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tRNA epitranscriptomic alterations associated with opioid-induced reward-seeking and long-term opioid withdrawal in male mice

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DNA cytosine methylation has been documented as a potential epigenetic mechanism of transcriptional regulation underlying opioid use disorder. However, methylation of RNA cytosine residues, which would drive another level of biological influence as an epitranscriptomic mechanism of gene and protein regulation has not been studied in the context of addiction. Here, we probed whether chronic morphine exposure could alter tRNA cytosine methylation (m^5C) and resulting expression levels in the medial prefrontal cortex (mPFC), a brain region crucial for reward processing and executive function that exhibits opioid-induced molecular restructuring. We identified dynamic changes in glycine tRNA ($tRNA^{Gly}_{GCC}$) cytosine methylation, corresponding to altered expression levels of this tRNA at multiple timepoints following 15 days of daily morphine. Additionally, a robust increase in methylation, coupled with decreased expression, was present after 30 days of withdrawal, suggesting that repeated opioid administration produces changes to the tRNA regulome long after discontinuation. Furthermore, forebrain-wide knockout of neuronal *Nsun2*, a tRNA methyltransferase, was associated with disruption of opioid conditioned place preference, and this effect was recapitulated by regional mPFC *Nsun2* knockout. Taken together, these studies provide a foundational link between the regulation of tRNA cytosine methylation and opioid reward and highlight the tRNA machinery as a potential therapeutic target in addiction.

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INTRODUCTION

Opioid use disorder (OUD) is a major continuing and worsening public health issue in the U.S., with a staggering 5.6 million people over the age of 12 diagnosed with OUD in 2021 alone [1] and contributing to over 70,000 deaths per year due to opioid overdose [2]. While a large segment of the population continues to receive opioids for pain management, with the potential for misuse and addiction, it is crucial to identify preventative measures or therapeutic approaches for OUD. Many studies suggest that molecular changes to the brain's reward circuit promote long-term functional regulation that underlies aberrant behavior in addiction [3]. Considerable work has been performed to identify patterns of transcriptional regulation in the reward circuit to identify biological domains affected by addictive drugs. However, most transcriptional changes are short-lived. Epigenetic mechanisms are thought to stamp in longer-term transcriptional regulation that drives persistent functional changes and behavioral patterns characteristic of addiction. Many studies have identified unique epigenetic transcriptional regulatory mechanisms in substance use disorders [4–6], although much of this work

has been focused on psychostimulants such as cocaine with less emphasis on opioids [7]. Further, while most examined canonical histone marks and DNA methylation, very little attention has been paid to other forms of epigenetic regulation, specifically the role of epitranscriptomics.

Epitranscriptomics has emerged as a novel field bridging RNA expression and protein synthesis in neuronal function to outline how chemical modifications to RNA can dynamically impact translational processes [8]. These modifications are persistent, lending the ability to induce long-term dysregulation of neurobiological function which may serve as a largely unexplored mechanism mediating addiction vulnerability. While some studies have investigated drug-induced changes in m^6A , a modification most prominent on mRNA [9, 10], no studies have explored the role of m^5C , a chemical modification most prominent on transfer RNAs (tRNAs), in addiction. tRNAs are integral for protein synthesis, a process altered by opioid exposure [11–13], and m^5C promotes tRNA stability in many mammalian tissues, including brain [14–17]. Surprisingly, early studies of morphine administration showed changes to other realms of tRNA functioning outside of the

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epitranscriptome (specifically, tRNA synthetase activity and amino acid binding to tRNAs) [18] and our recent study identified numerous changes in expression of tRNA synthetase genes throughout the brain reward system following heroin self-administration in the mouse [4]. Changes to other aspects of tRNA function have also been explored in brain and peripheral tissues, including tRNA fragment abundance following exposure to opioids [19] or other addictive drugs [20, 21]. Taken together, these studies point to a potential role for changes in tRNA function in addiction, including OUD.

Our laboratory has shown that altering the tRNA epitranscriptome through deletion of the m⁵C writer protein NSUN2 in neurons alters neuroplasticity and behaviors related to neuropsychiatric disease via glycine-specific changes in tRNA expression, amino acid abundance, and translation efficiency [22]. Notably, many of these changes overlap with processes relevant to addiction, including memory and synaptic plasticity. Here, we probed whether changes to the tRNA epitranscriptome and the functionality of NSUN2 within the medial prefrontal cortex (mPFC), a brain region heavily implicated in opioid reward-processing and addiction pathogenesis [23, 24], contribute to addiction-relevant behavior, and whether the observed opioid-induced changes persist through extended abstinence. To do this, we measured tRNA expression and m⁵C levels at previously identified candidate tRNAs [22] in mPFC tissue from wild-type mice that received chronic morphine injections followed by a brief or extended withdrawal period. Further, we investigated opioid preference in the morphine conditioned place preference test following knock-out of neuronal NSUN2. Our findings point to the tRNA epitranscriptome as a novel candidate for molecular mechanisms driving OUD and open a new line of research targeting tRNA machinery as a potential therapeutic target in addiction.

METHODS

Mice

All animal work was approved by the Institutional Animal Care and Use Committee of the Icahn School of Medicine at Mount Sinai. Mice were group-housed 2–5/cage with ad libitum access to food and water and a 12 h light/dark cycle (lights off at 7 pm) under constant conditions (21 ± 1 °C; 60% humidity). Mice bred in-house were weaned at ~P28, housed with same-sex littermates, and ear-tagged/genotyped. Male mice were used for the current study.

