miRNA-1-3p is an early embryonic male sex-determining factor in the Oriental fruit fly Bactrocera dorsalis

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Regulation of male sexual differentiation by a Y chromosome-linked male determining factor (M-factor) is one of a diverse array of sex determination mechanisms found in insects. By deep sequencing of small RNAs from Bactrocera dorsalis early embryos, we identified an autosomal-derived microRNA, miR-1-3p, that has predicted target sites in the transformer gene (Bdtra) required for female sex determination. We further demonstrate by both in vitro and in vivo tests that miR-1-3p suppresses Bdtra expression. Injection of a miR-1-3p mimic in early embryos results in 87–92% phenotypic males, whereas knockdown of miR-1-3p by an inhibitor results in 67–77% phenotypic females. Finally, CRISPR/Cas9-mediated knockout of miR-1-3p results in the expression of female-specific splice variants of Bdtra and doublesex (Bddsx), and induced sex reversal of XY individuals into phenotypic females. These results indicate that miR-1-3p is required for male sex determination in early embryogenesis in B. dorsalis as an intermediate male determiner.
Primary signals that initiate sex determination during development appear to be highly divergent among many taxonomic groups1-3. This is exemplified in dipteran insects, such as Drosophila melanogaster, in which the primary sex-determining signal consists of the dose of X chromosome-linked signal elements (XSE)4. A double XSE dose from XY diploid individuals leads to expression of Sex-lethal (Sxl), which directs female-specific splicing of transformer (tra) and doublesex (d sx) pre-mRNA whose gene products sequentially direct female differentiation. In the absence of Sxl expression from a single XSE dose in XY (or XO) individuals, male-specific tra pre-mRNA splicing results in nonfunctional TRA and the subsequent default DSX-male product that directs male differentiation5-7. Although the Drosophila Y chromosome is necessary for male fertility, it has no apparent role in determining somatic and germline sexual differentiation8,9. This is in contrast to other dipterans, including the mosquitoes Aedes aegypti, Anopheles gambicae and Anopheles stephensi, and the housefly, Musca domestica, in which a dominant male-determining factor (M-factor) is the primary determinant of male differentiation10-13. However, M-factors identified in these species have no obvious sequence conservation, nor is their mode of function necessarily conserved. For example, in M. domestica the paternally inherited male determinant (Mdmd) CW22 paralog in XY embryos prevents the zygotic activation of the positive Mdtra loop, resulting in the default male-specific splicing of Mdtra and Mdms pre-mRNAs leading to male differentiation10. The M-factors Nix, Yob/YG2 and Guy1 from A. aegypti, An. gambicae, and An. stephensi, respectively, exhibit early embryonic transcription and subsequent expression throughout development in their respective species, and are required to initiate male development but have no primary sequence homology11-13. Since a tra ortholog sequence appears not to be conserved, and is possibly lacking in mosquito genomes, the different M-factors could more directly effect male-specific splicing of d sx pre-mRNA.

Y-linked M-factors have been inferred for several dipteran insects, but have not yet been identified in telegeiid fruit flies, including the Oriental fruit fly, Bactrocera dorsalis (Hendel), which is one of the most devastating and highly invasive agricultural pests of fruits and vegetables throughout the world14. In contrast to Drosophila, the tra gene in B. dorsalis as well as other dipteran, hymenopteran and coleopteran species10,15-20 is able to autoregulate its female-specific splicing, analogous to the Drosophila Sxl gene. As maternal tra transcripts initiate a positive autoregulatory feedback loop in the early zygote, continuous female-specific functional TRA protein is provided and, thus, acts as a cellular memory, maintaining the female-determining signal15. For male development, the presence of an M-factor, which is required to prevent activation of the tra autoregulatory loop, leads to a male-specific nonfunctional TRA protein.

For B. dorsalis, and other telegeiid pests, the foremost biologically-based population control strategy is the sterile insect technique (SIT), which relies on the ability to generate sterile male-only populations for mass release into infested areas21. Understanding the genetic basis of sex determination and male fertility has therefore been central to the development of genetically modified strains for sex separation and sterility. Although Sxl is not involved in telegeiid sex determination, orthologs of the Drosophila tra, transformer-2 (tra-2), and d sx genes are functionally homologous22,23. Therefore, the discovery of M-factors, or other upstream factors, remains fundamentally important for telegeiid species. These factors may also include small RNAs as shown in the silkworm moth, Bombyx mori. In this lepidopteran species, a W chromosome-derived female-specific piRNA represses transcription of Masculinizer (Masc), leading to female-specific splicing of Bmdsx pre-mRNA in ZW females24. Thus, this piRNA acts as an essential primary feminizing factor in the ZW sex determination system. In view of this, we propose the hypothesis that small RNAs may also participate in sex determination by regulating the tra autoregulatory loop during early embryogenesis in B. dorsalis.

