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Dynamics of RNA m⁵C modification during brain development

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Abstract

Post-transcriptional RNA modifications have been recognized as key regulators of neuronal differentiation and synapse development in the mammalian brain. While distinct sets of 5-methylcytosine (m⁵C) modified mRNAs have been detected in neuronal cells and brain tissues, no study has been performed to characterize methylated mRNA profiles in the developing brain. Here, together with regular RNA-seq, we performed transcriptome-wide bisulfite sequencing to compare RNA cytosine methylation patterns in neural stem cells (NSCs), cortical neuronal cultures, and brain tissues at three postnatal stages. Among 501 m⁵C sites identified, approximately 6% are consistently methylated across all five conditions. Compared to m⁵C sites identified in NSCs, 96% of them were hypermethylated in neurons and enriched for genes involved in positive transcriptional regulation and axon extension. In addition, brains at the early postnatal stage demonstrated substantial changes in both RNA cytosine methylation and gene expression of RNA cytosine methylation readers, writers, and erasers. Furthermore, differentially methylated transcripts were significantly enriched for genes regulating synaptic plasticity. Altogether, this study provides a brain epitranscriptomic dataset as a new resource and

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Author contributions

H.X. conceived and designed the study; X.X. isolated and cultured E16.5 mouse cortical neurons and neural stem cells, constructed libraries for RNA-seq and RNA bisulfite sequencing; Z.J. and Y.L. performed the bioinformatic analyses; X.X., Z.J., Y.L., and H.X. interpreted results and wrote the manuscript. All authors have read and agreed to the published version of the manuscript.

Declaration of Competing Interest

The authors declare no conflict of interest.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.ygeno.2023.110604>.

lays the foundation for further investigations into the role of RNA cytosine methylation during brain development.

Keywords

Brain development; Neuron; Neural stem cell; RNA cytosine methylation; RNA bisulfite sequencing; RNA-seq

1. Introduction

Over the past decade, epitranscriptomics has emerged as a new field to study the post-transcriptional modifications of RNA bases on a transcriptome-wide scale [1]. Among the over 170 kinds of RNA modifications, RNA cytosine methylation (m^5C) has gradually been recognized as an important form regulating RNA metabolism. The m^5C modification on tRNAs regulates tRNA stability and protein translation [2], while cytosine methylation on rRNAs coordinates mitochondrial assembly and translation activity [3,4]. Despite their lower abundance, maternally derived m^5C -mRNAs are essential for embryonic development during the maternal-to-zygotic transition [5] and ablation of maternal m^5C -mRNAs results in developmental delay and embryonic defects [6]. Cytosine methylation on mammalian mRNAs is primarily catalyzed by NOP2/Sun RNA Methyltransferase 2 (NSUN2) [7,8] and is removed *via* m^5C oxidation by the Ten Eleven Translocation enzymes [9,10]. At DNA damage sites, the RNA methyltransferase, TRNA Aspartic Acid Methyltransferase 1 (TRDMT1), is recruited to introduce m^5C to mRNA in the form of DNA:RNA hybrids [11]. The methylated RNA in such hybrids was recognized by the m^5C readers RAD51/52 [11]. Another m^5C binding protein, Aly/REF Export Factor (ALYREF), mediates the export of methylated mRNAs from the nucleus to the cytoplasm [7]. mRNAs carrying m^5C sites can also be recognized by the Y-box binding protein 1 (YBX1), which recruits the poly(A) binding protein cytoplasmic 1a (PABPC1A) to achieve stabilization of methylated mRNAs [12].

Accumulating literature emphasizes the important roles of RNA post-transcriptional modifications in brain development and function. In neuronal stems cells, the loss of RNA methyltransferase NSUN2 results in decreased m^5C -tRNA levels, leading to aggregation of tRNA fragments, activation of a cellular stress response, and impairment of neuronal stem cell differentiation [13,14]. In mouse models, NSUN2 loss inhibits the neurogenesis of upper-layer cortical neurons inducing microcephaly and motor defects [13,15]. Later, targeted knockout of NSUN2 in the mouse prefrontal cortex was shown to have reduced levels of m^5C -tRNA and resulted in the disruption of neuronal synaptic signaling patterns and behavior [16]. Although the influence of NSUN2 loss on mRNA methylation remains unclear, methylated mRNAs in brain tissue were found to be enriched for genes involved in ion transport and synapse function [17]. In neuronal oxidative stress models, an elevation in mRNA methylation level was associated with the stress response and regulation of apoptosis [18]. In glioblastoma cells, methylated transcripts mediated by the m^5C writer NSUN6 were linked to the regulation of transcriptional and translational processes after alkylation treatment [19]. These findings indicate that m^5C -modified RNAs may influence brain cell

differentiation and function. However, no previous study has attempted to characterize m⁵C dynamics during mammalian brain development.

Brain development is driven by drastic transcriptome changes accompanied by an increase in cellular diversity and neuronal connectivity [20]. The expansion and migration of neuronal lineages occur at the pre-natal stage while post-natal brain development is characterized by active gliogenesis and rapid neuronal diversification [21,22]. Around postnatal day 17 (P17), the mouse brain enters a critical period of cortical plasticity that shapes neuronal circuits in response to early life experiences [23]. In this study, we performed RNA BS-seq to survey the landscape of mRNAs with methylated cytosines in the mouse brain at critical developmental time points, as well as neural stem cells and E16.5 cortical neuron cultures (Fig. 1A). In combination with bulk RNA-seq and single-cell RNA-seq data, we aimed to explore the links between the patterns of mRNA cytosine methylation and gene expression during brain development.

2. Materials and methods

2.1. Mice

C57BL/6 mice were maintained and bred in a 12-h light/dark cycle under standard pathogen-free conditions. Mouse brains were harvested at postnatal day 0, 17, and 6wk for total RNA extraction. Adult female mice were used for setting up timed pregnancy. Embryos were timed by checking vaginal plugs daily in the morning. Positive plugs were designated as E0.5. The experiments were approved by the Institutional Animal Care and Use Committee (IACUC) of Virginia Tech.

2.2. Mouse neural stem cell (NSC) culture

Mouse NSCs were isolated from the subventricular zone (SVZ) of the lateral ventricles as described previously [24]. NSCs were seeded on poly-ornithine and laminin-coated plates. The culture media was prepared by mixing Dulbecco's Modified Eagle Medium (DMEM) and Ham's F12 media at 1:1 volume ratio, supplemented with 2% B27, 2 mmol/L L-glutamine, 1× penicillin-streptomycin, 20 ng/ml epidermal growth factor (EGF, PeproTech), and 20 ng/ml basic fibroblast growth factor (bFGF, PeproTech).

