty (eg, conditional cash transfer, housing and public works programmes) alongside continuing research on more conventional vector control interventions.

INTRODUCTION

Arthropod-borne viruses (arboviruses) are transmitted between vertebrate hosts by haematophagous (blood-feeding) arthropod vectors, including mosquitoes and ticks.¹ Arboviruses with mosquito vectors, such as

dengue virus (DENV) and chikungunya virus (CHIKV), are of notable public health importance worldwide owing to their potential to cause explosive outbreaks and induce debilitating and potentially life-threatening disease manifestations.² In addition, congenital arboviral infections, such as with Zika virus (ZIKV), may result in severe congenital malformations with the potential to incur lifelong health and social costs for affected individuals and their families.¹⁻⁴

Infection due to arboviruses with mosquito vectors is becoming increasingly prevalent. The burden of DENV has grown dramatically in recent decades, with substantial impact on morbidity and mortality worldwide, and ZIKV, CHIKV and Yellow Fever virus (YFV) have re-emerged.⁵ Environmental factors, such as climate change (eg, rising temperatures) and habitat modification (eg, deforestation) along with social factors, such as increased international mobility, contribute to the global spread of competent vectors and arboviruses.^{6,7} In regions with established arboviral circulation, community-level factors, such as increased population density, inadequate water management, and poor housing, may exacerbate vector proliferation and elevate infection risks.⁸ This has been reported by several ecological studies, which have shown increased levels of arboviral infections in economically deprived areas at the population-level.⁹⁻¹¹ Furthermore, a recent systematic review employing descriptive synthesis reported a greater presence of *Aedes* mosquito vectors and associated arboviral diseases in regions with lower socioeconomic conditions in 50%–60% of evaluated studies.¹² As described in the early social epidemiology literature, steep inverse associations between social class and mortality from a wide range of diseases exist.¹³ To better understand individual- and household-level risk factors for arboviral infections, we conducted a systematic review and meta-analysis synthesising published evidence on the relationship between markers of socioeconomic position (SEP) and infection due to arboviruses with mosquito vectors.

METHODS

Search strategy and eligibility criteria

The protocol for this systematic literature review was registered in the International Prospective Register of Systematic Reviews (PROSPERO) as CRD42019158572 and was conducted in line with the 2009 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.¹⁴ We searched for studies measuring the association between SEP and arboviral infection published between 1 January 1980 and 30 June 2020 in MEDLINE (PubMed), Embase (Ovid) and LILACS (see online supplemental material 1), hypothesising that studies published more than 40 years prior to this work would lack relevance to current research. The search and full-text review were restricted to articles published in English, Portuguese, Spanish and French. Studies were eligible from any geographic location and

with individuals from any age group, and included peer-reviewed observational case reports, case series or studies that had a cross-sectional, case-control or cohort study design. Studies assessing the association between SEP and/or proxy measures of SEP (eg, individual social class, living conditions, education, employment, household income, race/ethnicity and asset ownership) at the individual-level or household-level and the occurrence of acute, recent or past arboviral infection, indicated by laboratory confirmation, were included. Laboratory confirmation of arbovirus infection was based on the presence of viral RNA, antigen and/or serological evidence (eg, IgM or IgG); the quality of assays used in the individual studies was not appraised. Studies from grey literature, using an ecological design, evaluating the economic burden of arboviral infections, or only describing the natural history of disease were excluded (online supplemental material 2).

Data extraction and meta-analysis

Data on the author, year of publication, study period, study type, source of population, data source, duration of follow-up (if applicable), geographic location, age, sex, individual-level and household-level socioeconomic characteristics, arbovirus infection type, comparison groups, confounders, frequency (number and percentage) and effect estimates (risk ratio (RR) or odds ratio (OR)) were extracted from studies and consolidated. Data screening was conducted in duplicate by four investigators (GMP, LQ, JMP and NSC) and extraction in duplicate by two investigators (GMP and AV). Discrepancies were resolved by consensus. Two reviewers (GMP and LQ) evaluated study quality by conducting a bias assessment using the Newcastle-Ottawa scale (NOS) for individual-level studies (NOS ranges from zero to nine). The NOS form for cohort studies was also used to evaluate data quality for cross-sectional studies; however, the maximum score is limited to six as it was not possible to demonstrate absence of infection at the start of these studies due to the lack of follow-up (online supplemental table 1). Evaluation was performed in duplicate, and discrepancies were resolved by consensus.

When effect estimates were provided for an indicator with comparable parameters in at least three cohort and/or cross-sectional studies, pooled effect sizes and the 95% CIs were calculated using random-effects meta-analyses. Since studies were highly heterogeneous, a random-effects model was preferred.¹⁵ Heterogeneity in RR estimates were assessed using I^2 statistics and Cochran's Q test p values. Case-control studies were not included in the meta-analyses since ORs with 95% CIs were calculated from these study data and, given the high frequency of infections in study populations, were considered to be not directly comparable with cohort and/or cross-sectional relative risk (RR) effect estimates. Further subgroup analyses were conducted for each virus within each of the meta-analyses. Analyses were performed using STATA

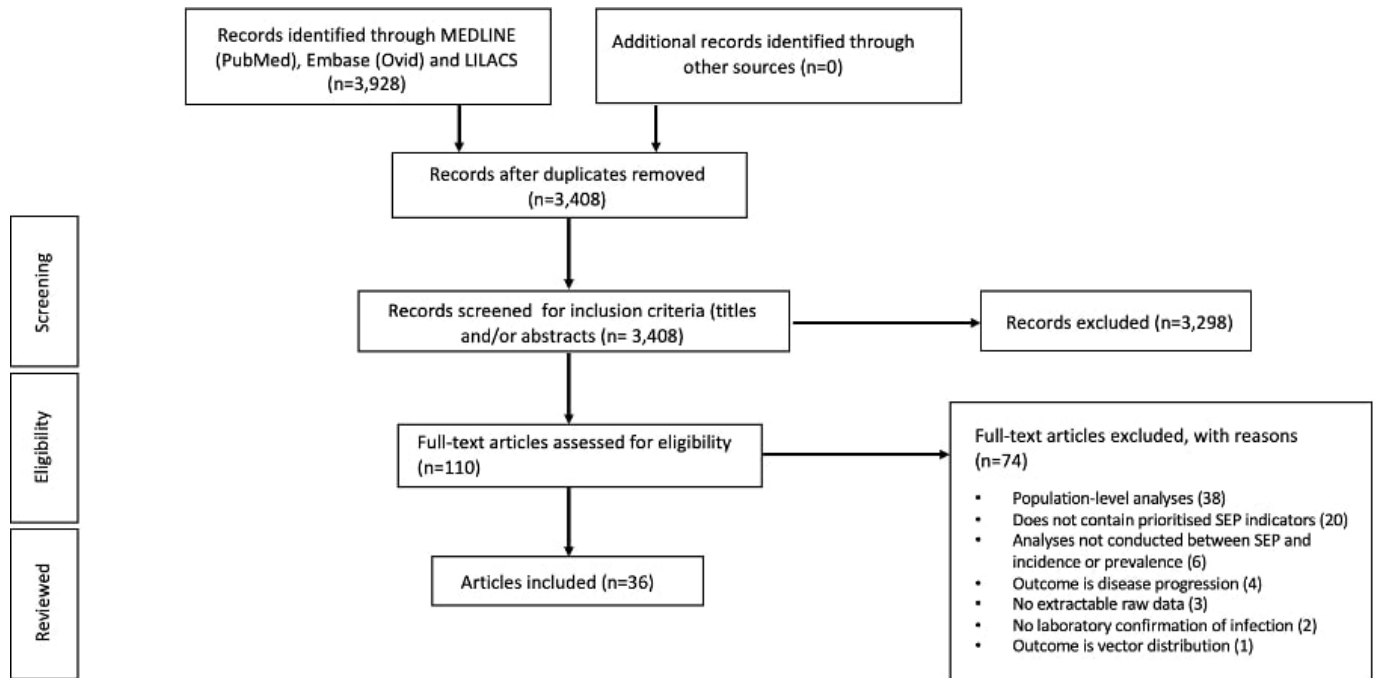


Figure 1 PRISMA flow chart illustrating selection of studies. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; SEP, socioeconomic position.

(V.14.0). A map indicating locations where studies were based was created using Tableau software.

Patient and public involvement

The patients and the public were not involved in the design, conduct or reporting of our research.

RESULTS

Our search generated 3928 published records. After screening titles and abstracts, 110 manuscripts were assessed for eligibility. Of these, 36 articles were deemed eligible for inclusion in this systematic review (figure 1).

