

## Review

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# DNA Methylation and Adult Neurogenesis

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**Abstract.** The role of DNA methylation in brain development is an intense area of research because the brain has particularly high levels of CpG and mutations in many of the proteins involved in the establishment, maintenance, interpretation, and removal of DNA methylation impact brain development and/or function. These include DNA methyltransferase (DNMT), Ten-Eleven Translocation (TET), and Methyl-CpG binding proteins (MBPs). Recent advances in sequencing breadth and depth as well the detection of different forms of methylation have greatly expanded our understanding of the diversity of DNA methylation in the brain. The contributions of DNA methylation and associated proteins to embryonic and adult neurogenesis will be examined. Particular attention will be given to the impact on adult hippocampal neurogenesis (AHN), which is a key mechanism contributing to brain plasticity, learning, memory and mood regulation. DNA methylation influences multiple aspects of neurogenesis from stem cell maintenance and proliferation, fate specification, neuronal differentiation and maturation, and synaptogenesis. In addition, DNA methylation during neurogenesis has been shown to be responsive to many extrinsic signals, both under normal conditions and during disease and injury. Finally, crosstalk between DNA methylation, Methyl-DNA binding domain (MBD) proteins such as MeCP2 and MBD1 and histone modifying complexes is used as an example to illustrate the extensive interconnection between these epigenetic regulatory systems.

**Keywords:** DNA methylation, adult neurogenesis, neural stem cell, neuronal differentiation, DNA methyltransferase (DNMT), Ten-Eleven Translocation (TET), and Methyl-CpG binding proteins (MBPs), MeCP2, MBD1

## INTRODUCTION

The term epigenetics was first used by Conrad H Waddington in 1942, before the age of DNA, to define the “study of those processes by which genotype gives rise to phenotype” [1]. Later iterations of this definition required that epigenetic modifications lead to heritable changes in gene expression and function that are maintained across cell divisions or generations [2]. Thus, epigenetic regulation is particularly relevant in proliferative cells, such as those involved in AHN. Recent advances have shown that certain

cell functions depend on dynamic regulation of epigenetic programs in addition to maintaining epigenetic programs through cell division. DNA methylation, one of the core epigenetic modifications along with histone modification, can be transmitted to daughter cells and was considered a largely permanent DNA modification until recent discovery of the Ten-Eleven Translocation (TET) enzymes in 2009 [3]. This discovery, coupled with the advent of more sophisticated techniques to measure and sequence DNA methylation and other derivatives has greatly expanded our knowledge of this epigenetic mark and the various roles it plays in the cell.

Likewise, the discovery and characterization of adult neurogenesis in humans and many other species over the past twenty years has caused a paradigm shift in the field of neuroscience: you are not born

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with all of the neurons you will ever have. In small regions of the brain stem cells continuously give rise to new neurons which play important roles in learning, memory and mood regulation. In addition, adult neurogenesis is also altered in many disease states—but can also be manipulated by pharmacological treatment and other interventions, making it a promising avenue for intervention. The role of DNA methylation in brain development is an intense area of research because the brain has particularly high levels of CpG and mutations in many of the proteins involved in the establishment, maintenance and interpretation of DNA methylation impact brain development and/or function. Epigenetic pathways, including DNA methylation, influence multiple aspects of neurogenesis from stem cell maintenance and proliferation, fate specification, neuronal differentiation and maturation, and synaptogenesis. In addition, DNA methylation during neurogenesis has been shown to be responsive to many extrinsic signals, both under normal conditions and during disease and injury.

## NEUROGENESIS

The epigenetic networks regulating neurogenesis are highly connected with each other and other signaling pathways and regulatory networks. To date, embryonic cortical neurogenesis and adult hippocampal neurogenesis (AHN) are the best-studied systems. Although they have many similarities and are regulated by many of the same networks, there are also many differences that warrant investigation. Embryonic brain development must integrate growth, patterning and differentiation signals to generate neurons, astrocytes, and oligodendrocytes (and sub-types of cells) in a spatial and temporally-specific manner while still maintaining populations of stem cells. aNSCs also generate multiple cell types; moreover they are responsive to environmental inputs, pharmacological interventions, and are a major source of plasticity in the adult brain [4]. Adult neurogenesis contributes to the behavioral problems and learning deficits observed in many neurodevelopmental, neurodegenerative, and injury-based disorders [5, 6]. Animal models have provided an invaluable resource to evaluate adult neurogenesis and the mechanisms that regulate it. Several promising results indicate that it may be possible to improve function through treatment at post-natal and adult stages, even for disorders with partial developmental etiology, such as fragile

X syndrome (FXS) and Rett syndrome (RTT) [7, 8]. Indeed, many anti-depressants and other medications are known to increase neural stem cell proliferation in animal models and humans. Because there is a distinct cellular progression of adult neurogenesis, and defined behavioral outcomes associated with altered neurogenesis, adult neurogenesis is an excellent model to study regulatory mechanisms, interventions, and functional outcomes.

Neurogenesis is defined as process that leads to the generation of new functional neurons. This process includes the proliferation and fate specification of neural stem cells and the differentiation and integration of newly generated neurons into the existing neural circuitry. Mammalian neurogenesis is divided into two phases: embryonic/developmental neurogenesis and adult neurogenesis. Embryonic neurogenesis encompasses the generation of neurons in the context of the formation of the central nervous system (CNS). In the adult brain, multipotent NSCs remain and continue to generate new neurons. There are two niches of the adult mammalian CNS confirmed to have ongoing neurogenesis: the subgranular zone (SGZ) of the dentate gyrus (DG) in the hippocampus and the SVZ bordering the lateral ventricles [9]. New neurons have also been observed in other regions of the brain, but these reports are more variable and may depend on the species and physiological or pathological state (reviewed in [10]). aNSCs in both the DG and the SVZ derive from embryonic NSCs. In mice, the DG begins to develop at E13.5 from NSCs located adjacent to the cortical hem and continues to develop until early postnatal stages [11]. Some NSCs retain their multi-potent properties and transform into radial glia like cells (RGLs), so named because their cell bodies reside in the subgranular zone (SGZ) and their radial processes extend through the DG molecular layer in manner reminiscent of RGs. RGLs, also known as type 1 cells, give rise to intermediate or transit-amplifying progenitors (type2a/b, type 3) which subsequently differentiate into neurons. A small number of astrocytes are generated from NSCs, but it is unclear if they arise directly from RGLs or if they pass through a non-committed transit amplifying intermediate [12, 13]. Cells in different stages of neurogenesis can be defined by a combination of morphology, cell-type specific markers, and proliferative capacity [14]. Throughout the process of neurogenesis, DNA methylation changes as cells proceed through different cell stages and respond to different inputs.

### *Function of adult hippocampal neurogenesis*

The hippocampus was recognized as being necessary for certain types of learning and memory based on cognitive studies of memory impairment in humans and hippocampal lesion studies in model animals [15]. Further studies determined that the hippocampus is required for spatial and contextual learning and memory [16]. Evidence for the importance of NSCs—and thus the formation of new neurons—in hippocampal learning and memory came from studies that ablated adult neural progenitors via genetic means, anti-proliferative drugs, or focal irradiation [17–22]. Hippocampal neurogenesis contributes to hippocampal plasticity and both play a role in hippocampal-dependent cognitive function.

