Methyltransferase DnmA is responsible for genome-wide N6-methyladenosine modifications at non-palindromic recognition sites in *Bacillus subtilis*

Taylor M. Nye, Lieke A. van Gijtenbeek, Amanda G. Stevens, Jeremy W. Schroeder, Justin R. Randall, Lindsay A. Matthews and Lyle A. Simmons

Department of Molecular, Cellular, and Developmental Biology University of Michigan, Ann Arbor, MI 48109-1055, USA

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ABSTRACT

The genomes of organisms from all three domains of life harbor endogenous base modifications in the form of DNA methylation. In bacterial genomes, methylation occurs on adenosine and cytidine residues to include N6-methyladenine (m6A), 5-methylcytosine (m5C), and N4-methylcytosine (m4C). Bacterial DNA methylation has been well characterized in the context of restriction-modification (RM) systems, where methylation regulates DNA incision by the cognate restriction endonuclease. Relative to RM systems less is known about how m6A contributes to the epigenetic regulation of cellular functions in Gram-positive bacteria. Here, we characterize site-specific m6A modifications in the non-palindromic sequence GACGmAG within the genomes of *Bacillus subtilis* strains. We demonstrate that the yeeA gene is a methyltransferase responsible for the presence of m6A modifications. We show that methylation from YeeA does not function to limit DNA uptake during natural transformation. Instead, we identify a subset of promoters that contain the methylation consensus sequence and show that loss of methylation within promoter regions causes a decrease in reporter expression. Further, we identify a transcriptional repressor that preferentially binds an unmethylated promoter used in the reporter assays. With these results we suggest that m6A modifications in *B. subtilis* function to promote gene expression.

INTRODUCTION

DNA methylation is pervasive across all three domains of life. In eukaryotes, 5-methylcytosine (m5C) modifications have been shown to function in development and the regulation of gene expression, with aberrant methylation implicated in human health, including cancer, autoimmune diseases, and metabolic disorders [for review, (1,2)]. m5C in promoter regions has been linked to the repression of downstream gene transcription, whereas gene body methylation has been positively correlated with gene expression [for review (3)]. A lesser-studied modification in the genomes of eukaryotes is N6-methyladenine (m6A). Recent studies have identified m6A in the genomes of *Chlamydomonas*, *Caenorhabditis elegans* and *Drosophila melanogaster* (4–6). In contrast to promoter m5C, m6A modifications appear to function in gene activation in the algae *Chlamydomonas* (4) and promoter m6A is also important in early *Drosophila* development (5). Further, m6A was positively correlated with gene expression in a diverse set of fungi (7). Thus, there is a growing recognition that m6A is critical for the regulation of gene expression in a broad range of eukaryotic organisms.

Bacterial genomes are known to harbor N4-methylcytosine (m4C) in addition to m5C and m6A [(8) and references therein]. All three modifications impart consequences to bacterial cells when methylation is lost (9). The most well understood example of DNA methylation in eubacteria is in the context of restriction-modification (RM) systems [for review (10,11)]. RM systems function as a bacterial host defense mechanism to prevent the invasion of foreign DNA, including phages and other mobile genetic elements (10,11). In organisms with RM systems, unmethylated foreign DNA is targeted for site-specific cleavage by a restriction endonuclease while the host chromosome is protected at the recognition sequence by site-specific DNA methylation (12). Methylation is achieved through the activity of DNA methyltransferases (MTases). MTases catalyze the transfer of a methyl group from the donor S-adenosylmethionine (SAM) to adenosine or cytidine residues in DNA (13,14). MTases that lack a cognate endonuclease and do not function in RM systems are referred to as ‘orphan MTases’ (15). In a limited set of...
Gram-negative bacteria, orphan MTases have been shown to function in critical processes including cell cycle control (16), origin sequestration (17,18), DNA mismatch repair (19–21), and the regulation of gene expression [for review (22)]. DNA methylation from orphan and RM-based MTases has also been shown to establish epigenetic inheritance through phase variation primarily in Gram-negative pathogens (23–25). While much work has been done to characterize RM and orphan MTases from Gram-negative bacteria, much less is known about how m6A contributes to the regulation of the cell cycle or gene expression in Gram-positive bacteria (26).

Until recently, tools for unbiased detection and functional characterization of DNA methylation were limited. Available tools for detection, such as methylation-sensitive restriction endonuclease treatment and bisulfitesequencing, are limited to the sequence context and modification type that can be detected (27). The recent development of the Pacific Biosciences (PacBio) Single Molecule, Real-Time (SMRT) sequencing platform allows for detection of modifications without a priori knowledge of their existence (28). SMRT sequencing enables the analysis of real-time DNA polymerase kinetics for inference of DNA base modifications. Base modifications in the template strand result in changes in DNA polymerase kinetics compared to their unmodified counterparts, allowing for reliable, sequence-context specific detection of methylated bases during sequencing reactions (29). While differences in kinetic signatures for m5C modified cytidine residues are modest, SMRT sequencing is adept for m6A and m4C detection (29).

Using the SMRT sequencing platform, a recent study of 230 diverse prokaryotes detected base modifications in 93% of the genomes surveyed (8). Of the genomes with detected modifications, 75% of the modifications were m6A, which is due in part to the robust signal of m6A modifications in SMRT sequencing relative to other modifications (29).

Given the high percentage of prokaryotic genomes with m6A detected and the contribution of m6A to the regulation of eukaryotic gene expression, it seems unlikely that the prevalent m6A modifications in prokaryotes are used exclusively in the context of regulating DNA cleavage by RM systems. As mentioned above, in Escherichia coli and Caulobacter crescentus m6A from orphan MTases occurs in palindrome recognition sequences and has been shown to mediate protein-DNA interactions (9,30), regulating important cellular processes including gene expression (31–34). Deletion of Dam methyltransferase (dam), which is responsible for m6A at GATC sites in E. coli, has severe pleiotropic effects (35,36). In C. crescentus deletion of the CcrM methyltransferase, which catalyzes the formation of m6A at GA(N)TC sites, is lethal when the CcrM-deficient strain is grown in rich media (16,37).

Much less is known about how m6A regulates cellular functions in Gram-positive bacteria. Recent work in Streptococcus pyogenes found that m6A from an active Type RM system regulates virulence gene expression in a clinical isolate, suggesting that m6A could have important roles for regulating gene expression in Gram-positive systems (26). Therefore, the importance of m6A in E. coli and C. crescentus and the pervasive occurrence of m6A in prokaryotes (8) highlights the importance of understanding how m6A regulates cellular functions in the numerous and diverse set of bacterial genomes that contain the modification.

Here, we characterize m6A modifications in the Gram-positive bacterium Bacillus subtilis strains PY79 and NCIB 3610. Using SMRT sequencing, we show that m6A is present at non-palindromic GACGm6A AG sites throughout the B. subtilis chromosome. Further, we characterize the methyltransferase, referred to herein as DnmA, as responsible for detectable m6A modifications in the B. subtilis genome of both strains. We found that DnmA does not function as part of an active, canonical Type I or Type II RM system. Moreover, we show that the promoter regions for a subset of genes contain the consensus sequence and that loss of methylation in these cis regulatory elements results in a decrease in gene expression. Further, we show that the transcriptional repressor ScoC preferentially binds a promoter region that is unmethylated. Together, our results show that m6A can function as an epigenetic signal in B. subtilis.