All studies included in this review were published between 1995 and 2020, the majority of which were published between 2015 and 2020 (n=28) and focused on DENV (n=21), CHIKV (n=6), Japanese encephalitis (JEV) (n=1), Sindbis virus (SINV) (n=1), West Nile virus (WNV) (n=1), ZIKV (n=1), DENV and JEV (n=2), DENV, CHIKV and Rift Valley fever virus (RVFV) (n=1) and flaviviruses in general with other arboviruses (n=2) (table 1, online supplemental table 2). There were no studies examining YFV. Included studies consisted of 2 cohort studies,^{16 17} 4 case-control studies,¹⁸⁻²¹ 27 cross-sectional studies,²²⁻⁴⁸ 1 nested cross-sectional study within a cohort,⁴⁹ 1 combined cross-sectional and cohort study⁵⁰ and 1 longitudinal serosurvey.⁵¹ Studies were conducted in 23 countries: 4 in low-income countries (Burkina Faso,⁴² Laos³⁵ and Sudan^{26 43}), 14 in lower-middle-income countries (Ecuador,⁴¹ India,¹⁹ Jordan,^{33 37} Kenya,^{17 36} Nicaragua,^{16 50} Nigeria,^{27 31 40} Pakistan,³⁹ Sri Lanka¹⁸ and Vietnam³⁴), 13 in upper-middle income (Brazil,^{23 30 45-47} China,^{20 24 38} Colombia,^{49 51} Malaysia,²⁵ Paraguay⁴⁴ and Thailand²⁹) and 5 in high-income countries/territories

(Mayotte (France),²⁸ French Guiana,²¹ Saudi Arabia,²² Sweden³² and USA⁴⁸) according to the Development Assistance Committee List of Official Development Assistance Recipients (figure 2).

Age and sex

Age and sex were investigated and/or adjusted for in 32 of the 36 studies on seven arboviruses (CHIKV, DENV, JEV, RVFV, SINV, WNV and ZIKV). These studies included three case-control, two cohort, 25 cross-sectional studies, one study comprising a cross-sectional and cohort investigation⁵⁰ and 1 cross-sectional nested in a cohort study, spanning 21 countries.

Of the 20 studies that evaluated the relationship between age and arboviral infection, 18 (90%) reported evidence of an association between increasing age and seropositivity for arboviruses, while four studies (20%) found statistical evidence of an association between age and past arboviral infection (DENV^{23 36 37} and CHIKV⁵⁰) in adjusted models.

All 36 studies considered the direct relationship between sex and arboviral infection or adjusted for sex in the model. Five (13.9%) of these studies reported evidence of higher prevalence of arboviruses among males in crude analyses.^{28 32 39 45 47} However, statistical analyses were not provided for every study, and just eight provided an adjusted point estimate.^{16 23 34 36 37 47 50 51} A study conducted in Sweden³² found a crude statistical association between being male and seropositivity for SINV; however, on adjusting for age and smoking in multivariate analyses, neither sex nor age were significant predictors of seropositivity for SINV. Twenty-four studies with 28 crude estimates comprising a total of 34 373

Table 1 Characteristics of included studies

Author (year)	Country/territory	Study period	Type of study	Population	Type of infection	Diagnostic test	Age range	Total size	Frequency measure	Cumulative incidence	NOS
DENV											
Brunkard <i>et al</i> (2007) ⁴⁸	USA	October 2004–November 2004	Cross-sectional	Probability-based, household selection stratified, multistage, cluster-sampling design	DENV	DENV IgM+; DENV IgG+	All ages	600	P	2%–7.3%; 40%–78%	5
da Silva-Nunes <i>et al</i> (2008) ⁴⁷	Brazil	2004–2006	Cross-sectional	Households in Ramal do Granada, were visited between March and April 2004. 466 dwellers <1–90 years of age (98.5% of the 473 areas permanent residents) were enrolled.	DENV	DENV IgG+	All ages	405	P	18.3%	6
Pessanha <i>et al</i> (2010) ⁴⁶	Brazil	June 2006–March 2007	Cross-sectional	All residents aged over 1 year in the three Belo Horizonte districts (Venda Nova, DS Leste and DS Centro-Sul)	DENV	Not specified	All ages	709	P	11.9% (95% CI 9.7% to 14.6%)	5
Kikutu <i>et al</i> (2015) ⁴⁵	Brazil	2009–2010	Cross-sectional	Individuals seeking medical care for acute febrile illness at the only public emergency health unit	DENV	DENV IgM+ and/or RT-PCR+	>5 years	2962	I	22.0%	5
Pereira <i>et al</i> (2015) ⁴⁴	Paraguay	2014	Cross-sectional	Inhabitants of three villages	DENV	DENV IgG+	All ages	418	P	24.2% (95% CI 20.2% to 28.6%)	5
Soghajer <i>et al</i> (2015) ⁴³	Sudan	2011	Cross-sectional	Randomly selected community population through multi-stage cluster sampling	DENV	DENV IgG+	All ages	540	P	9.4%	6
Fournet <i>et al</i> (2016) ⁴²	Burkina Faso	May 2004–September 2004	Cross-sectional	Children from Ouagadougou districts with different types and degrees of urbanisation	DENV	DENV IgG+	0–12 years	3015	P	22.7%	6
Kenneson <i>et al</i> (2017) ⁴¹	Ecuador	2014–2015	Cross-sectional	Individuals with DENV infections from sentinel clinics - as well as members of the same household and four neighbouring households located within 200 meters	DENV	DENV NS1 RDT+, All ages RT-PCR+ and/or IgM+	All ages	219	P	36.5%	5
Nasir <i>et al</i> (2017) ⁴⁰	Nigeria	May 2016–August 2016	Cross-sectional	Patients with febrile illnesses seeking medical assistance at hospital	DENV	DENV NS1 RDT+; DENV IgG+	1–49 years	171	P	8.8%; 43.3%	3
Khan <i>et al</i> (2018) ³⁹	Pakistan	2013–2015	Cross-sectional	DENV patient samples	DENV	DENV RT-PCR+	All ages	59 765	I	9.2%	4
Liu <i>et al</i> (2018) ³⁸	China	2013–2015	Cross-sectional	Samples selected from a 200,000-sample database holding serum collected from community residents living in Liwan and Yuexiu districts of Guangzhou	DENV	DENV IgM+; DENV IgG+	All ages	2085	P	3.98%; 11.8%	3
Obaidat and Roess (2018) ³⁷	Jordan	2015–2016	Cross-sectional	Healthy relatives of patients at governmental human health centres at 11 governorates	DENV	DENV IgG+	0–80 years	892	P	24.6%	6

Continued

Table 1 Continued

Author (year)	Country/territory	Study period	Type of study	Population	Type of infection	Diagnostic test	Age range years	Total size	Frequency measure	Cumulative incidence	NOS
Piedrahita <i>et al</i> (2018) ⁵¹	Colombia	2010–2012	Longitudinal serosurvey	School children	DENV	DENV IgG+	5–19 years	4385	I	53.8% (2010) to 64.6% (2012)	5
Udayanga <i>et al</i> (2018) ¹⁸	Sri Lanka	February 2017–April 2017	Case–control	Random selection of 200 households reporting past dengue incidence and 200 non-dengue reported households	DENV	N/A	All ages	4000	N/A	N/A	4
Al-Raddadi <i>et al</i> (2019) ²²	Saudi Arabia	2017	Cross-sectional	Residents of the four cities of all genders, age groups, and socioeconomic classes	DENV	DENV IgG+	All ages	6397	P	26.7%	6
Chiaravalloti-Neto <i>et al</i> (2019) ²³	Brazil	October 2015–March 2016	Cross-sectional	Residents of Vila Toninho neighbourhood	DENV	DENV IgG+	>10 y	1322	P	74.6%	8
Jing <i>et al</i> (2019) ²⁴	China	2015	Cross-sectional	850 participants from seven selected communities in Guangzhou with no reported dengue cases before 2014	DENV	DENV IgG+	1–84y	850	P	6.6%	6
Abd-Jamil <i>et al</i> (2020) ²⁵	Malaysia	2007–2010	Cross-sectional	Orange Asli populations residing in eight different villages in the forest or forest fringe areas of Peninsular Malaysia	DENV	DENV IgG+	All ages	491	P	17.0%	6
Eidigall <i>et al</i> (2020) ²⁶	Sudan	August 2017–May 2018	Cross-sectional	Eleven localities of Kassala state	DENV	DENV IgG+	All ages	600	P	11.4%	6
Omatola <i>et al</i> (2020) ³¹	Nigeria	2019	Cross-sectional	Visiting outpatients from the four hospitals in Anyigba	DENV	DENV IgG+	All ages	200	P	20.5%	3
Swain <i>et al</i> (2020) ¹⁹	India	2017	Case–control	Confirmed dengue patients within 1 year in six districts of the state	DENV	DENV IgM+	All ages	767	N/A	N/A	8
CHIKV											
Sissoko <i>et al</i> (2008) ²⁸	Mayotte	2005–2006	Cross-sectional	Household-based; complex multistage cluster sampling of population of Mayotte	CHIKV	CHIKV IgG+	≥2 years	1154	P	37.2%	6
Nakkhara <i>et al</i> (2013) ²⁹	Thailand	2008	Cross-sectional	Residents aged 18 years or more from three villages	CHIKV	CHIKV IgG+	>18 years	507	P	61.9%	5
Kuan <i>et al</i> (2016) ⁵⁰	Nicaragua	March 2015–April 2016	Cross-sectional; Cohort	Children aged 2–14 years enrolled in the Paediatric Dengue Cohort Study; Household recruitment	CHIKV	CHIKV total antibody+	2–14 years; >15 years	3362; 848	P	6.1% (2–14 years); 13.1% (>15 years)	9; 5
Rueda <i>et al</i> (2019) ⁴⁹	Colombia	2014	Cross-sectional nested in community cohort	548 suspected CHIKV patients from the COPCORD cohort	CHIKV	CHIKV IgG+	>18 years	548	P	53.8%	4
Anjos <i>et al</i> (2020) ³⁰	Brazil	2016–2017	Cross-sectional	All households of 3 contiguous valleys in Pau da Lima who are ≥5 years of age	CHIKV	CHIKV IgM+, CHIKV IgG+	All ages	1772	P	11.8%	4

