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| Citation         | Baldini, Francesco, Paolo Gabrieli, Adam South, Clarissa Valim, Francesca Mancini, and Flaminia Catteruccia. 2013. “The Interaction between a Sexually Transferred Steroid Hormone and a Female Protein Regulates Oogenesis in the Malaria Mosquito Anopheles gambiae.” PLoS Biology 11 (10): e1001695. doi:10.1371/journal.pbio.1001695. http://dx.doi.org/10.1371/journal.pbio.1001695. |
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| Published Version| doi:10.1371/journal.pbio.1001695                                                                                                                                                                                                                                           |
| Citable link     | http://nrs.harvard.edu/urn-3:HUL.InstRepos:11879031                                                                                                                                                                                                                       |
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The Interaction between a Sexually Transferred Steroid Hormone and a Female Protein Regulates Oogenesis in the Malaria Mosquito Anopheles gambiae

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Abstract

Molecular interactions between male and female factors during mating profoundly affect the reproductive behavior and physiology of female insects. In natural populations of the malaria mosquito Anopheles gambiae, blood-fed females direct nutritional resources towards oogenesis only when inseminated. Here we show that the mating-dependent pathway of egg development in these mosquitoes is regulated by the interaction between the steroid hormone 20-hydroxy-ecdysone (20E) transferred by males during copulation and a female Mating-Induced Stimulator of Oogenesis (MISO) protein. RNAi silencing of MISO abolishes the increase in oogenesis caused by mating in blood-fed females, causes a delay in oocyte development, and impairs the function of male-transferred 20E. Co-immunoprecipitation experiments show that MISO and 20E interact in the female reproductive tract. Moreover MISO expression after mating is induced by 20E via the Ecdysone Receptor, demonstrating a close cooperation between the two factors. Male-transferred 20E therefore acts as a mating signal that females translate into an increased investment in egg development via a MISO-dependent pathway. The identification of this male–female reproductive interaction offers novel opportunities for the control of mosquito populations that transmit malaria.

Citation: Baldini F, Gabrieli P, South A, Valim C, Mancini F, et al. (2013) The Interaction between a Sexually Transferred Steroid Hormone and a Female Protein Regulates Oogenesis in the Malaria Mosquito Anopheles gambiae. PLOS Biol 11(10): e1001695. doi:10.1371/journal.pbio.1001695

Academic Editor: David S. Schneider, Stanford University, United States of America

Received March 7, 2013; Accepted September 5, 2013; Published October 29, 2013

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Funding: This study was sponsored by the European Research Council FP7 ERC Starting Grant project ‘Anorep’ (grant ID: 260897; http://erc.europa.eu/), by the European Commission FP7 projects INFRAVEC (grant ID: 228421; http://cordis.europa.eu/fp7/home_en.html), by a Wellcome Trust grant (grant ID: 093553; http://www.wellcome.ac.uk/), and by Department funding from the Harvard School of Public Health to FC. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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

Abbreviations: 20E, 20-hydroxy-ecdysone; EcR, Ecdysone receptor; HR3, Hormone receptor 3; JH, Juvenile Hormone; Lp, Lipophorin; MAGs, male accessory glands; USP, Ultraspiracle; Vg, Vitellogenin; YPP, yolk protein precursor.

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Introduction

In many organisms, male–female molecular interactions occurring during sex shape reproductive success and may drive the rapid evolution of reproductive phenotypes [1]. While in species where females mate multiple times these reproductive interactions are often antagonistic due to the different reproductive strategies utilized by males and females [2–5], in monandrous species—that is, species where females mate a single time—they are believed to benefit both sexes [6]. Indeed this hypothesis has been proven experimentally in Drosophila melanogaster; removing sexual selection in this naturally promiscuous species through “imposed” monogamy induced the evolution of less antagonistic traits, where males became less harmful and females less resistant to induced harm [7].

In the malaria mosquito Anopheles gambiae, females rarely mate more than once during their lifetime [8]. As yet unknown male–female molecular interactions occurring during this single copulation regulate a series of postmating events that profoundly change the physiology and behavior of females. While in copula, females receive sperm, which are stored in a dedicated store organ named the spermatheca, and seminal secretions produced by the male accessory glands (MAGs). MAG secretions coagulate during mating to form a gelatinous mating plug that is transferred to the uterus (atrium), where it is digested in 1–2 d [9,10]. Following this copulation event, blood-fed females increase their egg production [11] and start laying eggs [12,13]. The regulation of egg production in A. gambiae is a particularly intricate process that depends on two main signals: one derived from blood feeding and one triggered by mating. While all females need to feed on blood to develop eggs, virgins in general have a pregravid state where they require two or more consecutive feedings to complete the first gonotrophic cycle [14–16]. This has profound implications for malaria transmission, as it increases the likelihood of contact with the human host. Pregravid behavior may be caused by insufficient metabolic reserves at emergence due to nutritional deprivation during larval stages [14,17]. This, in turn, may drive the need to optimize resource allocation between highly energy-demanding processes like flight and reproduction [18]. Indeed smaller A. gambiae mosquitoes tend to produce fewer eggs [19,20] and appear to feed as virgins [21], perhaps to build up energy reserves for mating.

