DNA methylation changes induced by long and short photoperiods in *Nasonia*

Mirko Pegoraro,1,3 Akanksha Bafna,1,3 Nathaniel J. Davies,1 David M. Shuker,2 and Eran Tauber1

1Department of Genetics, University of Leicester, Leicester LE1 7RH, United Kingdom; 2School of Biology, University of St Andrews, St Andrews KY16 9TH, United Kingdom

Seasonal timing is a key survival process for many organisms, especially in temperate regions. In most cases, light is the primary cue, and the seasonal change in day length is monitored by a so-called “photoperiodic clock” (Bradshaw and Holzapfel 2010). Despite intensive study of the photoperiodic clock for the last 80 years at the phenomenological level, the “black box” approach has provided almost no information about the underlying molecular mechanisms (Saunders et al. 2004). This is in marked contrast to our level of understanding of the circadian clock that regulates daily rhythms, in which studies in various model organisms have led to the discovery of general principles and a detailed understanding of the underlying molecules, which are highly conserved across diverse phyla (Hogenesch and Ueda 2011).

*Nasonia* is a typical temperate-zone insect, in which short photoperiods experienced by females lead to the induction of developmental arrest (diapause) in their progeny, allowing winter survival of the larvae. How female *Nasonia* control the developmental trajectory of their offspring is unclear. Here, we took advantage of the recent discovery that DNA methylation is pervasive in *Nasonia* and tested its role in photoperiodism. We used reduced representation bisulfite sequencing (RRBS) to profile DNA methylation in adult female wasps subjected to different photoperiods and identified substantial differential methylation at the single base level. We also show that knocking down DNA methyltransferase 1a (*Dnmt1a*), *Dnmt3*, or blocking DNA methylation pharmacologically, largely disrupts the photoperiodic diapause response of the wasps. To our knowledge, this is the first example for a role of DNA methylation in insect photoperiodic timing.

[Supplemental material is available for this article.]
DNA methylation in females and observe how these manipulations influence offspring diapause. We use two independent techniques that influence genome-wide methylation in different ways: (1) RNA-interference knockdown of Dnmt1a-c and Dnmt3; and (2) pharmacological knockdown using the methylation inhibitor 5-aza-2′-deoxycytidine.

**Results**

**The Nasonia methylome**

To identify DNA methylation changes induced by photoperiod, we quantified DNA methylation in genomic DNA extracted from whole female *Nasonia vitripennis* wasps kept in either long- or short-day conditions. We used reduced representation bisulfite sequencing (RRBS) (Gu et al. 2011) to achieve single-nucleotide resolution of DNA methylation of CpG dinucleotides distributed throughout the *Nasonia* genome at high coverage.

After removal of low-quality reads, we obtained 27,255,357 reads (1.06 Gbp) from the long-day sample (LD) and 30,818,609 reads (1.25 Gbp) from the short-day sample (SD). The mapping efficiency of these reads to the *Nasonia* reference genome was 84.1% (LD) and 83.2% (SD). The average coverage depth of the reads was 92.6× for the short-day sample and 86.4× for the long-day sample. By the same metric, 4.53% (1,270,716) of CpG sites in the genome were covered by both samples at an average depth of 95.72× for the SD sample and 88.97× for the LD sample. The average coverage depth of the reads was 92.6× for the short-day sample and 86.4× for the long-day sample. By the same metric, 4.53% (1,270,716) of CpG sites in the genome were covered by both samples at an average depth of 95.72× for the SD sample and 88.97× for the LD sample. The LD reads covered 1,304,087 CpG sites (4.65% of all genomic CpGs), whereas the SD reads covered 1,385,067 CpG sites (4.94%). We detected 1618 and 2018 methyl CpGs in the LD and SD samples, respectively, using a 1% false discovery rate (FDR) (see Methods), comprising 0.124% and 0.146% of the cytosines with sequence coverage that were analyzed.

