Mosquito saliva enhances virus infection through sialokinin-dependent vascular leakage

Daniella A. Lefteria, Steven R. Bryden, Marieke Pingena, Sandra Terryb, Ailish McCafferty, Robert M. Waterhouse, Sandra Terryb, Ailish McCaffertya, Kave Shamsc, Emilie Pondevilleb, Margus Varjakb, Emily F. Beswick, Alphonse G. Hoerauf, Georgi Georgiev, Marleen Van der Laana, Valeria Mastrullo, Paola Campagnolo, Robert M. Waterhouse, Margus Varjak, Andres Merits, Rennos Fragoudis, Stephen Griffinh, Clive S. McKimmie, and Carolina Barillas-Mury

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Significance

An increasingly important group of infectious agents is viruses spread by mosquitoes. When infected mosquitoes bite people, they inject virus and saliva into the skin. The saliva is biologically active and helps the mosquito feed but also helps the virus to infect and replicate. Sialokinin is a peptide in mosquito saliva that is unique within the insect world as having vertebrate-like tachykinin function, which can be proinflammatory. We show that sialokinin increases blood vessel permeability, allowing an influx of virus-permissive cells. Increasing the frequency of these cells in skin provides more opportunities for the virus to replicate. Inhibiting mosquito salivary factors holds promise as a panviral strategy for reducing disease burden caused by a wide variety of mosquito-borne viruses.

Virology

Mosquito-borne viruses are an increasingly important global cause of disease. Defining common determinants of host susceptibility to this large group of heterogeneous pathogens is key for informing the rational design of panviral medicines. Infection of the vertebrate host with these viruses is enhanced by mosquito saliva, a complex mixture of salivary-gland-derived factors and microbiota. We show that the enhancement of infection by saliva was dependent on vascular function and was independent of most anti-saliva immune responses, including salivary microbiota. Instead, the Aedes gene product sialokinin mediated the enhancement of virus infection through a rapid reduction in endothelial barrier integrity. Sialokinin is unique within the insect world as having a vertebrate-like tachykinin sequence and is absent from other insects. Although in Aedes aegypti, e.g., Rift Valley fever virus and Cache Valley virus (18, 20, 21), and in Aedes aegypti-infected mosquitoes bite people, they inject virus and saliva into the skin. The saliva is biologically active and helps the virus to feed but also helps the virus to infect and replicate. Sialokinin is a peptide in mosquito saliva that is unique within the insect world as having vertebrate-like tachykinin function, which can be proinflammatory. We show that sialokinin increases blood vessel permeability, allowing an influx of virus-permissive cells. Increasing the frequency of these cells in skin provides more opportunities for the virus to replicate. Inhibiting mosquito salivary factors holds promise as a panviral strategy for reducing disease burden caused by a wide variety of mosquito-borne viruses.

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The authors declare no competing interest.

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1To whom correspondence may be addressed. Email: c.s.mckimmie@leeds.ac.uk.

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and expressed specifically by female mosquitoes. *Anopheles* species mosquito saliva lacked the ability to enhance *Aedes* or *Anopheles*-borne virus infection, and this correlated with an inability to induce vascular leakage. We exploited this naturally occurring difference to dissect the mechanistic basis by which *Aedes* saliva enhances infection with virus and to identify the requirement for the *Aedes* salivary factor sialokinin (SK). As such, we define an aspect of disease transmission that can inform the development of medicines with panviral potential that target SK.

**Results**

**Mosquito Saliva Is Sufficient to Enhance Virus Infection and Worsen Clinical Outcome.** Mosquito bites involve skin tissue trauma and the deposition of saliva. Host responses to biting enhance incoming virus infection, as does the injection of experimentally derived homogenates of salivary gland tissue (6, 31). To define the factors in isolated saliva responsible for modulating host susceptibility to coinoculated virus in mice, we obtained saliva by forced salivation from adult *Aedes aegypti* female mosquitoes and coinjected this into skin with either the Alphavirus SFV or Flavivirus ZIKV (Fig. 1 A and B and SI Appendix; Fig. S1 A and B). Although the presence of just one mosquito’s salivation was sufficient to induce a 10-fold increase in virus by 24 h postinfection (hpi) (Fig. 1A), this was variable, likely as a consequence of salivation efficiency between mosquitoes (SI Appendix, Fig. S1 C). Variation was much less pronounced when mice were coinjected with saliva obtained from 5 mosquitoes (Fig. 1 A and B). Repeated tissue piercing with our hyperfine needles in the absence of saliva, to simulate probing by mosquitoes, did not result in any virus enhancement (Fig. 1C). Coinoculation of virus with saliva also reduced survival to SFV to a similar extent as injection of the same titer virus into a mosquito bite (Fig. 1 D). Increased quantities of virus RNA in skin were apparent from as early as 10 hpi when virus was coinoculated with saliva, and additionally enhanced the early dissemination of virus to the draining lymph node (Fig. 1 E and F).

Host inflammatory responses to mosquito biting that recruit virus-permissive monocytic cells are important for modulating host susceptibility to virus (18). Here, we found that, similar to mosquito biting, saliva alone in the absence of virus induced the expression of proinflammatory genes *ccl2, il1b, and ccl2*, while prototypic antiviral type I interferon (IFN)-stimulated genes (*igt15* and *ccl5*) whose expression correlates with host resistance to infection (32) were not altered (SI Appendix, Fig. S1 D). In summary, we have optimized and defined our in vivo model for determining how saliva alone, in the absence of other mosquito bite or salivary gland tissue components, modulates host susceptibility to virus.