AHN represents a unique source of plasticity in the brain. Plasticity, or the ability to respond and adapt to stimulus, is generated by AHN in the DG by two ways: 1) how many new cells are produced and 2) how these cells integrate into existing networks. First, increased NPC proliferation represents a major mechanism through which more neurons can be generated. The quiescence of RGLs, the cell cycle progression and differentiation of IPCs, and the maturation of new neurons are all tightly controlled by intricate molecular networks that consist of intrinsic genetic and epigenetic programs modulated by extrinsic physiological and pathological conditions [14, 23, 24]. The second mechanism driving adult neurogenesis-mediated plasticity is the integration of new cells to existing networks. These new neurons pass through a ‘critical period’ of enhanced synaptic and dendritic plasticity 3–6 weeks post-mitosis that is dependent upon the inputs cells receive [25–27]. Disruption of cells during this window of time disrupts hippocampal-dependent learning and memory [19, 21, 28, 29], revealing that cells in this stage are vital to the functional output of adult neurogenesis.

### **DNA METHYLATION**

Epigenetic mechanisms, including DNA methylation, are critical for establishing the diverse cell fates present in the central nervous system. Different regions of the human brain (cerebral cortex, cerebellum, and pons) each have a characteristic DNA methylation signature [30]. And even within brain regions like the hippocampus, global methylation varies between neuronal subtypes [31]. There is good support from genome-wide methylation studies that

DNA methylation globally represses neuronal genes in non-neuronal cells, supporting earlier studies of individual genes [32]. Multiple mechanisms likely contribute to gene repression by DNA methylation, including recruitment of repressive complexes by methyl-CpG binding proteins (MBPs) or through blocking the binding of pro-neuronal transcription factors (TFs). Though many studies have analyzed cell-type specific transcriptomes [33–37], the number of studies that combine transcriptome and genome-wide methylation is more limited. However, the growing feasibility of this type of study will allow researchers to ask questions about the role of DNA methylation in the cell. Such as, are DNA methylation patterns cell-type specific? Do they correspond to histone modifications or the binding patterns of other repressive complexes? Is methylation always associated with repression?

### *DNA methylation and DNMTs*

DNA methylation is well known for its role in long-term gene silencing; it serves as the basis of imprinting, X chromosome inactivation, and the establishment of cell fate [38–40]. DNA methylation involves the covalent addition of a methyl group from the cofactor SAM (S-adenosyl-1-methionine) to C5 of cytosine in CG dinucleotides (also referred to as CpG, 5mC, or mCG). This addition is catalyzed by a family of DNA methyltransferases (DNMTs). DNMT3a and DNMT3b establish *de novo* methylation, whereas DNMT1 maintains methylation patterns in newly synthesized DNA by recognizing hemi-methylated DNA and methylating the unmodified strand [41].

In addition to CG methylation, other dinucleotide pairs containing cytosine can be methylated, referred to as CH or CpH methylation, where H = A/C/T. Recent studies have shown that CH methylation (mCH) is high in the brains of humans and mice [42, 43]. And within the brain, non-CG methylation is much more prevalent in neurons than non-neuronal cells and is estimated to account for 25–38% of total methylation [44–46]. CH methylation has been shown to accumulate dramatically in neurons but not astrocytes during postnatal development, a critical period of neuronal maturation and synaptogenesis [46]. There is evidence that DNMT3A is responsible for the deposition of mCH, and that non-CG methylation is also associated with gene repression [46]. The growing awareness of non-CG methylation has the potential to yield novel insights into the role of

DNA methylation in regulating brain development and plasticity

Though many studies have analyzed cell-type specific transcriptomes [33–37], the number of studies that combine transcriptome and genome-wide methylation is more limited. However, the growing feasibility of this type of study will allow researchers to ask questions about the role of DNA methylation in the cell. Such as, are DNA methylation patterns cell-type specific? Do they correspond to histone modifications or the binding patterns of other repressive complexes? Is methylation always associated with repression?

#### *TET proteins and demethylation*

Until recently, methylation was thought to be a static DNA modification, with demethylation occurring only passively upon the reduction of DNMTs. However, the discovery of 5-hydroxymethyl cytosine (5hmC) and the subsequent elucidation of the cytosine demethylation pathway substantially changed the view of DNA methylation [47]. The regulation of DNA methylation and methylation derivatives is now known to be a dynamic and active process, though the biological functions of this process are not yet entirely clear [48]. Active DNA demethylation is a multi-step process in which the methyl group is modified before the entire base is replaced via base excision repair (BER) pathways (reviewed in [3, 49]). First, members of the TET family of proteins, including TET1, TET2, and TET3, catalyze the conversion of methylated cytosine to 5hmC and subsequently to other derivatives such as 5-formylcytosine (5fC) and 5-carboxylcytosine (5caC) which are removed by BER glycosylases [50]. A second pathway, which is still controversial, involves deamination of 5hmC by AID/APOBEC to 5hmU, followed by base excision repair [51, 52]. Although this second pathway may be important in certain situations, such as neuronal activity induced demethylation (described below), it is considered unlikely that AID and APOBEC are generally involved in 5hmC-dependent demethylation [3].

Mounting evidence indicates that 5hmC methylation may have biological function beyond acting as a chemical demethylation intermediate. For example, 5hmC has a unique distribution pattern across the genome, leading to the question of how it is deposited and maintained. Compared to 5mC, 5hmC is relatively abundant at CG islands (CGIs), promoters, and within gene bodies (exons), but low in intergenic regions [53, 54]. In addition, 5hmC is relatively abun-

dant in constitutively expressed exons and displays prominent 5hmC peak at the 5'splice site boundary [55, 56]. One of the key remaining questions is how 5hmC patterns are 'read' and interpreted by the cell. One possibility is through recruitment or exclusion of DNA-methyl binding proteins.

#### *DNA methylation readers: MBPs*

Three families of proteins are known to bind to methylated DNA, including the methyl binding domain (MBD) family, the zinc finger/Kaiso family, and SET and RING associated (SRA) domain family. In addition, recent work using quantitative proteomics has also allowed for the unbiased detection of proteins that interact with specific DNA sequences including methylated and hydroxymethylated sequences [57–59]. These methods have confirmed the binding properties of many MBPs and identified many novel methylated DNA binding proteins, such as RBP-J [58], a transcription factor within the Notch pathway. These findings highlight a growing appreciation of the contribution of DNA methylation to transcription factor binding, in addition to mediating the binding of the established MBP families. So far, only members of the MBD family have been implicated in human disorders affecting the brain [60], but many other MBPs are involved in cancer [61].

#### *MBD family*

DNA methylation-induced gene repression is primarily mediated by proteins containing a methyl-CpG binding domain (MBD) [62]. Among MBPs, MBDs were discovered first and remain the best-studied to date. This family includes MBD1–6 and MeCP2 (Methyl CpG Binding Protein 2) (Fig. 1) [62]. In addition to the MBD domain, members of the MBD family possess diverse functional domains that enable them to bind to chromatin modifying proteins and other DNA-methylation specific proteins (Fig. 1). For example, MBD1 possesses a transcriptional repression domain (TRD) that enables binding to multiple histone-related proteins (Table 1). Both MBD1 and MeCP2 are highly expressed in the brain and play important roles in neurodevelopment and plasticity [60]. MBD1 Members of the MBD family play significant roles in the regulation of adult neurogenesis, which will be discussed below.

A critical step in understanding the function of MBDs is to identify their binding specificity and preferred sequences. The initial description of

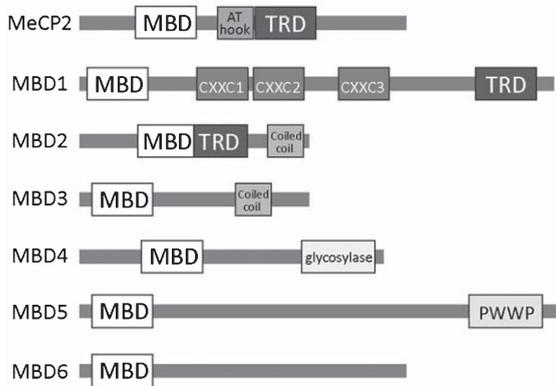


Fig. 1. A schematic overview of the MBD family of methyl binding proteins (MBPs) including known protein domains in (MBD, methyl-CpG binding domain; TRD, transcriptional repression domain; AT hook; CXXC, zinc finger Cys-x-x-Cys domain; PWWP, Pro-Trp-Trp-Pro).