Although most of the methylated cytosines were in a CpG context (Fig. 1A), a small fraction (<6.7%) of these were in non-CpG contexts (mCHG, mCHH, where H = non-G). A Venn diagram showing that a substantial proportion of the mCGs are methylated under both long- and short-day conditions in contrast to other sequence contexts that show negligible overlap (see text), suggesting that non-CpG methylation represents largely experimental noise. (C) The methylation level of CpG sites (percentage of covering reads that are methylated) shows a bimodal distribution. This bimodality is not seen in CHG or CHH (shaded yellow and orange) methylation contexts. (D) Fraction of mCgs identified in exon, intron, promoters (2-kb upstream of genes), and intergenic regions. (E) A snapshot of the methylome (751-kb region) is depicted overlaid with the raw long-day methylation data (scaled to 0%–2% methylation).

**Identification of differentially methylated genes**

Of the 12,055 genes with annotated mRNAs in the Nvit_2.0 build, 5972 (49.54%) had at least five covered CpG sites, and 5032 (41.74%) had at least 10 covered CpG sites. We used Fisher’s exact tests to compare methylation in long and short photo-periods and identified 51 differentially methylated CpG sites.
(DMCs; Benjamini-Hochberg multiple testing, FDR < 0.05) (Benjamini and Hochberg 1995), which were mapped to 37 unique genes (Fig. 2A; Supplemental Table S1). Approximately half of the DMCs (23/51) showed reduced methylation in long day compared to short day (i.e., hypomethylation), whereas the other showed the opposite trend. Because the annotation of the Nasonia genome is still in its infancy, we were able to assign gene ontology (GO) terms only to 25 genes in our list. The terms assigned to the genes represented a wide range of functions, such as GTPase activity, protein binding, and transferase activity (acyl and glycosyl groups). A few of these genes harbored multiple DMCs, including heixuedian (heix), an ubiA prenyltransferase domain containing protein 1 gene. Of the 42 CpG sites in this gene that were covered by our library, six were significantly more methylated in the short-day sample (Fig. 2B). In contrast, the gene misshapen (msn), encoding a protein kinase, was represented by seven CpGs sites in our reads, five of which showed increased methylation in the long-day sample (at the FDR-corrected \( q < 0.01 \)) (Fig. 2C). We have validated the differential methylation of specific CpG sites in the genes WDR36, (CG9799, WD repeat-

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**Figure 2.** Differential DNA methylation associated with photoperiod in Nasonia. (A) The methylation is shown across the five chromosomes and the unplaced (U) scaffolds. From the outer to inner circle, the data tracks are short-day methylated CpGs (red), long-day methylated CpGs (blue), methylation difference (red/blue), and significantly differentially methylated sites \( (q < 0.05, \text{dark gray}) \). The lack of differential methylation in the U scaffolds is probably because they encompass substantially fewer genes. (B) Differential methylation in the heix (LOC100118618) gene. Of 42 CpG sites that were covered, six (*) were significantly more methylated in short day \( (q < 0.05) \). (C) Differential methylation in msn indicates increased methylation in long day: There are seven covered CpG sites on this gene, of which five (*) are methylated at \( q < 0.01 \).
containing protein 36, putative), LOC100117390 (CG11148, PEQ, amino-acid rich with GYF domain-containing protein), and LOC100117821 (zinc finger protein 615) using a qPCR approach (Thomassin et al. 2004). In all three cases, we found a significant increase in methylation in long-day compared to short-day females, consistent with the RRBS analysis (three biological replicates, 2000 permutations, \( P < 0.0001 \)) (Supplemental Fig. S1).

During bisulfite conversion, 5-hmC has been shown to behave like its precursor, 5-mC (Huang et al. 2010). We have therefore sought to test for the presence of 5hmC in our data. Recent studies in mammals have shown that 5hmC is made from a 5mC precursor, a reaction which is mediated by ten-eleven translocation (TET) proteins (Tahiliani et al. 2009; Ito et al. 2010). We have analyzed the Nasonia genome and identified a single putative Tet ortholog. Although the overall level of identity to human TET2 is rather low (0.25), the 925-bp Tet motif (Pfam ID: TET_JBP) in the Nasonia gene results in a highly significant blast score \( (E = 4.9 \times 10^{-66}) \), suggesting it is a bona fide Tet ortholog.