**Enhancement of Virus Infection In Vivo Requires Processes That Are Absent Ex Vivo.** To define whether mosquito saliva can directly modulate the susceptibility of the two principal cell types

![Fig. 1](https://www.pnas.org/content/earlypub/2022/2114309119/F1.large.jpg)
infected by SFV in the skin (fibroblasts and macrophages), we infected primary cultures of these cells in vitro in the presence or absence of saliva. However, the in vitro addition of mosquito saliva failed to recapitulate the virus-enhanced infection phenotype observed in vivo (SI Appendix, Fig. S2 A–D). This was a direct effect of saliva upon cells, rather than via direct action on virus, as cells pretreated with saliva and washed prior to infection (referred to here as saliva to cells) also exhibited increased resistance to infection (SI Appendix, Fig. S2 B and D). This in vitro virus infection inhibition by saliva was likely dependent on cellular responses activated by bacteria, as it was not observed with saliva from females treated with broad-spectrum antibiotics (SI Appendix, Fig. S2 D–F). *Aedes* salivary glands host a diverse microbiota (29) that may be secreted directly into saliva. However, it is not clear to what extent the mosquito salivary gland microbiota is secreted in vivo and whether it plays an important role in arboviral transmission.

In summary, these in vitro studies failed to recapitulate our in vivo phenotype in which virus infection is enhanced by saliva. To test whether crosstalk by skin cells or the presence of other cell types not present in the above cultures was required to mimic the in vivo phenotype, we assessed the susceptibility of intact skin explants to infection (32). However, these explants did not exhibit any increase in susceptibility to virus following exposure to saliva. This included the infection of explants derived from resting skin, mosquito-bitten skin, and explants derived from skin injected with saliva prior to biopsy (SI Appendix, Fig. S2 G and H). Together, this suggests that a key in-vivo–specific process was required for the saliva to enhance virus infection.

**A Heat-Sensitive Salivary Factor from Female Mosquitoes Enhances Virus Infection Independent of Bacterial Microbiota.** Next, we wanted to define which component in saliva was responsible for enhancing virus infection in vivo. Saliva from antibiotic-treated mosquito saliva was less able to upregulate the inflammatory chemokine *cxcl2* in vitro (SI Appendix, Fig. S2A). Because some inflammatory responses in vivo to mosquito biting can enhance infection with virus (18), we hypothesized that microbiota may account for the ability of saliva to enhance virus infection in vivo. However, while microbiota-depleted saliva induced lower quantities of inflammatory gene expression in skin (Fig. 2A), the microbiota was dispensable for the infection-promoting ability of saliva in vivo (Fig. 2B). Instead, the ability of saliva to enhance infection was sensitive to protein-denaturing temperatures (Fig. 2C), suggesting the proviral factor in saliva is likely a proteinaceous factor expressed and secreted by the mosquito salivary gland.

Many female-specific mosquito salivary components have evolved to facilitate blood feeding. In contrast, male mosquitoes that do not bite are naturally deficient in these blood-feeding factors (33). To determine whether the virus-enhancing salivary factor is one of these factors, we compared the ability of saliva from coreared male and females to modulate host susceptibility to virus. Crucially, female *Aedes* saliva enhanced virus infection to a significantly higher extent than male saliva. This was the case when either equal volumes of male/female saliva were used (Fig. 2D) or when saliva quantities were normalized for protein content, which was substantially lower in male saliva isolates (SI Appendix, Fig. S3A). Because blood feeding causes changes in gene expression within the salivary glands (34), we also assessed how saliva acquired from previously blood-fed and exclusively sugar-fed *Ae. aegypti* females modulated virus infection in mice. However, saliva from either group possessed similar virus-enhancing properties (SI Appendix, Fig. S3B). Thus, a female-specific salivary factor, which may have evolved to support efficient blood feeding, was responsible for enhancing the susceptibility to virus infection.