In this study, we identify the key embryonic stages in which male sex determination is initiated in B. dorsalis, and show that a small autosomally derived RNA, miRNA-1-3p, induces male sex determination by suppressing Bdtra expression during this developmental stage. Embryonic injection of a synthetic miR-1-3p inhibitor, or disruption of miR-1-3p by CRISPR/Cas9, induces a sex reversal of XY chromosomal males into phenotypic females that exhibit female-specific splicing of both Bdtra and Bdms pre-mRNAs. Conversely, embryonic injection of a synthetic miR-1-3p mimic results in masculinization of XX individuals. These data support the conclusion that the miR-1-3p microRNA has a necessary intermediary function required for the induction of male sex determination in B. dorsalis.

Results

Analysis of small RNA libraries from embryonic stages. To identify factors that are involved in B. dorsalis sex determination, we first determined when Bdtra-f and Bdtra-m isoforms appear during embryogenesis. This was achieved by analyzing Bdtra sex-specific splicing isoforms in individual embryos at distinct hours post-oviposition (hpo) time points, thereby distinguishing the chromosomal sex of the embryos and their sexual fate. At 5 hpo, only female-specific Bdtra transcript was apparent in individual embryos, which was most likely due to maternal deposition of Bdtra-f in both XX and XY embryos. Male-specific Bdtra-m transcript was first detectable at 6 hpo while Bdtra-f transcript was not detectable at 7 hpo, presumably in XY male embryos. Embryos that expressed only Bdtra-f transcripts at 6 and 7 hpo were presumably XX (Fig. 1). These results suggest that the key stage for specifying male sexual development is between 5 and 6 hpo at 28 °C.

To identify candidate small RNAs that may be involved in sex determination, small RNA sequencing was performed using embryos of mixed sex at 5, 6, and 7 hpo (NCBI accession numbers: GSE117310). The three libraries produced a total of 6,178,809, 5,695,996, and 9,255,550, clean reads, respectively (Supplementary Table 1), and a total of 147 known and 103 novel miRNAs were identified in the combined data set (Supplementary Data 1 and 2). The majority of the small RNA reads ranged in size from 20 nt to 23 nt and had a peak at 22 nt, which is the typical length of mature miRNAs (Supplementary Fig. 1A). The first nucleotide position was enriched in uracil (U) with a frequency of

Fig. 1 The splicing patterns of the Bdtra pre-mRNA transcript. Shown in this figure are RT-PCR products from single B. dorsalis embryos at 5, 6, and 7 h post-oviposition (hpo). Bdtra-f and Bdtra-m are the female and male-specific isoforms of the transformer transcript, respectively. The sex of each embryo is distinguished by the presence of Bdtram that is only expected in XY embryos. The samples derive from the same experiment and that gels were processed in parallel.
67% (Supplementary Fig. 1B), consistent with one of the most fundamental features of miRNA sequences.

To study differential expression of known miRNAs within the three embryonic stages, the miRNA counts were normalized by transforming them into Reads Per Kilobase Per Million Mapped (RPKM). Sixty-five miRNAs were differentially expressed at 5, 6, and 7 hpo, according to a Fisher’s test and a chi-squared test (P-value < 0.05) (Supplementary Table 2). All 65 miRNAs, including miR-1-3p, are transcribed from autosomes and eight were found to potentially target the *Bdtra* gene (Supplementary Table 3). The putative target sequences of all eight miRNAs were found to be in the 3’ UTR of the *Bdtra* gene, with the smallest free energy value observed for miR-1-3p (Fig. 2; GenBank ID: JBF01001330.1).

### miR-1-3p regulates *Bdtra* transcript accumulation.

To test whether these eight miRNAs affect *Bdtra* expression in vitro, a dual luciferase assay was performed. A firefly luciferase reporter construct was made by inserting the 3’ UTR of *Bdtra* downstream of the *Renilla* luciferase coding sequence of the pSCECHECK-2 vector. To reduce interaction of endogenous miRNAs and mRNAs with our test system, we used a human cell line (HEK293T) for co-transfection with the dual-luciferase reporter and miRNA mimics. Whereas co-transfection with the miR-8-3p, miR-10-5p, miR-308-3p, miR-10-3p and miR-305-5p mimics did not significantly alter the luciferase signal, co-transfection with the miR-276a-3p, miR-1-3p and miR-12-5p mimics resulted in approximately 33%, 46% and 32% reductions in luminescence compared to controls, respectively (Supplementary Fig. 2). This suggested that only these three of the eight tested miRNAs targeted *Bdtra* in human cell lines.

To determine the interaction between miR-276a-3p, miR-1-3p, miR-12-5p and *Bdtra* in vivo, we injected newly oviposited embryos (0–1 h) with corresponding miRNA agomirs (mimics), antagomirs (inhibitors) (Supplementary Fig. 3) and a *Bdtra* dsRNA/antagomir. Following injection of the miR-1-3p agomir, *Bdtra* mRNA decreased significantly by 62%, similar to the 68% decrease resulting from dsBdtra injection in 10 h post-injection embryos (Fig. 3a and Supplementary Fig. 4A), whereas injection of the miR-1-3p antagomir inhibitor resulted in a 109% increase in *Bdtra* mRNA (Fig. 3b). Co-injection of antagomir-1-3p/ds*tra* decreased *Bdtra* transcript by 72%, whereas the control co-injection of antagomir-1-3p/ds*egfp* showed a similar increase in *Bdtra* transcript compared to the antagomir-1-3p treatments in 10 h embryos (Supplementary Fig. 4B). However, the expression level of *Bdtra* was not affected by miR-276a-3p and miR-12-5p antagomir and antagomir treatments (Fig. 3a, b). Together, these results indicate that miR-1-3p down-regulates the level of *Bdtra* mRNA.