2.3. Primary mouse cortical neuronal culture

Primary mouse cortical neurons were prepared as previously described [25] with some modifications. Briefly, C57BL/6 E16.5 mouse embryos were micro-dissected for cortex tissues and the cortex tissues were dissociated into single-cell suspensions using the neural tissue dissociation kit (Cat# 130–092–628) according to manufacturer's instructions. After dissociation, neuronal cells were filtered through a 70-µm strainer (Falcon) and spun at 300 *g* for 10 min. The cell pellet was resuspended in neuronal culture medium (Neurobasal medium containing 2% B27 supplement (Invitrogen), 1% Glutamax (ThermoFisher), and 1% penicillin-streptomycin (ThermoFisher) and seeded on laminin and poly-ornithine coated 10-cm dishes. Neurons were grown *in vitro* for 7 days with fresh medium changed on Day 3 and Day 6.

2.4. Immunostaining

Immunostaining was performed as previously described [26]. Briefly, NSCs or E16.5 mouse cortical neurons were seeded onto an 8-well chamber. The neurons were fixed with 4% paraformaldehyde in phosphate buffered saline (PBS) for 15min and permeabilized with 0.2% TritonX-100 in PBS for 10min. After being blocked with 5% Normal Goat Serum (Thermo Fisher) at room temperature (RT) for 1h, the cells were incubated with mouse anti-Nestin antibody (Millipore, MAB353) and rabbit anti-Sox2 antibody (Abcam, ab97959) for NSCs, or with mouse anti-Tuj1 antibody (Biolegend, 801,201) and rabbit anti-GFAP antibody (Sigma, HPA056030) for E16.5 cortical neurons at 4°C overnight. Then, the cells were incubated with Cy3 conjugated anti-rabbit IgG (A10520, Invitrogen) and Alexa Fluor 488 conjugated anti-mouse IgG (A10680, Invitrogen) secondary antibodies at RT in darkness for 1h. After washing 3 × 5min with 1 × PBS, cells were mounted with DAPI-Fluoromount-G™ Clear Mounting Media (Southern Biotech, 010020). Fluorescent images were acquired using a confocal microscope.

2.5. RNA BS-seq library construction

RNA bisulfite conversion was performed as previously described [27] with minor modifications. Poly(A) RNA was first mixed with spiked-in Xef1 unmethylated RNA at a ratio of 0.5%. The spiked-in unmethylated mRNA was transcribed from the pTRI-Xef plasmid supplied by the MEGAscript™ T7 Transcription Kit (Invitrogen) according to manufacturer's instructions. RNA bisulfite conversion was performed with an initial denaturation at 95 °C for 1 min, followed by three cycles of 70 °C for 10 min and 64 °C for 45 min using the EZ RNA methylation Kit (Zymo Research). The bisulfite converted RNA was subjected to the stranded RNA-seq library construction procedure using the TruSeq Stranded mRNA Library Preparation Kit (Illumina). We modified the procedure to skip the RNA fragmentation step and supply both random and ACT random hexamers during the first strand cDNA synthesis.

2.6. RNA BS-seq data analysis

RNA BS-seq data analysis was performed as previously described [27]. Raw sequencing reads were trimmed at the 5' and 3' ends by 6 bp to account for methylation bias and then filtered for low quality bases and adaptor sequences. After the removal of reads with short lengths, clean reads were mapped to the mm10 genome (Ensembl v.79) using meRanGh [28]. To exclude partially unconverted reads, mapped reads were further filtered by removing reads with >3 "C"s ("G"s on the cDNA strand). Methylation calling was performed using meRanCall and subjected to a series of filters to reduce false positive signals [27]. In addition to the thresholds applied on the methylation level (≥ 0.1) and read coverage (≥ 20) of m⁵C sites, additional filters were adopted including signal-to-noise ratio and the maximum number of m⁵C sites detected in a read to reduce false positive methylation calling [27]. The C-coverage threshold ranging from 6 to 10 was determined using Gini Index [8]. High-confidence sites were defined as the m⁵C sites passed all filters in both biological replicates.

2.7. Differential methylation analysis

Differential methylation analysis was performed using Fisher's exact test on high-confidence sites present in at least one sample. Benjamini-Hochberg p -value correction was used to correct for multiple comparisons, and sites with p -adjusted <0.05 were reported.

2.8. RNA-seq library construction

Stranded RNA-seq libraries were constructed using the TruSeq Stranded mRNA Library Preparation Kit (Illumina) following manufacturer's instructions. Briefly, after two rounds of poly(A) selection, the mRNA samples were fragmented and primed to synthesize first strand cDNA, followed by synthesis of the second strand cDNA. After Ampure XP bead purification, dA tailing was performed, and indexed adapters were ligated to both ends of the double-stranded cDNA. Adapter-ligated DNA fragments were enriched by PCR amplification for 12 cycles. After Ampure XP bead purification, the PCR products were size-selected with a range from 350 bp to 550 bp on 2% dye-free agarose gel using the pippin recovery system (Sage Science). The recovered libraries were sequenced on a HiSeq 4000 platform in the 150 bp paired end mode (Illumina).

2.9. RNA-seq data analysis

Trim Galore (version 0.6.5) was used to filter short reads, low quality reads, and trim adapter sequences from raw reads (https://www.bioinformatics.babraham.ac.uk/projects/trim_galore/). Clean reads were mapped to the mm10 genome and expression quantified using STAR (version 2.7.3a) [29]. Differentially expressed genes were identified using DESeq2 [30] using a fold change >1.5 and an adjusted p -value of 0.01 as cutoffs.

2.10. Gene ontology (GO) analysis

GO analysis was performed using DAVID (Database for Annotation, Visualization, and Integrated Discovery) [31]. The GO Direct terms, Biological Process (BP), cellular component (CC), and molecular function (MF) were identified. Significantly enriched terms were identified using Benjamini-Hochberg p -adjusted <0.05 .

2.11. Clustering analysis

The z-score of bulk RNA-seq expression of transcript per million (TPM) and differentially methylated site methylation levels were used as input for clustering analysis. Clustering analysis was performed with the Python package sklearn's Agglomerative Clustering. The 'complete' method was used with the number of clusters set to 5 for RNA-seq and RNA BS-seq datasets. Error bars represent the standard deviation of the z-score.

2.12. Availability of data and software

Data generated in this study were submitted to the NCBI Gene Expression Omnibus under accession number GSE207092. Analyses in this study were performed using the R v4.1.1, and Python 3.9.4 packages Biopython v1.78, matplotlib v3.3.4, Seaborn v0.11, and Pysam v0.16. The software package developed in this study is available in the GitHub repository (<https://github.com/zaustinj33/BrainDev>).