Continued

Table 1 Continued

Author (year)	Country/territory	Study period	Type of study	Population	Type of infection	Diagnostic test	Age range	Total size	Frequency measure	Cumulative incidence	NOS
Omatola <i>et al</i> (2020) ²⁷	Nigeria	2018	Cross-sectional	Febrile participants at five hospitals in Anyigba who test negative for typhoid and malaria	CHIKV	CHIKV IgM+, CHIKV IgG+	All ages	243	P	34.2%	3
JEV											
Luo <i>et al</i> (1995) ²⁰	China	June 1991–September 1991	Case-control	Active case finding in hospitals in Gusi County, Henan, China	JEV	JEV IgG+	>6 months - 10 years	150	N/A	N/A	8
SINV											
Ahlm <i>et al</i> (2014) ³²	Sweden	2009	Cross-sectional	Randomly selected from population registers	SINV	SINV IgG+	25–75 years	1729	P	2.9%	6
WNV											
Obaidat <i>et al</i> (2019) ³³	Jordan	November 2015–May 2016	Cross-sectional	Healthy relatives of patients seeking healthcare at health centres throughout Jordan.	WNV	WNV IgG+	15–50 years	801	P	8.6%	6
ZIKV											
Burger-Calderon <i>et al</i> (2018) ¹⁶	Nicaragua	August 2016–October 2016	Cohort	Laboratory-confirmed Zika index cases and their household members	ZIKV	ZIKV RT-PCR+	All ages	142	I	31.0%	8
Multiple arboviruses											
Bartley <i>et al</i> (2002) ³⁴	Viet Nam	April 1996–August 1997	Cross-sectional	Community and hospital-based subjects	DENV; JEV	DENV or JEV IgG+	All ages	308	P	66.0%	5
Conlan <i>et al</i> (2015) ³⁵	Laos	January 2009–March 2009	Cross-sectional	Random selection of 14 households per village and all household members over 6 years age asked to participate	JEV; DENV	NC; JEV HI+; DENV1 HI+; DENV2 HI+; DENV3 HI+; DENV4 HI+	≥6 years	1136	P	67.3% (Any flavivirus); 39.4% (JEV); 2.2% (DENV 1); 0.8% (DENV2); 0.8% (DENV3); 13.6% (DENV4)	4
Ochieng <i>et al</i> (2015) ³⁶	Kenya	2007	Cross-sectional	HIV-negative blood specimens from the 2007 Kenya AIDS Indicator Survey	CHIKV; DENV; RVFV	CHIKV IgG+; DENV IgG+; RVFV IgG+	15–64 years	1091	P	0.97%; 12.5%; 4.5%	3
Bonifay <i>et al</i> (2017) ²¹	French Guiana	March 2013–June 2014	Case-control	Group of patients infected with CHIKV in 2014 with a group infected with DENV	CHIKV; DENV	CHIKV RT-PCR+; DENV IgM+	>15 years and 3 months	336	N/A	N/A	6

Continued

Table 1 Continued

Author (year)	Country/territory	Study period	Type of study	Population	Type of infection	Diagnostic test	Age range	Total size	Frequency measure	Cumulative incidence	NOS
Horton <i>et al</i> (2019) ¹⁷	Kenya	December 2014–December 2015	Cohort	Acutely ill children presenting at one of four healthcare centres	Flavivirus, CHIKV; DENV	CHIKV IgG+; DENV IgG+	All ages	1604	P	3.7%	6

*The authors report it was not possible to distinguish between DENV and JEV IgG due to cross-reactivity. CHIKV, Chikungunya virus; DENV, Dengue virus; HI, Hemagglutination inhibition; I, Incidence; Ig, Immunoglobulin; JEV, Japanese Encephalitis virus; N/A, not applicable; NC, not clear; NOS, Newcastle-Ottawa scale; NS1, Non-structural protein 1; P, Prevalence; RDT, Rapid diagnostic test; SINV, Sindbis virus; WNV, West Nile virus; ZIKV, Zika virus.

individuals were included in the random-effects meta-analysis of the association of sex and arboviral infection. The crude combined RR for males was 1.1 (95% CI 1.0 to 1.2), with substantial heterogeneity between studies ($I^2=63.4%$) (figure 3A). Disease-specific pooled estimates indicated a RR of 1.1 (95% CI 1.0 to 1.3) and 1.0 (95% CI 0.9 to 1.2) in CHIKV and DENV subgroups, respectively.

Education and occupation

The association between education and arboviral infection was analysed in 1 cross-sectional study nested in a cohort, 2 case-control and 22 cross-sectional studies, spanning 18 countries and 6 arboviruses (CHIKV, DENV, JEV, RVFV, SINV and WNV). In these studies, education was classified in distinct ways depending on context, and included level of education,^{19 24 26 27 29 31–34 36 38–41 43 44 49} schooling age,²³ parental education,²⁰ the attainment of any formal education,^{25 37 42} length of education in years²⁸ and illiteracy.^{30 45}

Overall, there tended to be a higher risk of infection among less educated individuals in crude analyses. However, studies that developed multivariate models indicated weak or no statistical evidence of an association between education and arboviral infection after accounting for confounding factors.^{19 20 23 32 36 37} In addition, a cross-sectional study conducted in China presented evidence that fewer years of parental schooling was associated with increased risk of JEV infection;²⁰ however, on adjusting for JEV vaccination, there was very little evidence remaining. In the 17 investigations (n=15 760) included in the random-effects meta-analysis for education, the crude combined RR for lack of education was 1.5 (95% CI 1.3 to 1.9); however, there was considerable heterogeneity between studies ($I^2=83.1%$) (figure 3B).

Random-effects meta-analysis for disease-specific pooled estimates revealed that individuals with no education had a crude combined RR of 1.5 (95% CI 1.2 to 1.8) for DENV infections and 1.1 (95% CI 0.9 to 1.4) for CHIKV infections.

Occupation was assessed in 11 cross-sectional studies and 1 case-control study. Eleven of the 12 studies presented frequencies, 6 presented crude effect estimates and 2 presented adjusted effect estimates. The occupation-related variables analysed were employment status,^{25 26 30} location of work (inside or outside),²³ earnings (above the country's minimum wage or not),⁴¹ employment stability and occupation types.^{19 27–29 31 40 44} In a study conducted by Chiaravalloti-Neto *et al* in Brazil, there was a crude association between working outside and seropositivity for DENV, which was lost on adjusting for other socioeconomic and demographic covariates.²³ Swain *et al* indicated evidence to suggest that DENV infection was associated with occupations that required travel into certain parts of India.¹⁹ Collectively, in the six studies (n=4056) that were included in the random-effects meta-analysis for occupation, there was little evidence of an association between lack of employment and arboviral infection (pooled RR 0.9; CI 95% 0.7 to 1.3), with