MATERIALS AND METHODS

General bacteriology

The antibiotic concentrations used in this study are as follows: 5 μg/ml chloramphenicol, 0.5 μg/ml erythromycin, 100 μg/ml spectinomycin. Unless otherwise indicated, strains were grown in either LB (10 g/l tryptone, 5 g/l yeast extract, 10 g/l NaCl) or defined S750 minimal medium supplemented with 1% glucose (1 × S750 salts diluted from 10× S750 salts (104.7 g/l MOPS, 13.2 g/l ammonium sulfate, 6.8 g/l monobasic potassium phosphate; adjusted to pH 7 with potassium hydroxide), 0.1% potassium glutamate, 1% glucose, 40 μg/ml phenylalanine, 40 μg/ml tryptophan, 2 mM MgCl₂, 0.7 mM CaCl₂, 50 μM MnCl₂, 1 μM ZnCl₂, 1 μg/ml thiamine-HCl, 20 μM HCl, and 5 μM FeCl₃) at 30°C with shaking at 200 rpm.

Strain construction

The strains, plasmids and oligos used in this study can be found in Supplementary Tables S1–S3. Individual strain and plasmid construction can also be found in the Supplementary Materials and Methods. Deletions were created by ordering Bacillus subtilis 168 strains from the Bacillus Genetic Stock Center (http://www.bsgc.org/) where the respective genes were replaced with a loxP flanked erythromycin (erm) resistance cassette (BKE strains). Genomic DNA from the BKE strains was purified and used to transform B. subtilis strain PY79, and the erm resistance cassette was subsequently removed with Cre recombinase (38). Overexpression strains and all promoter GFP fusions were integrated in the PY79 amyE locus via double crossover (39). Three colonies containing the crossover were selected and colony purified on LB plates containing 100 μg/ml spectinomycin. Successful integration of the constructs was verified by PCR, Sanger sequencing, and screening for the ability to utilize starch.

Chromosomal DNA purification

Genomic DNA for Pacific Biosciences SMRT sequencing was purified as follows. Strains were struck out on LB
and grown overnight at 30°C. 500 ml LB cultures were inoculated at OD_{600} 0.05 and grown at 37°C. During mid-exponential phase (OD_{600} 0.6−0.8) an equal volume of methanol was added to each culture and centrifuged at 4000 rpm for 30 min. The supernatant was discarded and cells were resuspended in 12.5 ml of 10% sucrose Tris/HCl pH 8 buffer and transferred to Oakridge tubes. Resuspensions were then treated with 310 μl lysozyme (40 mg/ml in 10% sucrose Tris/HCl pH 8 buffer) for 30 min at 37°C and mixed every 5 min. 1.25 ml of 0.5 M EDTA was added to each tube and incubated on ice for five minutes followed by addition of 10 ml of freshly prepared lysis solution (0.1% Triton X-100, 62.5 mM EDTA, 50 mM Tris/HCl pH 8). Solutions were centrifuged at 15 000 rpm for 30 min and decanted into chilled graduated cylinders. To each lysate 0.95 g/ml of cesium chloride (CsCl) was added and dissolved followed by a 1/10 volume addition of 10 mg/ml ethidium bromide. Solutions were balanced and centrifuged at 44 000 (131 600 × g) rpm for 24 h. Chromosomal DNA was extracted and subjected to a second round of CsCl purification as described above. Solutions were centrifuged at 44 000 rpm (131 600 × g) for 48 h. Ethidium bromide was removed by extraction 4× with water-saturated butanol. The aqueous phase was transferred to an Oakridge tube and 1 volume of water and 2 volumes ethanol were then added. The solution was centrifuged at 15 000 rpm for 20 min and the supernatant was aspirated. The pellet was washed with 70% ethanol and resuspended in 1 ml TE buffer. In all other experiments, frozen strains were struck out and grown at 30°C. The plates were washed in S750 minimal media and 25 ml cultures were inoculated at an OD_{600} 0.05 and grown at 37°C with shaking to mid-exponential growth phase (OD_{600} 0.6−0.8). Genomic DNA was purified via phenol chloroform extraction method.

**PacBio SMRT sequencing and methylation analysis**

Chromosomal DNA was prepared for sequencing as described above. Library preparation and subsequent sequencing was performed as previously described (40,41). Modification and motif analyses were performed using RS_Modification_and_Motif_Analyses.1 version 2.3.0 with the appropriate B. subtilis reference genomes. The initial parameters used for modification analysis were performed using 0.75 minimum high quality reads, 50 bps minimum length, and a minimum ModQV call of 30. We also increased minimum high quality reads to >0.85 and minimum length to >1000 bp in subsequent analysis. Modification graphs were generated using functions from BaseModFunctions.v2.1.R available at: https://github.com/PacificBiosciences/Bioinformatics-Training/tree/master/basemods.

**Motif distribution analysis**

Motif distribution analysis was performed using the DistAMo web based server (42) available at http://computational.bio.uni-giessen.de/distamo searching the GACGAG motif for the B. subtilis PY79 genome via accession number NC_022898.1.

**Protein purification (DnmA, DnmA (Y465A) and YahB)**

Recombinant proteins were purified from E. coli BL21DE3 cells containing a pE-SUMO vector with the B. subtilis gene inserted (dnmA, dnmA (Y465A) or yahB). Cultures were grown in 4 l of terrific broth (2.4% yeast extract, 1.2% tryptone, 0.4% glycerol, 250 mM (NH4)2SO4, 500 mM KH2PO4, 1 × metals (1000 × metals: 2.5 mM FeCl3, 1 mM CaCl2, 0.5 mM ZnCl2, 0.1 mM CoCl2, 0.1 mM CuCl2, 0.1 mM NiCl2, 0.1 mM Na3MoO4, 0.1 mM Na2SeO3, 1 mM H2BO3) and 25 μg/ml kanamycin) at 37°C with orbital rotation for 2 h until reaching an OD_{600} of ~0.7. Overexpression was induced by adding IPTG to 1 mM and the cultures were grown for three additional hours at 37°C. Cells were then pelleted by centrifugation and frozen in liquid nitrogen to be stored at −80°C. Once thawed, the pellet was resuspended in lysis buffer (50 mM Tris–HCl pH 8, 300 mM NaCl, 10% sucrose, 10 mM imidazole, 1× protease inhibitors (Roche 1187358001)) and cells were sonicated on ice. Cell debris was pelleted by centrifugation. Supernatant was then poured through a 3 ml Ni2+-NTA agarose gravity-flow column. The column was washed with wash buffer (20 mM Tris–HCl pH 8, 10% glycerol, 20 mM imidazole, 2 M NaCl) and eluted with elution buffer (50 mM Tris–HCl pH 8, 150 mM NaCl, 400 mM imidazole). SDS-PAGE was performed to confirm the presence of desired protein. The sample was then dialyzed into anion exchange start buffer (50 mM Tris–HCl pH 8, 25 mM NaCl, 5% glycerol, 1 mM β-mercaptoethanol) and the sample was applied to a Q column (GE: 1715301) using an elution gradient of 50−750 mM NaCl. SDS-PAGE was performed and fractions containing desired protein were pooled and incubated with ULP1 protease at 25°C for 30 min. The digestion product was applied to another 3 ml Ni2+-NTA gravity-flow column, washed, and eluted using the same buffers as above. SDS-PAGE was again performed to confirm the SUMO tag was removed and the protein was concentrated and buffer exchanged into protein storage buffer (50 mM Tris–HCl pH 8, 150 mM NaCl, 50% glycerol), aliquoted, flash frozen in liquid nitrogen and stored at −80°C.