3. Results

3.1. Identification of high-confidence m⁵C sites in mouse NSCs, neurons, and brain tissues

To obtain RNA expression and m⁵C epitranscriptome profiles during brain development, we performed RNA-seq and RNA BS-seq for NSCs (Fig. 1B), neurons in culture (Fig. 1C), and brain tissues isolated from postnatal day 0 (P0), postnatal day 17 (P17), and 6-week-old (6 W) mice. Two biological replicates were generated for each condition with a total of 10 libraries obtained for RNA-seq and RNA BS-seq (Table S1). Approximately 20 million paired-end reads were generated for each RNA-seq library and aligned to the mouse reference genome (mm10). For these libraries, the average percentage of sequences uniquely mapped to the reference was around 92%. The normalized gene expression level of the samples was provided (Table S2). Since methylation calling requires a much higher read depth compared with those of RNA-seq libraries, five times more sequences were generated for RNA BS-seq libraries (~100 million pair-end reads for each library). Sequencing reads derived from RNA BS-seq libraries were processed following the procedure described in our recent study [27]. In brief, bisulfite sequencing reads were first trimmed to remove adaptors and bases with low quality scores. After quality control, reads were mapped to the C2T converted mm10 genome. Around 85% of bisulfite sequencing reads were uniquely aligned (Table S1). Methylation calling was performed using meRanCall [28] and putative m⁵C sites were subjected to a series of filters to reduce false positive signals [27].

A total of 2259 m⁵C sites were identified from the ten RNA BS-seq libraries. Among these m⁵C sites, 1758 sites passed all filters in one biological replicate but failed in the other; thus, these were denoted as “low-confidence” m⁵C sites. The remaining 501 m⁵C sites shared by both biological replicates were considered to be “high-confidence” in this study (Fig. 2A). The methylation levels of the high-confidence m⁵C sites were highly correlated between the biological replicates and had Spearman *R* values between 0.82 and 0.92 (Fig. 2B). Notably, 76.4% of these sites had a methylation level lower than 0.3 (Fig. S1). This result was consistent with previous reports that only a small percentage of RNA copies are methylated for a given gene [8,27,32].

To explore the factors contributing to the level of confidence in methylation calling, we first compared methylation levels and read coverages of the high-confidence and low-confidence sites. Across five conditions, the high-confidence sites showed average methylation levels in the range of 0.18–0.21, while the low-confidence sites had average methylation levels in the range of 0.06–0.12 (Fig. 2C). The average read coverages for the high-confidence sites were between 109 and 227 reads per site, while the average for the low-confidence sites was only 24–48 reads per site (Fig. S2A&B). We next examined the relationship between read coverage and methylation level (Fig. S2C). A very weak positive correlation was observed between the methylation level and the read coverage for both high-confidence (Spearman *R* < 0.19) and low-confidence sites (Spearman *R* < 0.08). Altogether, the high-confidence m⁵C sites tended to be found with a higher methylation level and more read coverage to enable them to survive all the stringent filters in both biological replicates.

Previous studies suggested that the process of library construction, the bisulfite conversion step in particular, could have a significant impact on methylation calling [8,27,32]. In this study, we generated both RNA-seq and RNA BS-seq data for the same pool of RNA samples.

This enabled us to perform correlation analyses for the read coverage of each methylated transcript in the paired libraries for RNA-seq and RNA BS-seq. Interestingly, compared with the low-confidence sites, the high-confidence sites showed higher correlations in read coverage with or without bisulfite conversion, except for P17 brain samples in which the two correlations are similar (Fig. S3A). We further examined the average length of methylated transcripts and the distribution of m⁵C sites in transcripts. No significant difference in transcript length was detected for transcripts containing either high or low-confidence sites (Fig. S3B). The distribution of high-confidence sites was generally biased towards the 3'UTR of transcripts, in particular for the P17 brain samples (Fig. S3C). This result indicates that low-confidence sites tend to reside in a transcript sensitive to bisulfite treatment, but such sensitivity may not have a link with transcript length.

3.2. High-confidence m⁵C sites differentially methylated in mouse NSCs, neurons, and brain tissues

We next focused on the 501 high-confidence sites to examine their sequence features and distribution across five conditions. Two common features of m⁵C sites on mRNAs have been reported: 1) they often localize upstream of the translation initiation sites (TISs), and 2) a “GGG” motif is frequently observed downstream from the m⁵C sites [8,27,32,33]. To determine the distribution of high-confidence m⁵C sites along the transcripts, we binned the lengths of methylated transcripts into the 5' - untranslated region (5'UTR: bins 1–5), coding sequence (CDS: bins 6–28), and 3' - untranslated region (3'UTR: bins 19–46). Two density peaks surrounding TISs and an additional strong peak right before the termination of the coding region were observed for m⁵C sites in all five conditions (Fig. S4A). A downstream “GGG” motif was determined for the high-confidence m⁵C sites in all five conditions (Fig. S4B), which may serve as potential targets of the NSUN2 enzyme [8].

Out of 501 high-confidence m⁵C sites, a total of 31 sites were identified as methylated in all conditions and 253 sites (50.8%) were shared by at least two conditions (Fig. 3A). Such a distribution indicates that at least some mRNAs are consistently methylated across multiple time points throughout brain development. Only 82 m⁵C sites were identified in NSCs while the libraries derived from the three-stage brain tissues reported a similar number of high-confidence sites ranging from 222 to 269. In addition, m⁵C sites in undifferentiated NSCs were largely over-lapped with those in neurons and brain tissues. To determine the effect of sequencing depth on methylation calling, we calculated the yield of m⁵C sites after normalizing to the number of total mapped reads for each library (Table S1). Despite more reads generated for NSC libraries, both NSC replicates yielded the fewest m⁵C sites regardless of high-confidence or low-confidence. The methylation level of high-confidence sites reported in the P17 brain were significantly lower than sites reported in the P0 brain (Fig. 3B). These results suggested that the diversity of mRNA cytosine methylation may increase during brain cell specification.

Since brain development is accompanied with substantial changes in gene expression, we selected a set of 50 high-confidence m⁵C sites that possessed at least 20× read coverage in all conditions. According to their average methylation levels, five clusters were identified with unique methylation patterns (Fig. 3C). The m⁵C sites in three clusters had relatively consistent methylation levels throughout all conditions (at high, low, and medium methylation levels, respectively). Such stable methylation patterns suggest that cells from neural lineage may share a regulatory mechanism to control the methylation of a small set of transcripts. Cluster C0 and C4 showed high methylation levels in neurons and P0 brain samples, respectively (Fig. 3D). We further determined their associated transcripts for GO enrichment analysis. Interestingly, transcripts highly methylated in neuronal culture (cluster C0) were associated with transcription, chromatin organization, and stem cell maintenance (Fig. 3E). For instance, genes including *Foxo3*, *RelA*, and *Rptor* are important mediators of neuronal cell functions including reprogramming and differentiation [34] and synaptic formation [35].