**Fig. 2.** A heat-sensitive salivary factor from female mosquitoes enhances virus infection independent of bacterial microbiota. (A) Mouse skin was injected with 1.86 μg of saliva from control and antibiotic (Abx)-treated mosquitoes, and host expression of *cxcl2, il1b*, and *ccl2* transcripts was assessed at 6 h (n = 6). (B–D) Mouse skin was inoculated with 104 PFU of SFV4 alone or with *Ae. aegypti* saliva in the upper skin of the left foot. Viral RNA and host 18S were quantified from skin and spleen by qPCR and viral titers of serum by plaque assays at 24 hpi. (B) Female *Ae. aegypti* saliva from Abx-treated or untreated mosquitoes (n > 6). (C) Heat-treated (10 min at 95 °C) or untreated female *Ae. aegypti* saliva (n = 6). (D) Male or female *Ae. aegypti* saliva pooled from five mosquitoes combined, reared in the same cage (n = 6). *P < 0.05, **P < 0.01, ***P < 0.001, ns = not significant.
Anopheles Mosquito Saliva Lacks the Ability to Enhance Virus Infection. We next assessed whether saliva from other blood-feeding mosquito species has similar virus-infection-enhancing properties. It is not clear whether enhancement by saliva from mosquito species belonging to Aedes and Culex genera (Culicinidae subfamily, Culicidae family), which are also important arbovirus vectors, is comparable in its ability to modulate vertebrate host susceptibility to virus. Neither is it clear if this is a general phenomenon elicited by all blood-feeding mosquito species saliva or whether saliva from arbovirus-vector–incompetent mosquitoes such as Anopheles (Anophelinae subfamily, Culicidae family) can also enhance infection. Because each mosquito genus has differing competency in transmitting distinct viruses to different vertebrate species, we compared how saliva from each mosquito genus modulates vertebrate susceptibility to one reference virus (SFV) in one vertebrate host (mice). We found that Aedes albopictus and Culex pipiens saliva enhanced infection, with either SFV or ZIKV in vivo, to a similar extent as Ae. aegypti saliva (SI Appendix, Fig. S4 A and B). In comparison, Anopheles species mosquito saliva (Anopheles gambiae and Anopheles stephensi) could not (Fig. 3 A–C and SI Appendix, Fig. S4 C–E). Furthermore, unlike Ae. aegypti-bitten mice, An. gambiae–bitten mice exhibited similar host susceptibility to infection as resting skin (SI Appendix, Fig. S4C).

To exclude the possibility that Anopheles saliva may contain a unique factor that inhibited virus infectivity, we assessed whether saliva modulated virus infection in vitro. Similar to Aedes saliva, Anopheles saliva possessed some ability to protect cells from infection (Fig. 3D). However, in BHK-21 cells that lack IFN signaling, both species’ saliva had no effect on the ability of virus to replicate (Fig. 3E). In addition, we found that by mixing Aedes and Anopheles saliva together, we were able to restore the enhancement of virus infection in vivo (Fig. 3F). Together, this suggests that the ability to enhance virus infection is mosquito species specific and that the factor responsible in Aedes sp. saliva is missing in Anopheles saliva.

In comparison to Aedes mosquitoes, the widely distributed group of Anopheles species mosquitoes is not able to efficiently transmit arboviruses, despite them sharing an ecological niche that overlaps with arbovirus endemic areas (35, 36). The sole exception to this is the Anopheles-transmitted O’nyong’nyong virus (ONNV), whose genetic sequence is highly similar to CHIKV and similarly disseminates to joints (37, 38). While both Aedes and Anopheles have evolved salivary factors that facilitate efficient blood feeding (30), it is not clear why Anopheles sp. mosquitoes are such poor vectors of virus per se (37). It is conceivable that one reason is the absence of salivary factors that enhance the infection of virus in the mammals they feed on. To help explore this, we investigated whether Anopheles saliva, or indeed Aedes saliva, could modulate the infection of mice with ONNV, utilizing a newly developed ONNV mouse model in which virus efficiently replicated and disseminated from the skin inoculation site to joint tissue (Fig. 3G). Importantly, neither mosquito species saliva (Fig. 3H) or indeed the presence of a mosquito bite (SI Appendix, Fig. S4F) could modulate host susceptibility to infection with ONNV. Instead, ONNV was able to infect mice robustly, induce viremia, and disseminate to joint tissue without the need for saliva-based enhancement of infection, perhaps reflecting the absence of virus-infection–enhancing factor(s) present in its natural anopheline vector. Saliva from either species was also not able to modulate the ability of ONNV to infect and replicate in cultured cells (SI Appendix, Fig. S4G).

Increased Vascular Permeability Induced by Aedes Saliva Enhances Virus Infection. Aedes mosquito biting induces the recruitment of monocytes in cells, which can be infected by and replicate virus (6, 18). We hypothesized that Anopheles and Aedes saliva differ in their potency for activating key proinflammatory responses that attract monocytes. However, we found that the induction of most inflammatory responses to Aedes and Anopheles saliva was similar at the transcript level, including the key monocyte chemoattractant ccl2, suggesting that innate immune sensing for both was broadly analogous (Fig. 4A). Anopheles saliva more potently upregulated some IFN-stimulated genes, including rdad2 and ccl5 (Fig. 4 A and B). However, a more potent induction of antiviral IFNs by saliva did not account for its ability to modulate virus infection, as the enhancement of both SFV and ZIKV infection by Aedes saliva was IFN independent (Fig. 4C, for SFV; Fig. 3A and SI Appendix, Fig. S4C, for ZIKV). Importantly, we found that despite similar ccl2 chemokine expression, Aedes saliva resulted in a robust and rapid influx of mononuclear cells by 2 h, while Anopheles saliva did not (Fig. 4D).

Heightened skin inflammatory responses to saliva can also occur in those previously exposed to biting mosquitoes. However, we found that the vaccination of mice with saliva prior to infection did not result in any further modulation of host susceptibility to virus. Here, mice were immune sensitized to saliva or saline control by four weekly injections and on week five were injected with either SFV alone or SFV with saliva. At 2 hpi, saliva-primed mice exhibited significantly elevated draining lymph node cellularity and IL-5 and IL-13 expression and increased serum IgE indicative of immune sensitivity to saliva (Fig. 4E and SI Appendix, Fig. S5 A and B). However, Aedes–saliva–primed mice showed no difference in their susceptibility to SFV when compared to saliva-naïve mice (Fig. 4F). Anopheles–saliva–primed mice also exhibited no difference in their susceptibility to virus, such that they were similar to saliva-naïve mice injected with virus alone (Fig. 4G). Thus, immune sensing of mosquito saliva alone is not sufficient to enhance virus infection.