We carried out hydroxymethylated DNA immunoprecipitation (hMeDIP), using Nasonia gDNA that was spiked with mouse DNA (Thomson et al. 2013), and tested for the presence of three candidate genes in the enriched 5-hmC fractions by qPCR (Supplemental Fig. S2). The percent recovery (enriched/total input) of the genes msn and heix was <1%, similar to the mouse negative control, whereas the gene Cact-2 was present at moderate level (1.13–2.33%) compared to the positive control. Thus, functional 5-hmC might exist in Nasonia but is unlikely to play a significant role in the photoperiodic response. The complete survey of 5-hmC in Nasonia will be published elsewhere.

Functional assays demonstrating a causative role of DNA methylation

To test the functional role of DNA methylation in mediating short-day-induced diapause, we knocked down Dnmt1a-c and Dnmt3 by injecting double-stranded RNA (dsRNAi) of these genes into female pupae of the wild-type N. vitripennis strain AsymC. Control females were injected with dsRNAi targeting the green fluorescent protein (GFP). We verified the knockdowns by qPCR of mRNA extracted from the injected females (Supplemental Fig. S3). Although the complete knockdown of Dnmt1a is lethal (Zwier et al. 2012), our protocol induced a partial knockdown (~60%) and resulted in the successful eclosion of adult females that were then tested for their photoperiodic responses. The diapause response of progeny of injected females is shown in Figure 3A. RNAi knockdown of Dnmt1a leads to females producing diapause offspring regardless of day length. Although control females exhibited the strong photoperiodic response typical of Nasonia, with increased diapause level in short days \( (t = 3.57, df = 28.6, P = 0.0012) \), the response of Dnmt1a dsRNAi-injected females was similar in both long and short days \( (t = 1.45, df = 36, P = 0.15) \), with a substantial increase in diapause induction in progeny of females maintained under long-day conditions. Silencing Dnmt3 (Fig. 3B) resulted in a nonsignificant difference in diapause response; although this result is close to significance, suggesting that the effect of Dnmt3 is much weaker or harder to resolve \( (t = 1.96, df = 36, P = 0.055) \). This effect of the dsRNAi was observed within 10 d of injection. For progeny laid 10–15 d after injection, diapause frequencies reverted to normal levels again (i.e., low diapause incidence in long days; data not shown). We have also attempted the knockdown of Dnmt1b and Dnmt1c, but the progeny of the injected wasps showed normal photoperiodic response (Supplemental Fig. S4).

We verified the effect of the RNAi knockdown on CpG methylation by using qPCR (MethyQuant). 9–10 d post-injection (5 d after eclosion) we analyzed the status of highly methylated CpG sites in three different genes (Supplemental Fig. S4: CG11148 (LOC100117390), WDR36 (CG9799)), and LOC100117821. Knockdown of Dnmt1 resulted in a significant reduction of methylation \( (P < 0.05) \), calculated by permutation, in both the WDR36 and LOC100117821 loci, whereas for Dnmt3 knockdown, the level of methylation was reduced in the treated group (DNMT3s-dsRNAi) compared to the control group (GFP-dsRNAi) in each of the tested genes \( (P < 0.05) \).

We also tested the impact of DNA methylation pharmacologically. We used the DNA methylation inhibitor 5-aza-2′-deoxycytidine (5-aza-dC), a potent inhibitor of DNA methylation (Christman 2002), and tested the female response to varying photoperiods. Although control females fed with sugar solution showed the normal photoperiodic response (again with a substantial difference between long- and short-day females: \( W = 338, P < 0.0001 \)) (Fig. 4A), wasps fed with 10 \( \mu \)M 5-aza-dC (a nonlethal dose), responded similarly to long and short days \( (W = 707, P = 0.47) \). Intriguingly, in this case, we observed a reduction of diapause in the progeny of wasps maintained in short day (median drop from 35% to 25%), accompanied by an increase in long-day progeny (8% to 20%). We verified the impact of the drug on DNA methylation of CpG sites in four different genes (Fig. 4B):
Discussion