The cascade of events triggered by blood feeding and leading to egg development, partially described in A. gambiae [22], has been well characterized in another mosquito species, the yellow fever and dengue vector Aedes aegypti. In these mosquitoes, after a blood
After mating the ovaries are released from their previtellogenic arrest and start vitellogenesis, the process of synthesis and secretion of yolk protein precursors (YPPs) by fat body cells. Upon secretion into the hemolymph, the YPP Vitellogenin (Vg) and the lipid transporter Lipophorin (Lp) become internalized into the ovaries via receptor-mediated endocytosis [23,24], leading to the maturation of 50–150 oocytes in approximately 2–3 d (reviewed in [25]). The transcription of YPPs is under endocrine regulation. After blood feeding the brain-secreted ovarian ecdysteroidogenic hormone (OEH) stimulates the ovaries to produce the steroid hormone ecdysone (E) [26,27], which in turn is hydroxylated into 20-hydroxyecdysone (20E) by the fat body cells. 20E synthesis releases the state of arrest of the fat body, activating the transcription of YPPs [25,28–30] by binding to the nuclear hormone receptor heterodimer Ecdysone Receptor (EcR)/Ultra-spireacle (USP), prompting it to function as a transcriptional activator [31]. A similar role of 20E in vitellogenesis after blood feeding has been demonstrated also in A. gambiae [22], where titers of 20E in blood-fed females correlated to Vg expression, suggesting a conservation of this pathway between Anopheles and Aedes mosquitoes.

No information is instead available on the factors regulating the mating-induced stimulation of oogenesis observed in A. gambiae. Mating increases the rate of egg production in a number of insects, and in some cases this effect has been attributed to the transfer of MAG secretions [reviewed in [32]]. The D. melanogaster Sex Peptide increases production of YPPs and oocyte maturation by inducing the female corpora allata to secrete the sesquiterpenoid Juvenile Hormone III-bisepoxide [JHB3] [33–35]. In Photoris fireflies, seminal secretions translocated to ovaries positively influence female fecundity [36]. In mosquitoes, a role of MAG products in egg development has been suggested by a number of studies where injections of MAG extracts into the hemolymph of Aedes females stimulated Vg synthesis and/or oogenesis [37–40]. In A. gambiae, indirect evidence suggests that MAG secretions act as master regulators of female postcopulatory behavior and physiology [41–43]. Thus far more than 100 A. gambiae MAG genes have been identified [46,47], and a number of them encode proteins that are...
by mated dsMISO females prompted us to analyze whether MISO plays a role in regulating the lipid transport to the oocyte. We therefore analyzed the expression levels of the viellagen-like lipid transporter \( Lp \) (AGAP007826) and the major VPP \( Vg \) (AGAP004203) in the fat body of blood-fed females at their peak of expression. In five different experiments, \( Lp \) transcript levels at 24 h after blood feeding were strongly reduced (34% mean reduction) in mated dsMISO females compared to mated controls, similar to virgin control levels (50% mean reduction) (Figure 2C) (Repeated Measures ANOVA, \( F_{2,4} = 8.142, p = 0.0118 \); Tukey’s Multiple Comparison post hoc test: virgin dsLacZ versus mated dsLacZ, \( p < 0.05 \); mated dsLacZ versus mated dsMISO, \( p < 0.05 \); virgin dsLacZ versus mated dsMISO, \( p > 0.05 \)). \( Vg \) instead was not significantly affected by MISO silencing (Repeated Measures ANOVA, \( F_{2,4} = 1.362, p = 0.3098 \) ) (Figure 2C). Taken together, these results indicate that mating increases the blood feeding–induced expression of \( Lp \) and that this regulation is dependent on MISO.

MISO Affects Male Ecdysteroid Titers in the Atrium and the Expression of 20E-Responsive Genes after Mating

In \( A. aegypti \) mosquitoes, the expression of the \( Vg \) and \( Lp \) after blood feeding is induced by the function of 20E, produced by the female [29,30], and a similar regulation occurs also in \( A. gambiae \) [22]. We tested whether the MISO-mediated upregulation in the expression of \( Lp \) in mated females after a blood meal was caused by an increased production of this hormone. We measured ecdysteroid levels secreted \textit{in vitro} by the ovaries of virgin and mated dsLacZ and mated dsMISO females before and 18 h after a blood meal, at their peak of secretion [48]. As expected, blood feeding strongly increased the steroidogenic capacity of the ovaries (Figure 3A) (one-way ANOVA, \( F_{1,4} = 11.17, p < 0.0001 \); post hoc Tukey’s multiple comparison, non-blood-fed versus blood-fed groups, \( p < 0.01 \)). However, no differences between virgin and mated females were observed, and silencing of MISO did not affect ecdysteroid secretion levels (\( p > 0.05 \) ) (Figure 3A).

Besides being produced by the female after blood feeding, in \( A. gambiae \) 20E is also synthesized in the MAGs and transferred to females during mating [48]. We therefore hypothesized that sexually transferred 20E may play a role in the MISO-mediated regulation of female physiology after mating. As a first step, we determined that the MAG-produced 20E is transferred to the female as part of the mating plug (Figure S3A). By 12 hpm, 20E localization was restricted to the anterior portion of the plug that is enclosed within the ampullae (Figure S3A), where MISO also localizes (Figure S1C). The amount of 20E detected in the MAGs corresponded to a mean of 632 pg (±17 pg), consistent with previous findings by others (Figure S3B) [48]. Interestingly, no 20E could be detected in the male reproductive tissues of two mosquito species, \( A. mosquito \) and \( A. aegypti \), which do not produce mating plugs (Figure S3B).