MeCP2 noted that it is localized to methylated DNA and pericentromeric heterochromatin [63]. Since then, site-specific analysis has shown that these initial observations hold true: MeCP2 binds throughout the genome and tracks with mCG density [64]. Genomic mapping of the MBD proteins MeCP2 and MBD1-4 revealed that MBD binding correlates with DNA methylation, peaking at regions of dense methylation, such as methylated CpG islands, except for

MBD3 which does not bind to methylated DNA [65]. A comparison of genome-wide promoter methylation and RNA Pol II enrichment showed that active transcription is opposite to that of MeCP2, MBD1, MBD2, and MBD4, confirming the general role of these MBDs in gene repression [65]. MBD1 has multiple splice isoforms with the longest form containing both MBD domain that can bind to methylated CpG and a CXXC zinc finger domain that can bind to unmethylated CpG. A shorter version of MBD1, called MBD1v3, contains only MBD domain and 2 CXXC domains that do not bind to DNA [66]. Thus far, MBD5 and MBD6 have not been found to bind to methylated DNA, although they are localized to heterochromatin [67]. This localization may be mediated by co-factors such as the PR-DUB Polycomb complex [68].

The discovery of 5hmC in the brain led multiple groups to question whether or not known methylated DNA binding proteins also bound to 5hmC. Multiple labs have shown that the MBD domain is specific for 5mC and that the presence of 5hmC reduces binding of MeCP2 and MBD1 [59, 69–72]. There is some conflict over the ability of MeCP2 to bind to 5hmC [71]; however, some binding may be mediated by hemi-hydroxymethylated DNA as MeCP2 is already known to bind to hemimethylated DNA [73, 74]. It

Table 1  
MBD1 interacting proteins

Reference	Interactor	Function	Methods	Required domains
Watanabe et al. [227]	MPG	DNA damage repair	Y2H (TRD) CoIP (HeLa)	TRD
Reese et al. [228]	CHAF1A	Histone octamer assembly on DNA during replication	Y2H CoIP (HeLa)	MBD
Fujita et al. [229]	MCAF1	Transcriptional co-factor	Y2H	TRD
Sarraf et al. [230]	SETDB1	H3K9 methyltransferase, PCR2 component	Y2H	TRD
Ichimura et al. [225]	CHAF1A	Transcriptional co-factor H3K9 methyltransferase, PCR2 component	Y2H	MBD
	MCAF2		GST pull-down Co-IP	TRD
	SETDB1		Co-IP	
Lyst et al. [222]	SP1	Zinc-finger TF, gene activation	Co-IP	
	HP1	Pericentromeric heterochromatin	Colocalization	
	PIAS1/PIAS3	E3 SUMO (small ubiquitin-like modifier)-ligases	Y2H See [230]	
Uchimura et al. [231]	MCAF1	Transcriptional co-factor	Y2H	
Villa et al. [221]	HDAC3	Histone deacetylase	CoIP	TRD
Sakamoto et al. [224]	HPC2	PRC1 component	Y2H (Cxxc domain) Co-IP (HeLa)	
	RING1B	PRC1 component		
	Xu et al. [232]	$\gamma$ H2AX MDC1	DNA-damage associated histone Cell cycle checkpoint	Co-IP (PANC1) Co-IP (PANC1)

CHAF1A, Chromatin Assembly Factor 1, Subunit A (P150), also known as CAF-1;  $\gamma$ H2AX, phosphorylated Histone H2A.X; HDAC3, histone deacetylase 3; HPC2 also known as CBX2; MCAF1, MBD1-containing chromatin-associated factor 1, also known as AFT7IP1 or AM; HP1, heterochromatin protein 1; MDC1, Mediator of DNA damage checkpoint 1; MPG, N-methylpurine-DNA glycosylase; MCAF2, MBD1-containing chromatin-associated factor 2, also known as AFT7IP2; PIAS1/3, protein inhibitor of activated STAT-1/3; SETDB1, SET Domain, Bifurcated 1; SP1, specificity protein 1; RING1B, Ring Finger Protein 1B.

is likely that the *in vivo* binding affinity of MeCP2 and other MBDs may depend on the local chromatin environment or DNA sequence.

Based on the observation that mCH increases specifically in the brain during postnatal development, Gabel et al. investigated how MeCP2, which also exhibits high postnatal expression, binds to different methylated dinucleotide pairs [75]. They found that MeCP2 binds to mCG, mCA and 5hmCA with relatively high affinity and to mCC and mCT with low affinity. Moreover, they found that mCA was enriched in longer genes, longer genes are more enriched in the nervous system, and that changes in gene expression in the absence of MeCP2 corresponds with both of these factors [75]. The correlation between MeCP2 loss and upregulation of long genes has also been confirmed in expression profiles from various neuronal subtypes [76]. Another study found that genes that acquire more mCH methylation after birth are more likely to be differentially expressed in MeCP2 models, adding further support to the idea that binding of MeCP2 to non-CG methylation is critical for gene regulation [77]. In addition, both mCG and mCH appear to be associated with nucleosome positioning [44, 46, 78], indicating that non-CG methylation may be linked to other epigenetic mechanisms. These studies indicate that it is important to consider more than just CG methylation when evaluating MBP binding or changes in DNA methylation. DNA methylation is increasingly viewed as a dynamic modification that can be interpreted by the cell in a variety of ways to mediate gene regulation. Binding by MBPs to DNA is a major mechanism that cells use to interpret methylation status.

### *Zinc finger family*

Kaiso (also known as ZBTB33) is a member of the BTB/POZ family of zinc-fingers (ZF) and contains three copies of a Krüppel-like C<sub>2</sub>H<sub>2</sub> zinc finger. Kaiso was originally identified as a binding partner of p120 catenin [79]. It was later found to be part of the MeCP1 complex—a methylated DNA binding complex that also contains MBD2 and NuRD [80]. Kaiso is unique in that it binds to mCpG sites in a methylation-dependent manner but also binds to a specific sequence (TCCTGCNA) similar to a transcription factor [80–83]. Recent crystallization of the zinc-finger added insight into how two different types of binding are mediated by the same domain [84]. There are conflicting reports on Kaiso's ability to bind to hydroxymethylated DNA as some groups

have found that it does not bind to 5hmC [85] while others have found that it binds to both 5mC and 5hmC [59]. Kaiso acts as a transcriptional repressor [80] and has been identified as a co-factor of N-CoR [83].

Other members in the Zinc finger/ Kaiso family include ZBTB38 (ZENON) and ZBTB4 (KIAA1538). The expression of ZBTB38 is restricted to the brain and specifically to differentiating neurons—it is highly expressed during development and continues in adult [86, 87]. ZBTB4 is expressed in multiple adult tissues, but not in embryonic stages [87]. Binding of both ZBTB38 and ZBTB4 is methylation dependent [87] and ZBTB4 also binds to the Kaiso non-methylated consensus sequence [88].