3.3. Gene expression dynamics of methylated mRNAs, m⁵C readers, writers, and erasers during brain development

We extended the analysis to identify differential methylation sites (DMSs) across five conditions. DMSs were defined as m⁵C sites with a methylation difference ≥ 0.05 between two conditions and a *p*-adjusted value ≤ 0.05 . Pairwise comparisons determined 83, 176, 62, and 172 DMS sites for NSCs vs neurons, P0 vs P17, P17 vs 6 W, and P0 vs 6 W, respectively (Fig. 4A). Interestingly, of the 83 DMSs identified between NSCs vs neurons, 96% showed increased methylation in neurons. Gene Ontology analyses indicated that differentially methylated sites in neurons compared to NSCs are significantly enriched for positive regulation of DNA-templated transcription, including RNA polymerase II transcription (Fig. 4B). The differentially methylated sites identified in postnatal brain tissues were significantly enriched for the regulation of GTPase activity and synaptic plasticity associated with brain development. For instance, the transcripts *Atxn1*, *Fgfr1*, and *Itsn1*, which are known to be critical for brain development, were found to be hypomethylated in the P0 brain.

To examine the relationship between RNA cytosine methylation and gene expression, we performed differential expression analysis of sample-matched RNA-seq libraries. These differentially expressed genes (DEGs) were defined as transcripts with an expression difference ≥ 1.5 -fold between two conditions and a *p*-adjusted value ≤ 0.01 . The majority of transcripts carrying DMSs were not differentially expressed between P17 and 6 W (Fig. 4C), while a number of genes showed significant changes in both gene expression and RNA methylation during early brain development. A moderate positive correlation ($R = 0.337$ in NSCs vs neurons) was observed between the changes in gene expression and RNA methylation (Fig. 4D). As aforementioned, transcripts carrying m⁵C sites tend to have a higher methylation level in neurons compared to those in NSCs. Of those differentially methylated transcripts, 52.3% exhibited an increased expression level in neurons. Notably in neuronal samples, the transcript *Psd*, which codes for a Plekstrin homology and SEC7 domain-containing protein, was over-expressed and hyper-methylated. SEC7 domains were conserved throughout many proteins and served to catalyze the guanine nucleotide exchange

factor initiating the formation of vesicle coating [36]. However, both positive and negative correlations between gene expression and RNA cytosine methylation were observed for pairwise comparisons of NSCs *vs* neurons, P0 *vs* P17, and P0 *vs* 6 W. For instance, the differentially methylated transcripts *Csf1r* and *Endod1*, which were hyper-methylated and under-expressed in the P0 brain, are markers for microglia and endothelial cell types, suggesting cell-type specific methylation of transcripts. Conversely, two transcripts, *Map6* and *Sdc3*, were consistently hyper-methylated and over-expressed in P0 samples compared to P17 and 6 W samples. *Map6* infers structural stability in developing neuronal microtubules, promoting interconnectivity and signaling [37,38], while *Sdc3* is a well-known mediator of neuronal lineage, promoting neuronal migration and synapse formation [39,40]. This result indicated that the regulation of RNA cytosine methylation and gene expression may be cell-type specific and is positively correlated for some transcripts.

Recently, significantly increased understanding has been gained about the writers, readers, and erasers of m⁵C in mRNAs [41–44]. To examine the expression patterns of these factors, we first identified the top 1000 variably expressed genes in our RNA-seq dataset and performed clustering analysis (Fig. 5A). Not surprisingly, NSCs showed distinct gene expression profiles from other samples. We further checked the expression profiles of the factors regulating RNA cytosine methylation. It showed highly dynamic expression patterns during neural cell specification and postnatal brain development (Fig. 5B). In the comparison between NSCs and neurons, it showed that NSUN2 and YBX1 are highly expressed in both, while NSUN4, ALYREF, and FMR1 are highly expressed in NSCs, TET1 and TET3 are highly expressed in neurons (Fig. 5B). In the comparison among the three postnatal stages (P0, P17 and 6 W), it underwent clear downregulation trend, with the P0 brain displayed the highest expression for most m⁵C readers/writers/erasers. Except that NSUN4 and RAD52 showed increasing upregulation during postnatal brain development (Fig. 5B). We further performed pairwise comparisons to determine the transitions in gene expression (Fig. 5C). With the cutoff of fold change >1.5 and an adjusted *p*-value <0.01, 9501 DEGs were identified in between NSCs and neurons, 6672 DEGs and 6997 DEGs were identified in the comparison P17 *vs* P0, and 6 W *vs* P0, respectively, while only 430 DEGs were identified in the comparison between 6 W and P17 (Fig. 5C). Consistent with the transcriptome-wide differences among the four comparisons, the 6 W *vs* P17 comparison yielded the fewest differentially expressed factors related to RNA m⁵C regulation (Fig. 5D). This suggests that m⁵C regulation is highly dynamic during early postnatal brain development and may reach homeostasis throughout the maturation process.

3.4. Differentially methylated transcripts have cell type-specific expression patterns and are temporally regulated in developing brains

During brain cell specification, more m⁵C sites (Fig. 3A) and increased methylation (Fig. 3B) were observed in differentiated neurons and maturing brains. This inspired us to further examine the temporal expression patterns of differentially methylated transcripts using bulk RNA-seq data publicly available for mouse embryonic/postnatal brain tissues including different neuronal subsets. We focused on 42 differentially methylated transcripts identified in the RNA BS-seq datasets, which also showed expression fold-change >2 across the five RNA-seq conditions (Fig. 6A). According to their expression profiles in bulk

RNA-seq, four distinct clusters were identified (Fig. 6B). Cluster 1 and cluster 3 were characterized with high expression levels in the adult brain and newborn brain, respectively. Of the Cluster 1 transcripts, *Endod1*, *Wbp2*, and *Rapgef4* were highly expressed in the mature 22-month-old brain samples and were found to be both over-expressed and hyper-methylated in our 6 W brain samples compared to the P0 brain. ENDOD1, an endothelial marker, was found in a previous study to have high expression in the cerebral cortex and hippocampus, while WW domain binding protein 2 (WBP2) and Rap guanine nucleotide exchange factor 4 (RAPGEF4) were mainly localized to the cerebral cortex [45,46]. WBP2 is a transcriptional co-activator of estrogen receptor α (ESR1), of which mutations induce abnormal glutamatergic synapse development [47].