We next investigated whether MISO affects the activity of 20E transferred by males during copulation. To this aim, we analyzed steroid hormone levels in the atra of dsLacZ and dsMISO females at five time points after mating (0.5, 6, 12, 18, and 24 hpm) to monitor 20E release from the mating plug over time. Immediately after mating (0.5 hpm), the atra of control and dsMISO females contained similar hormone titers (Figure 3B). Ecdysteroid levels in the atra of controls were statistically significantly decreased at the four later time points (Wilcoxon test, \( p < 0.001 \) ) and reached about

\begin{align*}
\text{MISO Influences Lipid Accumulation in Developing Oocytes by Regulating the Expression of the Lipid Transporter Lipophorin} & \\
\text{After assessing the role of MISO in the determination of the increase in oogenesis induced by mating, we next analyzed the progression of oocyte development in mated dsMISO and control virgin and mated females at two time points (24 h and 60 h) after a blood meal. At 24 h postblood feeding, dsMISO follicles showed delayed development compared to mated dsLacZ controls, similar to what observed in the ovaries of virgin dsLacZ females (Figure 2A). By 60 h postblood feeding, oogenesis was completed in all three groups (Figure 2B); however, dsMISO (and virgin dsLacZ) ovaries showed a number of undeveloped primary follicles (indicated by asterisks in Figure 2B) in agreement with the finding that MISO silencing reduces egg development. A time course of five time points (12, 24, 36, 48, and 60 h) after blood feeding in virgin and mated females confirmed that, similar to virgin dsLacZ controls, mated dsMISO females exhibited a statistically significant delay in egg development, and only achieved oocytes of the size exhibited by mated dsLacZ individuals at 60 h postblood feeding (Figure S2 and Table S2). These results suggest that the effects of MISO on egg development are due to delayed or impaired accumulation of lipids into the growing oocytes.
\end{align*}
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A

virgin dsLacZ

24h post blood feeding

mated dsLacZ

mated dsMISO

B

virgin dsLacZ

60h post blood feeding

mated dsLacZ

mated dsMISO

C

\[ Lp \]

\[ Vg \]

p<0.05   p<0.05

ns   ns

\[
\begin{array}{ccc}
\text{Lp relative expression} & \text{Vg relative expression} \\
\text{Virgin BF dsLacZ} & \text{Mated BF dsLacZ} & \text{Mated BF dsMISO} \\
\text{Virgin BF dsLacZ} & \text{Mated BF dsLacZ} & \text{Mated BF dsMISO}
\end{array}
\]
3 pg per individual by 24 hpm, suggesting that 1 d after copulation the steroids have been fully released from the mating plug and have circulated out of the atrium. Interestingly, ecdysteroid titers declined more slowly in the atria of dsMISO females (P-mixed effects model, \( p = 0.055 \)) (Figure 3B). No 20E was detected in the atria of virgin females (unpublished data), confirming that this hormone in the female is only produced after blood feeding. These results suggest that silencing of MISO impairs the release of ecdysteroids from the plug and/or their diffusion from the atrium, possibly affecting their function.

To confirm the latter hypothesis, we analyzed the transcription levels of five 20E-responsive genes at three time points after mating (6, 12, and 18 hpm) in the two RNAi-injected groups. If MISO impairs the release of 20E from the atrium, then the expression levels of these genes in surrounding tissues should be altered in dsMISO females compared to controls. Besides Vg and Lp [28,29], we analyzed Ecdysone Receptor (EcR, AGAP012211) [31], Ultraspiracle (USP, AGAP002095) [50,51], and Hormone Receptor 3 (HR3, AGAP009002) [52]. As mentioned above, EcR is a nuclear receptor that in conjunction with USP activates transcription of downstream genes upon binding of 20E [31,50,51], while HR3 is known to interact directly with EcR [52]. Three genes exhibited a significant reduction in postmating expression in dsMISO females over the time frame analyzed: HR3 was downregulated by 50% at 6 hpm (\( \text{t test, } t_6 = 2.431, p = 0.0256 \)), Vg was reduced by 54% at 12 hpm (\( \text{t test, } t_6 = 2.785, p = 0.0159 \)), while EcR was decreased by 44% at 18 hpm (\( \text{t test, } t_6 = 1.876, p = 0.0587 \)) (Figure 3C). The expression levels of Lp and USP did not significantly differ between control and experimental females (Figure 3C).

All together, these results show that MISO silencing impairs both the titers of 20E in the atrium and the expression of 20E-responsive genes after mating, reinforcing the hypothesis that

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**Figure 2.** MISO silencing alters the expression of the lipid transporter Lipophorin in developing oocytes after blood feeding. (A and B) Immunofluorescence experiments on ovaries dissected from virgin and mated dsLacZ and mated dsMISO females stained with the lipid-binding reagent Nile-Red (red). Scale bar: 200 μm. (C) qRT-PCR of Lp and Vg from the fat body of virgin and mated dsLacZ and mated dsMISO females 24 h after blood feeding (BF). Expression levels (shown in logarithmic scale) were normalized to the housekeeping gene RpL19. The box-and-whisker diagrams represent five replicates of pools of 6–10 tissues. doi:10.1371/journal.pbio.1001695.g002