Nsun2 conditional knockout mice

Nsun2 conditional knockout mice included forebrain *Nsun2* knockout (KO) (*Camk2a-Cre* +, *Nsun2*^{2lox/2lox} vs. wildtype controls (*Camk2a-Cre*-) and *Nsun2*^{2lox/2lox} mice injected with either AAV8^{hSyn1-CreGFP} vs. AAV8^{hSyn1-GFP} as controls. Both conditional knockout models were previously validated in Blaze et al. [22] and the *Camk2a-Cre* transgenic line used has no differences in organization of brain nuclei or cortical layers, and display no overt behavioral changes before recombination [25]. Mice with the mutant allele have exon 6 of *Nsun2* flanked by loxP sites, and presence of Cre-recombinase causes excision of exon 6 and a frame-shift resulting in early termination of *Nsun2* translation. *Nsun2*^{2lox/2lox} mice (2 copies of the mutant allele) were crossed with a *Camk2a-Cre*+ line to produce *Camk2a-Cre*+, *Nsun2*^{2lox/2lox} mice for knockout of *Nsun2* in forebrain neurons, which is associated with widespread neuronal Cre-mediated deletion across the forebrain by the third postnatal week [26–29]. Age-matched *Camk2a-Cre*-, *Nsun2*^{2lox/2lox} or *Nsun2*^{2lox/-} were used as wild-type controls.

Viral microinjections

For viral microinjection surgeries as described previously [22], we injected virus into adult male mouse mPFC (AAV8^{hSyn1-CreGFP} and AAV8^{hSyn1-GFP}; see Fig. 1a and Supplementary Fig. 2c for schematic of viral microinjection location and overlap with mPFC region of interest for rest of study), 10–12 week-old *Nsun2*^{2lox/2lox} mice were anesthetized with isoflurane and 1 µL of virus per hemisphere (bilateral injection) was injected at a rate of 0.25 µL/min using a Hamilton syringe (Reno, NV), a micropump (Stoelting) and a stereotactic frame (Stoelting). Stereotactic coordinates for mPFC injection

were as follows: 1.5 mm anterior/ posterior, ±0.5 mm medial/lateral and 1.5 mm dorsal/ventral. Control mice received 1 µl per hemisphere of AAV8^{hSyn1-GFP} using the same conditions. To confirm expression and location of viral CreGFP expression, we performed immunohistochemistry as previously described (Supplementary Fig. 2c) [22]. Imaging was performed using a Zeiss CLSM780 upright microscope and images were processed in NIH ImageJ software.

Morphine treatment

Morphine sulfate (from the National Institute on Drug Abuse) was diluted in saline and injected intraperitoneally at 10 mg/kg. For withdrawal studies, mice remained in the home cage for 2 h, 24 h, or 30 days after the last injection until sacrifice, at which point brains were rapidly extracted, sectioned (1 mm thick) using a brain matrix on ice, and mPFC was dissected using a 12 G punch and flash frozen on dry ice.

tRNA bisulfite sequencing

The tRNA amplicon bisulfite sequencing protocol was performed as described in Blaze et al. [22], and previously adapted from Bormann et al. [30]. mPFC punches (see Fig. 1a) were homogenized in Trizol reagent (Invitrogen) and RNA was extracted, purified, and quantified using the Qubit fluorometer. 500 ng of total RNA was used for bisulfite conversion with the EZ RNA Methylation kit according to manufacturer's instructions (Zymo Research) but with an additional 2 PCR cycles to denature tRNAs completely, as previously reported [15]. cDNA synthesis was performed on bisulfite-treated RNA using the reverse primer for each of 3 tRNA isodecoders (i.e., *GlyGCC-1* [22, 31], *GluCTC-1* [22], *AspGTC-1* [22, 30, 31]) followed by PCR amplification with the tRNA-specific forward primer. Amplicons were size-verified on a 4% agarose gel and PCR amplicons were extracted and purified (Qiaquick Gel Extraction Kit, Qiagen) before indexing (Nextera XT, Illumina), and pooled up to 32 amplicons in equimolar concentrations before running with 75 bp paired-end reads on the MiSeq (Illumina).

Data analysis. Bisulfite sequencing data was analyzed using BisAmp [30], a publicly available web server for quantification of methylated reads from Miseq output fastq files. tRNA bisulfite sequencing data from bisAMP (methylation percentages for each cytosine) was further analyzed by conducting a t-test for each cytosine with FDR correction using the two-stage linear step-up procedure of Benjamini, Krieger and Yekutieli.