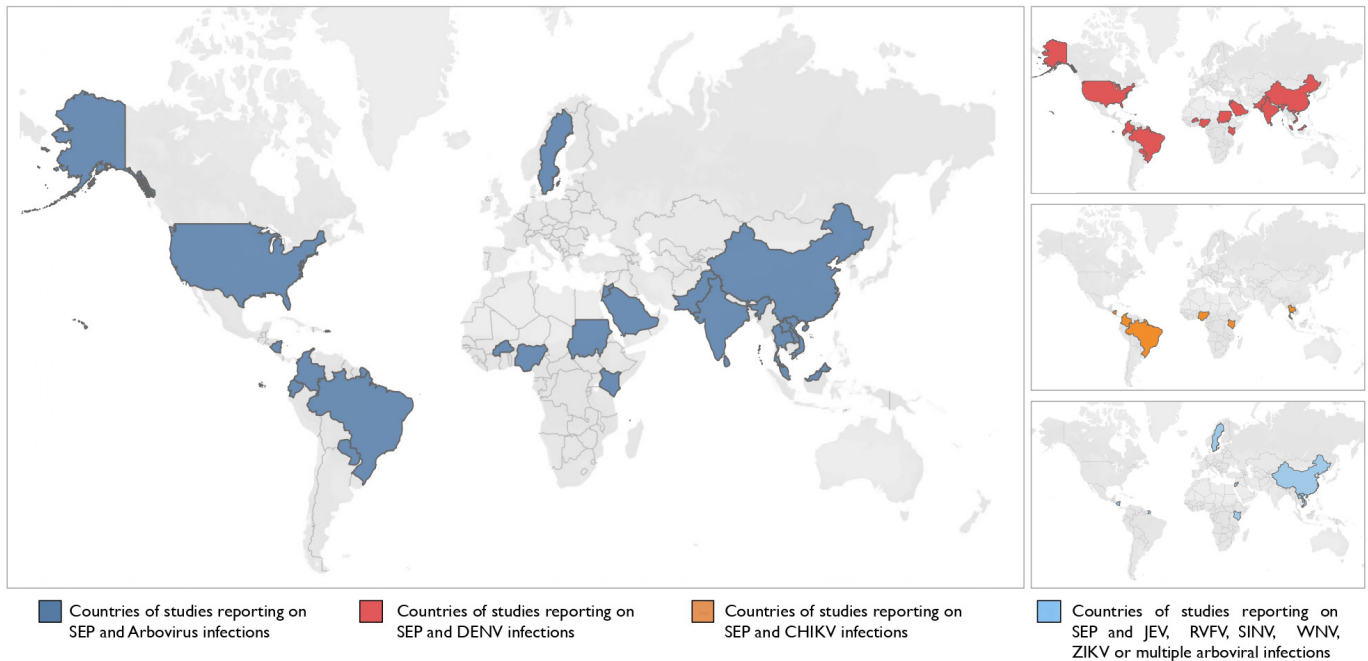


Figure 2 Geographic distribution of studies included in the systematic review. (A) All countries reporting SEP and arboviral infections, (B) Countries reporting SEP and Dengue virus (DENV) infections, (C) Countries reporting SEP and Chikungunya virus infections, (D) Countries reporting on SEP and Japanese encephalitis virus (JEV), Rift Valley fever virus (RVFV), Sindbis virus (SINV), West Nile virus (WNV), Zika virus (ZIKV) or multiple arboviral infections. SEP, socioeconomic position.

considerable heterogeneity between studies ($I^2=75.6\%$) (figure 3C).

Income poverty and social vulnerability

Variables indicating income poverty and social vulnerability varied considerably and thus were challenging to standardise; however, descriptive analyses indicate that lower income was a risk factor for arboviral infection, with limited empirical evidence.

The relationship between poverty or social vulnerability and arboviral infection was assessed in 1 cohort, 4 case–controls and 15 cross-sectional studies, across 16 countries and 4 arboviruses (CHIKV, DENV, JEV and WNV). Assessments were based on weekly or monthly household income,^{18 20 23 25 26 33 39 44–46 48 49} SEP categorised into groups,^{42 49 50} per capita income quartiles or quintiles.^{35 36 47} Health vulnerability was also assessed in two studies.^{21 46} This comprised estimating a health vulnerability index and health vulnerability through state or free care compared with social security and complimentary health insurance. Frequencies and/or effect estimates were extracted for 14. Four studies investigating DENV found evidence of a relationship between lower household income and increased arboviral infection.^{25 45 47 48} One case–control study, conducted in French Guiana, that specifically examined healthcare coverage status in relation to CHIKV and DENV infection, found that a lack of private health insurance was associated with higher CHIKV infection both in the crude and adjusted analyses. In contrast, however, DENV appears to affect a wealthier population.²¹ Since poverty indicators were not measured consistently between studies and study

contexts, a meta-analysis was not possible for income or social vulnerability factors in this study.

Household conditions

Four case–control, three cohort, one longitudinal serosurvey and 18 cross-sectional studies investigated the association between household characteristics and arboviral infections. These studies examined the type or size of residence,^{19 22–24 30 32 34 44 46} house appearance or quality,^{20 28 42} number of rooms,^{22 41} building density,⁴² household crowding,^{17 18 22 23 28 30 31 41 43 44 48 50} type or presence of walls,⁴⁷ wall gaps,⁴⁷ presence of screens,^{41 48} residential area,^{17 21 32 37} waste management^{42 45} and asset ownership (air conditioning,⁴⁸ refrigerator,¹⁶ television,³⁴ land tenure and home ownership^{23 41 47} and asset ownership index (presence of electricity, flush toilet, piped water and possession of a television set, radio or refrigerator).²⁸

Of the four studies that evaluated the association between type of residential area (urban vs rural) and arboviral infections,^{17 32 34 37} one reported higher risk of SINV infection in small, rural residential areas in Northern Sweden,³² one study showed that the risk of flavivirus infection was higher in urban residential areas or cities compared with surrounding rural areas and Southern Vietnam,³⁴ while a study in Kenya observed no difference in flavivirus infection between rural and urban areas but did note a higher seroprevalence among coastal compared with western study participants.¹⁷ In Jordan, a higher risk of WNV infection was reported for those living in Badia and the Jordan Valley regions (arid and hot climates) compared with those living in the

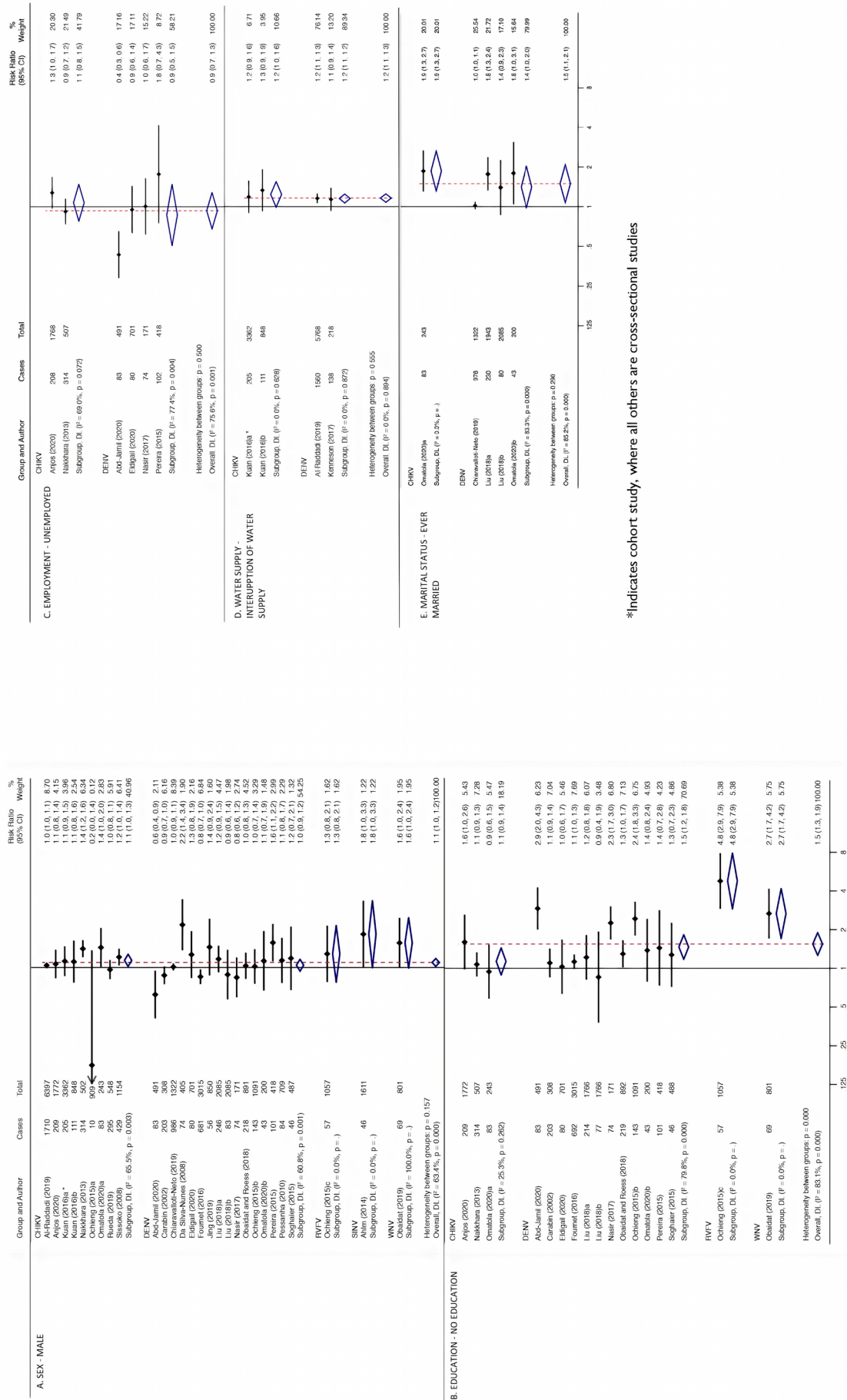


Figure 3 Meta-analysis for the association between socioeconomic risk markers and arboviral infections. Pooled estimates using random-effects meta-analyses are calculated by subgroups of socioeconomic markers, sex (A), education (B), employment (C), water supply (D) and marital status (E). Subgroups of arboviruses are additionally presented per risk marker. Error bars show the point RR with their 95% CIs on the log scale for each study. Diamonds show the combined point estimate. I² statistics and Q-test p values are reported. *Indicates cohort study, whereas all others are cross-sectional studies.