**Figure 3.** MISO silencing affects atrial 20E titers and reduces the activation of 20E-responsive genes after mating. (A) In vitro ovarian ecdysteroid secretion before and 18 h after a blood meal in virgin and mated dsLacZ and mated dsMISO. Graph shows data from eight individual ovaries. Data are represented as mean ± SEM. Means with the same letter are not significantly different (\( p > 0.05 \)). (B) Changes over time in the geometric mean of the ecdysteroid titer (natural logarithm) of dsLacZ (black solid line with circles) and dsMISO (green dashed line with triangles) females at 5 time points after mating (0.5, 6, 12, 18, and 24 hpm) with the mean trajectories estimated in regression mixed models (dashed and dotted lines). Nine replicates were performed using a pool of three atria each. (C) qRT-PCR of 5 20E-responsive genes (Vg, Lp, EcR, HR3, and USP) in dsMISO and dsLacZ females at different time points (0, 6, 12, 18 h) after mating. The levels of MISO after dsMISO injections are also shown. Four independent replicates were performed using a pool of 5–10 female abdomens. Expression was normalized to the housekeeping gene RpL19. Data are represented as mean ± SEM. One or two asterisks represent \( p < 0.05 \) and \( p < 0.001 \), respectively. doi:10.1371/journal.pbio.1001695.g003
MISO influences the function of male-derived ecdysteroids delivered by the mating plug.

**MISO Interacts with and Is Regulated by Male-Transferred 20E**

We next investigated whether the effects of MISO silencing on 20E titers and on the expression of 20E-responsive genes were caused by a possible interaction between MISO and 20E. To this aim, Western blot analyses were performed under native (i.e., nondenaturing) conditions. An anti-20E antibody detected a band of approximately 40 kDa in the atria of mated female (8 hpm) that was not detected in virgin extracts (Figure 4A). This band reacted also with anti-MISO antibodies, suggesting that the two factors are part of the same complex (Figure 4A). Moreover, immunoprecipitation of MISO in extracts of virgin and mated atria at 8 hpm followed by an ELISA coupled with anti-20E antibodies detected significant amounts of 20E co-immunoprecipitating in mated females, while no signal was observed in virgins (Figure 4B). All together these results suggest an interaction between MISO and 20E in the atrium of females after mating.

As 20E is known to regulate the expression of genes that are ultimately responsible for its function (reviewed in [33]), we next analyzed whether this steroid hormone plays a role in the expression of MISO in the atrium. To this aim, we injected three 10-fold dilutions of 20E into the hemolymph of virgin females, and analyzed MISO transcript levels specifically in the atrium (where the gene is not normally expressed in virgin females) at 24 h postinjection. At the highest concentration, 20E significantly induced MISO expression to levels similar to those achieved by mating (178- and 349-fold induction, respectively) (one-way ANOVA, F$_6,23 = 14.79$, p < 0.0001; post hoc Dunnett’s multiple comparison against virgins, p < 0.01), while the ethanol and cholesterol controls had no effect (Figure 4C). At lower dilutions, 20E injections increased MISO expression levels relative to controls, however this effect was not statistically significant. No effect on MISO expression was seen in tissues other than the atrium, confirming the tissue-specific restriction of expression of this gene (unpublished data). The expression of AGAP009584, an atrial gene that is not modulated by mating [10,49], was not induced by the injection of any of the 20E dilutions (Figure 4C) (one-way ANOVA, F$_6,23 = 0.5089$, p = 0.7947). Only the highest concentration of injected 20E achieved physiological atrial concentrations similar to those transferred during mating (Figure S4), explaining the observed titration-dependent upregulation of MISO expression.

Finally, to further confirm that 20E induces MISO expression in the atrium, we tested MISO induction levels in the absence of the 20E receptor EcR. We injected virgin females with dsRNA targeting EcR, and analyzed levels of MISO induction after mating. In four different experiments, injection of dsEcR (transcript mean reduction = 45%; one-sample t test, t$_s$ = 7.069, p = 0.0058, range 63%–36%) impaired MISO induction at 24 hpm by an average of 30-fold compared to injected controls (Figure 4D), reinforcing the notion that the expression of this gene after mating is regulated by male-transferred 20E. Interestingly, EcR silencing also reduced transcript levels of Lp$_e$ (24 hpm: t test, t$_s$ = 2.106, p = 0.0399), as expected as this gene is under the control of 20E and its expression is induced by both blood feeding and mating in A. gambiae (Figure 4D) [22,49]. These data demonstrate that the mating-induced expression of MISO is under the control of sexually transferred 20E, and that EcR mediates this regulation.

**Discussion**

In this study we unravel a major male–female molecular interaction that switches females to a mated state in terms of egg development and modulates their postmating physiology. We identify a female atrial protein, MISO, which is responsible for the increase in egg production after mating. Silencing of MISO reverses fecundity of mated females back to virgin levels, completely abolishing the effects of mating on oogenesis (Figure 1). Moreover we demonstrate that MISO is induced by and interacts with the steroid hormone 20E, transferred by the male (Figure 4). Sexually transferred 20E therefore acts as a “mating signal” that regulates female postmating physiology, and its interaction with MISO translates this signal into increased oogenesis in blood-fed females. To our knowledge, this is the first demonstration of an interaction between a male allohormone and a female protein in insects. The identification of this novel interaction in A. gambiae expands our knowledge of male–female molecular partnerships important for reproductive success, to date limited to few examples from *Drosophila* (reviewed in [34]).

The mating-induced increase in egg development seen in our experimental settings only partially reflects the deep impact that mating has on oogenesis in field conditions. Blood-fed virgins from natural mosquito populations rarely develop eggs after a single blood meal [14–16], presumably because of limited nutritional reserves from larval stages [17]. MISO may therefore represent a mating sensor that directs precious resources towards oogenesis only when females are inseminated. Indeed in two different phenotypic assays, MISO influenced pregravid behavior, and similar to virgin females, approximately 15% of dsMISO mated females completely failed to develop eggs compared to 4% of mated controls (Figure 1 and Table S1). It is reasonable to speculate that this effect would be much more pronounced in conditions of limiting resources such as those possibly available in field settings.