### miR-1-3p regulation of *Bdtra* is *BdDicer-1*-dependent.

We analyzed the influence of depletion of *BdDicer-1*, the core component of the miRNA biogenesis pathway, on miRNAs and *Bdtra* expression during embryogenesis. First, we analyzed the structure of *BdDicer-1* and assayed its expression. Structural analysis revealed that *BdDicer-1* contains an amino-terminal helicase domain, a PAZ domain, two RNAse III domains, and a dsRNA-binding domain (Supplementary Fig. 5). An expression profile of *BdDicer-1* transcript from different developmental stages showed that *BdDicer-1* is highly expressed in newly laid embryos of mixed sexes and showed increased levels at 6 and 7 hpo in early embryogenesis (Supplementary Fig. 6A, B).

To test the effect of *BdDicer-1* depletion on mature miRNA formation and *Bdtra* expression, we prepared a 643 bp dsRNA from the RNAse III domains of *BdDicer-1*. The *BdDicer-1* RNAi confirmed that transcriptional levels of the eight differentially expressed miRNAs were lower relative to *ds*egfp* controls in 10 h embryos of mixed sexes. Among them, RNA levels of miR-1-3p, miR-8-3p and miR-308-3p significantly decreased by 56%, 58% and 51%, respectively (Supplementary Fig. 6C), whereas *Bdtra* mRNA levels in dsBdDicer-1 treated embryos increased significantly by 137% compared to the *ds*egfp* controls (Fig. 3d). A rescue experiment using miR-1-3p agomir in *BdDicer-1* knockdown individuals showed that miR-1-3p transcript increased greater than 12-fold in ds*BdDicer-1*/agomir-1-3p treated embryos of mixed sexes (Fig. 3c). In those embryos

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**Fig. 2 miRNAs that potentially target the *Bdtra* gene.** a The position of miRNAs putative binding sites in the 3’ UTR of *Bdtra*, and (b) RNA-RNA duplexes of potential miRNA with the *Bdtra* gene as detected by RNA hybrids. miRNA seed sequences and their putative binding sites in the 3’ UTR are indicated by red. mfe: match free energy.
miR-1-3p is required for male sex determination in embryos.

Based on the detection of only Bdtra-m in single XY embryos, which is the major characteristic of sex-specific splicing of Bdtra pre-mRNA, we identified the sex of individual female and male embryos at 6, 7, 8, and 9 hpo. The expression pattern of the miR-1-3p gene in 0–9 hpo embryonic stages, including mixed embryos at 0, 1, 2, 3, 4, 5 hpo, and single female and male embryos at 6, 7, 8, 9 hpo, showed that miR-1-3p levels increased at 0 hpo to 6 hpo. Levels increased more so from 5 hpo to 6 hpo in mixed embryos (Supplemental Fig. 7A), and the relative expression levels of miR-1-3p in single male embryos were significantly higher than in single female embryos (Fig. 4a). The temporal profile in immature (IM), middle-aged (MA) and fully mature (FM) female and male adults confirmed that miR-1-3p has sexually dimorphic expression also during adulthood (Supplemental Fig. 7B). Our sequencing data indicated that miR-1-3p, having the Genbank ID: JFFB01001330.1, is transcribed from an autosome and is not X-linked (Supplementary Data 1). Furthermore, miR-1-3p PCR amplification from adult female and male genomic DNA also confirmed that miR-1-3p is not sex-specific and, thus, not linked to the male Y chromosome (Fig. 4b). These results indicated that a sexually dimorphic switch in miR-1-3p expression results in sex-specific splicing of tru between 5 and 6 hpo and, thus, has a role in inducing male sex determination in the XY embryo.

To investigate the role of miR-1-3p in determining male sex in early embryos, wild type embryos were injected with agomir-1-3p and antagonir-1-3p. We found that pooled embryos injected with agomir-1-3p resulted in progeny composed of 87–92% phenotypic males and pools injected with antagonir-1-3p resulted in 67–77% phenotypic females, whereas the female to male ratio of uninjected, agomir NC/antagomir NC control, and miR-1-3p mutant agomir/antagomir injected eggs was approximately 1:1 (Supplementary Table 4, 5). Two to 4% intersexual individuals exhibited a female dorsal phenotype, lacking an ovipositor, in agomir-1-3p injected pools and 2–6% intersexual individuals exhibited a male dorsal phenotype by the absence of bristles on the side of the third tergum in antagonir-1-3p injected pools (Fig. 5 and Supplementary Table 4). The low feminization rate of antagonir-injected embryos may be explained by the transient nature of embryonic artificial miRNA interference. A 76% reduction of miR-1-3p transcript levels was observed in embryos 10 h after antagonir-1-3p injection, whereas after 20 h, levels were comparable to those in control individuals, suggesting