Highlands and Plains regions (colder and higher precipitation areas).³⁷

The relationship between house or land ownership and arboviral infection was evaluated in three studies.^{23 41 47} A cross-sectional study conducted in Brazil showed little evidence of an association between home ownership and seropositivity in DENV, although living in a house compared with an apartment was positively associated with DENV seropositivity, after adjusting for socioeconomic and demographic covariates.²³ Crude analyses indicated evidence of a negative association between land tenure in rural Amazonia, Brazil, and DENV seropositivity; however, this association was weak in the adjusted analysis.⁴⁷

Of the seven studies that analysed building materials, three studies found an association between poor building materials or structures and arboviral infection.^{20 28 30} In addition, unstructured low building density households had higher prevalences of CHIKV and DENV.^{19 20 28 42}

Crowding, categorised by number of individuals per household,^{17 22 23 28 30 43 44 48 50} residents per room⁴¹ or residents per bed²⁷ was analysed in 11 studies, of which four found an association between crowding and arboviral infection.^{23 28 43 50} In a study conducted in Paraguay, DENV prevalence was higher for those who lived alone compared with those who lived with others.⁴⁴

Water supply and sanitation

Water supply or service consumption was investigated in eight studies^{16 22 37 41–44 50} and waste collection or sanitation in three studies.^{22 42 48} Having adequate water supply (ie, tap or piped water) was associated with lower DENV infection in Ecuador⁴¹ and Paraguay.⁴⁴ In addition, water supplied by water wells, onsite water storage and frequent/longer interruptions of water supply was associated with higher flavivirus seroprevalence in Burkina Faso,⁴² higher seropositivity for ZIKV in contacts of ZIKV index cases in Nicaragua,¹⁶ higher DENV infection in Ecuador⁴¹ and Saudi Arabia,²² and higher CHIKV infection in children in Nicaragua.⁵⁰

Improper waste management practices were also significantly associated with flavivirus IgG in different building density strata in Burkina Faso,⁴² while an association was found between lack of street drainage and higher DENV infection on the US/Mexico border.⁴⁸ The absence of sanitation was strongly associated with DENV infection in crude analysis in Saudi Arabia; however, this was not included in the multivariable analysis.²² The random-effects meta-analysis from three studies (one of which contained a cohort (A) and cross-sectional (B) study design) (n=10 196) revealed evidence of an association between interruption of water supply and arboviral infection (RR 1.2; 95% CI 1.1 to 1.3; I²=0.0%) (figure 3D).

Other (marital status, ethnicity and migration status)

A range of other sociodemographic factors that act as proxies for SEP were investigated by several articles identified in this review. Having been born overseas was

associated with greater risk of past arboviral infection, evidenced by one study,²¹ and crude analyses indicated individuals who identified as non-white or of a schedule caste in India, had a higher risk of arboviral infection.^{19 23 45 49} The evidence was limited, concentrated in six countries and largely focused on DENV or CHIKV.

Having been married, including currently or previously (ie, divorced or widowed), was associated with an overall increase in risk of arbovirus infection.^{23 31 38} Marital status and its association with DENV and CHIKV IgG and/or IgM antibody levels was investigated in four cross-sectional studies, conducted in Guangzhou, China,³⁸ São Paulo, Brazil,²³ Guinea Savannah, Nigeria,³¹ and Kogi state, Nigeria.²⁷ In São Paulo,²³ adjusted analyses showed that being single was a risk factor for DENV compared with being married, while in Guangzhou, China,³⁸ crude analyses showed that widowed or divorced individuals were at higher risk of infection compared with both their married and single counterparts. Adjusted analyses from these two studies, however, revealed no statistical evidence of an association. All four studies were included in the random-effects meta-analysis, which revealed statistical evidence that individuals who had ever been married, including currently married, divorced or widowed, had higher overall crude risks of arboviral infection (RR 1.5 95% CI 1.1 to 2.1; I²=85.2%) than those who were single (figure 3E).

Four studies examined race/caste as a correlate of arboviral infection, of which two were conducted in Brazil,^{23 45} one in Colombia⁴⁹ and one in India.¹⁹ The two Brazilian studies found that Black and non-white individuals were at increased risk of DENV^{23 45} and a case-control study conducted in Odisha, India, revealed higher odds of DENV infection in those considered a schedule caste or schedule tribe (official term given in India to those who have historically faced deprivation, oppression and marginalisation) compared with those considered non-schedule caste or non-schedule tribe.¹⁹ The crude analyses showed evidence of this association; however, this was lost on adjusting for unmentioned confounders. A meta-analysis was not performed due to the heterogeneity of study contexts and the countries' specific social constructions of race/caste.

Migration status, defined on the basis of the country of birth: French-born and Foreign-born, was investigated as a potential risk factor for arboviral infection in a case-control study conducted in French Guiana.²¹ This study found strong statistical evidence in crude analysis that individuals born abroad had over four times the odds of testing positive for DENV IgG than those born in French West Indies, French Guiana or Mainland France. One study additionally indicated that changing city within Brazil was not associated with an increase in DENV infection risk.⁴⁶

Quality evaluation

The quality scores of the 36 individual studies varied across study designs. For cross-sectional studies, scores

ranged from 3 to 6, with weaknesses related to selection bias of exposed cohorts and lack of adjustment for confounders. For the cohort studies, scores ranged from 6 to 9, with weaknesses related to no indication of absence of disease at the start of the study and to lack of adjustment for confounders (online supplemental table 1A). For case-control studies, scores ranged from 4 to 8, with weaknesses related to lack of adjustment for confounders (online supplemental table 1B).

DISCUSSION

In this systematic review and meta-analysis, we summarised published evidence linking markers of SEP and infection due to arboviruses with mosquito vectors. Descriptive results indicated lower education, income poverty, low healthcare coverage, poor housing materials, interrupted water supply, marital status (married, single, divorced or widowed), non-white ethnicities and migration status as potential risk factors for arboviral infection. Meta-analyses provided statistical evidence of an increased risk of infection due to arboviruses with mosquito vectors associated with lack of education, interruption of water and having ever been married.

Overall, the seroprevalence of arboviral-specific antibodies (in particular, to DENV) was shown to be highest in older age groups. This finding corroborates a number of studies that found a positive association between age and seropositivity for DENV and is assumed to be related to the longer period of exposure to DENV over time.⁵²⁻⁵⁸ No clear association between arboviral infection and sex was observed.

In addition, individuals with lower education were at greater risk of arboviral infection in both the descriptive summary and meta-analysis. Education is commonly used as a generic indicator for SEP, highlighting the accumulation of advantage and disadvantage over the lifecourse.⁵⁹⁻⁶⁰ It is associated with permanent income status, whereas income itself, for example, captures the level of income at the time of data collection and is thus, in general, volatile. These findings, therefore, might suggest that structural poverty is a relatively more important factor than transient poverty. Education is also argued to capture the knowledge and skill-related assets of an individual, which may contribute to the receptivity of health messaging and thus permitting more informed use of vector control activities to reduce risk of infection.⁶¹

The descriptive analysis for employment assessed several occupations and occupational exposure types, while the meta-analysis looked at unemployment compared with being employed. No overall statistical evidence for unemployment as a risk factor for arboviral infection was apparent. The unobserved effect is likely because the degree of vulnerability linked to unemployment is highly dependent on both the type of employment (indoor or outdoor occupations) as well as the country's overall economic circumstances.⁵⁹ Thus, this

indicator is limited when comparing across studies as well as geographic areas.

Poverty has long been considered a determinant of arboviral infections such as DENV and CHIKV; however, the scarcity of studies with consistent measures of income poverty and social vulnerability has meant that such a relationship has yet to be substantiated. Indeed, in this systematic review, a meta-analysis was not possible for the variables that indicated income poverty and social vulnerability, since contexts within which the data were collected for these were not standardised. Descriptive analyses, nonetheless, indicated that lower income appeared to be a risk factor, although with limited empirical evidence. This is additionally supported by the vast literature on social determinants of health.⁶² Income can influence a variety of material circumstances with direct implications for health and arbovirus exposure.⁶³ The conversion of money and assets into health-enhancing commodities or behaviours may be more relevant to understanding how this variable affects arboviral infection directly.⁵⁹

While a meta-analysis was not completed for the variables related to the constructs of race or caste, the descriptive analysis revealed that individuals who identified as non-white^{23 45} or of a schedule caste¹⁹ were at greater risk of arboviral infection. While there is no biological basis for an association between these constructs and health,⁶⁴ ethnicity, caste and race are proxies for the embodiment of xenophobia, casteism and racism in their structural, cultural and interpersonal forms.⁶⁵ Data from the US context, for example, observed that in areas where mortality rates are highest, the fraction of black residents is larger.⁶⁶ These findings may be extrapolated to the Brazilian context, where racial inequality and segregation are reflected in social disadvantage⁶⁵ and health inequities.