**ScoC purification**

Primers oTMN62 and 63 were used to amplify scoC from the B. subtilis chromosome and were subsequently combined with the pE-SUMO expression vector via Gibson assembly. Recombinant proteins were purified from E. coli BL21DE3 cells grown in 2 l of LB with 25 μg/ml kanamycin at 37°C with orbital rotation until an OD_{600} of 0.7 was reached. Overexpression was induced by adding 0.5 mM IPTG followed by culture growth for an additional three hours at 37°C with orbital rotation and cultures were subsequently pelleted via centrifugation and stored at −80°C. The pellet was re-suspended in lysis buffer and sonicated on ice as described for DnmA and YahB. Subsequent to centrifugation, the supernatant was applied to a 4 ml Ni2+-NTA agarose gravity-flow column. The column was washed with wash buffer (50 mM Tris–HCl pH 8, 25 mM imidazole, 2 M NaCl, 5% glycerol) and eluted with elution buffer (50 mM Tris–HCl pH 8, 150 mM NaCl, 50% glycerol), aliquoted, flash frozen in liquid nitrogen and stored at −80°C.
for 2 h at room temperature. The sample was then dialyzed into storage buffer (50 mM Tris–HCl pH 8, 150 mM NaCl, 5% glycerol) overnight at 4 °C. The dialyzed sample was then applied to another 4 ml Ni²⁺-NTA gravity-flow column to separate the recombinant protein from the SUMO tag. SDS-PAGE was performed to confirm the SUMO tag was removed. Glycerol was added to 25% and the protein was aliquoted and flash frozen for storage at -80 °C.

Methylation assays

All methylation reactions were performed in a buffer containing 50 mM Tris–HCl pH 8, 50 mM NaCl and 200 μM MgSO₄. The following substrates were annealed in the same buffer at 2.5 μM concentration by heating primers to 100 °C for 30 s and then cooling to room temperature on the bench top: dsDNA target (oTMN38, oTMN39); dsDNA non-target (oTMN40, oTMN41); and dsRNA (oJR270, oJR271). The H3-SAM (Perkin Elmer: NET155H001MC) was used at a concentration of 1 μM, and dNAM (V465A) was added to a concentration of 1 μM and all substrates were used at 0.25 μM in solution. The proteins were added in excess to determine if there was any off target methylation activity at higher protein concentrations. The total reaction solution came to 10 μl. All reactions were incubated at 37 °C for 150 min unless otherwise specified. Reactions were stopped using 450 μl of 10% TCA and placed on ice. The samples were filtrated using Glass microfiber filters (GE: 1822-025), washed with cold 70% ethanol, dried, and placed in a scintillation counter to measure mmol incorporation.

Growth curves

Strains were plated on LB and grown overnight at 30 °C. Plates were washed in LB and diluted to an OD₆₀₀ of 0.05 in 10 ml of LB in side-arm flasks. Cultures were grown in shaking water baths at 37 °C and optical density was measured using a Klett meter every half hour through late stationary phase. Growth curve experiments were done in triplicate and data was subsequently fit to a Gompertz growth function: 
\[
y(t) = A e^{\mu t} - e^{[e^{\mu t} - (\lambda - t) + 1]}
\]

where the parameters A, \(\mu\), and \(\lambda\) represent the time \(t\) when the growth rate equals zero (asymptote), the maximum growth rate, and the lag time, respectively, to obtain growth rate estimates (\(\mu\)) for each strain.

Transformation efficiency assays

Strains were plated on LB and grown overnight at 30 °C. Plates were washed with phosphate buffered saline (PBS) pH 7.4 and the cells were pelleted, the supernatant was aspirated, and a second PBS wash was completed before the cells were resuspended in PBS. The cells were used to inoculate a culture at an OD₆₀₀ of 0.05 into 1 ml of 1× MC media (10× MC media: 615 mM KH₂PO₄, 380 mM KH₂PO₄, 1.11 M dextrose anhydrous, 30 mM sodium citrate dihydrate, 840 μM ferrer ammonium citrate, 0.5 g casein hydrolysate, and 125 mM sodium aspartate monohydrate, to 50 ml with ddH₂O and filter sterilize) with 3 μl of 1M MgSO₄ and grown at 37 °C with aeration for 4 h. After 4 h, 3 μl of 1M MgSO₄ and 300 ng of pHPl3 purified from E. coli MC1061 cells was added to 300 μl of cells and grown for an additional 1.5 h at 37 °C. 10× serial dilutions were performed into PBS and appropriate dilutions were plated onto LB plates for colony forming unit (CFU) counts and chloramphenicol plates for transformation forming unit (TFU) counts. Transformation efficiencies were calculated as TFU/CFU and the average transformation efficiency for replicates performed over three separate days was plotted along with the corresponding standard errors.

Flow cytometry

Cells were grown overnight at 30 °C on LB plates containing 100 μg/ml spectinomycin. Exponentially growing colonies were washed from the plates using S7₅₀ medium, and washed two more times to remove residual LB agar before diluting the cells in pre-warmed S7₅₀ medium to an OD₆₀₀ of 0.05. Cells were grown to an OD₆₀₀ of 0.4 at 30 °C after which fluorescence of 200 000 cells was measured using an Attune™ Nxt Acoustic Focusing Cytometer (ThermoFisher Scientific) using the following settings: Flow rate, 25 μl/min; FSC voltage, 200; SSC voltage, 250; BL1 voltage, 250.

Streptavidin pull-down

5′ biotinylated primers were used to amplify the 233 bp region of the scpA promoter via PCR using genomic DNA from strains LVG087 and LVG102 as a template, which correspond to the GACGAG and GACGTG promoter, respectively. To obtain total cell lysate, 4 l of strain TMN85 (ΔdnmA) was grown in S7₅₀ medium at 37 °C with shaking until the culture reached an OD₆₀₀ of 1.0. After the cells were harvested the pellets were washed with 1× PBS (pH 7.5) and then subsequently washed with Pull-Down Binding Buffer (PDBB; 50 mM Tris–HCl pH 7.5, 0.5 mM EDTA, 100 mM NaCl, 0.01% (v/v) Triton X-100, 25% (v/v) glycerol and 1 mM DTT) and resuspended in ice-cold 20 ml PDBB supplemented with one tablet of eComplete™ EDTA-free Protease Inhibitor Cocktail (Roche, Mannheim, Germany). The cell suspensions were sonicated on ice (10 s on, 40 s off, 70 Hz) until the solutions cleared. Cell debris was removed from the lysate by two subsequent washing steps and the protein content of the supernatant was estimated using a Bradford assay (~20 mg/ml protein). For each pull-down experiment, 100 μl of Dynabeads™ M-270 Streptavidin magnetic bead slurry (ThermoFisher Scientific) was washed three times with 500 μl Pull-Down Wash Buffer (PDWB; 10 mM Tris–HCl pH 7.5, 1 mM EDTA, and 1 M NaCl). The beads were re-suspended in 250 μl PDWB, mixed with 200 pmol biotinylated probe DNA dissolved in 250 μl nuclease-free water, and incubated for 30 min at 25 °C with gentle rotation. The DNA-coated beads were washed three times with PDBB before 100 mg protein and 100 μg salmon sperm DNA (Millipore Sigma) were mixed and added to the DNA-bound beads. After 2 h of incubation at room temperature with gentle rotation, the beads were separated and washed once with PDBB, once with PDBB plus 100 μg salmon sperm DNA, and again with PDBB.
Bound proteins were eluted using Pull-Down Elution Buffer (PDEB; 50 mM Tris–HCl pH 7.5, 0.5 mM EDTA, 1 M NaCl, 0.01% (v/v) Triton X-100, 25% (v/v) glycerol and 1 mM DTT). The eluted proteins were desalted and concentrated using TCA precipitation and separated on a 4–20% Mini-PROTEAN TGX precast protein gel (Bio-Rad, Hercules, USA). Bands in the 20 and 40 kDa size range were excised from the gel followed by protein identification using mass spectrometry through the University of Michigan Proteomics Resource Facility, project PRF-2019-L-SIMM-29.