**Figure 3.** miR-1-3p targets the Bdtra transcript in vivo. a Levels of Bdtra in 10 h embryos injected with the miR-1-3p, miR-12-5p or miR-276a-3p agomir, (b) levels of Bdtra in 10 h embryos injected with the miR-1-3p, miR-12-5p or miR-276a-3p antagonir, (c) levels of miR-1-3p in 10 h embryos injected with dsBdDicer-1 and the miR-1-3p agomir, and (d) levels of Bdtra in 10 h embryos injected with dsBdDicer-1 and the miR-1-3p agomir. Error bars indicate the SEM of three independent biological replicates and asterisks (* and ** and *** indicate the statistically significant differences (P < 0.05 and P < 0.01 and P < 0.001) between the treatment group and control group based on Student’s t-test. Ns indicates no statistically significant difference between the treatment group and the control group.
a recovery of miR-1-3p expression (Fig. 6a). The expression of Bdtra increased 147% in embryos 10 h after antagomir-1-3p injection with no significant difference at 20 h after injection (Fig. 6b). The incompletely feminized intersexual individuals had immature ovaries with non-vitellogenic oocytes, whereas the normal females had normal vitellogenic ovaries with mature oocytes (Figs. 6c, d).

To further test whether miR-1-3p is required for male sex determination, miRNA-specific mutations were generated by CRISPR/Cas9 gene-editing (Supplementary Fig. 8). The 706 embryos injected together with Cas9-mRNA and miR-1-3p sgRNA resulted in 72 surviving adults comprised of 58 females, 11 males and 3 intersexual individuals, yielding a female to male sex ratio of 6:1. In contrast, 722 embryos injected with Cas9-mRNA/egfp sgRNA resulted in 85 adult survivors having a phenotypic female to male ratio of nearly 1:1 (41 females: 44 males) (Fig. 7a and Supplementary Table 6). A PCR-based analysis using DNA extracted from G0 individual adults, injected with Cas9/sgRNA as embryos, showed that 21 of the 58 G0 females and 3 intersexual individuals had indels in the targeted miR-1-3p region, whereas none of the 11 male adults exhibited miR-1-3p mutations, nor did the control injected individuals. Deletions ranged from 1 bp to 7 bp in the miR-1-3p mutated sequence (Fig. 7b) and a T7 Endonuclease I (T7EI) assay confirmed that PCR fragments spanning the miR-1-3p target sites could be cut into expected size bands in mutant individuals (Fig. 7c). This is consistent with a genomic disruption of miR-1-3p by the CRISPR/Cas9 system resulting in some chromosomal XY male individuals undergoing a sex reversal into phenotypic females.

However, we could not definitively determine the chromosomal sex of the 21 mutant phenotypic females (Fig. 7a), which was achieved by using the white/brown pupae strain developed for B. dorsalis sexing in the sterile insect technique28. In this strain, XX individuals homozygous for the recessive white pupae (wp−) mutation develop as non-pigmented white pupae, whereas XY individuals, having a wild type wp allele (wp+) translocated to the Y chromosome, develop as normally pigmented brown pupae (Supplementary Fig. 9). Thus, the CRISPR/Cas9 experiment was repeated in this strain to confirm the feminization of brown pupal XY males as a result of mutations in miR-1-3p (Fig. 8a, c). In the experimental group, 714 embryos injected with miR-1-3p sgRNA/Cas9 resulted in 62 surviving adults consisting of 26 phenotypic females and 36 phenotypic males (Fig. 8a and Supplementary Table 7). Of these adults, 47 developed from XY brown pupae, 11 of which were sexually reversed phenotypic XY females (Fig. 8a) that carried indel mutations in the targeted miR-1-3p region (Figs. 7b, c). They also expressed the female splice variants of Bdtra and Bddsx and had normal ovaries (Fig. 8b, d). The remaining 36 adults developing from XY brown pupae were normal phenotypic males. Among 15 adult females that developed from XX white pupae, three had miR-1-3p mutations and expressed the normal female splice variants of Bdtra and

**Fig. 4** The differential expression of miR-1-3p in female and male embryos. (a) Relative levels of miR-1-3p in single female and male embryos at 6, 7, 8, and 9 h post-oviposition (hpo). Female and male embryos were identified by the splicing patterns of Bdtra pre-mRNA. (b) PCR amplification of miR-1-3p from adult female and male genomic DNA. Error bars indicate the SEM of three independent biological replicates and asterisks (**) indicate the statistically significant differences (P < 0.01) between the treatment group and control group based on Student’s t test.