The interaction between MISO and 20E affects the function of the steroid hormone, as demonstrated by the effects of MISO silencing on 20E titers in the atrium and on the expression of a number of 20E-responsive genes (Figure 3B,C). Although the protein does not have any known functional domains that suggest a role as a sterol carrier, our data indicate that MISO facilitates the release of 20E from the mating plug and its diffusion from the atrium (Figure 3B). Further studies may help elucidating the mechanism by which this female atrial protein regulates 20E function. On the other hand, the finding that sexually transferred 20E induces the atrial-specific expression of MISO via the EcR receptor shows a remarkable mutual cooperation between the two factors (Figure 4C,D). Preventing males from producing and transferring 20E will clarify the full extent of the role that this ecdysteroid plays in regulating female postmating physiology and behavior.

A number of hypotheses can be formulated on the downstream events triggered by the interaction of MISO and 20E that lead to increased fecundity. One possibility is that this interaction may prime the fat body to respond to the female-derived ecdysteroids synthesized after a blood meal. This hypothesis is strengthened by the observations that mated dsMISO females experienced a reduced induction in Lp expression after blood feeding compared to controls, paralleled by delayed or impaired oocyte growth (Figure 2, Figure S2, and Table S2). The higher level of Lp expression seen in control mated females is not due to an increased release of ecdysteroids from the ovaries after blood feeding, as ecdysteroid titers were similar in control and dsMISO females (Figure 3A). Interestingly, MISO silencing affects the expression of
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Vg and Lp differentially: while the most prominent effect on Lp occurs after blood feeding (Figure 2C), Vg transcript levels are repressed only after mating (Figure 3C). This observation suggests a bimodal role for the MISO–20E interaction: a local effect on the expression of mating-responsive genes such as Vg that may regulate the function of reproductive tissues and possibly the remodeling of atrial cells observed after mating [49], and a later effect due to 20E release from the atrium that may control the response of the fat body to blood feeding, thereby affecting Lp transcript levels and egg development. Importantly, these results are consistent with a recent report that identified Lp rather than Vg as the factor most relevant for egg development in A. gambiae [55]. Another possible mechanism is that sexually transferred 20E may regulate resorption of ovarian follicles. In A. aegypti the interplay between JH and 20E influences the late of follicular resorption during the previtellogenic and vitellogenic stages [56]. Low JH titers during the previtellogenic stage result in higher follicular resorption that can be prevented by the application of methoprene, a JH mimic [57]. 20E can also stimulate resorption of “poor quality” follicles that express low levels of Vg and Lp receptors [58], probably by a caspase-mediated cell death mechanism [59]. In A. gambiae male transferred 20E may therefore act cooperatively with female-derived JH in determining correct follicular resorption. Alternatively, the large amount of 20E transferred from the MAGs, that as confirmed here exceeds the concentration produced by the ovaries after blood feeding [48], may increase the number of developing oocytes by causing yolk accumulation in secondary follicles already during the first blood meal. This process has been observed in A. aegypti [60] and A. stephensi [61] after 20E injection.

Mating does not modulate egg development in all anopheline species. For instance, oogenesis is not affected by copulation in the central American malaria vector A. albimanus [62], and interestingly, we could not detect any 20E in the MAGs of this mosquito species (Figure S3B). This result suggests that the effect of mating on fecundity in anophelines might be directly linked to the presence of 20E in the male reproductive tract. Intriguingly, secretion of lower 20E titers in A. gambiae compared to A. albimanus females after a blood meal [22,63] may be due to the availability of 20E from males in the former species. An increase in egg development following mating is also seen in A. aegypti [39], however the absence of 20E in the MAGs of this species suggests that this effect is caused by a different mechanism (Figure S3B). This increase may be regulated by MAG proteins stimulating the synthesis of growth hormones, as in the case of the stimulation of JH synthesis by Sex Peptide in Drosophila [33]. Indeed the existence of a Sex Peptide–like factor inducing postcopulatory changes in A. aegypti is supported by the observation that MAG extracts injected into virgin females trigger oviposition after blood feeding [64,65], contrary to A. gambiae where they have no effect [13]. Alternatively, hormones other than 20E produced by the male and transferred during mating may play this role. JH has been detected in the MAGs of A. aegypti [66], and the application of the JH analog methoprene to virgin A. aegypti females enhances oogenesis [39]. No evidence of JH synthesis exists in the MAGs of A. gambiae, and unlike A. aegypti, application of methoprene to blood-fed females inhibits egg maturation and vitellogenesis [22], suggesting differences in the mechanism of oogenesis in the two species. The analysis of the synthesis of 20E in the MAGs of other mosquito species, facilitated by the sequencing of an additional 16 anopheline genomes (http://www.vectorbase.com), will clarify the existence of a possible correlation between mating plug formation and 20E synthesis in the male, two reproductive features that are both present in A. gambiae but not in A. albimanus and A. aegypti, and between the sexual transfer of 20E and the occurrence of mating-induced oogenesis.