Substandard housing conditions are likely to lead to greater exposure to mosquitoes and thus increased risk of infection.⁶⁷ The association between poor quality housing conditions and arboviral infection was a common finding in many of the studies assessed. However, due to the diversity of indicators relating to household conditions, it was not possible to evaluate this in a meta-analysis. Poor living conditions are often also characterised by overcrowding. Indeed, household crowding appeared to be an additional risk factor for DENV infection. While the reasons behind this are unknown, it is likely due to the association between household crowding and income poverty as well as to the higher concentration of carbon dioxide and other chemicals in crowded houses which attracts a greater number of mosquitoes.⁶⁸ Furthermore, the meta-analysis conducted on water supply in this study provided evidence that interruption in water supply, likely resulting in storage of water in containers and creation of prime breeding spots for mosquitoes,⁶⁹ may increase risk of CHIKV and DENV infection.

The meta-analysis provided evidence that having been married, including currently or previously (ie, divorced or widowed), was associated with an increase in arboviral

infection risk; however, the descriptive analysis indicated that most of these associations diminish after adjusting for confounding. Age may be a particularly important confounder in this context. Migration was assessed in one study and presented descriptively in this analysis. Those classified as migrants were considered to be in a precarious social situation, since they did not have regular social security and health insurance and therefore were more at risk of arboviral infection.²¹

This review has strengths and limitations. First, it is among the first to conduct a systematic review and meta-analysis using diverse populations to assess SEP indicators that identify individuals at the highest risk of arboviral infection. Further research is required to understand the specific mechanisms by which these factors impact infection. The findings of this review should be interpreted with caution, since there were high levels of heterogeneity between studies, which is likely a result of differences in study design, study population and contexts within which these data were collected as well as differences inherent to the individual arboviruses and their mosquito vectors. While this review addressed several arboviruses that circulate in different ecological cycles and involve differences in vector-host preferences, local host abundances and herd immunity, assessing the social determinants of these arboviruses together allows for the analysis of distal risk factors, such as socioeconomic indicators, that have an overarching effect on all arboviral infections.⁷ However, we acknowledge that grouping findings from multiple arboviruses may obscure observations and the heterogeneity of the measures used to capture the range of socioeconomic factors analysed in these studies make it more difficult to delineate associations of interest. Furthermore, this review did not differentiate past infections from current infections and therefore changes in SEP, civil status and even location may have introduced misclassification bias.

CONCLUSION

Evidence from this systematic review suggests that indicators of lower SEP at the individual and household-levels are associated with increased risks of acquiring arboviral infection across a wide range of geographic and cultural contexts. Although not a sufficient determinant of arbovirus risk in itself, poverty is closely correlated with the risk factors for arbovirus infection identified in this review. Within settings experiencing a high burden of arbovirus infections, further work is required to delineate the roles of specific socioeconomic risk factors to inform locally relevant preventive activities. More broadly, the findings of this review underscore the importance of evaluating the arbovirus-related impacts of social protection policies that aim to reduce the consequences of poverty (eg, conditional cash transfer, housing and public works programmes) alongside continuing research on more conventional vector control interventions. To conclude, the findings of this review add to relatively sparse data

on the socioeconomic determinants of infection due to arboviruses with mosquito vectors and emphasise the need for further research to disrupt the cycle of poverty, vulnerability and arbovirus-related illness.

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Funding This work was supported by the British Council Newton Fund (527418645); Wellcome Trust & the UK Department for International Development (205377/Z/16/Z, <https://wellcome.ac.uk/>); Wellcome Trust (202912/Z/16/Z).

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval This study is a systematic review of available literature and did not involve direct access to participants of the primary research studies included. Research ethics approval was therefore not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as online supplemental information.

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Supplementary Material 1. Search strategy used to study socioeconomic factors associated with arboviruses**Pubmed**

("Arboviruses"[Mesh] OR "Arbovirus Infections"[Mesh] OR "Zika Virus"[Mesh] OR "Zika Virus Infection"[Mesh] OR "Dengue"[Mesh] OR "Severe Dengue"[Mesh] OR "Dengue Virus"[Mesh] OR "Chikungunya Fever"[Mesh] OR "Chikungunya virus"[Mesh] OR "Encephalitis, Japanese"[Mesh] OR "Encephalitis Viruses, Japanese"[Mesh] OR "Encephalitis Virus, Japanese"[Mesh] OR "Rift Valley Fever"[Mesh] OR "Rift Valley fever virus"[Mesh] OR "West Nile virus"[Mesh] OR "West Nile Fever"[Mesh] OR "Yellow Fever"[Mesh] OR "Yellow fever virus"[Mesh] OR zika[Title/Abstract] OR zikv[Title/Abstract] OR denv[Title/Abstract] OR dengue[Title/Abstract] OR chikv[Title/Abstract] OR chikungunya[Title/Abstract])

AND

("Social Conditions"[Mesh] OR "Socioeconomic Factors"[Mesh] OR "Social Class"[Mesh] OR "Poverty"[Mesh] OR "Poverty Areas"[Mesh] OR "Income"[Mesh] OR "Education"[Mesh] OR "Educational Status"[Mesh] OR "Ethnic Groups"[Mesh] OR "Race Factors"[Mesh] OR socioeconomic*)

Embase

exp Arbovirus/ OR (Arboviruses or Arbovirus Infections or Zika Virus or Severe Dengue or Dengue Virus or Chikungunya Fever or Chikungunya virus or Japanese Encephalitis Viruse or Rift Valley Fever or West Nile virus or West Nile Fever or Yellow Fever or zikv or denv or chikv).mp.
[mp=title, abstract, heading word, drug trade name, original title, device manufacturer, drug manufacturer, device trade name, keyword, floating subheading word, candidate term word]

AND

exp social class/ OR (Social Conditions or socioeconomic* or Social Class or Poverty or Poverty Areas or Income or Education or Educational Status or ethnic* or race).mp. [mp=title, abstract, heading word, drug trade name, original title, device manufacturer, drug manufacturer, device trade name, keyword, floating subheading word, candidate term word]

LILACS

Arboviruses or Arbovirus Infections or Zika Virus or Severe Dengue or Dengue Virus or Chikungunya Fever or Chikungunya virus or Japanese Encephalitis Viruse or Rift Valley Fever or West Nile virus or West Nile Fever or Yellow Fever or zikv or denv or chikv [Palavras]

AND

Social Conditions or Socioeconomic\$ or Social Class or Poverty or Poverty Areas or Income or Education or Educational Status or ethnic\$ or race or socioeconomic or pobreza or social or renda or educa\$ or raca [Palavras]

Supplementary Material 2: Inclusion and exclusion criteria

Inclusion Criteria	
Types of studies	Publication date: 1980 to 2020
	Studies from any geographical location.
	English, Spanish, Portuguese and French language.
	Studies using quantitative methods: Observational case reports, case series, cross-sectional, case-control and cohort studies.
Types of participants	All ages acceptable.
Types of exposure measures	Socioeconomic position and/or proxy measures of socioeconomic position at an individual level, such as social class, living conditions, education, household income, ethnicity if directly linked to socioeconomic status and asset ownership.
Types of outcome measures	Occurrence of infections due to arboviruses with mosquito vectors

Exclusion Criteria	
Types of studies	Grey literature / not published in a peer reviewed journal, ecological study designs/spatial analyses with no indication of individual risk factors
	Treatment guidelines documents, other systematic reviews
Dates of studies	<1980s
Types of outcome measures	Economic burden of arbovirus (e.g., economic evaluation of costs of disease to families or governments).

Supplementary Table 1. Bias assessment using the Newcastle-Ottawa scale (NOS) for individual-level studies for cross-sectional and cohort (A) and case-control (B)