**Fig. 5** Dorsal view of abdominal intersex phenotypes induced by embryonic injection. (a) Wild type female adult, (b) intersex of incomplete masculinization individuals exhibiting a female dorsal phenotype but lacking the ovipositor after miR-1-3p agomir injection, (c) dorsal overview of WT male adult, and (d) intersex due to incomplete feminization exhibiting a male dorsal phenotype without bristles on the side of the third tergum after the miR-1-3p agomir and sgRNA injection.
Bddsx. The backcross experiment of the 11 feminized XY females mated to normal XY males showed that eight (73%) mated normally, both the number of eggs per female and their hatching rate were similar to control injected (egfp sgRNA/Cas9) females (Supplementary Fig. 10), and the F1 adult sex ratios had no significant bias to either sex (Supplementary Table 8). The 729 control embryos injected with egfp sgRNA/Cas9 resulted in 75 non-mutant adult survivors (Fig. 7a and Supplementary Table 6) consisting of 17 phenotypic females and 58 phenotypic males, having an approximate 1:3 sex ratio (as observed in the untreated strain, consistent with Mendelian segregation of the dominant marker). The mRNA and protein levels of Bdtra were also investigated in normal and miR-1-3p mutant female and male individuals by qPCR and Western blots. The results showed that both mRNA and protein levels of Bdtra gene were female-specific in mutant feminized XY males, whereas they were hardly detectable in normal males (Figs. 9a, b). Taken together, these results show that CRISPR/Cas9 embryonic mutagenesis of miR-1-3p in XY individuals confers a complete male to female sex reversal consistent with its requirement for male development.

To further confirm that the miR-1-3p is required for masculinization, we investigated the influence of miR-1-3p overexpression on sexual differentiation by injection of agomir-1-3p into embryos of the white/brown pupae sexing strain. We observed 52 surviving adults consisting of 39 XY (wp+) brown phenotypic males, 5 phenotypic XX (wp-) white females, and 8 XX (wp-) white phenotypic males (Fig. 8a and Supplementary Table 9). The backcross experiments of the 8 masculinized XX males mated to the normal XX females, showed that five (63%) mated normally, and both the number of eggs per female and their hatching rate were similar to the control injected (agomir NC) treatment (Supplementary Fig. 11) and the majority of F1 progeny were female (Supplementary Table 10). For a control test, 697 white/brown pupae sexing strain embryos were injected with agomir NC, which resulted in 52 adult survivors consisting of 11 phenotypic females and 41 phenotypic males, with all males developing from brown pupae (Fig. 7a and Supplementary Table 9), consistent with Mendelian segregation of the dominant marker. Thus, the masculinization of XX white pupae by agomir-1-3p treatment indicates that the induced overexpression of miR-1-3p is sufficient to initiate male development in B. dorsalis in the absence of the Y chromosome.

Discussion

The primary molecular signals that determine sex are highly divergent in insects, but in several Orders they act on a functionally conserved key female-determining gene, transformer29-31. In this study, we showed that 5 to 6 hpo is a key stage for initiating male sex determination in B. dorsalis, during which the male-specific Bdtra-m transcripts appeared and the female-specific Bdtra-f transcripts degraded and became undetectable by 7 hpo at 28 °C. This timing is similar to that found in the other Bactrocera species, B. jarvisi and B. tryoni, where degradation of maternally deposited tra-f begins at 5–6 hpo and 9 hpo in male embryos at 25 °C, respectively32, whereas in C. capitata it occurs at 6-7 hpo at 25 °C. This time period for establishing the sex-determined state is coincident with the stage at which the proposed M-factor is likely to be active, for example, in maternal tra degradation32,33. The establishment of male sex determination during early embryogenesis in Bactrocera species, is also coincident with a similar sex-determining developmental stage in other insect species, including the dipterans, D. melanogaster34.
strain, in particular, showed that the miR-1-3p specific mutation in XY individuals results in female expression of Bdtra and complete adult feminization, whereas XX females developed normally. This proposed role for miR-1-3p in B. dorsalis male sex determination would be the converse of the proposed role for the Fem piRNA in B. mori, as the primary female-determining factor in the ZW/ZZ sex determination system. Here, the Fem piRNA represses Masc mRNA expression required for male Bmdsx pre-mRNA splicing and male sexual development. Nevertheless, the roles of miR-1-3p miRNA and Fem piRNA in the regulation of tra and Masc, respectively, indicate that small RNAs, in different orders of insects, have essential roles as post-transcriptional factors during the initiation of sex determination in early embryogenesis. However, miR-1-3p is a non-Y chromosome-linked miRNA with higher transcript levels in early XY embryos, which differs from Fem piRNA which is expressed from the female-specific W chromosome in B. mori. The dominant component of the silkworm W chromosome is female-enriched piRNAs gene clusters, whereas the major component of tephritid Y chromosome is repeat-rich sequences, as reported for C. capitata and Nasonia vitripennis. Indeed, the diversity of heterogametic sexual determinants between B. mori and B. dorsalis may explain, in part, the differing roles for small RNAs and mRNAs in regulating sex determination in the two species. Our study extends this pathway of interactions, at least for a tephritid species, by showing that miR-1-3p is a transducer of the Y-linked primary signal for male sex determination and may play a more direct role as a male-determiner by promoting male-specific tra pre-mRNA splicing and, hence, its functional inactivation in XY embryos.