Methylated transcripts in cluster 2 showed highest expression level in neurons. To further examine the expression patterns of these methylated transcripts, we integrated the epitranscriptome profiles generated in this study with an scRNA-seq dataset recently published for the P0 cortex [22] (Fig. 6C). Previous studies reported that the methylation of tRNAs is essential for proper development of cortical layers [13,15]. Intriguingly, scRNA-seq data also suggested that some genes with abundant neuronal expression were associated with a relatively high level of mRNA methylation (Fig. 6D). In neurons, the *Psd*, *Rundc3a*, and *Smardc3* transcripts were hyper-methylated, and their expressions were also broadly enriched for many neuronal subpopulations in cortical layers I–VI, interneurons 1–3, and striatal inhibitor neurons 1 and 2. For these genes, such a cell-type predominant expression was also reported in previous studies. For example, localized expression of *Rundc3a* was previously found to be concentrated within noradrenergic populations of multiple brain regions [45,46]. Taken together, these results indicate that highly methylated transcripts in mature neuron populations have temporal and cell-type predominant expression patterns.

4. Discussion

Cytosine methylation of mRNA has emerged as a critical regulator of mRNA transportation, stability and translation [5,7,33,48,49]. Despite distinct mRNA methylation profiles reported in brain and NSCs [17], no attempt has been made to characterize mRNA m⁵C methylation patterns in the developing mammalian brain. In this study, we provide both gene expression and mRNA cytosine methylation profiles for NSCs, mature neurons, and postnatal brain tissues.

RNA cytosine methylation profiling can be achieved with m⁵C-specific antibodies to enrich methylated transcripts followed by deep-sequencing [50,51]. Such an approach has resolution limitations and is highly dependent on the quality of antibody used. Despite the fact that bisulfite sequencing remains the gold standard for RNA cytosine methylation detection, our recent experiments [27], together with reports from other labs [8,52], indicated that both experimental procedure and methylation calling have a significant impact on the detection of RNA methylation. Thus, inconsistent methylation sites may be observed between biological replicates. In this study, we found that high-confidence m⁵C sites shared by two biological replicates tend to have a higher methylation level and more read coverage than those of low-confidence m⁵C sites identified in one biological replicate

only. In addition, low-confidence sites frequently reside in a transcript sensitive to bisulfite treatment.

Notwithstanding current technical challenges in the determination of RNA cytosine methylation, our analyses on high-confidence m⁵C sites led to a few interesting findings. Consistent with previous studies [7,8,17], methylated cytosines are preferentially enriched around the translation initiation sites of mRNAs and usually have a methylation level between 20 and 30% with a downstream “GGG” motif. Approximately 6% of high-confidence m⁵C sites identified in this study were methylated across all samples. However, the diversity of mRNA methylation increases in differentiated neurons. In addition, increased methylation in neurons were observed in 96% of the DMSs identified between NSCs vs neurons. Previous studies linked the methylation of both tRNA and mRNA to synapse formation [13,15]. Intriguingly, the DMSs identified in P17 vs P0 and 6 W brain tissues were significantly enriched for synaptic plasticity. This suggests that mRNA cytosine methylation may also play important roles during brain development.

In this study, we examined the expression profiles of RNA cytosine methylation readers, writers, and erasers. As a result of tRNA degradation, neuronal synapse function was impaired in the NSUN2 knockout mouse model [16]. Interestingly, most of the factors regulating m⁵C methylation were downregulated during development in the comparison between P17 and P0 but showed no significant change in the comparison between 6 W and P17. This result suggests the highly dynamic regulation of mRNA m⁵C methylation during early postnatal brain development, and it may reach homeostasis in the maturing brain. The integration of RNA-seq data with RNA BS-seq data allowed us to explore the correlation between the levels of gene expression and RNA methylation. A weak to moderate positive correlation was observed in general, but such a trend may not be true for all transcripts. This suggests that the regulation of RNA expression and methylation might be positively correlated and the methylated transcripts could be cell-type predominant, spatially, and/or temporally regulated. Future studies are needed to explore how cell-type predominant mRNA methylation may contribute to neuronal differentiation. Collectively, our study provided insight into the RNA m⁵C methylation dynamics of the developing brain and is a rich resource to facilitate further investigation on the role of brain mRNA cytosine methylation.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Data availability

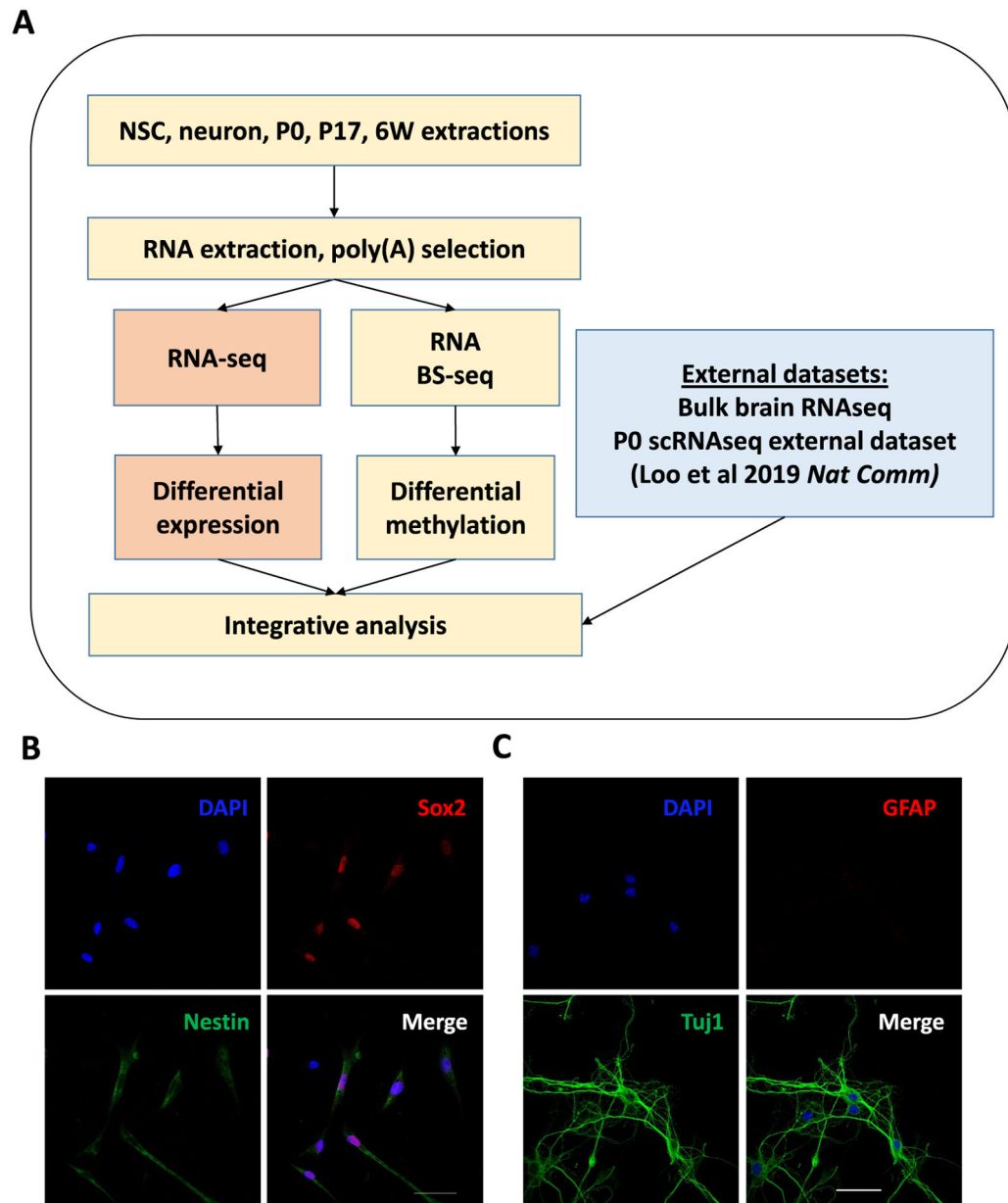
Data will be made available on request.