Here, we have shown that DNA methylation is associated with encoding the seasonal photoperiod in *Nasonia vitripennis*. Patterns of methylation across the genome change with photoperiod, and manipulation of the methylome via either the disruption of a key maintenance methyltransferase (*Dnmt1a*) or via a potent inhibitor of methylation (5-aza-DC) changes how female *Nasonia* respond to photoperiod. In addition, we have obtained a single-nucleotide resolution snapshot of the methylome of *Nasonia* that adds to the number of recently published insect methylomes, including the honeybee *Apis mellifera* (Lyko et al. 2010), the ants *Camponotus floridanus* and *Harpegnathos saltator* (Bonasio et al. 2012), the silk moth *Bombyx mori* (Xiang et al. 2010), and the locust *Schistocerca gregaria* (Falckenhayn et al. 2013). Moreover, the pattern of methylation in the wasp mirrors that of other insects, i.e., relatively low-density methylation, mostly mapped to gene bodies, particularly exons. Indeed, this seems to be an ancient property of eukaryotic genomes (Zemach et al. 2010), which is presumably involved in the regulation of alternative splicing (Lyko et al. 2010; Foret et al. 2012).

Profiling of DNA methylation by RRBS provides only a snapshot of the epigenome (Gu et al. 2011). It is therefore likely that the actual extent of differential methylation across the genome associated with the change in photoperiod is more substantial than that reported here (perhaps up to 20-fold higher, given our CpG coverage). It is also likely that our use of the whole body for methylation profiling hindered the detection of photoperiodic DMCs that may well be restricted to specific cell clusters in the brain or other tissues.

Of course, many of the DMCs we have identified may not necessarily be involved in the photoperiodic response (i.e., in inducing offspring diapause) because there are other physiological processes that might change with day length (for instance, involving changes in host-finding or resource allocation). Here, our goal has been to demonstrate that differential methylation with respect to photoperiod is present (although the link between a specific critical methylated locus and the photoperiodic circuit may be quite distant). Yet, some of the DMCs in our list may serve as candidates to follow. For instance, the gene *msn* has been shown to be involved in development in *Drosophila*, where it interacts with the JNK pathway; intriguingly, it is important for the neural development of photoreceptors (Treißmann et al. 1997) and therefore may play a role in photoperiodic light reception.

Crucially, the disruption of the normal photoperiodic response by the knockdown of *Dnmt1a* demonstrates the causative role of DNA methylation in the photoperiodic response (Fig. 3). The observed abnormal increase in diapause incidence under long-day conditions following the RNAi manipulation suggests that under normal conditions, methylation of key photoreceptor genes prevents diapause when days are long. However, the more pervasive pharmacological reduction of methylation by 5-aza-DC leads to diapause development under both long- and short-day conditions. In both cases, the photoperiodic response has been abolished, but not in quite the same way. This may be due to the fact that 5-aza-DC has a broad impact on DNA methylation (in contrast to *Dnmt1a* RNAi). As such, the differences in response in the two experiments may reflect multiple pathways and targets of DNA methylation with various effects on diapause, which are likely to be locus- and tissue-specific. Consistent with the fact that the change in day length is associated with both hyper- and hypo-methylation (Fig. 2), the 5-aza-DC blocking is expected to impact both long- and short-day response.

In mammals, the function of Dnmt1 is the maintenance of DNA methylation. If this function is conserved in *Nasonia*, then the effect of the knockdown would take place only in cells
undergoing DNA replication. The 5-aza-dC treatment would impact both maintenance and de novo methylation, however, which might explain the differences in the observed phenotypes (Fig. 3). The de novo methylation might be accomplished by Dnmt3, whose activity would be carried out both in dividing and nondividing cells. Dnmt3 has been shown to be important for caste determination in honeybees (Kucharski et al. 2008) via the regulation of DNA methylation levels and alternative splicing (Li-Byarlay et al. 2013), and its role in the photoperiod response merits further investigation.

In contrast to Dnmt1a, the RNAi knockdown of Dnmt1b-c did not affect the photoperiodic response. This may be due to different expression patterns of the genes or different catalytic functions for Dnmt1b-c, which would be rather interesting given their sequence similarity. Similar functional differences were suggested by Zwier et al. (2012), in which inhibition of Dnmt1a (but not of Dnmt1c) during early embryogenesis led to developmental arrest and lethality. It is also possible that differences between the Dnmts are caused by their noncatalytic activity. Such noncatalytic regulation of gene expression has been demonstrated in the vertebrate DNMT1 (Clements et al. 2012). The specific functional role of each of the Nasonia Dnmts again awaits further investigation.