### *SRA domain family*

UHRF1 and UHRF2 are 'hub' proteins that contain multiple epigenetic interaction domains including a ubiquitin-like (UBL) domain, tandem Tudor domain (TTD), plant homeodomain (PHD) finger domain, SET and RING associated (SRA) domain, and really interesting new gene (RING) finger domain [89]. The TTD and PHD domains are responsible for interactions with other epigenetic proteins such as DNMT1 and histones [90]. The SRA domain is responsible for recruitment to hemi-methylated DNA [91]. Crystallization of the SRA domain of UHRF1 revealed that binding of UHRF1 to DNA is not sequence dependent, as the SRA domain only makes contact with the hemimethylated CpG [92–94]. UHRF1 (also known as Np95) is essential in maintaining DNA methylation through cell division as it recruits DNMT1 to hemi-methylated sites [95, 96]. UHRF2, which is also known as Np95/ICBP90-like RING finger protein (NIRF), in contrast, does not maintain DNA methylation during DNA synthesis [97] but is involved in cell cycle progression and DNA damage repair. UHRF1 was shown to bind to 5mC and 5hmC with similar affinity [70]. This finding was confirmed by [59] who also showed that UHRF2 does not have any affinity for methylated DNA. Although further research is needed, this result is consistent with UHRF's role in the DNA damage response which is connected to the demethylation and BER pathways.

### *Methylation-specific transcription factors*

DNA methylation may also be interpreted/ influenced by transcription factors that are sensitive to DNA methylation, or that require DNA methylation to bind. Because many TF display cell and

developmental-stage specificity, they may represent a mechanism for fine-tuning the DNA methylation profile of different cell types. The relationship between TF and DNA methylation may function according to several different models (reviewed in [Reference 98]). Briefly, TFs can project specific sequences from DNA methylation. For example, deletion of the TF SP1, which binds to the consensus motif CCGCCC, has been shown to increase DNA methylation at a promoter with SP1 binding sites and to decrease gene expression [99]. Second, some TF complexes may promote DNA methylation. Although not a transcription factor itself, SETDB1, a H3K9 methyltransferase, interacts with DNMT3A and DNMT3B and is recruited to shared sites via TRIM28 and associated zinc-finger TFs [100, 101]. Third, there is some evidence that TF binding can reverse methylation. In REST knock out ES cells, sites surrounding REST binding motifs were methylated while they were not in WT cells, and reintroduction of REST into these cells removed the inappropriate methylation [102]. Finally, binding of TFs can reinforce repression of methylated regions. CpG methylation influences how E2F TFs bind to their recognition sites, which contain two possible methylation sites (T/CTTC/GG/CCGC/G); bi-methylation inhibits binding of all E2F TFs, but methylation of one site inhibits binding of E2F1 but not of E2F2-E2F5 [103]. E2F TFs are well known as regulators of the cell cycle, but during neurogenesis E2F3 and E2f4 bind to the promoters of many pathways involved in cell fate and differentiation such as members of the Notch, Wnt, and Fgf pathways [104].

## DNA METHYLATION AND NEUROGENESIS

The epigenetic landscape of the brain is unique: it has a specific pattern compared to other tissues [105] characterized by high levels of non-CpG methylation and hydroxymethylation, which are exclusive to the brain [106]. In addition, many epigenetic factors, such as MBPs, are highly and/or uniquely expressed in the brain during development and beyond. These observations indicate that establishing, maintaining and interpreting DNA methylation serves an important function in the brain. These roles can be divided into two general processes: 1) to help establish the diverse cell fates found in the central nervous system and 2) to alter gene expression programs based on inputs received by the cell. These two processes

occur in neurogenesis during development and in the adult. Technological advancements, such as the ability to detect DNA modifications beyond classical CpG methylation, have greatly expanded our knowledge of the function of DNA methylation in the brain.

Mounting evidence indicates that altered neurogenesis contributes to many pathological conditions [5, 6]. Adult neural stem/progenitor cells play a major role in normal brain functions and the brain's response to injury and disease. Many neurodevelopmental, psychiatric, and neurodegenerative disorders are characterized by symptoms that have been linked to reduced adult neurogenesis, such as depressive behaviors and memory and learning deficits [107]. Knockout and conditional deletion models have been used to elucidate how the loss of DNA methylation-related proteins effects neurogenesis. Although the picture is not entirely complete, it is known that MBDs, DNMTs and TET proteins play a role in some stages of AHN ranging from stem cell maintenance and proliferation to neuronal differentiation and maturation (Fig. 2, Table 2).

### DNA methylation

Neurogenesis occurs before astrogenesis during embryonic brain development; therefore, suppressing astrocytic genes in NSCs is a vital step in the temporal fate specification of these cells. DNA methylation is a major mechanism through which this occurs. For example, during neurogenic phases the STAT3 binding sites in the promoters of *Gfap* (Glial Fibrillary Acidic Protein) and *s100 $\beta$*  are highly methylated, but at later stages that corresponding to astrogenesis they become demethylated, allowing STAT3 to bind [108, 109]. A similar phenomenon occurs in ESCs: the *Gfap* promoter is highly methylated but becomes demethylated in ESC-derived astrocytes [110, 111]. Binding of NFIA (Nuclear Factor I/A), a downstream mediator of Notch and JAK/STAT activation, at the *Gfap* promoter has been shown to lead to DNA demethylation and gene activation via dissociation of DNMT1 [112].

The early embryo undergoes large-scale DNA methylation remodeling, but the changes that occur in DNA methylation during brain development are less clear. A large study that used *in vitro* neural differentiation showed that the DNA methylation state of a cell undergoes large changes as mouse embryonic stem cells (mESCs) transition to restricted neural progenitor cells and differentiated neurons [113]. Newly-methylated genes are associated with pluripo-

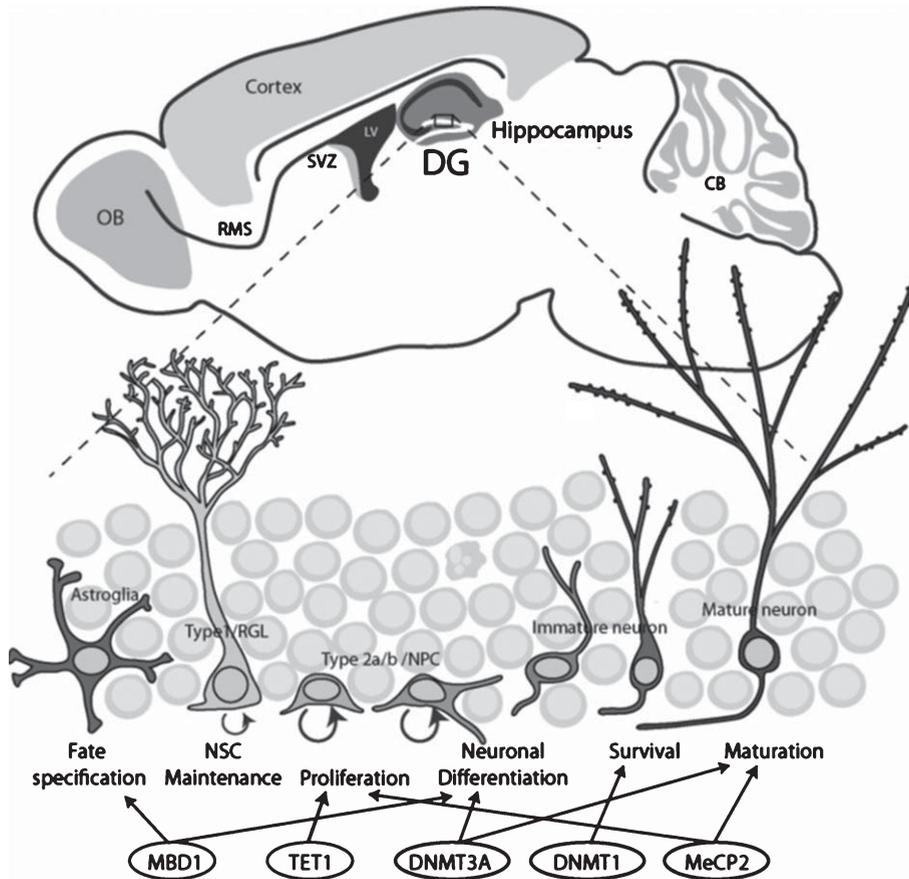


Fig. 2. Schematic drawing illustrates the two neurogenic regions in the adult rodent brains and the stages of neurogenesis regulated by DNA methylation-related epigenetic proteins. Top, Adult neurogenic niches in the brain include the dentate gyrus (DG, blue) and the subventricular zone (SVZ, purple). Bottom, the stages of DG neurogenesis and regulation by epigenetic proteins.

tency, early embryonic development, or germ line development while the majority of the demethylated genes are brain-related, supporting the hypothesis that DNA methylation in general acts to repress genes. Another important finding of this study is that patterns of DNA methylation correlate more strongly with histone modifications than with the underlying genetic code, highlighting the interplay between these two systems [114].