Author(s)	Year of	Selection (4)			Comparability (2)			Outcome (3)			Total Score	Study design	Study description
		Representativeness of the Exposed Cohort	Selection of the Non-Exposed Cohort	Ascertainment of Exposure	Demonstration That Outcomes of Interest was Not Present at Start	Comparable in Main Factor (age)	Comparability in other controlled factors (i.e. sex)	Assessment of Outcome	Sufficient period of follow up	Adequacy of follow up			
Abd-Jamil, I., et al.	2020	1	1	1	1	1	1	1	1	1	6	Cross-sectional	This present study was performed to investigate the seroprevalence of dengue among the Orang Asli (OA) residing at the forest fringe areas of Peninsular Malaysia and determine the factors that could affect the seroprevalence of dengue.
Ahlin, C., et al.	2014	1	1	1	1	1	1	1	1	1	6	Cross-sectional	Seroprevalence study to determine seroprevalence of SIVV and associated risk factors.
Al-Raddadi et al	2019	1	1	1	1	1	1	1	1	1	6	Cross-sectional	To estimate the seroprevalence of dengue in these regions and the risk factors associated with acute serology.
Anjos et al	2020	1	1	1	1	0	0	1	1	1	4	Cross-sectional	Investigate factors associated with prior CHIKV infection.
Bartley, L.M., et al.	2002	0 (only typhoid patients)	1	1	1	1	1	1	1	1	5	Cross-sectional	Study assesses associations between sociodemographic factors and dengue and Japanese encephalitis seroprevalence in Southern Vietnam.
Brunkard, J. M., et al.	2007	0 (more females than male)	1	1	1	1	1	1	1	1	5	Cross-sectional	Cross-sectional serosurvey used to assess dengue seroprevalence in the southern Texas-Mexico border and assess associated risk factors.
Burger-Calderon, R., et al.	2018	1	1	1	0	1	1	1	1	1	8	Cohort	Study assessing the prevalence of ZIKV and its social determinants in Nicaragua.
Charavallot-Neto, F., et al.	2019	1	1	1	0	1	1	1	1	1	8	Cross-sectional/cohort	Seroprevalence study to determine seroprevalence and incidence of DENV and identify if SES and demographic covariates are associated with seropositivity.
Comlan, J. V., et al.	2015	1	1	1	1	0	0	1	1	1	4	Cross-sectional	Seroprevalence study to determine seroprevalence of Flaviviruses (JEV and DENV) and associated risk factors.
Da Silva-Nunes, M., et al.	2008	1	1	1	1	1	1	1	1	1	6	Cross-sectional	Seroprevalence study to determine seroprevalence of DENV in Amazonian region of Brazil and associated risk factors.
Edigal, M. H., et al	2020	1	1	1	1	1	1	1	1	1	6	Cross-sectional	In the present investigation, a cross-sectional study was conducted to advance an understanding of the prevalence of DENV and associated risk factors were determined in Kozhikode State, India.
Foumet, F., et al.	2016	1	1	1	1	1	1	1	1	1	6	Cross-sectional	Seroprevalence study to analyse Flavivirus prevalence relative to the socioeconomic, demographic, health and environmental data concerning children, their family and household and the district.
Hortton, J., et al.	2019	0 (only acutely ill patients in hospital)	1	1	1	0	0	1	1	1	6	Cohort study	This seroprevalence study aimed to investigate the frequency of alphavirus and flavivirus incident infections in two regions in Kenya and identify potential risk factors.
Hing, Q., et al.	2020	1	1	1	1	1	1	1	1	1	6	Cross-sectional	A cross-sectional serosurvey using a stratified random sampling method among individuals aged 1-84 years-old in 7 communities in Guangzhou with no reported dengue cases before 2014 was performed.
Kemneson, A., et al.	2017	0 (2/3 of cases are referred from MDH health facilities)	1	1	1	1	1	1	1	1	5	Cross-sectional	The authors conducted a household-level study to identify KAP and social-ecological risk factors associated with acute or recent DENV infections in the city of Machakos, Kenya.
Khan, J., et al.	2018	1	1	1	1	0	0	1	1	1	4	Cross-sectional	Conducted enhanced, community-based surveillance in the only public emergency unit in a slum in Salvador, Brazil to identify acute febrile illness (AFI) patients with laboratory evidence of dengue.
Kikuti, M., et al.	2015	0 (only acutely febrile patients)	1	1	1	1	1	1	1	1	5	Cross-sectional	Conducted enhanced, community-based surveillance in the only public emergency unit in a slum in Salvador, Brazil to identify acute febrile illness (AFI) patients with laboratory evidence of dengue.
Kuan, G., et al.	2016	1	1	1	1	1	1	1	1	1	9	Community based cohort (0-14)	Two studies were conducted to analyse the seroprevalence of CHIKV after the first chikungunya epidemic in a community-based cohort of children ages 2-14 years and a cross-sectional survey of persons over 15 years old in the same area of Managua, Nicaragua.
		0 (more females)	1	1	1	1	1	1	1	1	5	Cross-sectional (>15 yo)	
Liu, J., et al.	2018	0 (more females)	1	1	1	0	0	1	1	1	3	Cross-sectional	This cross-sectional study explored the seroprevalence of dengue virus infection in Guangzhou.
Nakkhara, P., et al.	2013	0 (more females)	1	1	1	1	1	1	1	1	5	Cross-sectional	
Nair, I. A., et al.	2017	0 (patients with febrile illness)	1	1	1	0	0	1	1	1	3	Cross-sectional	
Obaidat, M. M. and A. A. Roess	2018	1	1	1	1	1	1	1	1	1	6	Cross-sectional	Seroprevalence study to understand the prevalence of DENV in Jordan and assess risk factors that may be associated with increased seropositivity.
Obaidat, M. M., et al.	2019	1	1	1	1	1	1	1	1	1	6	Cross-sectional	
Ochieng, C., et al.	2015	0 (only HIV negative samples)	1	1	1	1	0 (did not adjust for sex)	1	1	1	3	Cross-sectional	Seroprevalence study to understand the prevalence of DENV, CHIKC and RVFC in Kenya and associated risk factors.
Omatola, C. A., et al.	2020	0 (Only included patients who had fever and suspected dengue or malaria)	1	1	1	0	0	1	1	1	3	Cross-sectional	This study identifies past exposure to DENV among people in Anyigba, located in the Guirga-Saramba region, Nigeria.
Omatola, C. A., et al.	2020	0 (Only patients with febrile illness were included)	1	1	1	0	0	1	1	1	3	Cross-sectional	This study identifies recent CHIKV infection in Anyigba, Nigeria.
Pereira, Y., et al	2015	1	1	1	1	1	0	1	1	1	5	Cross-sectional	Study to establish the seroprevalence of infection by the dengue virus in a district of the Paranaíba region, Brazil.
Pessanha, J.E.M., et al	2010	1	1	1	1	1	0	1	1	1	5	Cross-sectional	Study to determine dengue seroprevalence for to different viral serotypes in three districts in Belo Horizonte, Brazil.
Piedrahita, L. D., et al.	2018	1	1	1	1	1	0	1	1	1	5	Cross-sectional	This longitudinal serological survey and spatial analysis study estimated dengue virus (DENV) transmission in schoolchildren (aged 5-19 years) in Medellín from 2010 to 2017.
Rueda, J. C., et al	2019	1	1	1	1	0	0	1	1	1	4	Cross-sectional	The objective of this study was to describe the demographics and clinical characteristics of suspected chikungunya cases in six Colombian cities.
Sisooko, D., et al.	2008	1	1	1	1	1	1	1	1	1	6	Cross-sectional	Household-based cross-sectional serosurvey to investigate the association between CHIKV seropositivity and risk factors.
Soghater, M.A., et al.	2015	1	1	1	1	1	1	1	1	1	6	Cross-sectional	The objective of this study was to identify socio-demographic factors associated with the prevalence of dengue serotypes in Katsina State in the eastern part of Kaduna State, Nigeria.

Supplementary Table 1. Bias assessment using the Newcastle-Ottawa scale (NOS) for individual-level studies for cross-sectional and cohort (A) and case-control (B)
NOS

Author(s)	Year of publication	Total	Study design	Study description
Bonifay, T., et al.	2017	6	Case (CHIKV)-control (DENV)	Study to describe the socioeconomic indicators of individuals infected with CHIKV and compare to those infected with DENV and the local population.
Luo, D., et al.	1995	8	Case control	Study examines children with Japanese encephalitis and compares them with neighborhood controls matched by age and sex in terms of several social and environmental variables.
Swain, S., et al.	2019	8	Case-control	The study aims to identify the social and ecological factors associated with emerging dengue in Odisha, India.
Udayanga, L., et al.	2018	4	Case-control	Evaluation of demographic, socio-economic and other associated risk factors affecting the

Study ID	Year	Age	Country	Region	Study design	Study population	Study period	Sample size	Sex	Age	Outcome		Outcome		Outcome		Outcome		Outcome		Outcome		Outcome	Outcome
											HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI		
1	2002	65-74	England	Europe	Cohort	Individual	2002-2010	100	50%	65-74	HR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
											95% CI	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2		
2	2002	65-74	England	Europe	Cohort	Individual	2002-2010	100	50%	65-74	HR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
											95% CI	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2		
3	2002	65-74	England	Europe	Cohort	Individual	2002-2010	100	50%	65-74	HR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
											95% CI	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2		
4	2002	65-74	England	Europe	Cohort	Individual	2002-2010	100	50%	65-74	HR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
											95% CI	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2		
5	2002	65-74	England	Europe	Cohort	Individual	2002-2010	100	50%	65-74	HR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
											95% CI	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2	0.8-1.2		