These results lead us to propose an updated model for the sex determination pathway in B. dorsalis (Fig. 10), that might be extended to related tephritid species. Since the miR-1-3p gene is present in both sexes, albeit only transcribed at high levels in male embryos during a developmental period critical to sex determination, and is thus not Y-encoded, this model proposes that its expression is activated in early embryogenesis by a Y-linked primary signal in male embryos. This primary signal, is the putative male determining factor (M-factor), whereas miR-1-3p has a more direct role in inhibiting maternal Bdtra expression. The inhibition of maternal Bdtra expression would result in male-specific splicing of zygotic Bdtra pre-mRNA, producing a truncated BdTRA unable to maintain the threshold level needed to sustain the feedback loop of Bdtra expression. Thus, a lack of BdTRA protein in XY would lead to female sexual differentiation. However, due to low expression levels of miR-1-3p in female embryos, insufficient inhibition of maternal Bdtra transcript would allow it to translate into functional BdTRA protein to initiate the female-specific autoregulatory loop in the zygotes. BdTRA, in concert with Bdr-TRA-2, would then result in female-specific splicing of the downstream Bddxs gene pre-mRNA, thus promoting female sexual differentiation.

In summary, we have shown that miR-1-3p is essential in determining the male sex in B. dorsalis through a combination of miR-1-3p knockdown, knockout and over expression experiments. This is the first report of a direct role for a miRNA-mediated sex determination mechanism in insects, and the second finding of a role for a small RNA, although the B. mori piRNA and miR-1-3p act conversely. This suggests that small RNAs may have an expanded and varying repertoire of regulatory functions for sex specification in insect species, and perhaps other organisms. For B. dorsalis in particular, manipulation of miR-1-3p may be used for sexing in SIT, for example, by producing male-only populations by its conditional ectopic overexpression in XX females during embryogenesis.
Fig. 8 miR-1-3p is sufficient to initiate male development in B. dorsalis. (a) The sexual phenotype of the white/brown pupae sexing strain (wp+/−) adults having an miR-1-3p knock-out deletion (miR-1-3p−) mediated by CRISPR/Cas9 and after embryonic injection of the miR-1-3p mimic (miR-1-3p+). wp+ are genetic males (XY) and wp− are genetic females (XX). The red bar represents the number of miR-1-3p mutant adults, and the blue bar represents the number of phenotypic male adults which developed from the white pupae (XX wp−) embryos treated with miR-1-3p agomir. NC is the control group injected with the agomir negative control. (b) Sex-specific transcripts of Bdtra and Bddsx in WT females and males and XY miR-1-3p mutant phenotypic females. −1, −2, −4, −5, and −7 represent different types of miR-1-3p mutant females. (c) Phenotypes of XX wp− female (left) and XY wp− male (middle) adults from the white/brown pupae sexing strain, and an XY wp+ miR-1-3p mutant phenotypic female (right) induced by CRISPR/Cas9. (d) Mature vitellogenic ovaries from an XX wp− female (left) and an XY wp+ miR-1-3p mutant phenotypic female (right).

Methods

Insect rearing. B. dorsalis wild type and white/brown pupae sexing (provided by Dr. Eric Jang, USDA-ARS, Hilo, HI) strains were reared at the Institute of Horticultural and Urban Entomology at Huazhong Agricultural University (Wuhan, China). Adult flies were fed an artificial diet consisting of yeast extract and sugar and cultured at 28 °C under a 12 h light: 12 h dark photoperiod in cages, and hatched larvae were maintained on bananas18. The sex ratio of normally bred adult females to males is approximately 1:1 for wild type and 1:3 for the sexing strain (provided by Dr. Eric Jang, USDA-ARS, Hilo, HI) strain. This sex ratio was confirmed by sexing flies from each wild-type and sexing strain experiments. The sex ratio of normally bred adult females to males is 1:1 for wild type and 1:3 for the sexing strain. We have complied with all relevant ethical regulations for B. dorsalis testing and research.

Sample preparation. Embryo total RNAs were extracted using RNAiso Plus reagent (TaKaRa, Japan) according to the manufacturer’s protocol. First strand cDNA was synthesized from 1ug total RNA using PrimeScript® Reverse Transcriptase (TaKaRa, Dalian, China) with the oligo (dT) adapter primer. Female and male transcripts from Bdtra were amplified by RT-PCR from 5, 6, 7, 8, and 9 h post-oviposition (hpo) embryo cDNA with Bdtra sex-specific exon primers (Supplementary Table 1). Embryos at 5, 6 and 7 hpo were collected and stored in RNAlater® Solution (Ambion, Austin, TX, USA). To identify the small RNAs involved in B. dorsalis primary sex determination, three small RNA libraries were constructed from the 5, 6 and 7 hpo samples. Total RNAs were extracted using RNAiso Plus reagent (TaKaRa, Dalian, China) according to the manufacturer’s protocol, with the quantity and purity examined using a Bioanalyzer 2100 (Agilent, CA, USA).

Small RNA library construction and deep sequencing. Small RNA libraries were generated from the three RNA samples using the TruSeq Small RNA Sample Prep Kits, following manufacturer’s guidelines (Illumina, San Diego, USA). The 18-30 nt length RNA fractions were separated by gel electrophoresis and ligated sequentially to 3′ and 5′ adaptors. Ligation products were then reverse transcribed using a primer to the 3′ adaptor with 15 PCRs amplified for the first strand synthesis production with both adapter sequence primers. Ultimately, PCR products were purified, validated and sequenced using an Illumina HiSeq 2500 System (LC-Sciences, Hangzhou, China).