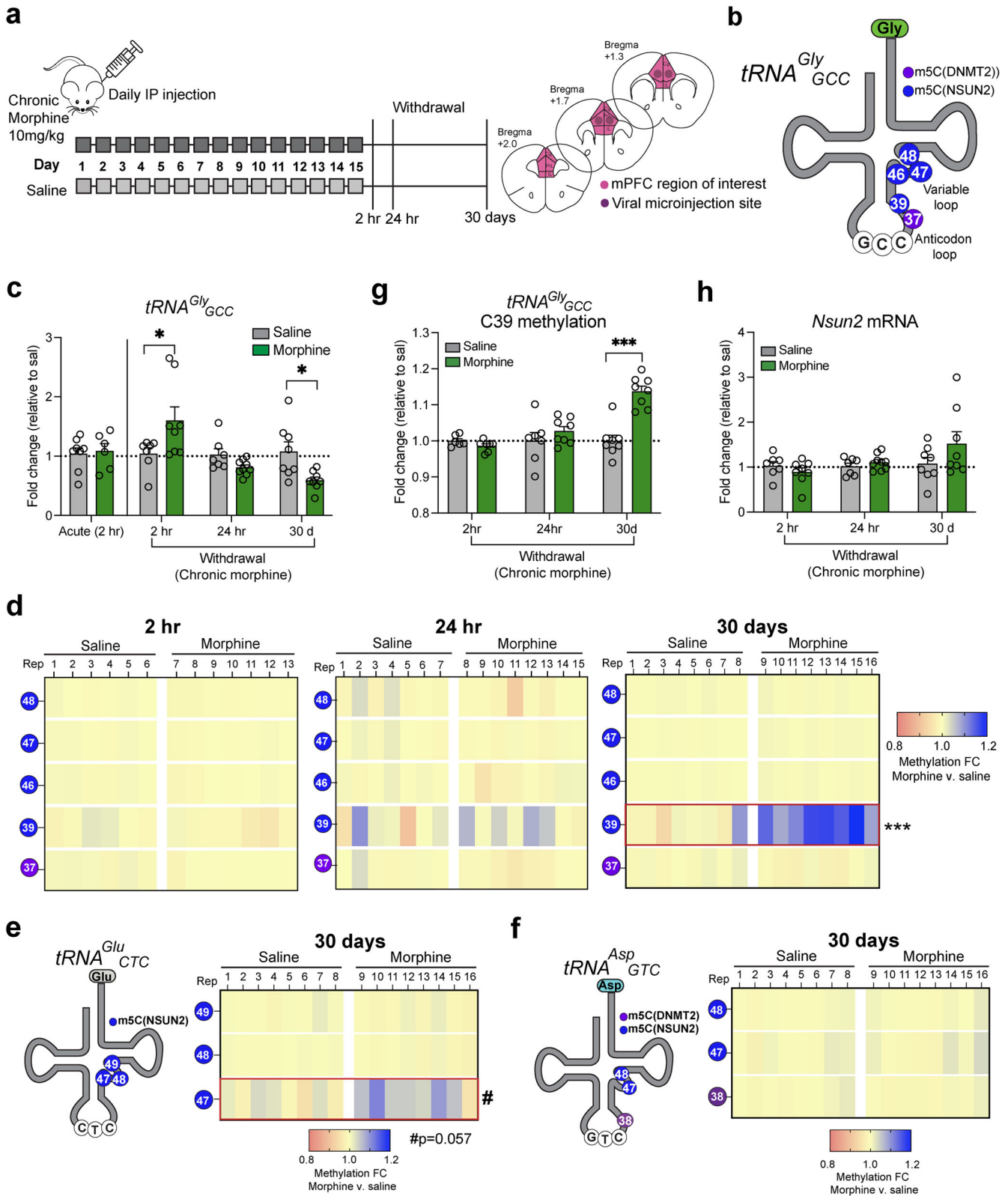
YAMAT/UMI seq

Y-shaped Adapter MAture tRNA (YAMAT) sequencing was performed as previously described [22] and was adapted from Shigematsu and colleagues [32] to include unique molecular identifiers (UMIs) in order to avoid overamplification of individual tRNA isodecoders. Briefly, 5 µg total RNA was isolated from frontal cortex of morphine- or saline-treated mice. RNA was then deacylated to remove amino acids from 3' ends and demethylated to remove m1A, m1C, and m3C modifications using a proprietary demethylation mix (Arraystar, Inc). 40 µm of YAMAT forked linkers were then incubated with pure demethylated and deacetylated RNA followed by addition of 10X annealing buffer (50 mM Tris HCl pH 8, 100 mM MgCl₂, 5 mM EDTA) and then overnight incubation with T4 RNA ligase 2. Linker-ligated RNA was then incubated with RT Primer (TruSeq Small RNA Library Prep Kit, Illumina) and reverse transcription was performed with Superscript III RT (Invitrogen) followed by bead purification. Libraries were amplified using Phusion Hotstart II Polymerase (Thermo Scientific) with primers and indexes from the TruSeq Small RNA kit (Illumina) for 16 PCR cycles. Amplified libraries were then bead-purified and run on the Agilent Bioanalyzer for confirmation of library size and quantified with the Qubit Fluorometer (Invitrogen). 8 libraries (*n* = 4 morphine, 4 saline) were pooled at equimolar concentrations and run with 75 bp paired-end reads on the MiSeq (Illumina).

Data analysis. Raw sequencing reads were processed using cutadapt to trim adapter sequences and UMItools to extract UMIs from reads and perform deduplication. Paired end reads were merged using Pear and reads were filtered before differential expression analysis was performed using Deseq2, with significant differences in morphine vs. saline tRNA expression denoted at FDR-adjusted *p* < 0.05.

Real-time qPCR

For real-time qPCR to identify changes in *Nsun2* mRNA expression, we used 1 µg total RNA to generate cDNA followed by Taqman qPCR using Taqman



Universal Master Mix and Taqman probes for *Nsun2* (Assay ID: Mm01349532_m1) and *Gapdh* as a control gene (Assay ID: Mm99999915_g1; Applied Biosystems). For qPCR to detect change in tRNA expression, we used the cDNA described above and pre-validated PCR primer sets specific to the mature tRNA sequence for *GlyGCC-1* or *5S*

tRNA as a housekeeping gene (Arraystar, Inc.) using SYBR-based qPCR. Data were analyzed using the comparative Ct method and normalized to the housekeeping gene and saline-injected controls. An unpaired t-test was used to compare treatment groups and significance was denoted at $p < 0.05$.

Fig. 1 Dynamic tRNA^{Gly} expression and methylation following acute morphine and withdrawal in wild-type mice. **a** Adult wild-type male mice were injected IP with 10 mg/kg morphine daily for 15 days, followed by 2 h, 24 h, or 30 days of withdrawal and collection of medial prefrontal cortex (mPFC) tissue. See overlap of mPFC region of interest and viral Cre injections used in behavioral experiments. **b** Schematic of tRNA^{Gly}_{GCC} depicting methylated cytosines. **c** qPCR for tRNA^{Gly}_{GCC} expression after a single acute morphine injection ($n = 8$ saline/6 morphine) or morphine withdrawal (2 h: $n = 7$ saline/8 morphine, 24 h: $n = 7$ saline/9 morphine, 30d: $n = 8$ saline/8 morphine; $*p < 0.05$). **d** Bisulfite amplicon sequencing for tRNA^{Gly}_{GCC} after 2 h, 24 h, and 30 days of withdrawal. Note the increase in methylation for morphine-treated mice at cytosine 39 at 30 days (t-test; $***p < 0.001$). **e** Left, schematic representation of tRNA^{Gly}_{CTC} and cytosines methylated by Nsun2. Right, Bisulfite amplicon sequencing for tRNA^{Gly}_{CTC} showed a trending increase in methylation for morphine-treated mice at cytosine 47 (t-test $\#p < 0.057$; $n = 8$ /group). **f** Left, schematic representation of tRNA^{Asp}_{GTC} and cytosines methylated by NSUN2 and DNMT2. Right, Bisulfite amplicon sequencing for tRNA^{Asp}_{GTC} showed no changes in methylation at any sites between saline and morphine-treated mice ($n = 8$ /group). **g** Cytosine 39 methylation at each timepoint shown as a fold change relative to the average of each group's saline control ($***p < 0.001$). **h** qPCR for Nsun2 mRNA levels at all 3 withdrawal timepoints.