Finally, the identification of a previously uncharacterized reproductive pathway in A. gambiae has promise for the development of tools for the control of malaria-transmitting mosquito populations. The effects of the 20E-MISO partnership are likely to be more prominent in field mosquitoes, where nutritional resources are limited and egg development rarely occurs in virgins. Manipulation of this interaction with specific inhibitors or with genetically manipulated males impaired in 20E synthesis
might therefore offer an attractive option for reducing the reproductive output of natural *Anopheles* populations. Moreover, interfering with the mating-induced pathway of oogenesis may have an effect on the development of *Plasmodium* malaria parasites. A recent study has shown that the expression of Vg and Lp reduces the mosquito *Plasmodium*-killing efficiency mediated by TEP-1, the principal antiparasitic factor in *A. gambiae* [55]. As VgPs are regulated after a blood meal via a MISO-dependent mechanism, the 20E-MISO interaction may play a role in the modulation of *Plasmodium* development in *A. gambiae*.

Materials and Methods

Mosquito Procedures

Mosquitoes from a laboratory colony of the *A. gambiae* G3 strain were reared under standard conditions [26–28°C, 65–80% relative humidity, 12 h:12 h Light/Darkness (L:D) photoperiod]. For mating experiments, mosquitoes were separated by sex as pupae and raised in cages supplied with sucrose *ad libitum*. Matings were performed as described previously, and couples were captured in *copula* [49].

RNA Interference

A 397 bp region corresponding to the coding sequence of MISO (AGAP0092620) was amplified from atrial cDNA 24 hpm using specific primers FW: 5’GGGTGGCCATTGTGTGTGT-3’ and REV: 5’AGTACTCGGCCAGCTGAATG-3’ and cloned into the pLl10 plasmid [67]. A 435 bp region corresponding to AGAP012211 (ER) was amplified from female abdomen cDNA using the primers FW: 5’CTGCCCTCAGTGAGGAGTGATGA-3’ and REV: 5’GGCCACGTTCAGTTCTTCAG-3’, while a 495 bp portion of the eGFP control gene was amplified using the primers FW: 5’TGTTCTGCTGTTAGTGTTGCG-3’ and REV: 5’ACGTAAACGGCCACAAGTTC-3’; both amplicons were cloned into pCR2.1 (Invitrogen). These constructs were then used to synthesize dsRNAs targeting the different genes, following established protocols [10,67,69]. Females were sexed as pupae and injected with 69 nl of dsRNA (5 mg/ml) within 24 h of eclosion. Surviving females were allowed to mate with 4-d-old virgin males. After mating, mated females were then used for phenotypic assays or dissected for qRT-PCR analysis. RNA extraction, cDNA synthesis, and SYBR-green based qRT-PCR were performed as described previously [49] using the primers listed in Table S3. The ribosomal protein gene Rpl19 (AGAP004422) was used for normalization, using previously described primers [49].

Oviposition, Egg Development, and Fertility Assays

Three days after dsRNA injections, females were captured during mating and kept in isolation until blood feeding. Females were blood fed *ab libitum* on human blood. Partially fed or unfed mosquitoes were removed. For oviposition and fertility assays, 3 d after the blood meal, females were put into individual oviposition cups for 4 nights. After completion of oviposition, eggs were counted under the microscope and those that hatched into a larva were scored as fertile. For the egg development assay, abdomens were dissected 3 d after blood feeding, and eggs developed inside the ovaries were counted under the microscope.

Polyclonal Anti-MISO Antibodies

Affinity-purified polyclonal antibodies against MISO were raised in rabbit against the peptide epitope CSNGPSSSYGPPRNT by a commercial supplier (GenScript Corp., Piscataway, NJ).

Immunoblots

Female tissues were homogenized in 20 μl RIPA buffer (10 mM Tris/HCl pH 7.6, 100 mM NaCl, 10 mM EDTA, 0.5%, Nonidet P40, 0.5% Triton ×100, 1× protease inhibitor from Roche). Samples were centrifuged at 13,000 rpm for 15 min at 4°C. The supernatant was diluted into NuPAGE reducing agent and sample buffer (Invitrogen), heated at 70°C for 10 min, and applied to precast NuPAGE gels (Invitrogen) under reducing conditions according to the manufacturer’s instructions. For native conditions, protein extraction was performed by homogenizing the tissues in a hypotonic solution (10 mM Tris/HCl pH 7.6, 10 mM NaCl, 10 mM EDTA, 1× protease inhibitor from Roche) followed by centrifugation at 13,000 rpm for 15 min at 4°C. The soluble phase was then loaded onto an acrylamide gel in the absence of SDS. Proteins were transferred to a Hybond ECL membrane using the XCell II Blot module (Invitrogen). Membranes were immunostained using standard protocols with the following primary antibody titres: anti-MISO, 0.96 mg/ml; anti-20E (1:10 dilution, Cayman Chemicals); and anti-β-actin (1:1,000 dilution, Santa Cruz Biotechnologies). HRP-conjugated secondary antibodies (Santa Cruz Biotechnologies) were used at a dilution of 1:10,000. Bands were visualized using ECL Western blotting detection reagents (GE Healthcare). Reprobing with additional primary antibodies was performed after incubating membranes in stripping solution (10 mM Tris/HCl pH 6.8, 100 mM DTT, SDS 2%) at 50°C for 30 min. Before adding the new primary antibody, incubation with the secondary antibody used in the first analysis was tested by ECL to exclude any signal from the previous incubation.

Immunofluorescence and Confocal Analysis

MAGs or female reproductive tracts from 3–4-d-old mosquitoes (virgin and mated) were dissected on ice, fixed in 4% formaldehyde, washed in PBS, then blocked and permeabilized in PBS with 1% BSA and 0.1% saponin. Samples were incubated in either 3 mg/ml anti-MISO or a 1:10 dilution anti-20E (Cayman Chemicals), then a 1:1,000 dilution of anti-rabbit Alexa-Fluor 488 (Invitrogen). Alternatively, ovaries were stained with 1:1,000 dilution of Nile-Red (10 mg/ml in DMSO, Sigma-Aldrich). Tissues were then mounted in DAPI-containing Vectashield medium (Vector Laboratories, Inc.) and visualized using a Point Scanning Confocal microscope Nikon TE2000 or a Zeiss Axio Observer inverted fluorescent microscope with apotome.