ScoC EMSA

5′ IR dye end-labeled substrates oTN67/oTN68 and oTN70/oTN71, corresponding to the GACGAG and GACGTA oligos, respectively, were annealed at a concentration of 50 nM by heating at 95°C for 1 minute and then snap-cooled on ice. Care was taken to avoid subjecting the IR dye labeled oligos to light. Annealed oligos were mixed at a final concentration of 5 nM with indicated concentrations of purified ScoC in 1× EMSA reaction buffer (5× EMSA reaction buffer: 250 mM Tris×HCl pH 8, 5 mM EDTA, 150 mM KCl, 10 mM MgCl2, 5 mM DTT, 1% Tween 20, 125 μg/ml sheared salmon sperm DNA) to a final volume of 10 μl. Reactions were incubated at 37°C for 15 min and subsequently loaded onto and resolved via 6% Native-PAGE, which was performed covered and on ice for 60 min at 100 V. The samples were visualized with the LI-COR Odyssey imager. The intensity of the shifted band was normalized to the no protein control for each sample to calculate the percent band shifted. Three replicates were completed and quantified across separate days and the average and standard errors for percent band shifted was reported in Figure 6.

RESULTS

Characterization of B. subtilis PY79 and NCIB 3610 methylomes

It was previously published that B. subtilis does not have m6A at the E. coli Dam MTase recognition site, GATC, and that ectopic expression of Dam in B. subtilis induced the DNA damage response (44,45). However, until recently it remained unknown if B. subtilis contains m6A in another sequence context because the detection of m6A without a priori knowledge of the sequence context would require a new experimental approach. PacBio SMRT sequencing was used to determine if DNA modifications were present in the genome of several B. subtilis strains with the results deposited on the publicly available web resource REBASE maintained by New England Biolabs. This resource reports m6A occurring in various sequence motif contexts in 19 of 23 B. subtilis strains where SMRT sequencing was used. Among the B. subtilis strains analyzed, methylation at GACGAG sites was reported in four of the 23 strains (http://rebase.neb.com). Previously, our group performed PacBio sequencing on the widely used B. subtilis laboratory strain PY79 for whole-genome assembly (41). As part of our effort to study DNA methyltransferases, we used PacBio sequencing to characterize the PY79 methylome. We purified genomic DNA from the wild type (WT) B. subtilis strain PY79 and analyzed our results using the SMRT sequencing platform, allowing for genome-wide base modification detection in sequence-specific contexts (29).

SMRT sequencing of the B. subtilis PY79 chromosome showed that the second adenosine residue within the sequence context 5′-GACGmAG showed high modification quality values (modQVs), which indicates a statistically significant difference in DNA polymerase kinetics from the expected background at particular loci (Supplementary Figure S1A, Table 1). The interpulse duration (IPD) ratios, which are a comparison of DNA polymerase kinetics at a base within a particular sequence context compared to an unmethylated in silico control, were far higher for the second adenosine residue in the GACGmAG motif compared to any other modified motifs in the B. subtilis chromosome (Table 1, Supplementary Figure S1A). Thus, we identify m6A in the sequence context 5′-GACGmAG in the chromosome of B. subtilis PY79, herein referred to as the m6A motif.

We found that 99.7% of m6A motifs (1215/1219) were called as methylated in the PacBio SMRT sequencing analysis at the 3′-adenosine during exponential growth in defined minimal medium. While our sequencing analysis identified other motifs in the B. subtilis PY79 chromosome, the average modQVs, IPD ratios, and the percentage of motifs called as modified were far lower compared to m6A identified within the GACGmAG sequence (Supplementary Table S4 and Figure S2). It is likely that most of the other motifs called represent DNA secondary structures that affect DNA polymerase kinetics or sequencing noise instead of genuine nucleic acid modifications (Supplementary Table S4). For completeness we chose to report all motifs called during analysis of the SMRT sequencing data (Supplementary Table S4).

Of the 1219 m6A motifs that occur in the B. subtilis PY79 genome, 1118 (91.7%) occur in protein coding regions. Intergenic regions, which compose 11.2% of the genome, contain 7% (85 motifs in 76 regions) of the m6A motifs. With the exception of only a few sites, the majority of m6A sites had >75% of sequencing reads called as methylated independent of genome position or occurrence on the plus or minus strand of the chromosome (Supplementary Figure S2 and Supplementary Table S4).

B. subtilis PY79 is a commonly used laboratory strain, however selection in the lab has caused PY79 to lose many

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Table 1. Relevant modified motifs detected in B. subtilis by PacBio SMRT sequencing

| Motif        | Type | %Detected | Mean QV | Mean Cov | Mean IPD ratio |
|--------------|------|-----------|---------|----------|---------------|
| GACGAG       | m6A  | 99.7      | 388     | 286      | 6.72          |
| CTCGAR      | m5C  | 70.8      | 74      | 270      | 1.89          |
| WT PY79     | m6A  | 94.7      | 362     | 313      | 4.84          |

**a**All motif calls by SMRT sequencing are reported in Supplementary Table S4.

**b**Modification type confirmed via methylation sensitive restriction endonuclease digest as described in the supporting document.
of the robust phenotypes associated with ancestral strains of *B. subtilis* (46). To determine whether m6A is present in the ancestral strain, we purified genomic DNA from *B. subtilis* strain NCIB 3610 (40) for SMRT sequencing and found m6A within the same GACGAG sequence context (Supplementary Figure S1B and Table 1). In NCIB 3610, 94.7% (1208/1275) of m6A sites were called as methylated in the PacBio SMRT sequencing analysis. The chromosome of the ancestral strain is considerably larger than PY79 and harbors an 84-kb plasmid, both of which account for the increased number of m6A motifs (40). The decrease in the percentage of motifs called as modified between PY79 and NCIB 3610 (99.7% → 94.7%) could be the result of biological variation, such as an increase in protein binding or other factors that may obscure methylation of recognition sites. The decrease in motifs called could also be due to technical variation in sequencing reactions. We note that we also detected many additional motifs in the ancestral strain that did not appear in the lab strain PY79, with each motif called listed in supplementary Table S4. Further, m6A at GACGAG sequences has also been reported for three *B. subtilis* strains other than PY79 and NCIB 3610 on REBASE.

In addition to m6A modifications, SMRT sequencing of the PY79 genome identified cytidine modifications in the sequence 5′CTCGARB (where R represents a purine and B either a cytidine or a guanosine). These results are described in the supplementary results section, where we show using methylation-sensitive restriction digest that m5C formation occurs in the *B. subtilis* PY79 genome through the BsuMI RM system (Supplementary Figure S3) previously described for *B. subtilis* Marburg (47).