Behavior

Morphine conditioned place preference (CPP). CPP was carried out using an unbiased design in three-chamber CPP boxes with overhead monitoring captured by a camera and analyzed using Ethovision (Noldus) to track the animal's location within the box. The two end chambers differed in visual (stripe vs. gray) and tactile (small grid flooring vs. large grid flooring) cues to create distinct contexts. Dim white overhead light was used throughout testing. In the first session (pretraining), mice were placed in the CPP chamber and allowed to explore all three compartments freely for 20 min. Pairing sides were then determined for each mouse such that the group average of time spent was not biased for either chamber. Subsequently, mice received three days of morphine-context conditioning. On each conditioning day, mice were treated with saline and confined to one chamber for 45 min in the morning, and, in the afternoon, received morphine (10 mg/kg IP) and were confined to the other chamber for 45 min. On the fifth day, CPP was assessed by placing mice in the chamber and allowing them to explore freely. A place preference score was generated by subtracting the time the animal spent in the morphine-paired chamber vs. time spent in the saline-paired chamber. Locomotor activity measurements presented in Supplementary Fig. S2 were obtained as distance traveled measures (in cm) in the CPP arena during pretest or posttest phases of the conditioning procedure.

Statistical analyses

Statistical analyses were performed using Graphpad Prism 8.4.3 software. For tRNA/mRNA expression and tRNA methylation data with multiple timepoints, we used a two-way ANOVA (timepoint \times condition) followed by Bonferroni's posthoc test when appropriate. For comparisons with one timepoint and two conditions, we used unpaired two-tailed t-tests, and FDR-correction was used for tRNA bisulfite amplicon sequencing. We used simple linear regression for correlation analysis. Statistical significance was denoted by $p < 0.05$. All bar graphs are presented as the mean with error bars representing standard error of the mean (SEM) and include all individual data points.

RESULTS

mPFC tRNA^{Gly} expression is dynamically altered during 30 days of morphine withdrawal

To assess whether a clinically relevant model of opioid addiction and withdrawal can alter tRNA expression and tRNA cytosine methylation in the rodent brain, wild-type adult male mice were injected intraperitoneally (IP) acutely (single dose) or daily for 15 consecutive days with 10 mg/kg morphine (or saline as control) and left undisturbed until sacrifice at one of three timepoints post-injection: 2 h, 24 h, or 30 days (Fig. 1a). The 30-day timepoint was chosen to coincide with a period of abstinence known to promote high levels of craving that drive relapse [33], which we have shown to promote robust transcriptional regulation throughout the reward circuit [4]. In this opioid treatment procedure, 10 mg/kg daily morphine has been shown to promote psychomotor sensitization and other reward- and addiction-related behavior, and many of these phenotypic changes have been linked to the mPFC, a crucial part of the well-known addiction circuitry in rodent models and human subjects [23, 24]. At each timepoint, mPFC, including infralimbic and prelimbic cortex (refer to Fig. 1a), was collected, and RNA was extracted to query tRNA levels and

tRNA cytosine methylation following acute or chronic morphine treatment or withdrawal.

As a starting point, we chose to quantify tRNA^{Gly} levels because this tRNA family is highly sensitive to alterations in tRNA cytosine methylation. This sensitivity is likely due to the fact that brain is one of the organs with the highest levels of tRNA cytosine methylation, altogether encompassing 5 cytosine methylation sites [22], the most of any tRNA family and not seen for any of the other tRNAs assigned to the other 23 canonical amino acids. Specifically, of the four tRNA^{Gly} isoacceptors (ACC, CCC, GCC, TCC), tRNA^{Gly}_{GCC} (Fig. 1b) is the most highly expressed in brain and sensitive to disruptions in cytosine methylation [22]. Thus, we assessed tRNA^{Gly}_{GCC} expression in the mPFC of morphine- and saline-treated mice by qPCR. Indeed, there was a dynamic change in tRNA^{Gly}_{GCC} expression levels throughout the 30 day withdrawal period (2-way ANOVA treatment \times timepoint interaction; $F(3,53) = 6.118$, $p = 0.001$; acute: $n = 8$ saline/6 morphine, 2 h: $n = 7$ saline/8 morphine, 24 h: $n = 7$ saline/9 morphine, 30d: $n = 8$ saline/8 morphine), defined by a transient increase followed by a significant decline in the long-term. Specifically, tRNA^{Gly}_{GCC} was increased 1.6-fold in morphine-treated mice 2 hours after the last injection ($t(53) = 3.063$, $p = 0.014$; $n = 7$ saline/8 morphine) but showed no change at 24 hours ($t(53) = 1.215$, $p = 0.919$; $n = 7$ saline/9 morphine). Conversely, after 30 days of withdrawal, morphine-treated mice showed a 0.6-fold decrease in tRNA^{Gly}_{GCC} expression ($t(53) = 2.756$, $p = 0.032$; $n = 8$ saline/8 morphine) (Fig. 1c), suggesting a dynamic pattern of tRNA^{Gly}_{GCC} expression throughout the withdrawal period after repeated morphine exposures. In contrast, no alterations were observed after a single dose of morphine ($t(53) = 0.243$, $p > 0.999$; Fig. 1c), suggesting that dynamic changes in tRNA^{Gly}_{GCC} expression are driven by chronic opioid exposure.