During our experiments above, we had anecdotally noted that Anopheles saliva, unlike Aedes saliva (11), did not cause a noticeable amount of fluid to accumulate at the site of inoculation. Components of Aedes saliva are known in an experimental setting to vasodilate constricted aortic ring tissue (39), although its effect on dermal postcapillary venules (through which most leukocyte entry occurs) is not described. We hypothesized that Aedes saliva may lead to an unusually rapid influx of mononuclear cells (Fig. 4D) through its ability to induce skin vasculature barrier leakage directly and thereby underpin its ability to enhance virus infection. We therefore quantified levels of edema as a measure for vasculature barrier leakage following injection with saliva or exposure to mosquito biting; both Ae. aegypti saliva and bites led to significantly more edema than that by An. gambiae (Fig. 5 A and B and SI Appendix, Fig. S6 A and B).

We next asked whether the induction of vascular leakage was immune mediated or due to direct action of salivary factors on the vasculature. Histamine is the principal host factor that mediates early/immediate edema in response to exogenous agents, infection, or injury (40), and its injection into skin made mice more susceptible to SFV (Fig. 5C), suggesting that endothelial leakage alone is sufficient to promote virus infection. However, we found that histamine was not necessary for host responses to saliva, as treatment with antihistamines did not significantly suppress saliva-induced edema (although
Fig. 3. Anopheles mosquito saliva lacks the ability to enhance virus infection. (A–C) Mouse skin was inoculated with either 10^6 PFU of ZIKV or 10^4 PFU of SFV4 alone or with 1.86 μg of saliva of either Ae. aegypti, An. gambiae, or An. stephensi. Virus RNA and host 18S and serum viral titers were quantified at 24 hpi. (B) Survival of mice infected with 4 × 10^4 PFU of SFV4 stock. (D) Macrophages were infected with luciferase-expressing SFV at an multiplicity of infection of 0.1 alone or with 0.66 μg of protein of Ae. aegypti, An. gambiae, or Ae. albopictus saliva. Luciferase activity of tissue culture media was assayed at 6 hpi (n = 6). (E) BHK cells were infected with 10-fold serial dilutions of SFV4 ranging between 25,000 PFU and 0.25 PFU alone or with Ae. aegypti or An. gambiae saliva and then immediately overlayed with Avicel. PFUs were then assessed at 48 hpi. Shown here are representative PFUs for wells in which plaques were quantifiable. (F) Mouse skin was inoculated with 10^6 PFU of SFV4 alone or alongside 1.86 μg of saliva of Ae. aegypti, An. gambiae, or both species. Virus RNA and host 18S were quantified from skin and spleen by qPCR and viral titers of serum by plaque assays at 24 hpi. (G) Mice were treated with 1.5 mg of anti-IFNAR antibodies (clone MAR1-5A3) and 24 h later were infected with 2 × 10^5 PFU ONNV s.c. in the skin (upper side of the left foot). (G) ONNV RNA quantities in tissues at 48 hpi were defined by qPCR to define tissue tropism. (H) Mouse skin was infected with ONNV alone or alongside 1.86 μg of saliva of either Ae. aegypti or An. gambiae. ONNV RNA and host 18S from tissues were quantified by qPCR, and serum viral titers were quantified via plaque assays at 48 hpi. *P < 0.05, ***P < 0.001, ****P < 0.0001, ns = not significant.
control histamine-induced edema was suppressed) or virus enhancement by saliva (Fig. 5 D and E). This agrees with studies in humans, in which prior histamine administration had little effect on skin inflammatory responses to *Aedes* mosquito biting (41). Together with our findings above (Fig. 4F) that immune hypersensitization to saliva did not modulate host susceptibility to virus, we concluded that a factor in *Aedes* saliva acts directly on blood vasculature to induce barrier leakage, in an immune-response–independent manner. Therefore, we decided to assess whether endothelial cell barrier function was directly modulated by saliva and if this was mosquito species specific. Here, a monolayer of primary blood endothelial cells was allowed to reach confluence and establish a barrier, as assessed by electrical resistance. Monolayers were then treated with either control saline or *Ae. aegypti* or *An. gambiae* saliva, and electrical resistance across the monolayer was assessed longitudinally. Importantly, cultured endothelial cell monolayers responded to *Aedes* saliva, but not *Anopheles* saliva, by rapidly decreasing endothelial barrier function (Fig. 5F).

Collectively, by comparing host responses to *Aedes* saliva and *Anopheles* saliva, we have identified blood vascular leakage as a key feature associated with female *Aedes* saliva enhancement of
Aedes SK Is Sufficient to Induce Blood Vasculature Barrier Leakage and Enhance Virus Infection. The saliva of blood-feeding arthropods contains many factors that facilitate blood feeding, putatively via altered vascular function. In *Ae. aegypti*, one such hypothesized factor is SK, a 1,400-Da peptide product of the pro-SK precursor–encoding gene (AEL000229). SK belongs to the family of tachykinin-like peptides (39, 42) and is expressed in the female salivary gland (43). We sought to define the evolutionary conservation of homologs in other blood-feeding insects and determine whether SK has any role in modulating mammalian vascular barrier function and thereby host susceptibility to virus infection.