Study ID	Year	Country	Study Design	Exposure	Outcome	Duration	Sample Size	Age Group	Sex	Ethnicity	Setting	Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value	Quality Score	Bias Risk	Notes							
																					Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value	
Study 1	2018	USA	Cohort	Physical activity	Mortality	10 years	10,000	65-75	Male	White	Urban	Physical activity	Sedentary	Mortality	Relative risk	0.85	0.75-0.95	0.001	12	Low	No selection bias, no confounding, no measurement bias, no reporting bias						
																						Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value
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Study 2	2019	UK	Cohort	Dietary intake	Mortality	15 years	15,000	45-65	Mixed	White	Urban	Dietary intake	Low fiber	Mortality	Relative risk	1.25	1.10-1.40	0.001	12	Low	No selection bias, no confounding, no measurement bias, no reporting bias						
																						Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value
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Study 3	2020	Canada	Cohort	Physical activity	Mortality	8 years	8,000	55-75	Mixed	White	Urban	Physical activity	Low	Mortality	Relative risk	0.90	0.80-1.00	0.001	12	Low	No selection bias, no confounding, no measurement bias, no reporting bias						
																						Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value
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Study 4	2021	Australia	Cohort	Dietary intake	Mortality	12 years	12,000	60-80	Mixed	White	Urban	Dietary intake	High	Mortality	Relative risk	0.70	0.60-0.80	0.001	12	Low	No selection bias, no confounding, no measurement bias, no reporting bias						
																						Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value
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																						Intervention	Comparator	Outcome Measure	Effect Size	95% CI	P-value

Study ID	Year	Country	Study Design	Intervention	Comparator	Outcome	Effect Size	95% CI	P-value	Quality of Evidence	Bias	Confounding	Reporting	Other	Overall		Subgroup	
															Mean	SD	Mean	SD
Study 1	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 2	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 3	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 4	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 5	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 6	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 7	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 8	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 9	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 10	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 11	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 12	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 13	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 14	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 15	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 16	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 17	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 18	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 19	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85
Study 20	2018	Kenya	Randomised controlled trial	Vitamin A	Control	All-cause mortality	Relative risk	0.85	[0.75, 0.96]	0.001	Low	No	No	No	0.85	0.85	0.85	0.85
															0.85	0.85	0.85	0.85

Study ID	Year	Country	Study Design	Intervention	Comparator	Outcome	Population	Sample Size	Follow-up	Risk of Bias	Quality of Evidence	Effect Size		95% CI	NNT	IC95%	P-value	I ²	Heterogeneity	Publication Bias	Subgroup
												Mean	SD								
Study 1	2012	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 2	2013	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 3	2014	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 4	2015	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 5	2016	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 6	2017	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 7	2018	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 8	2019	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 9	2020	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	
Study 10	2021	India	Randomized Controlled Trial	Hand hygiene	Control	Incidence of diarrhoeal illness	10-14 years	100	12 months	Low	High	0.15	0.05	0.05-0.25	10	95%	0.001	0	None	None	

Study ID	Year	Country	Study Design	Population	Intervention	Comparator	Outcome	Effect Size	95% CI	P-value	Quality of Evidence	Risk of Bias		Sensitivity Analysis		Subgroup Analysis		Heterogeneity	Publication Bias												
												Low	High	Low	High	Low	High			Low	High										
Study 1	2018	India	Randomized Controlled Trial	Adults	Intervention	Control	Mortality	Relative Risk	0.85	[0.75, 0.95]	0.001	High	Low	High	Low	0.001	0.001	0.001	0.001	0.001											
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.85	[0.75, 0.95]	0.001	High	Low	0.001	0.001	0.001	0.001
Study 2	2019	India	Randomized Controlled Trial	Adults	Intervention	Control	Mortality	Relative Risk	0.75	[0.65, 0.85]	0.001	High	Low	High	Low	0.001	0.001	0.001	0.001	0.001											
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.75	[0.65, 0.85]	0.001	High	Low	0.001	0.001	0.001	0.001
Study 3	2020	India	Randomized Controlled Trial	Adults	Intervention	Control	Mortality	Relative Risk	0.65	[0.55, 0.75]	0.001	High	Low	High	Low	0.001	0.001	0.001	0.001	0.001											
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.65	[0.55, 0.75]	0.001	High	Low	0.001	0.001	0.001	0.001
Study 4	2021	India	Randomized Controlled Trial	Adults	Intervention	Control	Mortality	Relative Risk	0.55	[0.45, 0.65]	0.001	High	Low	High	Low	0.001	0.001	0.001	0.001	0.001											
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001
																					Intervention	Control	0.55	[0.45, 0.65]	0.001	High	Low	0.001	0.001	0.001	0.001

Resumo

Introdução

Os arbovírus possuem notável importância em saúde pública em todo o mundo devido ao seu potencial de causar grandes surtos e de gerar manifestações clínicas debilitantes ou possivelmente fatais. Esta revisão sistemática e meta-análise tem como objetivo avaliar a relação entre indicadores de posição socioeconômica (SEP, sigla em inglês) e infecção por arboviroses com mosquitos vetores.

Métodos

Realizamos uma busca sistemática nas bases de dados Pubmed, Embase e LILACS para identificar estudos publicados entre 1980 e 2020 medindo a associação entre marcadores de SEP e infecção por arboviroses. Incluímos os estudos observacionais, sem realizar restrições por localização geográfica ou idade. Excluímos estudos da literatura cinzenta, revisões e ecológicos. Os dados dos estudos foram extraídos, resumidos e utilizada a meta-análise com efeitos aleatórios para obtenção das estimativas combinadas de efeito.

Resultados

Identificamos 36 estudos observacionais usando dados referentes a 106.524 participantes do estudo em 23 localizações geográficas que examinaram empiricamente a relação entre indicadores socioeconômicos e infecções causadas por sete arbovírus (vírus da Dengue, Chikungunya, Encefalite Japonesa, Febre do Vale do Rift, Sindbis, Febre do Nilo Ocidental e Zika). Embora os resultados tenham variado, a síntese

descritiva apontou um maior risco de infecção por arboviroses associado a indicadores de SEP mais baixos, incluindo menor escolaridade, menor renda, baixa cobertura de saúde, materiais de habitação precários, abastecimento de água interrompido, estado civil (casado, solteiro, divorciado ou viúvo), etnias não brancas e status migratório. As estimativas brutas combinadas indicaram um risco aumentado de infecção por arboviroses associado à baixa escolaridade (RR = 1,5 IC 95%: 1,3, 1,9); I²=83,1%), interrupção do abastecimento de água (RR = 1,2; IC 95%: 1,1,1,3; I² = 0,0%) e ser casado anteriormente (RR = 1,5 IC 95%: 1,1, 2,1; I²=85,2%).

Conclusão

As evidências dessa revisão sistemática sugerem que pior SEP aumenta o risco de adquirir infecção por arbovírus. No entanto, houve grande heterogeneidade entre os estudos. Mais estudos são necessários para definir a relação entre indicadores específicos de SEP a nível individual, domiciliar e comunitário e a infecção por arbovírus para informar intervenções direcionadas de saúde pública.

Palavras-chave: posição socioeconômica, equidade em saúde, infecção por arbovírus, mosquitos vetores, revisão sistemática

Resumen

Introducción

Los arbovirus son de notable importancia para la salud pública global por su potencial de causar brotes explosivos además de manifestaciones clínicas debilitantes y potencialmente letales. Esta revisión sistemática y meta-análisis tiene como objetivo la evaluación de la relación entre indicadores de posición socioeconómica (PS) e infecciones por arbovirus transmitidos por mosquitos vectores.

Métodos

Realizamos una búsqueda sistemática en las bases de datos Pubmed, Embase, y LILACS para identificar estudios publicados entre 1980 y 2020 que median la asociación entre marcadores de PS e infección arboviral. Incluimos estudios observacionales sin restricciones sobre la localidad geográfica o edad de los participantes. Excluimos estudios de la literatura gris, revisiones y estudios ecológicos. Los hallazgos de los estudios fueron extraídos y resumidos y se realizaron meta-análisis de efectos aleatorios para obtener estimaciones combinadas de efecto.

Resultados

Identificamos 36 estudios observacionales con datos pertenecientes a 106,524 participantes de 23 localidades geográficas que examinaron empíricamente la relación entre factores socioeconómicos e infecciones causadas por siete arbovirus (Dengue, Chikungunya, Encefalitis Japonesa, Fiebre del Valle de Rift, Sindbis, Fiebre del Nilo Occidental, y el Zika). Mientras que los resultados fueron variados, la síntesis descriptiva señaló un riesgo mayor de

infección arboviral asociada con marcadores de PS más bajos, incluyendo menor nivel educativo, escasez de ingresos, baja cobertura de saneamiento, materiales de viviendas de baja calidad, interrupciones del suministro de agua, estado civil (casado, soltero, divorciado o viudo), grupos étnicos no caucásicos y estatus migratorio. Las estimaciones agrupadas brutas indicaron un riesgo aumentado de infección arboviral asociado con menor nivel educativo (RR = 1.5 95% CI: 1.3, 1.9); $I^2=83.1\%$), interrupciones del suministro de agua (RR = 1.2; 95% CI: 1.1,1.3; $I^2 = 0.0\%$) y haber estado casado (RR = 1.5 95% CI: 1.1, 2.1; $I^2=85.2\%$).

Conclusión

Esta revisión sistemática señala que el tener una PS inferior aumenta el riesgo de adquirir infección arboviral, sin embargo hubo una gran heterogeneidad entre los estudios. Más estudios son necesarios para mejor definir la relación entre indicadores de PS individuales, a nivel de hogar, y a nivel comunitario y la infección arboviral para mejor diseñar intervenciones de salud pública dirigidas.