Because tRNA^{Gly}_{GCC} showed a significant decrease in morphine-treated mice after 30 days of withdrawal in chronic morphine-exposed mice, we asked whether other tRNA^{Gly} isoacceptors show similar types of alterations. We quantified tRNA^{Gly}_{TCC} and observed a marginal, non-significant decrease in mPFC expression levels after 30 days of morphine withdrawal (Supplementary Fig. 1a, b; unpaired two-tailed t-test, $t(13) = 1.398$, $p = 0.186$; $n = 8$ saline, 7 morphine), suggesting that tRNA^{Gly} may have isoacceptor-specific regulation following repeated exposure to opioids and long-term withdrawal. Next, we examined whether these effects of morphine withdrawal on tRNA^{Gly} expression were specific to the infralimbic and prelimbic regions of the mPFC (which are firmly implicated in opioid addiction circuitry) or alternatively representative of changes broadly affecting the entire rostral cortex (Supplementary Fig. 1a, c, d). Therefore, we repeated the morphine exposure and 30-days withdrawal procedure in a different cohort of mice and dissected out the entire rostral cortex (see Supplementary Fig. 1c for exact location). There were no differences between treatment groups for tRNA^{Gly}_{GCC} levels by qPCR ($t(18) = 1.136$, $p = 0.271$; $n = 10$ saline/10 morphine) or by YAMATseq (Supplementary Fig. 1e; $n = 4$ saline/4 morphine), an unbiased screen of the tRNAome using next-generation sequencing (see Methods).

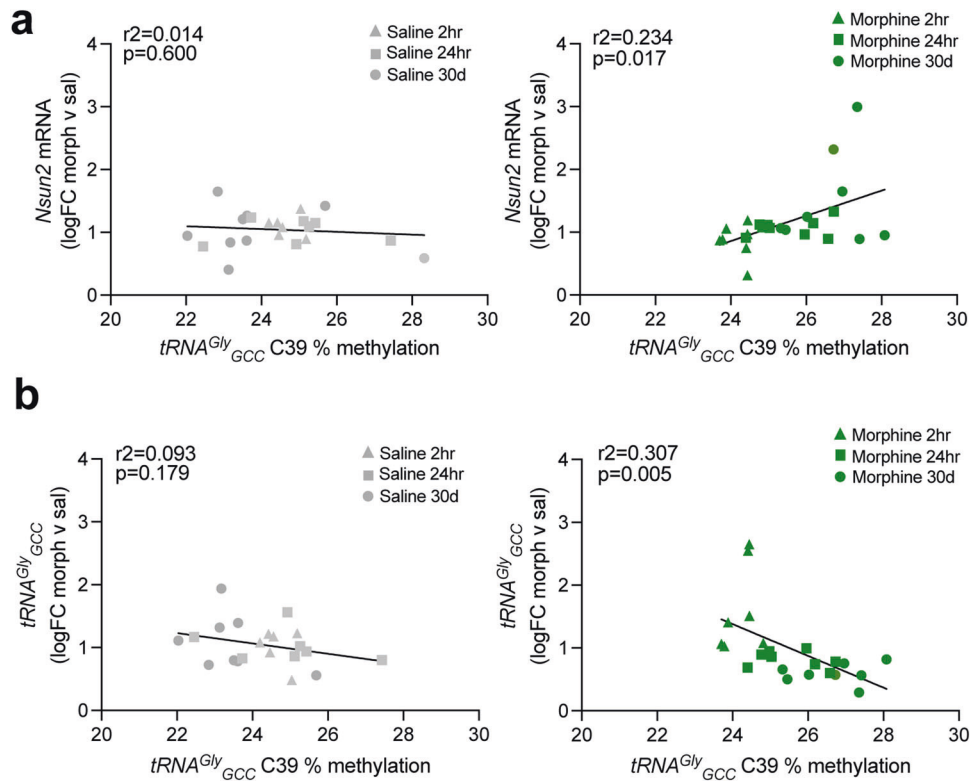


Fig. 2 Dynamic changes in morphine-induced tRNA hypomethylation are associated with shifts in *Nsun2* and tRNA abundance. Correlations between tRNA^{Gly}_{GCC} methylation and (a) *Nsun2* mRNA levels and (b) tRNA^{Gly}_{GCC} expression levels. Left, saline groups show no significant correlation or anticorrelation, while the morphine groups (right) show significant relationships between expression levels of tRNA and *Nsun2* and C39 methylation.

Notably, the input required for YAMATseq was prohibitive for conducting this analysis on mPFC specifically. We conclude that morphine withdrawal-induced changes in tRNA^{Gly}_{GCC} are specific to the mPFC without affecting other areas of rostral cortex.

tRNA methylation in mPFC is increased following 30 days of morphine withdrawal

Having shown that withdrawal from opioids affects tRNA^{Gly}_{GCC} expression, we then wondered if a mechanism for these types of fluctuation in tRNA levels are associated with tRNA epitranscriptomic dysregulation, given that previous studies in peripheral cell lines have uncovered a strong link between tRNA expression levels and cytosine methylation changes following stress [14, 15]. Because we have previously shown decreased methylation is directly linked to decreased tRNA^{Gly}_{GCC} expression and our data here point to dynamic alterations in tRNA^{Gly}_{GCC} in mPFC after 2 h and 30 days of morphine withdrawal, we queried the methylation status of tRNA^{Gly}_{GCC} in mPFC of saline- and morphine-injected mice at all three morphine withdrawal timepoints. We used tRNA bisulfite sequencing with approximately 40,000 reads per sample to measure tRNA^{Gly}_{GCC} cytosine methylation levels in mPFC of mice that experienced morphine withdrawal (or saline) at 2 h, 24 h, and 30 days and identified a significant 1.15-fold increase in methylation only after 30 days for morphine-treated mice at cytosine 39 ($t(14) = 6.330$, FDR adj. $p < 0.0001$; $n = 8$ saline/8 morphine), while methylation at all other cytosines remained unaltered by morphine exposure (Fig. 1d; Supplementary Table 1). Notably, methylation at C39 (and other cytosines) were unchanged by morphine treatment at earlier timepoints of 2 h and 24 h into withdrawal, showing a temporal specificity for methylation changes that only appear after protracted abstinence (Fig. 1d; Supplementary Table 1). Because we saw a significant increase in tRNA^{Gly}_{GCC} methylation at 30 days, we also measured

cytosine methylation at other tRNA families that contain fewer cytosine methylation sites. First, tRNA^{Glu} contains 3 methylcytosines (Fig. 1e), and bisulfite sequencing of tRNA^{Glu}_{CTC} revealed a near significant methylation increase at cytosine 47 for morphine-treated mice after 30 days (Fig. 1e; $t(14) = 2.654$, FDR adj. $p = 0.057$; $n = 8$ saline/8 morphine; Supplementary Table 1). We also examined tRNA^{Asp}_{GTC} which likewise contains 3 methylcytosines and found no morphine-induced changes after 30 days of withdrawal (Fig. 1f; Supplementary Table 1).