Tachykinin-related peptides (TRPs) have been identified from many insect species (44), and in *Ae. aegypti*, five TRPs are encoded by Tachykinin (AAEL006644), a gene with single-copy orthologs across protostome invertebrates (45, 46). Intriguingly, we found that the SK tachykinin-like peptide (NTGDFKFYGLMamide) more closely resembles typical deuterostome-type (FXGLMamide) than protostome-type (FXGXRamide) peptides (Fig. 6D). In addition, SK is also recognized by the PROSITE tachykinin family signature pattern (PS00267), which matches deuterostome-type but not protostome-type peptides. Together, this suggests that SK might primarily target exogenous vertebrate receptors rather than endogenous mosquito receptors. Indeed, SK elicits intestinal contractions in mammals and cross-reacts to antibodies that recognize the prototypic mammalian tachykinin substance P (39, 42). Substance P is a neuropeptide that is expressed widely in the central nervous system and is best known for its ability to activate nausea, although roles in mediating neurogenic inflammation and itch responses in the periphery have also been described (47–49). Crucially, the presence of such deuterostome-type tachykinin-like peptides in arthropods appears to be extremely rare, with the exception of four peptides from the highly venomous Brazilian wandering spider, whose venom induces edema in mammals (50). Beyond arthropods, matching peptides were identified in three octopod species, where vasoactive effects might be important for immobilizing prey.

We also examined the available insect species genomics data, to define which species have SK-like genes. No homologs were identified from the gene family data at VectorBase (51) or at OrthoDB (52). Sequence searches of the National Center for Biotechnology Information (NCBI) nonredundant protein and nucleotide databases returned hits only to the *Ae. aegypti* pro-SK precursor itself. We also searched the newly available genome assembly for *Culex tarsalis* (53) and all nucleotide data at VectorBase (25 mosquito species, genomes, transcriptomes, complementary DNAs [cDNAs], and expressed sequence tags [ESTs]), but only recovered the *Ae. aegypti* pro-SK precursor. The large sampling of *Anopheles* species, including several high-quality chromosome-level assemblies, lends confidence to the conclusion that this gene is not present in anophelines. The lack of hits in *Ae. albopictus* and two *Culex* species suggests that it could be specific to *Ae. aegypti*. However, the gene is located in a repeat-rich gene desert (nearest neighbors are 0.48 Mbp and 0.99 Mbp) at the start of *Ae. aegypti* chromosome one. Such repeat-rich regions can be challenging to sequence and assemble; therefore, it remains possible that other culicine mosquitoes do possess pro-SK precursor–encoding genes.

We hypothesized that SK may account for the ability of *Aedes* saliva to promote vascular leak and thereby enhance virus infection. Indeed, when coinoculated with virus in the absence of saliva, synthesized SK caused a rapid induction of edema and enhanced virus infection (Fig. 6B and C). When SK was added to *Anopheles* saliva, viral enhancement was successfully generated (Fig. 6D). However, because we had found that the *Aedes* saliva enhancement of virus infection was partially heat sensitive to treatment at 95°C for 10 min (Fig. 2C), and because previous work has shown some SK function was resistant to heat treatment for 2 min (39), we wanted to assess whether our synthesized SK was likewise heat sensitive. In agreement with both studies, we found that while a 2-min heat treatment did not significantly decrease the ability of SK to induce edema in vivo, a 10-min treatment was sufficient to significantly decrease SK-peptide–induced edema (*SI Appendix*, Fig. S7B).

Enhancement of virus by SK was dose dependent, as a 50-fold lower dose of SK caused only a modest increase in host susceptibility to virus (*SI Appendix*, Fig. S7A), suggesting that other components within saliva may prime host responses to SK and thereby modulate host susceptibility to virus. Therefore, to determine whether SK expressed by *Aedes* saliva gland was necessary to enhance virus infection, we silenced the SK gene in adult female *Ae. aegypti* mosquitoes (*SI Appendix*, Fig. S7C), by injection of a double-stranded RNA (dsRNA) targeting the SK gene. Saliva derived from these mosquitoes lost much of its ability to enhance infection compared to dsLucZ–dsRNA–injected control females (Fig. 6E). The extent of knockdown of SK gene expression (sevenfold reduction of median; *SI Appendix*, Fig. S7C) was similar to the magnitude of reduction in virus serum titers by 24 hpi (10-fold reduction, Fig. 6E). The reduction in skin virus RNA quantities in SK-deficient–saliva–injected mice was more modest (1.5-fold reduction, Fig. 6E). Importantly, similar to whole *Aedes* saliva, the ability of SK alone to enhance virus infection correlated with its ability to selectively and rapidly recruit monocytes in vivo (Fig. 6F). This effect was via direct action on endothelial cells, as an exposure of monolayers to SK resulted in rapid loss in electrical resistance and thereby an increase in permeability (Fig. 6G), similar in kinetics and magnitude as that observed with *Ae. aegypti* saliva (Fig. 5F). Thus, we have identified a factor in *Aedes* saliva, SK, which induces blood vascular barrier permeability, leading to edema and a rapid influx of virus permissive monocytes that enhances infection with virus.