Small RNA sequence analysis. After removing low quality reads and adapter sequences from the raw data, large amounts of clean small RNAs were analyzed by
Differentiation. The components that are inactive are shown in gray. In XX embryos low expression levels of miR-1-3p results in insufficient inhibition of maternal Bdtra gene, allowing it to translate into sufficient functional BdTRA protein to initiate the female sexual fate autoregulatory loop in zygotes. BdTRA protein then activates the female-specific splicing of the downstream Bddsx pre-mRNA, thus, promoting female sexual differentiation. The components that are inactive are shown in gray.

**Comparison of differentially expressed miRNAs.** As male sex determination occurs within 5, 6, and 7 hpo, the expression patterns of miRNAs during these three key embryonic stages were compared to identify differentially expressed miRNAs of *B. dorsalis* (NCBI Assembly# ASM78921v2) and hairpin RNA structures containing sequences were predicted from *B. dorsalis* full length sequence of the 3′ UTR of *Bdtra* and miRNA sequences were submitted to the miRBase database for searching of miRNA precursors/mature miRNA sequences and previously identified miRNAs of *B. dorsalis* with the matched or single mismatch sequences, were identified as known miRNAs. Unmapped sequences were aligned to the whole genome sequence (WGS) of *B. dorsalis* (NCBI Assembly# ASM78921v2) and hairpin RNA structures containing sequences were predicted from flanking 80 nt sequences using RNAfold software (http://rna.tbi.univie.ac.at/cgi-bin/RNAfold.cgi) complying with criteria from pre-miRNA statistics in miRBase to identify potentially novel miRNAs.

**Target gene prediction.** For the prediction of miRNAs targeting the *tra* gene in *B. dorsalis*, three different target prediction programs were used: miRanda, RNAhybrid, and TargetScan. To characterize the interaction of the *Bdtra* gene (GenBank ID: KP769599.1) with differentially expressed miRNA candidates, the full length sequence of the 3′ UTR of *Bdtra* and miRNA sequences were submitted on-line to RNAhybrid, an algorithm taking into account the free energy level of RNA-RNA duplexes and degree of miRNA target sequence complementarity using default parameters.

**Cell culture and dual luciferase reporter (DLR) assay.** HEK293T cells provided by Prof. Shengbo Cao (Laboratory of Animal Virology, College of Veterinary Medicine, Huazhong Agricultural University) were cultured and maintained in Dulbecco’s Modified Eagle’s Medium (Gibco, Carlsbad, CA, USA) supplemented with 10% fetal bovine serum (Gibco, Carlsbad, CA, USA), and 1% penicillin/streptomycin (Gibco, Carlsbad, CA, USA) at 37°C in 5% CO2.

The *Bdtra* 3′ UTR (332 bp) was amplified by PCR from *B. dorsalis* cDNA and inserted into the psiCHECK-2 vector (Promega, Madison, WI, USA) which contains the firefly and renilla luciferase gene using Not I and Xho I sites, respectively (see sub-cloning PCR primers in SI Appendix, Table S8). After growth in 96-well plates with serum-containing medium for 12 h (104 seeded cells per well), cells were transfected with 50 nM miRNA agomir or the negative control (NC) and 100 ng luciferase reporter vector psiCHECK-2, containing the *Bdtra* 3′ UTR, in each well using 0.5 μl FuGENE® HD (Promega, Madison, WI, USA), 48 h after transfection, cells were lysed and the activity of firefly and renilla luciferase was determined with the Dual-Luciferase® Assay System (Promega, Madison, WI, USA) according to the manufacturer’s protocol. Firefly luciferase values were normalized to renilla, and the ratio of firefly/renilla was presented.

**miRNA agomir/antagomir and dsRNA embryo microinjection.** A 598 bp fragment of the *Bdtra* gene and a 643 bp fragment of the *BdDicer-1* gene (RNase III coding region) were produced by PCR, and a control *egfp* fragment was amplified with primers from the *egfp*-containing vector pPBSiLsEGFP® (Supplementary Table 1) according to the transformation procedure described by Sharma et al.®. Equal volumes of NCs and *egfp* were injected for controls. Ten hours post-injection, ten embryos from each treatment group were assayed for expression of miR-1-3p and *Bdtra* that was replicated at least three times for each treatment group. In miR-1-3p agomir/antagomir injection experiments embryos were incubated in an oxygen-saturated culture chamber at 28 °C. Surviving larvae were reared and placed on banana media until eclosion. Adults were collected and their emergence rate, sex ratio, and morphological changes were recorded.