**Distribution of m6A sites across the *B. subtilis* genome shows enrichment on the lagging strand of the left chromosomal arm**

To begin to understand the function of m6A in *B. subtilis*, we used the motif enrichment program DistAMo (42) to determine the location of m6A sites on the *B. subtilis* chromosome. This was done to determine if m6A sites are uniform or showed areas of enrichment and de-enrichment throughout the chromosome (Figure 1). We present the analysis using sliding windows of 50 kb to 500 kb over the length of the chromosome by the rings from outside (large) to inside (small) scaling in 50 kb increments. Over (red) and under (blue) enrichment are colored by z-scores in the scale as shown. From the analysis we determine that the locations of m6A sites are certainly not uniform across the chromosome and instead show patterns of enrichment in particular regions. We find that several areas are largely devoid of m6A sites, including the terminus region and the origin of replication (Figure 1). Analysis of enrichment shows that locations in the *B. subtilis* chromosome with high z-scores includes the right and left chromosomal arms with the largest enrichment on the lagging strand of the left chromosomal arm (Figure 1C). With these results we suggest that m6A is unlikely to function in origin sequestration or DNA mismatch repair as described for Dam methylation in *E. coli* (17,18) due to our finding that the origin does not contain m6A sites and because m6A is non-palindromic and not uniform across the *B. subtilis* chromosome. To be certain, we empirically test if m6A contributes to replication timing, mutagenesis, or recombination in the supplementary results and show no effect (Supplementary Figures S4, S8 and Supplementary Table S5).

**Methyltransferase YeeA is necessary for m6A formation in vivo**

DNA methylation is catalyzed by DNA methyltransferases (MTases) (48). To identify putative MTase(s) responsible for the observed m6A modification, we searched all protein coding sequences for the conserved DNA m6A MTase catalytic motif (D/N/S)PPY (48). This search yielded two uncharacterized MTases, coded for by the genes yabB and *yeeA* (*dnmA*) (41). We created clean deletions of the *ΔyabB* and *ΔyeeA* (*dnmA*) coding regions as well as a *ΔyabBΔyeeA* double deletion. Each of these strains was viable and none of the deletions conferred a growth defect on *B. subtilis* under the conditions used here (Figure 3A, described later in the results).

To identify the MTase responsible for genomic m6A, DNA was harvested from each strain when cultures reached an OD600 of ~0.7 followed by SMRT sequencing. Subsequent methylation analysis revealed that chromosomal DNA from *ΔyeeA* (*dnmA*) cells lost all detectable methylation at the m6A motif previously identified in WT cells in both PY79 and NCIB 3610 strain backgrounds (Table 2, Supplementary Figure S5, and Supplementary Tables S6, S7 and S8 for all PY79 *ΔyeeA* (*dnmA*) GACGAG sites). Expression of *yeeA* (*dnmA*) from an ectopic locus in the *ΔyeeA* (*dnmA*) background restored methylation at the m6A site (Supplementary Figure S5C and Table 2). Computational analysis from sequencing data posted on REBASE also predicted YeeA (DnmA) as the MTase responsible for m6A detected in strains of *B. subtilis* with modifications at the m6A motif described here.

Genomic DNA from *ΔyabB* cells retained the methylation at m6A sites (Supplementary Figure S6, Supplementary Table S9) whereas detectable modifications at the m6A site were lost in the double deletion strain (Supplementary Figure S6B, Supplementary Table S9). Interestingly, while methylation is maintained at the m6A site in the *ΔyabB* strain, we noticed additional motifs not present in the WT or *ΔyeeA* (*dnmA*) strains that were detected upon loss of yabB in the single or double deletion strains (Supplementary Table S9). These additional motifs are likely to result from sequencing noise and/or DNA secondary structure given the low IPD ratios (Supplementary Table S9). With these results we show that *yeeA* (*dnmA*) is necessary for genomic m6A formation in the sequence context GACGAG in vivo and we refer to YeeA herein as DNA methyltransferase A (DnmA), with the formal name of *M. BsuPY79I* and *M. Bsu3610I* for strains PY79 and NCIB 3610, respectively. For simplicity, we will collectively refer to *M. BsuPY79I* and *M. Bsu3610I* as DnmA in the work presented below.

**DnmA is sufficient for methylation of m6A sites in double stranded (ds)DNA in vitro**

DNA MTases typically use SAM to catalyze the transfer of a methyl group to a DNA base (9). DnmA (*M. BsuPY79I*),...
YabB, and a DnmA catalytically inactive variant (Y465A) were purified (Figure 2A). In addition to the predicted ~120-kDa band corresponding to the DnmA monomer, a high molecular weight species was observed in the DnmA purifications. The slower migrating protein was analyzed by mass spectrometry identifying it as multimer of DnmA. We speculate that the DnmA multimer is caused by irreversible disulfide bonding or another crosslink that forms between two purified DnmA monomers during isolation (Supplementary Table S10).

A time course methylation experiment was performed to determine if DnmA is sufficient to catalyze methylation of the m6A motif in DNA (Figure 2B). The purified proteins were incubated with tritiated SAM and an oligonucleotide sequence from the *B. subtilis* addA locus containing the m6A (target) motif. Incorporation of the labeled methyl group over time indicates that DnmA is indeed sufficient for methylation at m6A motifs in dsDNA (Figure 2B). With the results from the time course methylation experiment we suggest that purified DnmA does not have significant activity...
DnmA showed activity on the dsDNA substrate with the incorporation of methylation from tritiated SAM. As expected, there was no appreciable incorporation of the methyl group by DnmA to the non-target sequence (m6A motif). We also tested DnmA for methylation of ssRNA, ssDNA and ssRNA bearing the target sequence. The results show little to no methylation for any of these substrates with the exception of ssDNA, which yielded only weak methylation activity relative to dsDNA (Figure 2D). Together, these results provide strong evidence that DnmA is specific for dsDNA containing the m6A motif.

To determine if the lack of methylation at the non-target sequence was caused by an inability of DnmA to bind DNA, an electrophoretic mobility shift assay (EMSA) was performed on 5' end-labeled target (GACGAG), non-target (GACGTTG), and a degenerate sequence where the entire target sequence had been removed. Incubation of DnmA with the target, non-target, and degenerate sequences each resulted in a shift, indicating that the methylation specificity is not due to a loss of DNA binding at other sequences (Supplementary Figure S7). Additionally, the Y465A catalytically inactive variant still bound the target sequence, suggesting that this variant is only dysfunctional for methyltransferase activity (Supplementary Figure S7). We conclude that DnmA is necessary and sufficient to methylate dsDNA that carries the GACGAG sequence in vivo and in vitro and that Y465 is an important residue for activity.

**DnmA does not function as part of an active RM system**

We next asked if DnmA functions as part of an active RM system. DnmA shares 38% identity and 57% similarity with the MmeI enzyme, which is a bifunctional protein with a methyltransferase domain and a PD-ExK endonuclease motif in the amino terminal domain. MmeI belongs to a subgroup of Type II RM systems that use DNA hemimethylation for host chromosome protection. DnmA was included in a set of MmeI homologs that lack the endonuclease motif in the amino terminal domain but are flanked by conserved genes similar to yeeB and yeeC, which are immediately downstream of dnmA. It was found that under the conditions tested for other MmeI homologs DnmA lacked endonuclease activity, however it is important to note that the downstream yeeB and yeeC gene products are annotated as a putative helicase and an endonuclease, respectively. Deletion of dnmA does not result in a growth defect (Figure 3A), which would suggest that yeeB or yeeC lacks endonuclease activity associated with typical Type II RM systems, where endonuclease activity is achieved independent of the MTase.