### DNMTs

The establishment of DNA methylation during development is essential as all DNMT null mutations (except DNMT2 and DNMT3L) are lethal in embryonic stages or shortly after birth [115]. DNMT1-KO mice develop neural tube defects (NTDs) and die by E9 [116]. Likewise, homozygous mutations in DNMT3B, which is highly expressed in the embry-

onic neural ectoderm, leads to NTDs and death by E9.5 in mice [117]. DNMT3B expression in the brain is high at embryonic day 10.5–13.5, corresponding with peak neurogenesis, but is not detectable after E13.5 [118]. In contrast, in human ESCs RNAi-mediated knockdown of DNMT3B resulted in accelerated maturation of the neuroepithelium and precocious expression of neuronal markers such as NEUROD1 coupled with loss of Polycomb repressive complex (PRC) component EZH2 at the promoters of early neuronal genes [119]. Does DNMT3B promote or suppress neuronal programs? Are the effects specific for developmental windows? DNMT3A-null mice appear normal at birth, but die at around 4 weeks of age [117]. The specific roles of DNA methyltransferases in development has been investigated using cell-type specific deletion models.

Deletion of DNMT1 in the entire CNS by E9–10 via crosses with Nestin-cre mice results in extensive

Table 2  
DNA methylation related proteins in AHN

	Mouse model	AHN Stage	Direction	Details	Ref
DNMT1	Nestin-CreER <sup>T2</sup> cKO	NSC maintenance	No change	Type 1 cells, 36 dpi Tamoxifen	[124]
		Proliferation	No change	Ki67+, BrdU+(24 hrs), 36 dpi Tamoxifen	
DNMT1/DNMT3A	Synapsin1-Cre CamK-Cre dKO	Survival	↓	BrdU+, 28 dpi BrdU	[126]
		Maturation	No change	DCX+	
DNMT3A	KO, 4 week-old mice	Maturation	↓	Reduced hippocampal volume and impaired spatial memory	[123]
		Proliferation	No change	Ki67+	
TET1	KO	Neuronal differentiation	↓	DCX+	[141]
		Neuronal survival	No change	TUNEL+	
TET1	KO and Nestin-CreER <sup>T2</sup> cKO	NSC maintenance	No change	Type 1 cells	[141]
		Proliferation	↓	Nestin-GFP+, BrdU+Ki67+	
TET1	Nestin-CreER <sup>T2</sup> cKO	Neuronal differentiation	↓	6 weeks pi Tamoxifen: DCX+BrdU+ (7 dpi), NeuN+BrdU+ (21 dpi)	[142]
		Behavior	↓	Learning and memory MWM	
TET1	KO	Neuronal diff/survival	No change	NeuN hippocampal density	[142]
		Behavior	↓	Memory extinction in MWM and contextual fear conditioning	
MeCP2	Nestin-Cre cKO	Maturation	↓	Reduced neuronal cell body	[166]
MeCP2	Retrovirus sh-MeCP2	Maturation	↓	Reduced dendritic branching and spine development	[170]
MeCP2	MeCP2 <sup>S421A</sup>	Proliferation	↓	Ki67, BrdU (24 hrs)	[174]
		Neuronal Differentiation	↑	NeuN+BrdU+ (4 weeks pi)	
MBD1	KO	Fate specification	↓ neurons	BrdU+NeuN+, BrdU+GFAP+ (4 weeks pi)	[163]
			↑ astrocytes		

MWM, Morris Water Maze.

hypomethylation in cells across the CNS [120]. These mice display precocious astrocytic differentiation linked to hypomethylation of astrocytic genes and activation of the JAK-STAT pathway [121]. This result mirrors the findings of the role of DNA methylation in astrocyte promoter activation and fate specification. Likewise, mice with conditional deletion of DNMT3A in nestin-expressing cells are apparently normal but lose DNA methylation at the *Gfap* promoter [122]. These mice die during young adult hood, most likely due to defects in the neuromuscular junctions of the diaphragm that cause respiratory failure [122].

#### DNMTs and adult neurogenesis

The observation that DNMT3A-null mice appear normal at birth but die by 4 weeks of age, suggests a role for DNMT3A in postnatal de novo methylation. A group that evaluated postnatal neurogenesis in the DG and SVZ found that the number of DCX+ immature neurons is greatly reduced, though there are no changes in proliferation [123]. This group found that DNMT3A promotes the transcription of a number of neurogenic genes by antagonizing PRC2-mediated repression. DNMT1 is also involved in adult neu-

rogenesis: deletion of DNMT1 in DG aNSC does not change cell proliferation, but does reduce new neuron survival [124]. Retinal-specific *Dnmt1* deletion from the onset of neurogenesis by Chx10-Cre also leads to defective neuronal differentiation and eventual wide-spread neuronal death [125]. These results indicate that DNA methylation in postnatal neurogenesis is required for neuronal differentiation and survival, as opposed to embryonic neurogenesis where DNA methylation contributes to cell fate specification. However, differences between *in vivo* and *in vitro* models may also contribute to some of these differences. Further exploration of how DNA methylation is altered in these mutants is needed to understand the mechanisms regulating neurogenesis.

DNA methylation also likely contributes to neuronal maturation. In addition DNMT1 and DNMT3A may have overlapping functions in post-mitotic neurons. Only Camk2a-cre double KO, which affects neurons in the hippocampus, resulted in an observed phenotype [126]. These mice had smaller hippocampi, impaired spatial memory, increased expression of genes associated with synaptogenesis, and reduced DNA methylation [126]. DNMT-mediated methylation in post-mitotic neurons is also important in other regions of the brain. For example,

Emx1-cre *Dnmt1* conditional knockout mice fail to generate the correct structures in the somatosensory barrel cortex and do not establish long term potentiation (LTP) [127]. Deletion of *Dnmt3a* driven by Sim1-Cre (single-minded1) causes obesity and overgrowth phenotypes in mice [128]; SIM1 is expressed in a subpopulation of neurons of the paraventricular hypothalamus (PVH) and amygdala that are known to contribute to energy homeostasis [129]. Because some phenotypes of DNMT3A overgrowth syndrome (discussed below) are recapitulated by loss of DNMT3A in SIM1 neurons, it suggests that de novo methylation may be more critical to the function of specific subpopulations of neurons which may contribute to the disease phenotype in different ways. Interestingly, deletion of MeCP2 in SIM1 expressing cells also causes obesity, highlighting the importance of both the establishment and interpretation of DNA methylation in this specific population of neurons [130]. In Purkinje neurons, DNMT3B is responsible for de-novo methylation of protocadherin genes that results in stochastic and monoallelic expression of different cadherin isoforms in individual neurons, a mechanism hypothesized to contribute to dendritic arborization by self-avoidance, synaptogenesis and circuit formation [131]. These studies indicate that de novo DNA methylation is required in post-mitotic neurons, though the extent to which this is required for maturation or integration of input signals particularly in the hippocampus remains to be determined.