**In vitro synthesis of Cas9-miRNA and sgRNA.** A miRNA-targeted genomic mutation was created by CRISPR/Cas9 using a miR-1-3p specific sgRNA according to the methods of Ling et al.® and Zhang et al.®. To design the miR-1-3p specific sgRNA, miR-1-3p mature regions were searched with the protospacer-adjacent motif (PAMs) of the recognition site for Streptococcus pyogenes (SpCas9) and predicted the target sequences for the Cas9 endonuclease digestion products were purified using the E.Z.N.A.®, Cycle-Pure Kit (Omega, Norcross, GA, USA). In vitro transcription of sgRNA and Cas9-miRNA were carried out following the instructions of the MEGAscript® Kit (Ambion, Austin, TX, USA). The sgRNA and Cas9-miRNA were purified with the MEGAclear® (Promega, Madison, WI, USA) and stored at -80 °C until use. Embryos were microinjected with a mixture of purified Cas9-coding miRNA and miR-1-3p sgRNA according to the microinjection protocol described above. Final concentrations of Cas9-coding miRNA and miR-1-3p sgRNA were 300 ng/μl and 150 ng/μl respectively and Cas9-coding miRNA and *egfp* sgRNA were used as controls. Surviving larvae were placed on banana media until adult eclosion at which time G0 adults were collected with their emergence rate, sex ratio and morphology changes recorded.

**Fertility tests and mutagenesis analysis.** Phenotypic females that developed from chromosomal male (*y*, wp®) brown pupae and phenotypic males that developed from chromosomal female (*wp−*) white pupae were reared and tested for fertility under standard laboratory conditions. Each female was backcrossed to three virgin normal XY males and each male was backcrossed to three virgin normal XX females in separate cages. Matings were observed at 30 min intervals during the dark photoperiod, and eggs from each cage were collected and counted continuously for 5 days. The newly hatched larvae were counted and then transferred onto banana until adult emergence. To molecularly characterize CRISPR/Cas9 induced mutations, genomic DNA was extracted from G0 individuals following the E.Z.N.A.® Insect DNA Kit protocols (Omega, Norcross, GA, USA). The miR-1-3p target loci were amplified by PCR using the following conditions: 94°C for 3 min, 35 cycles of 94°C for 30 s, 55°C for 30 s, and 72°C for 30 s, followed by a 7-min extension at 72°C.
final extension at 72 °C for 10 min. PCR products were cloned into pEASY-T1 vector (TransGen Biotech, Beijing, China) and sequenced using GenScript (Nanjing, China). The PCR products were also used for the T7 endonuclease I (T7EI) assay according to the manufacturer’s instructions (V-soldi, Beijing, China). For expression analysis of the Btdra and Bddxs genes, total RNA was extracted and cDNA prepared from a G0 individual. Primers that were designed to detect mutagenesis in targeted sites and sex-specific expression of Btdra and Bddxs are listed in Supplementary Table S11.

Quantitative real-time PCR of miRNA and miRNA. Quantitative real-time PCR (qPCR) was used to assay the developmental transcript expression profiles of Bdl-Dicer-1 in 0, 5, 6, and 7 hpo embryos, 1st, 2nd, and 3rd instar larvae, early and late stage pupae, and 1 and 10 d female and male adults. qPCR was also used to test transcript expression levels of miR-1-3p and Bdra in miR-1-3p agomir/antagomir and ds/tras/dicer-1 treated individuals. Total RNA was extracted using RNAiso Plus reagent (TaKaRa, Dalian, China), with 200 ng for each sample subjected to reverse transcription of mRNA using the PrimeScript™ RT Master Mix (TaKaRa, Dalian, China). Reverse transcription reactions for mature miRNAs were conducted with 1 ug total RNA using miRNA specific stem-loop primers (Supplementary Table S11). The reverse transcription products from miRNAs and miRNA were used for qPCR using primers listed in SI Appendix, Table S11. qPCR was performed using the SYBR Green qPCR mix following the manufacturer’s instructions in a real-time thermal cycler (Bio-Rad, Hercules, CA, USA) using the test technological blot detection was performed using an ECL Luminol Substrate Reagent (Frdbio, China).

Statistical analysis. All experiments were repeated at least three times, except for the small RNA sequencing, and analyzed using GraphPad Prism 5.0 (GraphPad Software, San Diego, CA, USA) or Microsoft Excel (Microsoft, Redmond, WA, USA). Results are expressed as the mean ± SEM. Data was compared with either a two-way ANOVA, with subsequent t tests using Bonferroni post-tests for multiple comparisons, or with the Student’s t test. For all tests, differences were considered significant when P < 0.05.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability. The miRNA-seq data presented in this article are deposited in GEO database: GSE117310. All relevant data are available from the authors. The source data underlying Figs. 1, 3a–d, 4a, b, 5a, 6a–d, 7c, 8b–d and 9a, b and Supplementary Figs 2, 4a, b, 6a–c, 7a, b, 10a, b and 11a, b are provided as a Source Data file.

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**Author contributions**

H.Z. and W.P. conceived of and designed the study, and W.P. conducted all experiments related to the functional analysis of miR-1-3p, with the assistance of S.Y. in embryo microinjections. A.M.H. instructed W.P. on the embryo microinjection technique. G.S. and Z.J.T. analyzed the data and Z.X. provided the white/brown pupae sexing strain. W.P., H.Z., and A.M.H. wrote the manuscript with intellectual input from all authors.

**Competing interests**

The authors declare no competing interests.

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