References

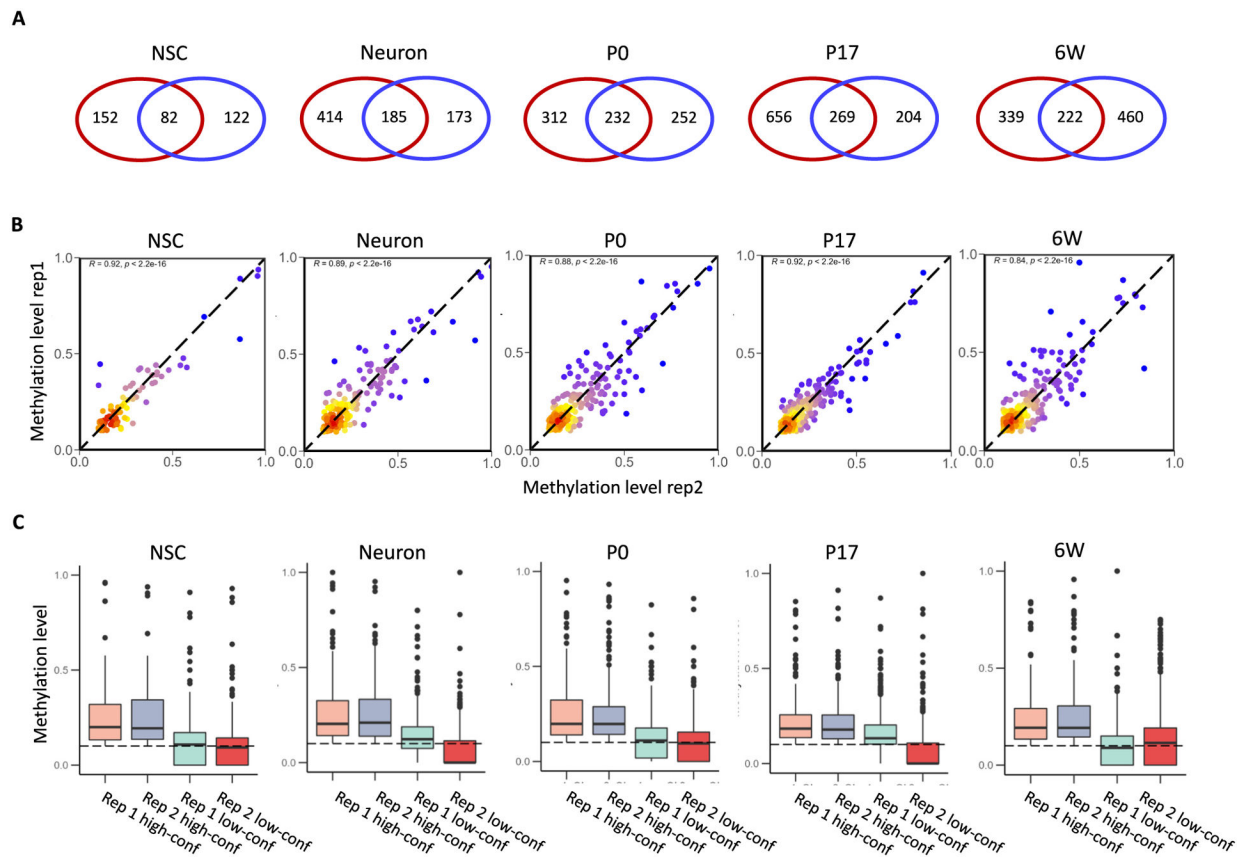
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**Fig. 1.**

Experimental Design. (A) Overall experimental design of this study. (B) Mouse neural stem cells (NSCs) were double stained with the neural progenitor markers Nestin (cytoplasmic, green) and Sox2 (nuclear, red); nuclei were counterstained with DAPI (blue). Scale bar: 50 μ m. (C) E16.5 mouse cortical neuronal culture was double stained with the neuronal marker Tuj1 (green) and glial marker GFAP (red); nuclei were counterstained with DAPI (blue). Scale bar: 50 μ m.

**Fig. 2.**

Reproducibility of replicates among RNA BS-seq samples. (A) Overlap of m⁵C sites identified between biological replicates. The red circle indicates replicate 1 and the blue circle indicates replicate 2. (B) Methylation correlation (Spearman's R) of the overlapping m⁵C sites between biological replicates. Color code indicates the density of sites, ranging from red (high density) to blue (low density). (C) Distribution of methylation levels for the overlapping and non-overlapping m⁵C sites identified between biological replicates.