Also noteworthy is that our methylome data show that photoperiodic transitions induce both up- and down-regulation of methylation, suggesting a complex regulation of methylation involving various methylating (and possibly demethylating) agents. Likewise, in honeybee caste determination, the methylation level in the brains of workers and queens shows both up- and down-regulation in a gene-specific manner (Lyko et al. 2010), which cannot be explained by merely the up-regulation of Dnmts in one of the castes.

In summary, our experiments clearly demonstrate a significant role of DNA methylation in photoperiodic timing. How these methylation changes underlie the maternal transfer of the photoperiodic response to the progeny remains to be determined. The fact that DNA methylation is involved in the photoperiodic clock echoes the role of epigenetic regulation in the circadian clock (Masri and Sassone-Corsi 2010; Ripperger and Merrow 2011). Indeed, most of those studies (carried out in mammals) underscored the role of histone modifications, which was also recently shown to be important for insect photoperiodism (Lu et al. 2013).

Whether photoperiodic timing is itself relying on the circadian clock is still a subject of debate (O’Brien et al. 2011), but some evidence suggests that this might be the case in Nasonia (Saunders 1974). If such a link between the two clocks does exist, it may involve light-induced epigenetic changes of Clock or Clock-controlled genes. Revealing the genes and cells that underlying the Nasonia circadian and photoperiodic clocks will soon allow a better understanding of the epigenetic mechanisms associated with these systems.

Methods

Wasp strains
We used the wild-type strain, AsymC, whose genome has been sequenced (Warren et al. 2010). To test for diapause, newly eclosed females were kept individually in glass vials containing two blowfly pupae (Phaenicia cuprina; hosts) at 18°C under various photoperiods (using LED illumination) for 10 d. Hosts were replaced twice, after 5 and 9 d, and kept in darkness at the same temperature for an additional 20 d. At this stage hosts, were dissected and diapause status of the progeny was determined (diapause larvae are easily identified through their morphology).

RRBS library preparation
Newly eclosed females were maintained in either a long (18 h) or short (6 h) photoperiod. After 10 d, 2 h after lights on, genomic DNA was isolated from 10 females using the DNeasy kit (Qiagen). The RRBS libraries were prepared as previously described (Gu et al. 2011). Briefly, the gDNA (300 ng) was digested using MspI and the resulting CpG-terminated fragments were purified, and 3′ A-overhangs attached using Klenow fragment (3′-5′ exo-nuclease). The blunted end fragments were then ligated to methylated Illumina adapters (15 μM; Integrated DNA Technologies) that also included a 6 nt unique barcode at the 3′ end, allowing sequencing the samples in a single Illumina lane (see below). To obtain DNA fractions in the 40- to 200-bp range of MspI-digested products, the adapter-ligated fractions were excised from a 3% NuSieve gel. Bisulfite conversion was conducted using the EpiTect kit (Qiagen), following the manufacturer’s instructions. The final libraries were generated by PCR amplification using PfuTurbo Cx Hotstart polymerase (2.5 U/μL; Agilent Technologies). The RRBS libraries were analyzed by an Agilent 2100 Bioanalyzer (Agilent Technologies).

RRBS sequencing and data analysis
The libraries were sequenced using an Illumina HiSeq 2000 analyzer (BGII). The raw sequencing data was obtained through two Illumina sequencing runs on the same libraries, yielding 117,646,358 raw 50-bp reads in total. FastQC version 0.10.1 (http://www.bioinformatics.babraham.ac.uk/projects/fastqc) was used to provide data quality information at each step of the quality control process. The data were taken through several preprocessing steps before being aligned to the Nasonia genome (version Nvit_2.0) using Bismark v.0.7.7 (Krueger and Andrews 2011). These steps included de-multiplexing and barcode trimming, as well as trimming of low-quality bases and Illumina adapter sequences from the 3′ end of the reads, using the Trim galore v0.2.5 package. Low quality reads (cutoff of 20 Phred score) were removed using FASTX-Toolkit (0.0.13; http://hannonlab.cshl.edu/fastx_toolkit/), which was also used to remove potential noise (i.e., reads that did not begin with YGG [where Y = C or T], as all RRBS reads are expected to begin with either CGG or TGG, in the nonconverted and converted cases, respectively). To facilitate downstream analysis, the resulting Bismark SAM output was sorted by chromosome and position using SAMtools version 0.1.18 (Li et al. 2009).