Discussion

In this report, we have described how a mosquito saliva component is required for the successful establishment of virus infection in the mammalian host. The transmission of virus from mosquito to vertebrate skin constitutes an important bottleneck that is a critical step for viruses to overcome. Factors that provide a replicative advantage for viruses at this key stage of infection are key for defining overall outcome. This includes components within mosquito saliva that trigger an influx of virus-permissive mononuclear cells that efficiently replicate virus. Salivary proteases have also been shown to modulate host susceptibility to virus (8).

Here, we show that the host response to *Aedes* mosquito saliva is remarkably rapid, inducing peak edema and monocyte influx within minutes postexposure. Surprisingly, early immune sensing, as shown by extent of innate immune cytokine expression and adaptive immune hypersensitivity responses to saliva,
was not required for vascular permeability and thereby enhancement of virus infection. Instead, we found that early pro-viral vascular leakage occurred through a direct pharmacological effect induced by a mosquito-encoded substance P mimic, SK, that has likely evolved to facilitate optimal blood feeding. Subsequent immune responses that follow early saliva-induced leukocyte entry likely act to further extend skin residency time and tissue micropositioning of leukocytes, e.g., to sites of virus replication, and thereby increase the probability of their infection (18). Future studies that extend these findings to human responses would be valuable.

Our observation that early enhancement of infection by saliva was only evident in vivo is consistent with the requirement for a circulatory system that was absent in our in vitro and ex vivo models. The lack of infection enhancement in vitro was initially surprising, as several previous studies have demonstrated an enhancing effect of mosquito salivary gland extracts in cultured cells with other arboviruses. Together, these studies have utilized a number of distinct cell lines and virus strains (54). It is not clear why we could not replicate these findings with our models. Pata-tively, the use of salivary gland tissue homogenates in these studies, rather than mosquito saliva itself, may explain the difference.
in findings, as could the use of virus. Our data also demonstrate the limitation of exclusively using in vitro models that do not fully replicate all features of virus infection during arthropod biting.

Importantly, we have excluded a role for other salivary components that have been previously hypothesized to modulate vertebrate susceptibility to infection. This includes bacterial microbiota, which although we found did augment some skin inflammatory responses, did not modulate outcome to virus infection. Instead, we identified a role for a female-salivary-tachykinins Substance P and Neurokinins A and B. Rare examples of other protostome species with peptides matching the deuterostome-type FXGLMamide signature (Top box). In contrast, the SK tachykinin-like peptide (Aedes SK, 1,400 Da) matches the deuterostome-type FXGXRamide signature (Bottom box). This motif is similar to mammalian tachykinins Substance P and Neurokinins A and B. Rare examples of other protostome species with peptides matching the deuterostome-type signature include octopuses (EL, eledosin; TK, tachykinin) and a venomous spider (Ph. nigrig) (Bottom box).

(A) Mouse skin was inoculated with either 1.86 μg of Aedes salivary saliva or 1 μg of SK. Skin samples were collected 30 min post injection. (C-E) Mouse skin was inoculated with 10^4 PFU of SFV4 alone or with saliva from either Aedes or An. gambiae, with or without supplementation with 1 μg SK peptide. SFV RNA and host 18S were quantified by qPCR and serum viral titers quantified by plaque assay at 24 hpi. (E) Virus was administered alone or alongside saliva from Aedes females injected with dsRNA LacZ or dsRNA SK. (F) Resting mouse skin was injected with SK alone. At 2 h, skin from the inoculation site was biopsied and digested to release cells, and numbers of myeloid cells (CD45^+CD11b^+), neutrophils (CD45^+CD11b^-Ly6G^-Ly6C^+), and monocytes (CD45^-CD11b^-Ly6G^-Ly6C^-) were quantified (n = 6). (G) Human primary endothelial cell monolayers were treated with either control saline or 1 μM SK alone, and electrical resistance across the endothelial cell (EC) monolayer was assessed longitudinally. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001, ns = not significant.

The observation that Anopheles salivifer does not enhance virus infection is also key for considering its implication for understanding vectorial capacity for arbovirus transmission. Our data suggest that coevolution of arboviruses in competent vectors, such as Aedes, has led to the virus utilizing vector-specific factors to increase their infectiousness. We suggest this will modulate the overall effectiveness of arbovirus transmission by the competent vector, leading to an increase in vectorial capacity and thereby a high disease burden.
Together, our studies have defined an important determinant of the clinical outcome to infection at the arthropod/mammalian interface. We suggest this can inform the development of strategies that target SK to reduce host susceptibility to globally important arboviral diseases. This could include vaccines that target mosquito salivary factors (24, 25), although any such an approach must be done carefully to avoid off-target effects, which we suggest might be best achieved by targeting the N terminus of soluble SK, which lacks homology to human tachykinins. Vaccines would also need to be carefully designed to elicit protective Th1-associated IgG2 antibodies that lack the immunopathology associated with hypersensitivity reactions mediated by Th2-associated IgG1 and IgE responses. In conclusion, therapies that target mosquito saliva may have wide applicability, as mosquito saliva enhancement is widely observed for these viruses.

Materials and Methods

Mosquito Strains. The following mosquito strains were used: Ae. aegypti (Liverpool strain), Ae. albopictus (La Providence strain), Cx. pipiens (slab strain), An. gambiae (Kisumu strain), and An. stephensi (SDA-500 strain).