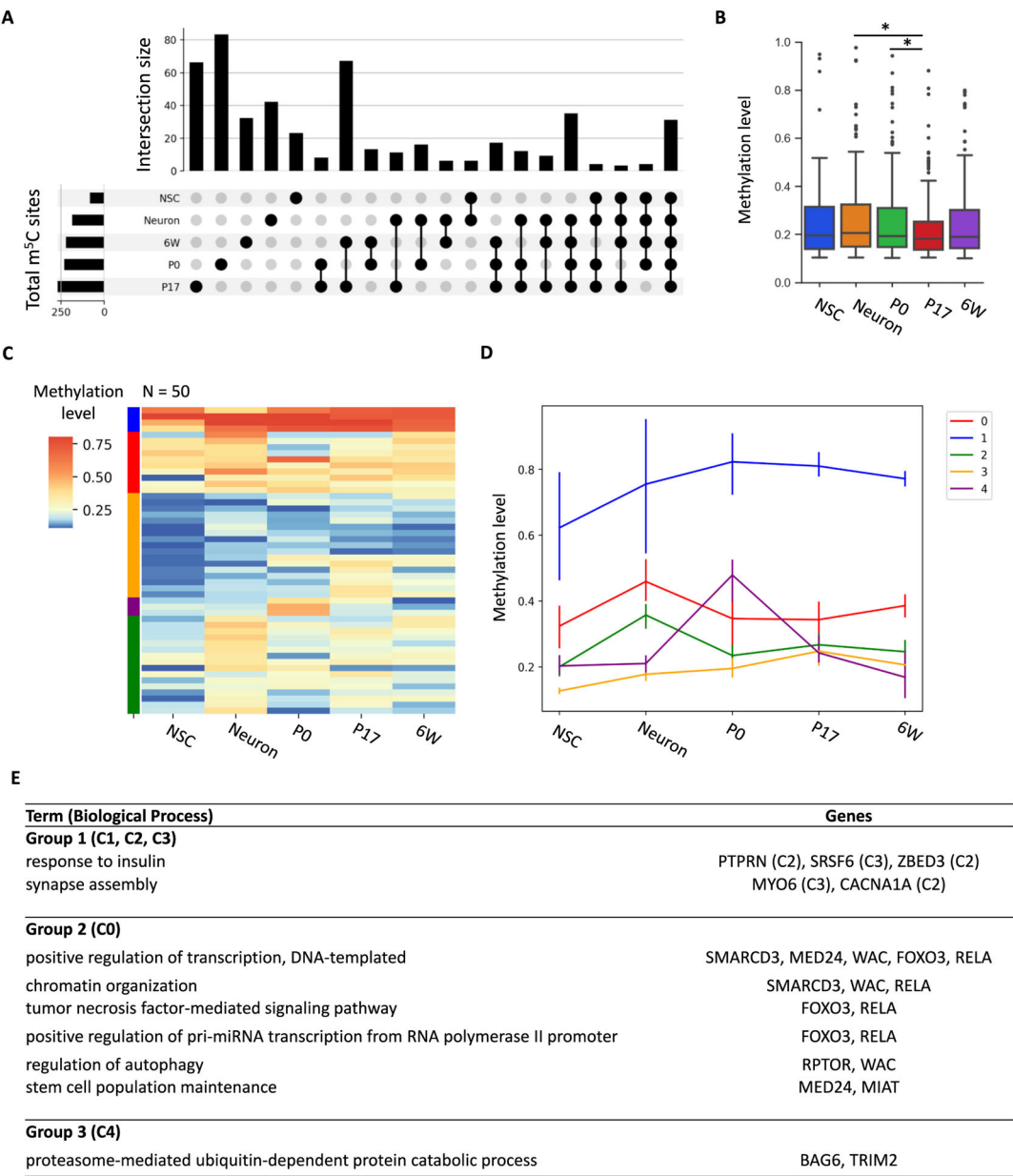
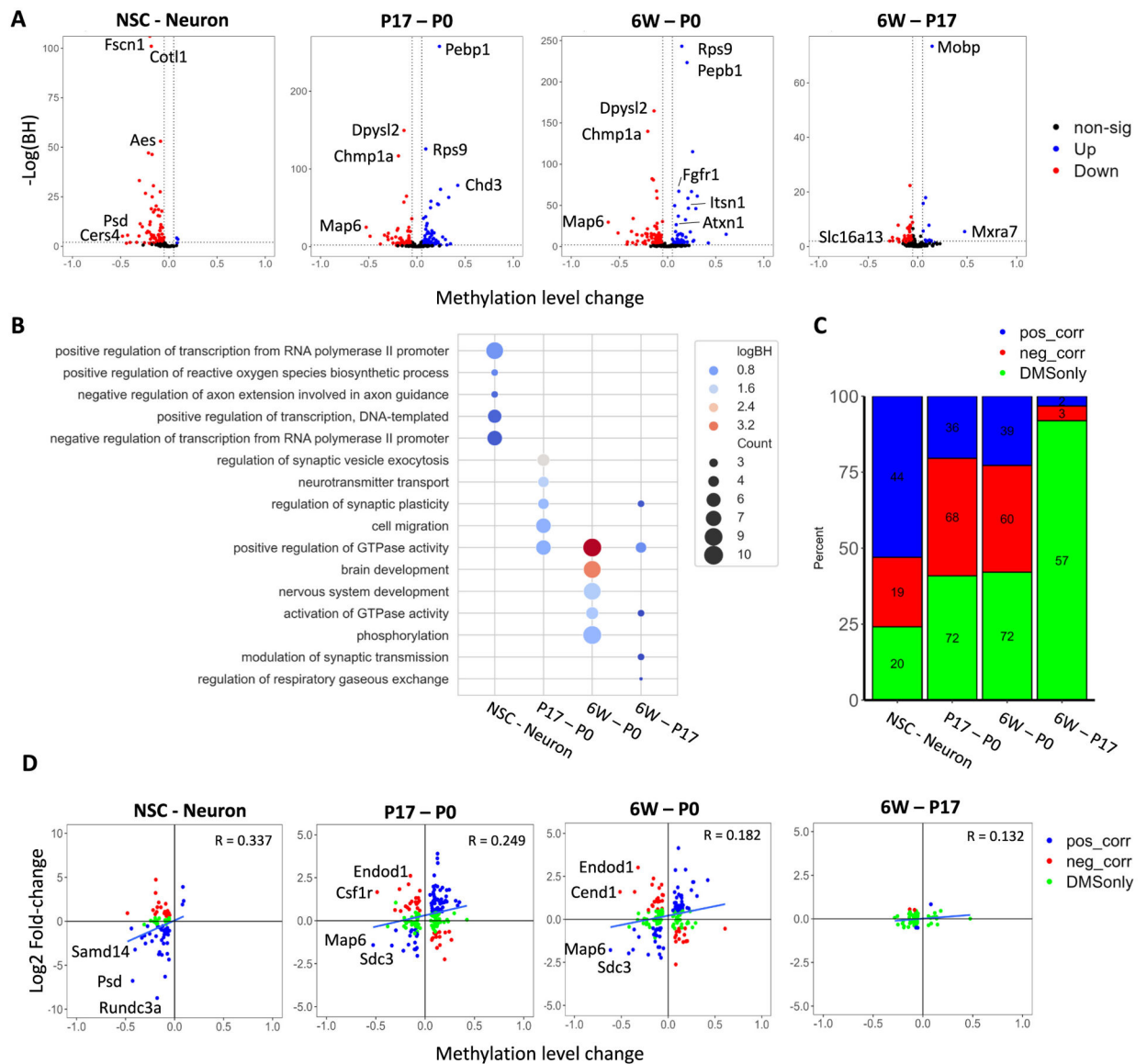