In Vitro Ovarian Culture

Ovaries of dsRNA-injected females were dissected from virgin and mated mosquitoes before or after 18 h after a blood meal. Blood feeding was performed 1 h after mating. Ovaries of mated non-blood-fed females were dissected 19 h after copulation. After dissection in Schneider medium (Sigma-Aldrich), individual pairs of ovaries were separately transferred to 50 μl of Schneider medium and incubated for 5 h at 25°C. After incubation, culture medium was stored at −80°C until ecdysteroid quantification.

20E ELISA

Atria from groups of three virgin females or from groups of three mated females at different time points after mating, previously injected with dsMISO or dsLacZ, were placed in 50 μl methanol and frozen at −80°C. Alternatively, MAGs or testes from 10 *A. gambiae, A. albimanus*, and *A. aegypti* males were dissected and placed in 50 μl methanol. Tissues were then homogenized and loaded into separate wells of a 96-well plate pre-coated with mouse anti-rabbit IgG (Cayman Chemical). For the analysis of the *in vitro* ovarian ecdysteroid secretion, 50 μl of Schneider medium
where the ovaries have been incubated were directly loaded into the gel. A standard curve was prepared from 18 ng 20E (Sigma-Aldrich) in methanol or Schneider medium (Sigma), with a series of seven 3-fold dilutions. After evaporation of the methanol, 50 μl of each of the following solutions were added: Enzyme Immuno-Assay Buffer (0.1 M phosphate solution containing 0.1% BSA, 0.4 M sodium chloride, 1 mM EDTA, and 0.01% sodium azide); 20E acetylcholinesterase (AChE) Tracer, which is a covalent conjugate of 20E and AChE; and anti-20E rabbit IgG (Cayman Chemical). The plate was incubated with the solutions overnight at 4°C, washed with PBS 1× containing 0.05% TWEEN20, incubated with 200 μl Ellmans reagent (5,5'-dithiobis-(2-nitrobenzoic acid)) (Cayman Chemical), and finally developed for 90–120 min and measured in an ELISA reader at 420 nm.

20E Injections
Three-day-old females were injected with different quantities (2.5 μg, 0.25 μg, and 0.025 μg) of 20E (158 nl of 10% ethanol solution). As controls, either the same volume of 10% ethanol or 0.25 μg of water-soluble cholesterol (which is the maximum soluble concentration) (Sigma-Aldrich) were injected. Female lower reproductive tracts (LRT, atrium, spermatheca, and parovarium) were dissected 24 h after injection and analyzed by qRT-PCR. Three replicates were performed containing 6–8 tissues per replicate. LRTs were also dissected and analyzed by qPCR from noninjected virgin females and from mated females at 24 hpm.

Immunoprecipitation Experiments
Fifty atria from virgin and mated (8 hpm) females were dissected and homogenized in 15 μl of hypotonic solution (10 mM Tris/HCl pH 7.6, 10 mM NaCl, 10 mM EDTA, 1× protease inhibitor from Roche) and centrifuged at 13,000 rpm for 15 min at 4°C. The soluble phase was then incubated for 1 h at 4°C under gentle rocking with 2 mg of anti-MISO rabbit IgG that had been previously linked to Dynabeads protein A (Invitrogen) in a 10 min incubation at 25°C under gentle rocking followed by three PBS 1× washes. The immunoprecipitate was washed three times with PBS 1× and split in two aliquots: one-third of the total volume was utilized in a Western blot incubated with anti-MISO, while the remaining two-thirds were diluted with 100 μl of methanol, to extract 20E, and kept at –80°C. The methanol solution was then analyzed with an anti-20E ELISA. As controls, 25 ng of 20E were incubated under the same conditions with 2 mg of Rabbit anti-MISO linked to Dynabeads protein A to measure the unspecific binding of 20E to the antibody or to the Dynabeads. All samples were also immunoprecipitated using pre-immune rabbit IgG to control for unspecific bindings. ELISA quantification was performed normalizing the signal to anti-MISO rabbit IgG-Dynabeads protein A incubated in methanol.

Statistical Analysis
To examine the effects of MISO on oviposition and egg development, we utilized a generalized linear model approach where the number of eggs was modeled with a log link function and Poisson distribution function using SAS Proc GenMod (SAS, Inc., Cary, NC). Replicate was also included as a covariate in each of these analyses. Post hoc comparisons for fecundity were made using the Bonferroni Multiple Comparison Procedure in SAS (SAS, Inc.).