#### *5hmC and TET*

Multiple studies have shown that 5hmC is much more abundant in brain tissues relative to other tissues of the body [132–134]. In adult cells, 5hmC is abundant in the neurons relative to other cell types or cancer cells, constituting 0.6% of all nucleotides in Purkinje neurons and 0.2% in cerebellar granular neurons [135]. The abundance of 5hmC increases as neurons differentiate and mature in both embryonic and adult models [136, 137]. Multiple studies indicate that genes that gain or are enriched for 5hmC are important for neuronal differentiation [56, 136]. 5fC and 5caC, derivatives of 5hmC, also accumulate in differentiating human ESCs (hESCs), in mouse embryos during peak neurogenesis (E11.5–13.5), and during initial differentiation of aNPCs [138].

TET proteins have garnered significant interest in recent years for their role in activity-mediated demethylation, but they appear to be involved in early neuronal development as well. All TET proteins are

expressed in the brain; TET3 is expressed at the highest level, followed by TET2 and TET1 [132]. About half of TET1-KO; TET2-KO mice survive to adulthood and are fertile; but those that die exhibit defects in head development, indicating that TET1 and TET2 are likely involved in neural development [139]. Deletion of TET3 in mice causes perinatal lethality early in development, but it is not known whether or not neurological defects contribute to this outcome [140]. Overexpression of TET2 or TET3 by in-utero electroporation enhanced cortical neuronal differentiation whereas knock-down via sh-RNA reduced the progression of neuronal differentiation and resulted in the accumulation of progenitor cells [136]. As described previously, other studies have found that DNA methylation is lost on neurogenic genes during neuronal differentiation, which is consistent with a role for TET proteins in this process.

#### *TET and adult neurogenesis*

Even though TET1 deletion is compatible with life, multiple groups have shown that its loss impacts adult neurogenesis. In TET1-KO mice there are no changes in the proliferation of RGLs but proliferation of nIPCs is reduced, leading to the generation of fewer new neurons [141]. TET1-KO animals also have learning and memory deficits [142]. Multiple studies indicate that TET1 controls the expression of neuronal-activity related genes such as *Bdnf* (Brain-derived neurotrophic factor) and *c-Fos* and that 5hmC is a key intermediate in this regulation [51, 142–144]. However, other labs have found that demethylation is not necessarily associated with increased transcription of neuronal genes during neuronal differentiation [136].

There is growing evidence that 5hmC, irrespective of its role in DNA demethylation, may be involved in transcriptional regulation, maintenance of bivalent chromatin states and neuronal differentiation. TET1 most likely does not bind to methylated DNA on its own: it likely recruits or is recruited by other factors. So far, three mechanisms have emerged that might couple 5hmC distribution, TET1 binding patterns, and gene expression. First, multiple reports have shown that TET1 and Polycomb (PcG) binding overlaps significantly genome-wide and that TET1 binds to PcG proteins targets [145–148]. PcG proteins play a critical role in regulating pluripotency genes and TET1 and TET2 knock-down leads to decreased expression of pluripotency genes [53], indicating that DNA demethylation may also be involved in

resolving bivalent chromatin. Second, TET1 may be involved in recruiting the NURD (Nucleosome Remodeling and Deacetylase) repressive complex. There is evidence indicating that the NuRD complex is targeted to specific genomic loci by TET1 and 5hmC. MBD3, which is part of NuRD complex, co-localizes with TET1 across the genome, MBD3 knock down affects expression of genes with 5hmC, and localization of MBD3 is TET1 dependent [149]. PRC2 and the MBD1-NURD complex also co-regulate a set of common targets that are characterized by bivalent chromatin state (H3K27me3 and H3K4me3), highlighting the possible interplay of multiple epigenetic pathways [150]. Third, a recent report indicates that TET1 may be recruited to a portion of its genomic loci in by Lin28Aa protein which was previously thought of as an RNA binding protein [151]. Nor are the interacting partners of TET1 limited to the mechanisms described above. TET1 Co-IP identified methyl-CpG binding proteins MeCP2 and UHRF1, histone modifying proteins including several HDACs, SIN3A and LSD1 (lysine specific demethylase), and PCNA (proliferating cell nuclear antigen) [148]. Binding competition at 5mC may be one mechanism of TET1 regulation; one study found that MeCP2 inhibited TET1 activity [152]. In addition 5hmC levels in the mouse cerebellum have been correlated with MeCP2 gene dosage [153]. In summary, the precise role of TET proteins and DNA demethylation at different stages of development is still unclear. In addition, the crosstalk between TET proteins and MBPs in the context of neural development, whether it is facilitative (MBD3) or antagonistic (MeCP2), warrants further exploration.

### MBDs

MeCP2, MBD1 and MBD5 are highly expressed in the brain. MeCP2 is highly expressed in neurons throughout the brain of adult rodents and primates and expression correlates with age [154–157]. MeCP2 is expressed in some nestin-positive precursor cells from the neuroepithelium [158] and its expression increases with neuronal differentiation [159]. Although initial reports focused on MeCP2 was expression in neurons [158, 159], subsequent reports have identified it in astrocytes and oligodendrocytes [160] and confirmed MeCP2 is not expressed in microglia [161]. MBD1 is expressed in the brain by P7 (postnatal day 7), but it is not known if it is expressed earlier [162]. In adult mice, MBD1 is expressed in neurons throughout the brain, though its

expression is particularly high in the DG and CA1 of the hippocampus [163]. MBD5 is highly expressed in the brain relative to other tissues [67]. This expression pattern is strongly indicative of a role of MBDs in brain development and function.

### MBDs and neurogenesis

MeCP2 performs dual complementary functions during embryonic fate specification of NPCs to neuronal and astrocytic fates. MeCP2 represses astrocyte genes during neurogenesis, and this repression must be released during astrogenesis. For example, during neurogenic phases, the *s100β* promoter is highly methylated, but at later stages that correspond to astrogenesis, it becomes demethylated, releasing MeCP2 during astrogenesis [109]. Other evidence suggests that the presence of MeCP2 supersedes other fate specification cues because the expression of MeCP2 in transplanted neuroepithelial progenitors promotes neuronal differentiation even when cells are transplanted into non-neurogenic brain regions [164]. In neurons, astrocyte induction cues were not sufficient to increase GFAP expression because the promoters of astrocytic genes *Gfap* and *s100β* were found to be highly methylated and repressed by MeCP2 [165]. It is still unclear if MeCP2 and other MBPs inhibit astrocyte genes directly, or if they also regulate the major pathways that promote gliogenesis. It is also possible that many of these experiments may be of limited biological relevance because the expression of MeCP2 is relatively low during embryonic development relative to its postnatal expression.

However, the majority of MeCP2 research involves the loss of MeCP2, which is more relevant for neuronal maturation rather than fate specification. NSCs derived from the cortex of E13.5 MeCP2-KO mice do not display altered fate specification for neurons, astrocytes or oligodendrocytes relative to controls [159]. Nor do MeCP2-null brains display obvious structural deficits associated with altered cell fate specification [166]. Rather, MeCP2-KO mice, similar to humans with RTT [167–169], have neurons with smaller nuclei, reduced dendritic branching and immature spines [170].

Both MBD1 and MeCP2 play a role in aNSC fate specification. In NPCs, oligodendrocytes, and astrocytes *in vitro*, both MeCP2 and MBD1 are localized to the RE1/NRSE region of the mGluR2 promoter, which is suppressed in non-neuronal cell-types [171]. This suggests that MBD1 and MeCP2 are involved in suppressing neuronal genes in astrocytes,

which is distinct from MeCP2's function in embryonic NSCs. A role for MBD1 in fate specification is supported by differentiation analysis of MBD1-KO animals injected with DNA analog BrdU (5-Bromo-2'-deoxyuridine): 4 weeks after BrdU injection, the number of BrdU-labeled neurons decreases greatly, while the number of astrocytes increases slightly [163]. These mice also show signs of depression, and exhibit impaired spatial learning, and reduced long-term potentiation believed to be a result of decreased adult neurogenesis [172]. However, in the function of MBD1 in NSCs is unexplored and it is not clear what stage of NSC development leads to reduced neurogenesis. In summary, many questions remain about how MeCP2 and other MBDs regulate stem cell maintenance, proliferation, or fate commitment.