It has been suggested that DnmA, along with YeeB and YeeC, comprise a Type I-like RM system, where restriction endonuclease activity requires the MTase subunit and DNA cleavage would not occur efficiently in the absence of DnmA. To test this possibility, we performed a transformation efficiency assay in WT and ΔdnmA cells with the plasmid pH13, which is a 4.7 kb plasmid containing three m6A sites as the donor DNA (Figure 3B). Plasmid purified from E. coli cells was used to transform competency deficient (ΔcomK), hyper-competent (Δrok), WT and ΔdnmA strains followed by selection for transformants conferring resistance to chloramphenicol. We found that compared to ΔcomK and Δrok strains, with transformation efficiencies of less than 1 × 10⁻⁸ and 177 × 10⁻⁵ (SE 13.2 × 10⁻⁵), respectively, the transformation efficiencies of WT [7.33 × 10⁻⁵ (SE 3.30 × 10⁻⁶)] and ΔdnmA [9.44 × 10⁻⁵ (SE 1.25 × 10⁻⁵)] were nearly indistinguishable (Figure 3C). We show that DnmA, YeeB and YeeC do not function to restrict DNA update during natural genetic competence. Based on the transformation results and the conservation of these three genes clustering together, we suggest that DnmA, YeeB and YeeC could be part of an inactive or inefficient Type I-like RM system or perhaps a noncanonical RM system. We also cannot exclude the possibility that restriction activity could be measured under some other circumstance, such as phage predation.

**Proximity of m6A sites to -35 boxes of housekeeping sigma factor SigA regulates promoter activity**

Due to the enrichment of m6A within particular genomic locations (Figure 1), we considered a role for m6A in regulating gene expression. Several prior studies have shown

| Motif | Type | %Detected | Mean QV | Mean Cov. | Mean IPD ratio |
|-------|------|-----------|---------|-----------|----------------|
| ΔdnmA WT PY79 | CTCGAR | m5C<sup>b</sup> | 46.7 | 51 | 120 | 2.00 |
| ΔdnmA WT 3610 | None<sup>c</sup> | | | | |
| ΔdnmA, amyE::Pspac dnmA PY79 | GACGAG | m6A | 99.7 | 213 | 152 | 6.32 |
| ΔdnmA, amyE::Pspac dnmA WT 3610 | CTCGAR | m5C | 52.7 | 59 | 149 | 2.00 |

<sup>a</sup> All motif calls by SMRT sequencing are reported in Supplementary Table S6.
<sup>b</sup> Modification type confirmed via methylation sensitive restriction endonuclease treatment as described in the supporting document.
<sup>c</sup> GACGAG and CTCGAR were not detected in NCIB 3610 ΔdnmA. All other motifs called are reported in Supplementary Table S6. The average coverage is reported for each spurious motif detected.

Table 2. Relevant modified motifs detected in B. subtilis by PacBio SMRT sequencing.
that DNA methylation from RM systems can also regulate gene expression (23,25,26). Accordingly, DNA MTase targets that occur within promoter or operator regions have the potential to influence transcription (50). Thus, we hypothesized that DnmA-dependent methylation might exhibit a similar function in *B. subtilis*.

To identify genes that might be affected by DnmA-dependent methylation, we used the list of transcribed regions 5′ of *B. subtilis* 168 open reading frames (ORFs) reported previously (51) to prioritize the subset of promoters in *B. subtilis* with m6A sites located on the left chromosomal arm where we observed m6A enrichment. The promoters chosen for analysis included those of non-coding and anti-sense RNAs as well as promoters embedded inside transcriptional units and we excluded promoters where the target site occurs downstream of the transcriptional start site (Supplementary Table S11). *B. subtilis* PY79 contains 32 transcribed regions 5′ of ORFs with the m6A motif in the vicinity of known or predicted sigma factor binding sites (Supplementary Table S11). To examine if m6A in promoter regions influences gene expression in *B. subtilis*, we constructed a series of transcriptional fusions where a *gfp* allele was introduced downstream of the respective m6A motif-containing promoter (Figure 4A). All transcriptional fusions were introduced at the ectopic *amyE* locus to separate the promoter from other potential *cis*-acting regulatory elements or chromosome structure contexts that could affect expression (Figure 4B). Promoter activity was monitored in WT and ΔdnmA strains using fluorescence as a reporter in single cells during mid-exponential growth by flow cytometry (please see Materials and Methods).

We found that loss of m6A in a subset of *B. subtilis* promoters, specifically those that contain an m6A motif in or slightly downstream of the –35 region of the SigA-
Figure 3. Loss of DnmA does not affect growth rate or transformation efficiency of foreign methylated DNA. (A) Growth curves for WT, \(\Delta dnmA\), \(\Delta yabB\), and \(\Delta dnmA\Delta yabB\) were performed in triplicate and fit to a Gompertz growth model (43) to calculate growth rate. Growth rate and the corresponding 95% confidence interval for each strain are indicated. (B) Plasmid map of pHP13 with the location of each m6A site shown. The orange carrots indicate the relative position and strand orientation for each site. (C) Transformation efficiency assays were performed using pHP13 plasmid purified from E. coli as donor DNA in WT, \(\Delta dnmA\), \(\Delta rok\), and \(\Delta comK\) recipient strains. The average transformation efficiency and standard error for each strain is indicated.

binding box (PscpA, Phbs, PrnhC, PyumC, PzapA), consistently resulted in decreased activity from the unmethylated promoter relative to the methylated counterpart (Figure 4C and D). The m6A sites in the promoter region for PscpA, Phbs, PrnhC, PyumC, PzapA in PY79 are identical to the promoter regions in B. subtilis strain NCIB 3610.

We did not observe this trend for the promoter fusions that contained m6A sites away from the -35 box. For example, the activation level of the SigB-inducible rsbV-rsbW-sigB-rsbX promoter (PrsbV), with an m6A site directly upstream of the –10 box, was not influenced by the presence of methylation during normal growth or even after stressing the cells with 4% ethanol for 1-hour as described (52) (Figure 4C and D). Similarly, we did not observe differences in gfp expression with the PcomEA, PwprA or PyloA fusions in the \(\Delta dnmA\) background relative to WT.

The m6A motif was present just upstream and overlapping the -35 region of the SigA binding box for PzapA (transcription unit: zapA-yshB-polX-mutSB-yshE) and PyumC, respectively, and both reporters showed a decrease in activity in \(\Delta dnmA\) cells relative to WT (Figure 4C and D). ZapA is involved in FtsZ ring assembly and YumC
Figure 4. Methylation of DnmA motifs in proximity of -35 boxes affects downstream gene expression. (A) Schematic overview of the promoter regions containing DnmA sites that were selected for analysis using transcriptional GFP fusions. Indicated are the locations of the predicted sigma factor -35 and -10 boxes with respect to the DnmA motifs. U numbers correspond to the transcribed regions 5′ of ORFs identified by Nicolas et al. (51). (B) The location of the studied promoters on the PY79 chromosome map with respect to the amyE site used for integration and analysis of the promoter-GFP constructs. (C) Histograms depicting the GFP fluorescence in 200,000 WT (blue) or ΔdnmA (red) cells in three biological replicates that were grown in S750 medium to an OD600 of 0.5 at 30°C and measured using flow cytometry. For U0374/PsigB, an additional experiment was performed in which the cells were treated with 4% EtOH an hour before analysis with flow cytometry. The standard deviations are represented as shaded areas. Promoter regions that appear methylationsensitive are shown in green. (D) Scatter dot plots, with indicated mean and standard deviation, depicting the median fluorescence of each strain taken from the histograms shown in (C) and appended with similar measurements taken on at least one different day. A standard T-test was performed to evaluate differential GFP expression between WT and ΔdnmA for each promoter. P-values: *P < 0.05, ***P < 0.005, ****P < 0.001.
is an essential ferredoxin/flavodoxin reductase (53,54). The m6A site for the remaining three promoter fusions that showed decreased expression upon loss of m6A, PscpA (transcription unit: scpA-scpB-ypuI), Phbs (transcription unit: S861-hbs), and PhnC, was located just downstream of the -35 region of the SigA binding box. Interestingly, the gene products for two of the differentially expressed promoter regions, scpA and hbs, have important roles in chromosome segregation, chromosome structure, and organization (55–60). The changes in Phbs activity were mild, which is likely due to the fact that Phbs contains two SigA-binding boxes, of which the m6A-positive box is the least dominant of the two promoters (61). Another promoter fusion that exhibited a DnmA-dependent increase in expression was PnhiC, which codes for RNase HIII, an enzyme important for cleavage of RNA-DNA hybrids (62,63). One type of RNA-DNA hybrid is an R-loop, which could affect local chromosome structure and transcription (64). Together, decreased expression from PscpA, Phbs and PhnC could have impacts on global chromosome structure, altering the expression of other genes.