Once aligned to the genome, methylation sites were mapped by identifying C to T conversions produced by the bisulfite conversion. The error rate of methylation calling was estimated for each of the samples. Because the protocol uses unmethylated Cs for the filling in of the MspI restriction site, Cs at this site should always appear as Ts in the sequencing data. Any Cs that do appear in the sequencing data at this position are thus due to incomplete bisulfite conversion or T > C sequencing error. Only reads that contained the Illumina adapter and barcode at their 3′ end with an overlap of 11 bases or more were used to estimate the error because this was judged to be a good indication that the filled-in position could be located within these reads. The error estimate (E) due to incomplete bisulfite conversion and sequencing error is given by E = nC/[(nC + nT)], where nC is the number of Cs found at the filled-in position; and nT is the number of Ts found at the filled-in position. The error estimates for each sample were 0.0068 (LD) and 0.00622 (SD).
To classify a cytosine as methylated (mC), the binomial distribution was used to calculate the probability of getting k successes (methylated Cs) in N trials (read depth) with probability E of a success (Lister et al. 2009). The P-values that had been generated in this way for each site were adjusted using the Benjamini-Hochberg method (Benjamini and Hochberg 1995), setting the FDR at 1%.

Differentially methylated Cs were identified using methylKit (Akalin et al. 2012) that uses the raw Bismark output data as an input. For each CpG, the proportions of methylated reads in the two photoperiods were compared using the Fisher’s exact test, and the P-values were corrected to genomic-wide false discovery rate (FDR)-based Q-values by using the SLIM method (Wang et al. 2011). The methylated sites were mapped to genes using custom Perl scripts.

To test for 5-hydroxymethylation, we used the previously published hmeDIP protocol (Thomson et al. 2013). Briefly, genomic DNA (5 μg) was sonicated for 10 cycles, vacuum dried, and was spiked with mouse liver DNA (300 ng). Immunoprecipitation was carried with rabbit polyclonal antibody against hydroxymethylcytosine (Active motif #39769) and purified (Qiagen). Enrichment of 5-hmC was tested using SYBR green qPCR in a 25-μL reaction volume using a two-step cycling program (15 min at 94°C and 45 min at 60°C) for 45 cycles. The target Nasonia genes were tested along mouse Gapdh and Tex19.1 (negative and positive controls, respectively). Information about the tested genomic regions and primer sequences are in Supplemental Table S2.

Validation of differential methylation by qPCR

We used the MethylQuant method (Thomassin et al. 2004) to validate specific differentially methylated CpGs in candidate genes. Following bisulfite conversion, the DNA was preamplified for 15 PCR cycles using nondiscriminatory (ND) primers (Supplemental Table S2). Three independent DNA samples were tested from long and short day for RNAi validation, and two independent replicates were used in testing the effect of the drug. The product was purified and used in qPCR (25 μL reaction volume) using discriminatory (D) primers harboring the methylation status-specific nucleotide at the 3’ end (Supplemental Table S2). The fluorescence data was analyzed using the qPCR library of the R statistical software (Ritz and Spiess 2008). The PCR efficiencies were averaged across replicates using the sliding window method, and ratios between the two conditions (long/short day) of a gene-of-interest were calculated using normalization against the nondiscriminatory qPCR data. Statistical significance for the ratios is calculated by a permutation approach (n = 20000) of randomly reallocated versus non-reallocated data (Ritz and Spiess 2008).

Functional assays

The protocol for generation of Dnm1Δ, Dnm3Δ, and GFP double-strand RNA and injection of pupae was previously described (Lynch and Desplan 2006). For the oligonucleotide sequences, see Supplemental Table S1. We verified that the dsRNA fragments do not match the transcripts of any other genes (off-targets) using a custom-made off-target prediction tool, which we have made available online (http://WaspAtlas.com) (Davies and Tauber 2015). The tool dices the RNA sequence into all possible 19-mers, which are then matched against the published WaspAtlas (http://WaspAtlas.com), our new Nasonia genomic database (Davies and Tauber 2015).

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