Although MeCP2 seems to play a more prominent role in neuronal maturation, it may also function in NSCs and other cell types. MeCP2 expression in astrocytes has recently been shown to have significant impact on neuronal function and RTT pathology [160, 173]. Since astrocyte-"specific" manipulations utilize the promoter of *Gfap*, a gene that is also expressed in NSCs, it is possible that some of these effects might be, at least in part, due to changes in NSCs. In addition, the phosphorylation-incompetent MeCP2 mutant S421A reduced proliferation of and enhanced neuronal differentiation in the adult dentate gyrus, indicating that MeCP2 function is important in NSC/NPCs [174].

MeCP2 has been shown to play an important role in translating neuronal activity and signaling cascades into epigenetic gene regulation. Because phosphorylation is a dynamic modification, MeCP2 phosphorylation can integrate activity-induced signaling and gene expression [175]. For example, Chen et al. 2003 found that depolarization of neurons led to phosphorylation of MeCP2 and decreased occupancy at the *Bdnf* promoter [176]. A complementary study by Martinowich et al. linked activity-dependent changes in DNA methylation with reduced MeCP2 binding at the *Bdnf* promoter which corresponded to increased BDNF expression [177]. BDNF signaling via its receptor, TRKB (Tropomyosin receptor kinase B), is also essential for AHN [178]. MeCP2 phosphorylation also integrates Notch signaling with neurogenesis: phosphorylation-incompetent MeCP2 mutants S421A exhibited deficits associated with reduced Notch signaling and could be rescued by overexpression of the Notch intra cellular domain (NICD) [174].

## DNA METHYLATION IN DEVELOPMENT AND DISEASE

Given the unique methylation landscape in the brain and the contributions of epigenetics to brain development and function, connections between the factors responsible for the regulation of DNA methylation and human disease are of great interest. Indeed, mutations in a proportion of DNA methylation associated-genes have devastating impacts on brain development and function. Deficits in AHN may contribute to the etiology of some of these disorders [179]. However, mutations in others may be related to other brain regions and functions. But even in situations where there is no direct link between an epigenetic factor and AHN in humans, studies of AHN can be used to elucidate the function of a protein in different cells ranging from stem cells through new neurons. Moreover, AHN is a useful model system because aNSCs generate a relatively homogenous population of neurons that pass through well-defined developmental stages with consistent timing. Information gained from such studies may provide important mechanistic insights into disorders that effect the brain, but which are not primarily caused by defects in AHN, such as neurodegeneration caused by mutations in DNMT1 (discussed below). The scope of disorders linked to DNA methylation offers insight into the specific role of DNA methylation in the brain compared to other cell types and tissues.

### DNMTs

Mutations in DNMT3A have recently been associated with human disease. Researchers searching for genes responsible for overgrowth disorders identified heterozygous *de novo* mutations DNMT3A in approximately 10% of overgrowth patients, all of whom also displayed moderate intellectual disability [180]. Mutations in DNMT3A are also frequently found in acute myeloid leukemia (AML) [181] and other hematological cancers like myelodysplastic syndromes (MDS) [182]. Mutations that cause DNMT3A overgrowth syndrome have been found in all functional domains of the protein and modeling suggests that many mutations interfere with histone binding [180]. Interestingly, other epigenetic proteins seem to play a dual role in growth syndromes and myeloid neoplasms. Mutations in EZH2, NSD1, and MLL (mixed lymphomic leukemia) cause Weaver, Sotos, and Wiedemann-Steiner syndromes,

respectively, and are also associated with hematological cancers [180, 183]. This suggests that loss of de novo methylation or the inability to interpret the methylation state contributes to a cell's oncogenic transformation.

In addition, DNMT3A was identified as an autism spectrum disorder candidate gene by genome-wide association from whole-genome sequencing [184]. One study suggests that DNMT3A overgrowth syndrome and possibly the intellectual disability associated with this disorder may be comorbid with autistic symptoms: a study that integrated two large copy number variation (CNV) data sets identified small de novo deletions of DNMT3A in 3 patients with autism who also had heights and weight characteristic of over-growth syndromes [185].

Mutations in DNMT3B results in ICF1 syndrome which is about 50–60% of the cases of ICF syndrome (immunodeficiency, centromere instability and facial abnormalities syndrome), a rare autosomal recessive disorder that leads to CG hypomethylation of pericentromeric satellite regions of chromosomes 1, 9 and 16 [186–188]. About half of ICF1 patients also display a high incidence of intellectual disability [189], indicating that de-novo methylation mediated by DNMT3B is important for proper brain function and development.

Mutations in DNMT1 have been linked to two rare neurodegenerative syndromes: autosomal dominant cerebellar ataxia-deafness and narcolepsy (ADCA-DN) and hereditary sensory neuropathy with dementia and hearing loss (HSN1E) [190, 191]. Mutations that cause HSN1E prevent binding of DNMT1 to heterochromatin during the G2 phase of the cell cycle, leading to global DNA hypomethylation and selective hypermethylation [190]. Whole genome bisulfite sequencing of HSN1E patients revealed that differentially methylated regions (DMR) are associated with neurological disorders and NAD<sup>+</sup>/NADH metabolism [192]. Defects in DNA methylation have been identified in many neurodegenerative diseases including Alzheimer's disease, Parkinson's disease and Huntington's disease, reviewed in [193].

### MBPs

Mutations in MBDs have been linked to multiple neurodevelopmental disorders that manifest with varying degrees of severity. A common thread between MBPs is autism-like symptoms, with disruptions to MeCP2 representing the most severe and

MBD1 the most mild [194]. The similar yet distinct phenotypes associated with MBDs suggest that overlapping but distinct mechanisms contribute to disease etiology.

Mutations in *MeCP2* are the cause of the neurodevelopmental disorder Rett syndrome (RTT), but they are also associated with a number of other neurological disorders, including cases of Angelman syndrome, Prader-Willi syndrome, autism, and non-syndromic mental retardation [195]. Mouse models have replicated key phenotypes of RTT and revealed that MeCP2 plays a critical role in neuronal maturation and synaptogenesis, in part through the regulation of dendritic morphology, synaptic transmission, and long-term plasticity [196, 197]. Both clinically and in mouse models of RTT, deletion or loss of function MeCP2 and duplication both result in similar pathology, highlighting the necessity for its precise control [195].

As detailed above, changes in adult neurogenesis have been described for MeCP2 mouse models [174, 198]. A potential exciting avenue of research is the modulation of adult neurogenesis by various pharmacological and non-pharmacological interventions. Because adult neurogenesis is responsive to a variety of inputs, it may be possible to counteract some behavioral deficits by increasing or normalizing adult neurogenesis. For example, treatment of FXS mice with a GSK3 $\beta$  inhibitor was capable of restoring hippocampal-dependent learning and rescuing neurogenesis in adult mice [8]. Reactivation of MeCP2 in post-natal neurons also partially rescues multiple pathologies observed in MeCP2-null mice [7]. These studies suggest that some aspects of neurodevelopmental-associated pathology may be reversible and may offer useful information in the development of therapies.