To further investigate how m6A methylation affects transcription, the m6A site within the PscpA-GFP promoter was mutated to GACGAG, ensuring loss of methylation at this site in both the WT and ∆dnmA backgrounds. The GACGAG containing promoter adopted the same activity as observed in the ∆dnmA strain, indicating that m6A at the fifth position of the motif stimulates gene expression (Figure 5A and B). Interestingly, an A → T at the fifth position of the m6A site (GACGTG) made PscpA-GFP behave as if it were m6A (GACGmAG) in both WT and ∆dnmA backgrounds (Figure 5A and B middle panel). The reason for how thymidine in the fifth position of the motif stimulates gene expression to the same extent as m6A is unclear. To further test how integrity of the motif modulates PscpA activity, the motif was subsequently changed to GACGAC so that the fifth position was unchanged but the DnmA recognition site was lost. The promoter adopted the same activity as quantified in the ∆dnmA strain, indicating that m6A or T at the fifth position of the motif stimulates gene expression for the scpA promoter (Figure 5A and B). With these data we suggest that m6A is capable of regulating gene expression when located near the -35 binding site for SigA with methylation promoting gene expression from a subset of promoters in B. subtilis.

Transcriptional repressor ScoC binds GACGAG sites

The mechanism for m6A-dependent promotion of gene expression could be explained by an increase in SigA binding at methylated promoter regions or a less direct mechanism, such as competition for SigA binding with a methylation-sensitive transcriptional regulator. To determine if proteins in B. subtilis differentially associate with unmethylated DNA, we performed a pull-down in cell extracts using two different oligos. We amplified biotinylated oligos corresponding to the PscpA promoter region containing the GACGAG site. We could not obtain complete methylation of the substrate in vitro using purified DnmA. Therefore, we amplified the region and introduced a mutation in the m6A motif to GACGTG, which behaved like the WT methylated promoter in our reporter assay using the same promoter region (Figure 5A, B, middle panel). We isolated protein lysates from exponentially growing B. subtilis cells, incubated the lysates with our biotinylated oligos, performed a streptavidin pull-down, and visualized the proteins from each pull-down experiment via SDS-PAGE. We noted differences in the protein bands for the GACGAG relative to GACGTG oligo in the 20 and 40 kDa molecular weight range. These regions were excised from the gel and the proteins were identified using mass spectrometry. Of the top four most abundant proteins across the samples, the transcriptional regulator of the transition state, ScoC (65,66), was the only protein that did not appear in both pull-down experiments (Figure 6A). We found that ScoC was only present in the pull-down with the oligo that contained the GACGAG site, representing the unmethylated promoter state. No peptides corresponding to ScoC were identified in the pull-down of the GACGTG control site (Figure 6A).

To directly test if ScoC binding is affected by the A → T mutation, we purified ScoC (Figure 6B) and performed electrophoretic mobility shift assays (EMSAs). We used labeled oligos representing the PscpA promoter that only differed in the GACGAG and GACGTG sites, which overlap the –35 box but occur just outside of the ScoC consensus binding site (Figure 6C). The intensity of the shifted band was quantified and normalized to a no protein control for three independent experiments across a range of protein concentrations and the percent band shifted was compared at 250 nM and 500 nM ScoC. Consistent with the results from our pull-down experiment, we observed a 33.4% (S.E. ± 2.6) and 14.7% (S.E. ± 1.1) percent band shift at 250 nM ScoC for the GACGAG and GACGTG oligos, respectively (Figure 6D-E). We also observed percent band shifts of 70.6% (S.E. ± 9.0) and 45.7% (S.E. ± 5.1) at 500 nM ScoC for the GACGAG and GACGTG oligos, respectively (Figure 6D, E). The increased binding of ScoC to the oligo with the GACGAG site compared to the oligo with the GACGTG site (Figure 6E) and the decrease in expression we observed from the GACGAG promoter region compared to the GACGTG or GACGmAG promoter (Figure 5) supports the model that ScoC is a transcriptional repressor (65,66) and that ScoC shows preferential binding to an unmethylated promoter with the m6A site proximal to the ScoC binding site. With these results we suggest that ScoC binds to unmethylated GACGAG sites in promoter regions repressing transcription. When the GACGAG site overlaps or is adjacent to the ScoC binding site we suggest that methylation or A → T mutation at the fifth position could weaken ScoC binding leading to an increase in gene transcription.

DISCUSSION

We report that DnmA (M.BsuPY79I or M.Bsu3610I) is responsible for endogenous m6A modifications that promote gene expression in B. subtilis strain PY79. We have shown that m6A in B. subtilis occurs at non-palindromic GACGmAG sites in the chromosome with enrichment on the left chromosomal arm. In B. subtilis PY79 there are only 1,219 chromosomal m6A sites in contrast to the ∼20 000
Figure 5. Mutating the DnmA recognition motif is sufficient for differential gene expression in the PscpA promoter. (A) Analysis of the effect of mutating WT GACGAG to GACGCG (first graph), GACGTG (second graph), or GACGAC (third graph) on the activity of PscpA-GFP in WT (teal) and ΔdnmA (orange) cells. (B) Scatter dot plots, with the indicated mean and standard deviation, of the median GFP fluorescence of each strain taken from the histograms shown in (A) and appended with measurements from a similar experiment taken on a separate day. The median values were tested against each other, including the median values from the strain expressing PscpA-GFP in WT cells, for differential expression using a one-way ANOVA post-hoc Tuckey test. P-values: *P < 0.05, ***P < 0.005, ****P < 0.001, ns = not significant.

and ~4500 palindromic m6A sites in E. coli and C. crescentus, respectively (67,68). While non-palindromic sites have been described (8) and have been shown to affect gene expression (25), the palindromic nature of m6A sites in E. coli and C. crescentus is necessary for function in DNA mismatch repair, origin sequestration, and cell cycle control (67). During these processes, protein binding or activity is dictated by full versus hemi-methylated states of m6A motifs, which determines the downstream regulatory role (67,69). Here, we have shown that loss of m6A at the non-palindromic GACGAG sites in B. subtilis also affects the regulation of gene expression, with loss of methylation resulting in decreased expression of genes, including scpA and hbs, which code for proteins important for chromosome structure, organization, and maintenance (55–60) (Figure 4C and D). Our data indicate that the presence of m6A promotes the expression of a subset of genes in PY79 that could have important downstream effects on gene expression and chromosome structure.