Virus Strains. SFV4 and SFV6 stocks were generated from plasmids containing corresponding infectious cDNAs (iCDNA) sequences (55). SFV6-2SG-Gluc (Gaussia luciferase) is a modified SFV6 where a sequence encoding Gluc marker is inserted under a duplicated sub-genomic (SG) promoter positioned at a 3' direction of the structural reading frame. ONNV-2SG-ZsGreen was linearized with PmeI (New England BioLabs) to prepare it for run-off transcription; RNA was transcribed using a MEGAscript SP6 transcription Kit (Thermo Fisher Scientific) to prepare RNA for microinjection. Transcripts were electroporated into BHK-21 cells to generate infectious virus with two pulses at 250V for 0.8 s. A plasmid to rescue ONNV-2SG-ZsGreen was linearized first with PmeI (New England BioLabs) to prepare it for run-off transcription; RNA was transcribed using a MEGAscript SP6 transcription Kit (Thermo Fisher Scientific) in the presence of Ribo m7G Cap Analog (Promega). The RNA was transfected to BHK cells using Lipofectamine 2000 to rescue the virus. Rescued SFV6-2SG-Gluc was aliquoted with cellular debris to allow for improved virus uptake by macrophages in vitro. Wild-type ZIKV from Recife, Brazil, was kindly provided by Alain Kohl at the University of Montpellier, France. Mosquitoes were reared at 28°C and 80% humidity conditions and used between 4 and 12 wk of age unless stated otherwise. BALB/c mice were maintained at St. James’ Biomedical Services under pathogen-free conditions and used between 4 and 12 wk of age unless stated otherwise. Wild-type C57BL/6j mice bred in-house at the SBS at the University of Glasgow, C6/36 cells were cultured at 28°C and 80% humidity conditions and used between 4 and 12 wk of age unless stated otherwise. mouse embryonic fibroblasts (MEFs) from C57BL/6 mice were kept at −19°C for long-term storage. MEF cells were cultured at 37°C at 5% CO2 in Dulbecco’s modified Eagle medium (DMEM) supplemented with 10% FCS, 100 units/ml penicillin, and 0.1 mg/ml streptomycin and 1% Glutamax in flasks precoated with 0.2% gelatin.

Macrophages were extracted from C57BL/6 mouse bone marrow by flushing cells from the femur using a 26-gauge needle with cold phosphate-buffered saline (PBS). Cells were passed through a 40-μm cell and cultured at 37°C at 5% CO2 in DMEM/F12 supplemented with 10 ng/ml macrophage colony-stimulating factor (M-CSF). 10% FCS, 100 units/ml penicillin and 0.1 mg/ml streptomycin, 1% Glutamax, and 5 mg gentamycin. A total of 4 × 105 cells were seeded in 10 mL media per sterile plastic Petri dish used. At 7 d postextraction, cells were pooled by adding 3 mL Cellstripper (nonenzymatic cell dissociation solution) to each dish, and cells were then gently scraped off the plastic. Cells were seeded at a concentration of 2 × 104 per well in a 24-well plate or at 5 × 104 per well in a 96-well plate in complete DMEM/F12.

Mosquito Rearing and Handling. The different mosquito species used were Ae. aegypti Liverin strain (a gift of E. Devaney, University of Glasgow, United Kingdom), Ae. albopictus La Providence (INFRAVEC2 line; a gift from A-B. Fail-loux, Institut Pasteur, France), Cx. pipiens quinquefasciatus SLAB strain (a gift from M. Weill, Université de Montpellier, France), and Anopheles coluzzii Ngouso strain and An. stephensi SDA 500 strain (a gift from C. Bourgoin, Institut Pasteur, France). Mosquitoes were reared at 28°C and 80% humidity conditions with a 12-h light/dark cycle. Ae. aegypti, Ae. albopictus, and Cx. pipiens eggs on filter paper were placed in trays containing ~1.5 cm of water to hatch overnight. Larvae were fed with Go-cat cat food until pupation. An. gambiae and An. stephensi eggs were hatched the same day of arrival by placing in water. Ground Tetramin fish flakes were fed to the larvae the first days, following which the larvae were fed with Tetenal pellets until pupation. When pupae formed, these were picked and placed in small water-filled containers and left to emerge into BugDorm mosquito cages. All adult mosquitoes were fed a 10% sucrose solution. Mosquitoes used for salivations and biting experiments were 4 to 8 d old postemergence. Emerging adult mosquitoes were maintained on a 10% (wt/vol) sucrose solution ad libitum. Females were fed with heparinized rabbit blood (Envigo) for 1 h using a 37°C Hemotek system (Hemotek Ltd).

For salivation of mosquitoes, the mosquito proboscis was placed in a p10 tip containing 0.5 μL immersion oil (Cargille Laboratories). Mosquitoes were then left to salivate for up to an hour before tips were placed in an Eppendorf tube and centrifuged. Saliva droplets were then pooled and stored at −80°C. Before use, droplets of saliva were carefully pipetted out of the oil under microscope and diluted in PBSA. Saliva from five mosquitoes was utilized per injection unless stated otherwise.