In brain, tRNA cytosine methylation is mainly deposited by NSUN2, which methylates ~80% of tRNAs, while the other RNA methyltransferase, DNMT2, methylates a much smaller subset of tRNAs. Among the methylcytosines within tRNA^{Gly}_{GCC}, 4 out of 5 are methylated by NSUN2 (C39, C46-48, Fig. 1b), while one site is DNMT2-dependent (C37). Conversely, tRNA^{Glu}_{CTC} has only 3 NSUN2-dependent methylation sites (C47-49; Fig. 1e), and tRNA^{Asp}_{GTC} carries 2 NSUN2-mediated cytosines and 1 DNMT2-mediated cytosine (Fig. 1f). We noticed that the subset of cytosines altered after morphine withdrawal (tRNA^{Gly}_{GCC} C39 and tRNA^{Glu}_{CTC} C47) are both methylated by NSUN2, suggesting that tRNAs with more NSUN2-dependent cytosine methylation sites may be selectively vulnerable to changes associated with long-term withdrawal from opioids. Therefore, we quantified NSUN2 levels by qPCR at each withdrawal timepoint in mPFC and found a positive correlation between methylation levels at tRNA^{Gly}_{GCC} cytosine 39 (the only cytosine modified by morphine withdrawal) and *Nsun2* mRNA levels only for the morphine-treated mice (Fig. 2a; saline: $r^2 = 0.014$, $p = 0.600$, $n = 22$ mice, 7-8 per timepoint; morphine: $r^2 = 0.234$, $p = 0.017$, $n = 24$ mice, 7-8 per timepoint), thereby suggesting that dynamic changes in morphine-induced tRNA hypermethylation could, at least in part, be linked to corresponding adaptations in *Nsun2* expression. We did not detect significant changes in *Nsun2* mRNA levels between

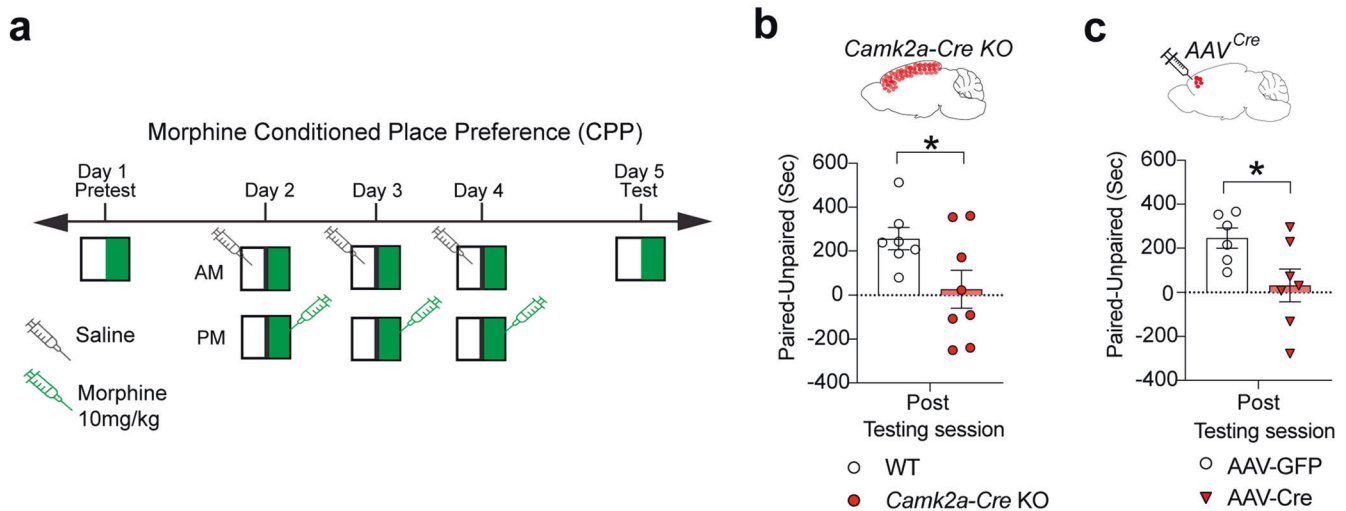


Fig. 3 Morphine conditioned place preference in *Nsun2* conditional knockout mice. **a** For CPP, mice received a 20-min pre-test wherein all compartments of the chamber could be explored. Subsequently, mice underwent three 45-min conditioning sessions in which saline was injected in the morning restricted to one side of the chamber (“unpaired”), and 10 mg/kg morphine was injected in the afternoon on the other side (“paired”). Conditioning to morphine was assessed in a subsequent 20-min session with access to both sides of the chamber, with positive conditioning indicated by a higher preference on the morphine-paired side. **b** *Camk2a-Cre* *Nsun2* KO mice ($n = 7$ WT, 8 KO) and **c** PFC AAV-Cre *Nsun2* KO ($n = 6$ AAV-GFP, 7 AAV-Cre KO; see Fig. 1a for AAV localization schematic) mice both show a reduction in time spent in the morphine-paired chamber compared to the unpaired chamber, indicating a blunted reward response to morphine (t-test $*p < 0.05$).

morphine and saline-treated mice in the mPFC at 2 h, 24 h, or 30 days (Fig. 1g, h; 2-way ANOVA, $n = 7$ -8 saline/8-9 morphine; all p 's > 0.05), however we did not measure NSUN2 protein levels directly in the current study. Additionally, tRNA^{Gly}_{GCC} cytosine 39 methylation levels negatively correlated with tRNA^{Gly}_{GCC} expression levels only in morphine-treated mice (Fig. 2b; $r^2 = 0.307$, $p = 0.005$, $n = 24$ mice, 7-8 per timepoint) while no correlation was observed in saline-treated mice (Fig. 2b; $r^2 = 0.093$, $p = 0.179$, $n = 24$ mice, 7-8 per timepoint). Taken together, our data showing increased tRNA^{Gly}_{GCC} cytosine methylation potentially driven by NSUN2 coupled with decreased tRNA^{Gly}_{GCC} expression suggest that long-term morphine withdrawal induces enduring changes in the mPFC tRNA regulome, including NSUN2-mediated tRNA cytosine methylation and expression.