Supporting Information

Figure S1 MISO is strongly induced in the atrium after mating and is secreted in the ampullae. (A) Quantitative RT-PCR (qRT-PCR) showing MISO expression in three conditions: virgin females after a blood meal (VBf), mated females (M), and mated females that have been blood fed immediately after mating (MBf). Atria, ovaries, and the rest of the body (carrass) were analyzed at different days (1, 3, and 6 d) postmating and/or blood feeding, and in age-matched virgin females. Expression levels (shown in logarithmic scale) were normalized to the housekeeping gene RpL19. The analysis was performed in three replicates on pools of 5–10 tissues, and data are represented as mean ± SEM. (B) Immunoblot analysis of MISO using a polyclonal antibody raised against a peptide fragment of the protein. Atria were dissected from different groups of females: virgins (Vf); mated (M) at 24 hpm; virgin blood fed (VBf) dissected at 24 h post-blood-feeding; mated blood fed (MBf), dissected at 24 h postmating and blood feeding; and MBf dissected after egg laying (EL). Immunoreactive bands (arrow) corresponding to the predicted 15 kDa size of MISO were detected in M, MBf, and EL atria. Actin was used as loading control. (C) Confocal analysis of MISO (green) in the atrium of virgin and mated females. The images next to the bright field (BF; scale bar: 100 μm) are magnifications (xy section, scale bar: 50 μm) of the regions indicated in the inset. At 12 hpm the mating plug is visible in the atrium (arrowhead). Cell nuclei (blue) are labeled with DAPI. (D) cDNAs from 15 independent replicates of dsMISO injections in virgin females analyzed by qRT-PCR at 24 hpm. RpL19 relative expression levels were compared between dsMISO- and dsLacZ-injected females (dotted line). Data are represented as a box-and-whisker diagram. (E) Immunoblot analysis of the efficacy of MISO silencing in protein extracts from atria, ovaries, and eggs. Atria and ovaries were dissected from virgin or mated females at 24 hpm that were injected with either dsMISO or dsLacZ, as indicated. Eggs were collected 1–4 h after oviposition. Actin was used as loading control. The arrow indicates the expected size for MISO. (TIF)

Figure S2 MISO silencing induces a delay in ovarian development. Immunofluorescence of oocyte development in ovaries dissected from dsMISO or dsLacZ-injected virgin or mated females at five points (12, 24, 36, 48, and 60 h) after blood feeding.
Nile-Red (red) and DAPI (blue) were used to stain lipids and cell nuclei, respectively. Scale bar: 50 μm.

**Figure S3** 20E localization in MAGs and atrium and quantification in male reproductive tracts from three mosquito species. (A) MAGs dissected from virgin males (MAGs) and atria dissected from virgin (V) and mated females at two time points after mating (0.5 hpm and 12 hpm) were dissected and incubated with anti-20E antibody (green). Cell nuclei (blue) are labeled with DAPI. Scale bar of the bright field (BF): 100 μm. The images next to the bright field (BF) are a magnification (xy section) of the region indicated by the inset (scale bar: 50 μm). (B) ELISA quantifications of 20E levels in MAGs and testes from either *A. gambiae* virgins or *A. aegypti* males. A pool of 10 tissues was used for each of three replicates. Data are represented as mean ± SEM.

**Figure S4** 20E quantification in the atrium after injection. ELISA quantification of 20E levels in female atria was performed prior or post injection (at 0.5 h, 6 h, and 24 h postinjection) of different 20E dilutions in the hemolymph of virgin females, or at the same time points after mating. Three 1:10 dilutions starting from 2.5 μg per mosquito were injected. Ethanol injections were used as a control. A pool of 10 atria was used for each of three replicates. Data are represented as mean ± SEM.

**Table S1** Summary of phenotypic analysis of dsMISO-injected females. MISO knockdown results in higher proportion of females that fail to develop eggs in both the oviposition and the egg development (oogenesis) assay (dsLacZ vs MISO; P<0.0083; dsLacZ vs mated; P=0.0594). Among females that completed oogenesis, injections of dsMISO reduced the number of developed eggs (oviposition: t test: t₁₂₁ = 0.9944, P = 0.1594; fecundity: one-way ANOVA: F₂,95 = 7.196, P = 0.0009; Tukey’s multiple comparison post hoc test: virgin dsLacZ vs mated dsLacZ, P<0.01; mated dsLacZ vs mated dsMISO, P<0.001; virgin dsLacZ vs mated dsLacZ, P>0.05). One, two, and three asterisks indicate P<0.05, P<0.01, and P<0.001, respectively.

**Table S2** Oocyte length in mated dsMISO females compared to virgin and mated dsLacZ controls after blood feeding. Oocytes showing lipid accumulation (as estimated by Nile-Red) were measured in ovaries dissected from dsMISO or dsLacZ virgin or mated females at five points (12, 24, 36, 48, and 60 h) after blood feeding. Oocytes from dsMISO and virgin females are consistently smaller than oocytes from dsLacZ females throughout development, and the three groups reach the same size only at 60 hpm (one-way ANOVA: 12 h, F₂,95 = 10.34, P<0.0001; 24 h, F₂,95 = 132.0, P<0.0001; 36 h, F₂,95 = 169.2, P<0.0001; 48 h, F₂,95 = 82.29, P<0.0001; 60 h, F₂,95 = 1.024, P = 0.03627). At each time point, means with different letters are significantly different (Tukey’s multiple comparison post hoc test: P<0.001).

**Table S3** List of primers and concentrations used for qRT-PCR.

**Acknowledgments**

The authors are grateful to Elena A. Levashina for useful discussion, to Evdokia Kakani and Perrine Mercenac for help with laboratory procedures, and to Emily Lund and other members of the Catteruccia laboratory for help with mosquito maintenance and for careful reading of the manuscript. The Nikon Imaging Center at the Harvard Medical School was utilized for confocal microscopy analysis.

**Author Contributions**

The author(s) have made the following declarations about their contributions: Conceived and designed the experiments: FB PG FM. Analyzed the data: FB PG AS VC FC. Contributed reagents/materials/analysis tools: FC. Wrote the paper: FB FC.

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