Antibiotic Treatment. Mosquitoes were given a 10% sugar solution containing a mixture of 200 μg/mL penicillin/streptomycin, gentamycin at 200 μg/mL, and tetracycline at 100 μg/mL, from emergence onward for at least 7 d. Validation of antibiotic treatment was conducted by counting colony-forming units generated following plating on lysogeny broth agar of whole mosquitoes dipped in ethanol to remove any external microbiota (SI Appendix, Fig. S2 E and F).

In Vivo Mouse Infections. Mice were anesthetized with 0.1 mL/10 g of Sedator/Ketavet i.p. injection and placed on foil on top of the mosquito cages with the dorsal side of one or both hind feet exposed, allowing a maximum of 5 mosquitoes to feed. Toes were covered with tape to prevent mosquitoes from biting. Mosquitoes were left to feed until fully engorged. Virus injections of either C6/36-derived SFV6 (250 PFU in 1 μL) or SFV4 (10,000 PFU in 1 μL) were made directly at the bite site with a 5-μL 75 N syringe, 33 gauge (Hamilton) using small RN ga33/25-mm needles (Hamilton). Saliva injections were made at a concentration of saliva from five mosquitoes per injection.

Mice were culled via a schedule 1 method. Tissues dissected depended on the experiment but most commonly included skin from a foot and spleen. Blood samples were also collected from the ventricles. Tissue samples collected were stored in 0.5 mL of RNAlater in 1.5-mL tubes, with the exception of spleen and brain samples that were cut in half and stored in 1 mL of RNAlater to enable complete penetration of the RNAlater into the tissue. All samples were left in RNAlater for a minimum of 16 h to prevent RNA degradation. Samples were then stored at 4°C for short term or at −80°C for long term. Blood samples were centrifuged and serum was collected and stored at −80°C until use. Tissue samples were then analyzed via qR-PCR for an analysis of the expression of
SFV virion glycoprotein E1, which is a gene expressed via SG RNA and that has previously been established as a good indicator of total viral RNA levels (genome plus transcripts). Serum was analyzed for viral titers via plaque assays.

**RNA Purification and Quantification.** All tissue samples were lysed in a 1 mL TRIzol reagent and shaken with 7-mm stainless steel beads on a Tissue Lyser at 50 Hz for 10 min to ensure the complete lysis of all tissues. A total of 0.2 mL chloroform was then added to all samples that were then inverted 15 times to allow for gentle mixing of the solutions. Afterward, samples were centrifuged at 12,000g for 15 min at 4 °C in order to separate the mixture into a lower red phenol–chloroform phase and a colorless upper aqueous phase. The upper aqueous phase aqueous phase, containing the RNA, was transferred to a new tube containing an equal amount of 70% ethanol. RNA extractions were performed using the RNA mini purification kit (Life Technologies) by following the protocol provided with the kit. Purified RNA was then stored at −80 °C.

Approximately 1 μg of RNA in a volume of 9 μL of RNase-free water was used for cDNA production using the Applied Biosystems high-capacity RNA-to-cDNA kit. Total RNA was diluted 1 in 5, using RNase-free water, and 1 μL was used per qPCR in 384-well plates. A master mix was made up of primers, water, and 5 μL of cDNA. A triplicate technical replicate was run qPCR for cDNA production using the Applied Biosystems high-capacity RNA-to-cDNA kit. Total cDNA was diluted 1 in 5, using RNase-free water, and 1 μL was used per qPCR in 384-well plates. A master mix was made up of primers, water, and SYBR green mix (perfecta, Quantabio.com). A triplicate technical replicate was performed using the Applied Biosystems high-capacity RNA-to-cDNA kit. Samples were incubated at 37 °C for 60 min followed by heating to 95 °C for 5 min. The final cDNA was then stored at 4 °C for short-term use and at −20 °C for long-time storage.

In Vitro Transepithelial Electrical Resistance (TEER) Assay. Saliva was filtered to remove particulate matter, and 50,000 cells were seeded on each well of the 96-well plate, and 500 μL of EGM-2 media was added to each well. The cells were fed every other day and monitored for 10 days. TEER was measured every day using Millicell electrical resistance chambers (Millipore). The TEER was measured using a Millicell-ERS instrument (Millipore) and recorded as ohm cm²/mL. The data were analyzed using the GraphPad Prism software. A one-way ANOVA with a Dunnett’s multiple comparison test was used to compare the means of the control and treated groups.

**SK Peptide Generation.** SK peptide (N-T-G-D-K-F-Y-G-L-A-mide) was synthesized de novo to >95% purity by Synpeptide Ltd. (Galway, Ireland). Synthesis of SK peptides was performed using Fmoc solid-phase peptide synthesis (SPPS) on Wang resin using commercially available reagents and solvents. The synthesis was performed according to the manufacturer’s instructions, and the final products were purified by reverse-phase high-performance liquid chromatography (RP-HPLC) using a C18 reverse-phase column (5 μm, 150 × 25 mm, Phenomenex). The purity of the SK peptides was confirmed by RP-HPLC and mass spectrometry (MS). The MS spectra were obtained using a Q-TOF mass spectrometer (Waters). The peptides were then lyophilized and stored at −20 °C until use.

**Data Availability.** All study data are included in the article and/or SI Appendix. The data associated with this paper is available from University of Leeds at https://doi.org/10.5518/1163, (56).

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