*MBD5* was identified as the critical region contributing to 2q23.1 microdeletion syndrome [199, 200]. Microdeletions of that disrupt *MBD5* cause a spectrum of neurodevelopmental phenotypes that share some behavioral, developmental, and neurologic features with autism, Rett syndrome, and other disorders [200]. In addition to being a hotspot for microdeletions, mutations in *Mbd5* have been identified in ASD patients [185, 201–203]. Also, duplications of *MBD5* have been identified in patients with similar features as those with the gene disruption, indicating that gene dosage of MBD5, like MeCP2, is critical for neural development [204, 205]. Mice with MBD5 deletions display craniofacial abnormalities and decreased motor function and altered social interaction [206]. The contribution of

developmental or adult neurogenesis to these phenotypes has not been examined but may be affected.

In humans, mutations or polymorphisms in *MBD1* have been identified in sporadic cases of autism spectrum disorder (ASD) [202, 207]. *MBD1* is also contained within the region of del (18) (q12.2q21.1) syndrome, which has also been identified in some cases of atypical Rett syndrome [208, 209]. Mice with *MBD1* deletion (*MBD1*-KO) exhibit behavioral deficits associated with ASD, including learning impairment, increased anxiety, reduced social interest, and impaired sensorimotor gating [163, 172]. These phenotypes are consistent with a role for *MBD1* in the fate specification and differentiation of aNSCs.

So far, Kaiso has only been associated with cancer [210] and little is known about its function in the brain. Kaiso KO mice show no overt developmental phenotype, nor does its loss effect neural stem cell fate decision or stem cell maintenance [211]. However, another group that analyzed the behavior of the same mice found that the KO mice had increased locomotion, exploratory behavior and sensorimotor gating as well as shrunken ventricles [212]. A double KO of Kaiso and MeCP2 does not worsen the MeCP2-KO phenotype [213], indicating that Kaiso is not functionally redundant to MeCP2. It is unknown if ZBTB38 and ZBTB4 have a brain-specific function.

UHRF2 is expressed in the adult brain at comparable levels to other tissues in the body, while UHRF1 is expressed primarily in pluripotent and proliferative cells [214, 215]. Consistent with its function in DNA damage and repair, UHRF2 is frequently upregulated in cancer [89]. During embryogenesis, UHRF1 is highly expressed in proliferating NPCs around E11. In the adult hippocampus it is also expressed in proliferating NPCs, suggesting that it is involved in maintaining DNA methylation in this proliferating stem cell populations in the brain at multiple stages [216].

## CROSTALK BETWEEN EPIGENETIC PATHWAYS

DNA methylation is heavily interconnected with other epigenetic and regulatory systems such as histone modifying complexes, and transcription factor binding. The proteins that mediate the deposition, maintenance, interpretation and removal of DNA methylation are also highly connected with each other. For example, MeCP2 has been found to inter-

act with diverse proteins including, but not limited to NCOR-SMRT (nuclear receptor co-repressor), HP1 (heterochromatin protein 1), SOX2, TET1, p300, and CREB1 (cAMP responsive element binding protein 1) [197]. In addition, it is becoming increasingly clear that signaling pathways and transcriptional regulation pathways are highly connected to epigenetic regulation. Understanding how disrupting one aspect of the network influences other pathways is vital in understanding the system as whole.

*MBD1* is important in linking DNA methylation to the regulation of gene expression in combination with other epigenetic and regulatory pathways. *MBD1* binds directly to the promoter and regulates the expression of *Fgf-2*, a molecule that promotes proliferation and is commonly used to expand NSC populations. Furthermore, a DNA methylation inhibitor blocked the effects of *MBD1* in forebrain-derived aNSCs and increased *Fgf-2* expression, underscoring the important role the methylation state of the promoter plays [217]. In part, *MBD1* regulates neurogenesis by silencing miR-184; inhibiting miR-184 hinders proliferation and promotes the differentiation of hippocampal DG adult NSCs *in vitro*. In turn, miR-184 binds the 3' UTR of Numbl (Numbl), a signaling protein required for differentiation in adult neurogenesis, and targets it for degradation [218]. Numbl is known to inhibit the Notch pathway [219], which has a significant impact on neurogenesis. Inhibition of miR-195, a microRNA repressed by *MBD1*, increases neuronal differentiation *in vivo*, similar to miR-184 [220].

Similar to MeCP2, *MBD1* has been found to interact with multiple epigenetic co-factors and it is highly likely that *MBD1*-mediated regulation of adult neurogenesis requires cooperation from other epigenetic machinery. *MBD1*-interactors have been identified primarily in cancer cell lines and it is not known if these interactions are relevant to *MBD1* regulation in the context of adult neurogenesis (Table 1). HDAC3 can facilitate the regulation of gene expression by *MBD1* in cancer cells [221] but it is not clear gene repression is mediated through the recruitment of HDACs by *MBD1* as the HDAC inhibitor trichostatin A (TSA) does not always activate *MBD1*-repressed genes [222, 223]. However, MeCP2, *MBD1*, and HDACs have also been shown to target the RE1 REST binding site in a subset of neuronal genes, reflecting the possibility that both DNA methylation and histone remodeling are required to mediate gene repression [171]. In addition, *MBD1* has been shown to interact with PRC1 components HPC2 and RING1B [224],

and PRC2 component SETDB1 [225], indicating a possible role in the resolution of bivalent chromatin states. In addition, in the absence of MBD1, the levels of activating and repressive histone marks are altered at the promoter of miR-184 in aNSCs, suggesting that MBD1 can influence the recruitment of histone-modifying proteins [218]. However, exactly how MBD1 interacts with other epigenetic pathways remains unknown.

## CONCLUSION

The brain is a hot-spot of epigenetic regulation: the generation of heterogeneous cell populations from shared progenitors requires many levels of gene regulation. Many epigenetic factors are highly expressed or highly prevalent in the brain, such as many MBPs and DNA modifications. Recently, improved technologies for detecting DNA modifications beyond classical CpG methylation has greatly expanded our knowledge of the existence and possible function of mCH, 5hmC, other modifications, and demethylation pathways. Epigenetic pathways are increasingly found to be highly interconnected and DNA modifications are poised to be at the center of networks linking transcriptional regulation, chromatin states, and environmental inputs. The proteins that deposit, read and remove DNA methylation are vital for the function of the cell. The contributions of these proteins in the development of the nervous system are particularly interesting given how they are already known to regulate neurogenesis. However, there are many outstanding questions that remain regarding the function of DNA methylation in adult neurogenesis. The fields of epigenetics and AHN are starting to address what roles methylation-specific proteins play at each stage of neurogenesis (Fig. 2), but the functions in all cell types are not all known. One example is MBD1, which we have been studying for the past decade. Does MBD1 play a role in NSC maintenance and proliferation? Does it regulate neuronal maturation? What pathways and mechanisms control these possible functions? What are the cofactors or facilitators for MBD1 regulation of neurogenesis? Does MBD1 have a role in neurogenesis during juvenile period which may have important role in developing social ability [226]? Beyond neurogenesis, questions about the basic mechanisms of these proteins still remain. What is the binding specificity of MBDs in physiologically relevant cell types? How are MBDs recruited to methylated DNA? Do some MBDs have specific affinity for each type of non-

CG methylation? Can MBDs compensate each other? Do MBDs interact with each other? As the field of adult neurogenesis moves increasingly towards an integrated network-based understand of regulatory networks and disease mechanisms, understanding the how epigenetics and other pathways intersect is becoming increasingly important.

## ACKNOWLEDGMENTS

This work was supported by grants from the NIH to X. Z. (MH080434, MH07897, R21NS095632), a center grant from the NIH to the Waisman Center (P30HD03352), and a NIH Molecular Biosciences Training Grant to E.M.J (MBTG: T32 GM07215).

## CONFLICT OF INTEREST

The authors have no conflict of interest to report.

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