Neuronal *Nsun2* deletion impairs opioid preference

Given our findings linking NSUN2-regulated tRNA^{Gly}_{GCC} methylation and expression following long-term morphine withdrawal in the mPFC, we then tested whether deletion of NSUN2 impacts addiction-related behavior. Early studies revealed alterations in protein synthesis in synaptosomal fractions following acute and chronic morphine exposure [11, 12], suggesting that protein translation in neurons, and potentially regulatory molecules such as tRNAs in neurons, are involved in opioid addiction phenotypes. Therefore, we generated mice with conditional neuron-specific knockout of *Nsun2* in forebrain (*Camk2a-Cre*⁺, *Nsun2*^{2lox/2lox} mutant mice) driven by *Ca*²⁺/*calmodulin-kinase II* (*CamK*)-*Cre* promoter, which produces widespread neuronal *Cre*-mediated deletion across the forebrain before postnatal day 18 as previously described. We conducted a standard morphine CPP procedure using the same dose as the aforementioned long-term morphine exposure procedure (Fig. 1a), which is a well-established task to measure reward seeking and has translational relevance to the drug-environment associations in human OUD [34]. Intact drug-cue interactions and reward seeking in CPP require expression of addiction-related proteins, but it is unknown whether dysfunction of the neuronal tRNA regulome is involved in CPP performance. To test this, *Nsun2 Camk2a-Cre* conditional knockout (cKO) mice and wild-type controls underwent a traditional morphine CPP task (see Methods; Fig. 3a). *Nsun2 Camk2a-Cre* cKO mice spent significantly

less time in the morphine-paired side compared to wild-type controls (unpaired t-test, post-test: $t(13) = 2.194$, $p = 0.047$; pre-test: $t(13) = 0.506$, $p = 0.621$; $n = 7$ WT, 8 KO; Fig. 3b, Supplementary Fig. 2a). Interestingly, *Nsun2 Camk2a-Cre* cKO mice exhibited heightened locomotor activity in the pre-test phase of the CPP (Supplementary Fig. 2b; pre-test: $t(13) = 3.968$, $p = 0.002$) which was no longer present during the post-test phase (post-test: $t(13) = 0.784$, $p = 0.447$). These results suggest that deficits in CPP cannot be explained by generalized reductions in locomotor activity but do indicate a potential sensitivity of *Nsun2 Camk2a-Cre* cKO to novel stimuli or environments.

Because forebrain deletion of *Nsun2* may affect numerous psychological and behavioral processes that broadly impair function, we examined whether the observed impairment in opioid preference after *Nsun2* knockout could be recapitulated by specifically targeting the mPFC. To test this, we conducted the same type of CPP experiment in 12–16-week-old conditional *Nsun2*^{2lox/2lox} mice that received a highly localized bilateral mPFC microinjection of AAV8^{hSyn1-CreGFP} (AAV-Cre), or of AAV8^{hSyn1-GFP} as a control (AAV-GFP) for region- and cell-type-specific knockout of neuronal *Nsun2* (see Fig. 1a for overlap with mPFC region of interest used in molecular studies, Supplementary Fig. 2c). Previous studies have shown that this area of mPFC is essential for opioid-induced reward behavior but not for contextual memory. Deletion of *Nsun2* in adult neurons of the PFC produced a similar impairment in morphine CPP (AAV-GFP vs. AAV-Cre unpaired t-test, post-test: $t(11) = 2.337$, $p = 0.039$, pre-test: $t(11) = 0.258$, $p = 0.801$; $n = 6$ AAV-GFP, 7 AAV-Cre; Fig. 3c, Supplementary Fig. 2d), while having no impact on basal locomotor activity (Supplementary Fig. 2e; pre-test: $t(11) = 0.247$, $p = 0.810$, post-test: $t(11) = 0.114$, $p = 0.911$). These results are consistent with the role of prelimbic cortex in mediating the expression of context-dependent drug-seeking [35, 36]. Together, these findings suggest that mPFC *Nsun2* expression, and possibly NSUN2-mediated tRNA methylation, in this discrete brain region is necessary for opioid preference.

DISCUSSION

We show here that chronic morphine exposure elicits dynamic changes in expression of tRNA^{Gly}_{GCC} in the rodent mPFC,

characterized by an early increase in expression and ultimately leading to a decrease in expression in morphine-treated mice after 30 days of withdrawal. Furthermore, changes in expression are tightly linked to site-specific NSUN2-mediated cytosine methylation levels at tRNA^{Gly}_{GCC}. We also demonstrate that loss of neuronal NSUN2 in forebrain or more locally within a discrete area of the mPFC impairs opioid preference, suggesting the importance of NSUN2, and potentially the m⁵C epitranscriptomic system, in regulating addiction phenotypes. As the first investigation of tRNA cytosine methylation in the context of addiction, we now provide a previously unexplored regulatory system, the tRNA epitranscriptome, as an important mediator of opioid action.

The epitranscriptome in addiction

There is a wealth of evidence that chronic exposure to opioids induces dynamic and long-term changes to transcriptional machinery via epigenetic modifications to chromatin in the brain's reward circuitry [4–7], but very few studies have investigated the epitranscriptome in relation to addiction. To our knowledge, the only investigation into CNS epitranscriptomic changes in relation to opioids focused on m⁶A, showing that chronic morphine administration altered m⁶A profiles in spinal cord for many circular RNAs (circRNAs), non-coding RNAs that are single-stranded and covalently bound in a closed loop, which may be a molecular change contributing to the effects of morphine tolerance [10]. The m⁶A epitranscriptome was also implicated in alcohol use disorder (AUD), with a small cohort of human postmortem brains showing AUD-specific patterns of methylation at various mRNAs, long non-coding RNAs, and miRNAs [9]. The current study investigates the m⁵C tRNA epitranscriptome in the context of addiction. Because tRNAs are crucial components of protein translation in the ribosome, and many studies have pointed to protein synthesis abnormalities in the brains of morphine-exposed rodents [11–13], it is surprising that there has been such little investigation into tRNA regulatory mechanisms in opioid addiction.