One mechanism by which m6A regulates gene expression is through dictating transcription factor binding to promoter regions. In prototypical E. coli the methylation state of recognition sites for Dam methyltransferase in promoter regions has been shown to affect expression of a subset of genes, including virulence factors (67,69). One such example is the agn43 promoter, where methylation at the promoter blocks binding of the redox sensitive repressor OxyR, thereby stimulating production of Agn43, which is important for non-fimbrial adhesion (70). Also, uropathogenic E. coli use phase variation to evade the host immune system by altering the expression of the pyelonephritis-associated pilus (pap) in a Dam methylation-dependent manner (24). In the Gram-negative pathogen Neisseria meningitidis non-palindromic m6A sites from an active Type III RM system also function in phase variation (25). The Gram-negative bacterium C. crescentus has a transcriptional activator, GcrA, which associates with RNA polymerase-σ70 and recognizes a subset of promoter regions that are methylated at palindromic recognition sites by the CcrM MTase (71).

Here, we have demonstrated that m6A regulated promoters in B. subtilis PY79 contain the methylation site at or slightly downstream of the -35 region of the housekeeping
Table 1

| Protein  | Function                              | MW [kDa] | # PSMs GACGAG | # PSMs GACGTG |
|----------|---------------------------------------|----------|---------------|---------------|
| FabL     | Enoyl-acyl carrier protein reductase  | 27.2     | 105           | 110           |
| RpsE     | Ribosomal protein                     | 17.6     | 29            | 22            |
| RpsG     | Ribosomal protein                     | 17.9     | 23            | 11            |
| ScoC     | Transcriptional regulator of transition state | 23.7     | 16            | 0             |

**Figure 6.** Transcription factor ScoC binds the scpA promoter with an unmodified GACGAG site. (A) Top protein hits identified in the pull-down of the biotinylated scpA promoter regions with GACGAG and GACGTG sites. The #PSMs indicates the total number of peptide spectra identified for each protein using the indicated oligo in the lysate pull-down assay. (B) SDS-polyacrylamide gel of ScoC overexpressed and purified from E. coli and stained with Coomassie. (C) Schematic of the scpA promoter region. The ScoC binding consensus sequence is shown in blue, the m6A site is in red, and the -35 box is also indicated. (D) ScoC binding to 5' IR dye end-labeled scpA promoter region containing a GACGAG or GACGTG site was determined via EMSA. Representative electrophoretic mobility shift assay (EMSA) of ScoC binding to scpA promoter regions is shown. The concentration of ScoC is shown with (−) indicating the absence of ScoC from the reaction. Oligos containing the GACGAG or GACGTG site are also indicated at the top of the gel. The DNA substrates used in the reaction are otherwise identical. (E) Quantification of the percent band shifted using 250 and 500 nM concentrations of ScoC for the GACGAG and GACGTG oligos as indicated on the graph. The percent band shifted was normalized to the no protein control for each substrate. Three replicates were completed with the error bars representing the standard error between reactions.
SigA binding site (72). We have shown that, in the absence of modification at the m6A site, we observe increased binding of the transcriptional repressor ScoC in the promoter region for the gene scpA (Figure 6A–E). The increased binding of the transcriptional repressor ScoC at the scpA promoter containing a GACGAG site relative to the GACGTG site supports our reporter results, showing that the GACGAG site phenocopied the higher expression levels in a wild type strain relative to the ΔdnmA strain (Figure 5A, B). We speculate that increased binding of the ScoC repressor to unmethylated GACGAG sites is responsible for the decreased gene expression we observe from the scpA promoter, representing one mechanism by which m6A could regulate gene expression in B. subtilis PY79.

While m6A-mediated binding of ScoC represents one mechanism by which m6A regulates gene expression, we find it likely that many other mechanisms exist. The methylation-responsive promoters identified in the current study do not share an obvious ScoC consensus binding sequence. Future work will be necessary to determine the additional regulatory mechanism(s) that result in increased gene expression at methylated promoter regions in B. subtilis PY79 and 3610.

Each of the promoter fusions tested was ectopically expressed at the amyE locus, which allowed us to assay for the effect of promoter methylation status independent of the effects of chromosomal location and local chromosome architecture. This experimental design allows for identification of promoter region activities that were affected by loss of methylation at the m6A site but did not account for other factors. Interestingly, as shown (Figure 4B), the genes for many of the downregulated promoter fusions occur toward the terminus (hbs, scpA, rnhC and zapA) and on the left arm of the chromosome, whereas the amyE locus is origin proximal and occurs on the right arm of the chromosome. Thus, we are able to conclude that methylation at the m6A site in B. subtilis PY79 promotes gene expression for a subset of genes but we cannot rule out other factors that control gene expression at the endogenous loci or indirect regulatory functions of m6A elsewhere in the chromosome.

In addition to its direct regulatory function at select promoter regions, m6A may have indirect effects on gene expression. It has been shown that m6A can increase the curvature of the DNA that may, in turn, influence protein binding and chromosome architecture (73,74). Alternatively, m6A might directly influence the expression of DNA binding proteins that contribute to chromosome architecture. Consistent with this hypothesis, we observe slight but significant downregulation of the hbs gene, which codes for the essential and highly abundant histone-like protein HBsu (Figure 4C). A potential decrease in HBsu levels concomitant with the preference of HBsu for highly curved regions of DNA creates the possibility for an m6A-dependent mechanism for changes in overall DNA topology and chromosome architecture. Thus, loss of m6A may affect protein occupancy throughout the chromosome to influence chromosome architecture in such a way that results in more changes to gene expression. It is important to note that both direct and indirect models of m6A-dependent changes are possible and that they are not mutually exclusive.

Genomic m6A from orphan and active RM system MTases has been shown to function in the regulation of gene expression [e.g. (23–26)]. Here, we demonstrate that loss of MTase DnmA does not affect the natural transformation efficiency of foreign methylated DNA from a plasmid with multiple recognition sites in competent cells. Therefore, we suggest that DnmA is an MTase from an inefficient or inactive RM system. We have also discovered that DnmA-dependent m6A in the promoter regions of a subset of genes promotes gene expression in B. subtilis PY79 and we show that transcriptional repressor ScoC binds unmethylated DNA. In addition to influencing ScoC binding, we find it interesting that m6A promotes expression of several genes involved in chromosome structure and maintenance, which could in turn have effects on the expression of other genes. In total, we have shown that DNA methylation from DnmA has an effect on gene expression, prompting further investigation of RM systems and their possible regulatory contribution outside of DNA restriction.

DATA AVAILABILITY

The SMRT sequencing data for PY79 and 3610 are available (40,41) with accession number CP006881 for PY79 and accession numbers CP020102 for the NCIB 3610 chromosome and pBS32 plasmid CP020103. The SMRT sequencing data for PY79 strains ΔdnmA, ΔyabB, P spac ΔdnmA; ΔdnmA, ΔdnmAΔyabB, and NCIB 3610 strain ΔdnmA is available [GEO GSE130695] at https://www.ncbi.nlm.nih.gov/geo. The raw data, equations, and descriptions of calculations used to generate Supplementary Figure S4F and Supplementary Table S5 have been deposited to http://figshare.com DOI 10.6084/m9.figshare.8070995 and are publicly available.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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