Fig. 3. Characteristics of high-confidence m⁵C sites identified in NSCs, neuron, and brain samples. (A) High-confidence m⁵C sites shared among conditions. (B) Methylation distribution of the high-confidence m⁵C sites. Wilcoxon rank-sum test was performed to assess significance (**p* < 0.05). (C) Clustering of m⁵C sites according to methylation changes. (D) Average methylation levels of clusters identified in Fig. 3C. Bars indicate standard deviation. (C) and (D) share the same color code. (E) Gene ontology analysis of methylated transcripts.

**Fig. 4.**

Methylation dynamics throughout brain development. (A) Differential methylation analysis of neuron and brain samples. Blue and red points represent significantly hyper and hypo methylated transcripts, respectively. Significance was determined using Fisher exact test followed by Benjamini-Hochberg correction for multiple comparison ($p < 0.05$), requiring a minimum difference of 0.05 in methylation. (B) Gene Ontology analysis of differentially methylated transcripts identified in (A). (C) Correlation analysis of changes in RNA expression and methylation for transcripts carrying differentially methylated sites. Blue and red bars represent positive and negative correlation between methylation level difference and expression fold-change, respectively. Green bars represent transcripts without expression fold-changes above 2 and p -adjusted < 0.01 . (D) Dot plot representation of transcripts to indicate the changes in methylation and expression. Overall correlation was determined using Spearman's R . Transcripts with methylation difference over 0.3 are noted.

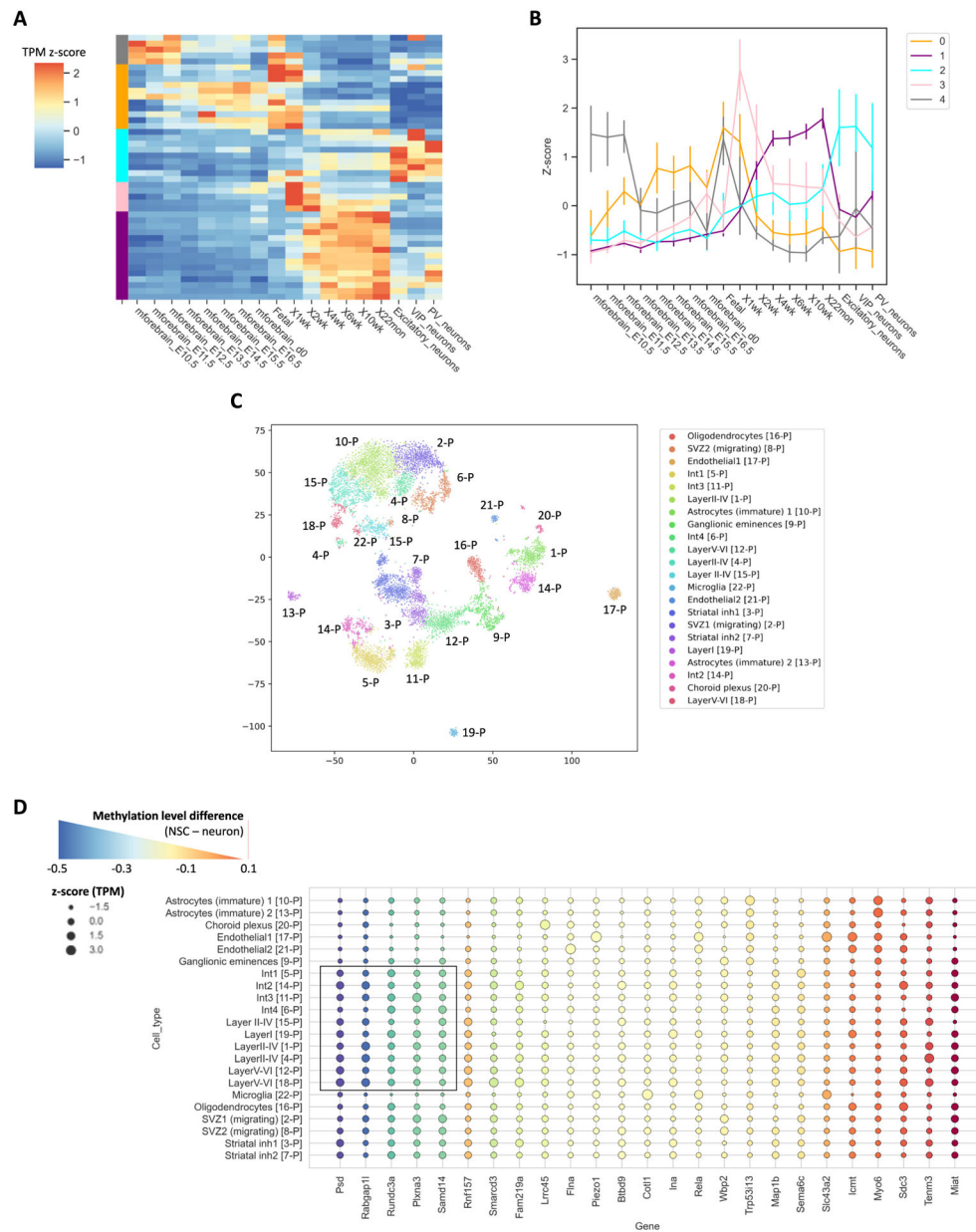


Fig. 6. Temporal and spatial expression of differentially methylated transcripts. (A) Clustering analysis of expression profiles for differentially methylated transcripts. Z-score of TPM was used to perform clustering. (B) Linear representation of clusters identified in (A). Error bars represent standard deviation. (A) and (B) share the same color code. (C) tSNE clustering of scRNAseq P0 brain tissue from Loo et al., 2019 [22]. p is denoted as postnatal. (D) TPM z-score of differentially methylated transcripts using scRNA-seq data generated for P0 brain. Color of bubble plot indicates methylation level differences between NSC and neuron RNA BS-seq samples.