Full length tRNAs and cleavage products altered by addictive drugs

While we profiled cytosine methylation and full-length tRNA expression, there are reports of drug-induced changes to tRNA fragments of varying sizes originating from various tRNA isoacceptor families. In the rat nucleus accumbens, tRNA fragments were altered following methamphetamine self-administration, including a decrease in the fragment originating from the 5' end of tRNA^{Gly}_{GCC} [21]. Taken together with our findings of NSUN2 and cytosine methylation correlations in brain following opioid exposure, the results suggest that the tRNA regulome has a drug- and tissue-specificity that may contribute differentially to addiction phenotypes.

Furthermore, there is evidence that tRNA-mediated effects of addictive drugs are not limited to the nervous system. For example, in the male reproductive system, including sperm and epididymosomes, which are tissues that have been well-characterized as carrying large amounts of tRNA fragments that are sensitive to stressful conditions including low-protein diet [37]. Additionally, chronic ethanol exposure in rodents altered NSUN2 expression and increased the abundance of tRNA fragments originating from tRNA^{Glu}_{CTC} and tRNA^{His}_{GTG} in the male reproductive tract, while fragments from tRNA^{Ser}_{AGA} were decreased in sperm [20]. Conversely, another study measured tRNA fragments in sperm of human opioid users and found a decrease in various tRNA fragments, with the most marked effect being at tRNA^{Gly}_{GCC} [19].

Importantly, changes in NSUN2-mediated tRNA methylation in the mature brain selectively decrease mature tRNAs [22]. For this reason, in the present study we chose to focus on mature tRNA expression and m⁵C in the brains of morphine-exposed mice, but

future studies could link our findings with tRNA^{Gly}_{GCC} fragmentation. The downstream molecular or behavioral sequelae of these changes have not been elucidated. In peripheral tissues and cultured cells, it has been established that impairments to the tRNA methylation machinery via ablation of NSUN2 causes accumulation of tRNA fragments, which activates cellular stress pathways and globally decreases protein synthesis [14, 15].

Remaining questions and future directions

Here we identified 30 days after morphine withdrawal a significant morphine-dependent increase in tRNA^{Gly}_{GCC} C39 methylation that correlated with an increase in tRNA^{Gly}_{GCC} expression. The question remains: How can an increase in tRNA expression modulate the addiction circuitry to confer vulnerability to future drug-seeking? While most studies demonstrate loss of methylation following environmental perturbations [14, 15, 38], we show here a site-specific experience-dependent increase in methylation of m⁵C. Therefore, the mechanism by which increased methylation drives tRNA abundance is unknown. Our group has previously shown that viral overexpression of NSUN2 in wild-type mouse mPFC selectively increases methylation at tRNA^{Gly}_{GCC} C39 and also confers a depressive-like phenotype [22]. While this provides an additional link between increased tRNA^{Gly}_{GCC} C39 methylation and downstream behavioral consequences, future studies will identify if virally overexpressing *Nsun2* also increases tRNA^{Gly}_{GCC} abundance and how this contributes to addiction-related or depressive-like behavior, which involve overlapping circuitry.

In the current study we identified a deficit in morphine preference in the CPP following deletion of neuronal *Nsun2* in both the postnatal forebrain and the adult mPFC. We note here that this deficit may not be specific to morphine or other opioids, but instead could be relevant to a broader impairment of the reward circuitry or impairments of reinforcer-context associations. Future studies will identify if these effects extend to other addictive drugs or even other rewarding stimuli, including food reward or social contact. Additionally, while we speculate that *Nsun2*-mediated nuclear-encoded tRNA methylation and expression play a role in morphine CPP deficits, we cannot rule out other molecular targets of *Nsun2* enzymatic activity, including (but not limited to) mitochondrial tRNAs [39], lncRNAs, miRNAs, and rRNAs that may be partially mediating these behavioral effects [40]. Sex differences and brain region specificity will also be probed in further studies, as our current study used male mice and pooled adjacent mPFC subregions (IL, PL) for tRNA assays following the morphine withdrawal procedure due to input restrictions, but we note that the well-established molecular and functional differences in these areas, and between males and females, may produce differential effects on *Nsun2* levels, tRNA expression, or methylation.

tRNAs can sense environmental cues and modulate the epitranscriptomic landscape in an experience-dependent manner. While this has mostly been demonstrated in the context of toxin exposure [38] or cellular stress via UV radiation [14, 15], there is evidence of global tRNA m¹A levels increasing following non-associative learning in the cerebral ganglia of *Aplysia californica* [41]. We now add to this concept by showing drug- and withdrawal-dependent dynamic changes in the tRNA m⁵C epitranscriptome in mammalian brain. Not only are these observed molecular changes dynamic, but the presence of a robust tRNA phenotype at 30 days after the last morphine injection provides new insight into mechanisms by which the brain may be primed for relapse or reinstatement after prolonged abstinence, including potential changes to protein synthesis linked to tRNA-specific codon content. While we have shown here two converging lines of evidence that tRNA regulation is linked to opioid withdrawal and opioid preference, it will be crucial in future work to functionally link the biochemical and

behavioral assays by investigating tRNA changes occurring during the CPP training itself. Although the current study is unable to describe exactly how Nsun2 function is affecting behavioral responses to morphine, linking Nsun2 manipulations, the tRNA regulome, and translational outcomes to behavioral changes that arise after long term opioid exposure (e.g., withdrawal behavior, relapse, drug seeking) is of interest as a future direction of this work.

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AUTHOR CONTRIBUTIONS

J.B. and C.J.B. conceptualized and designed experiments. C.J.B. and R.F. performed mouse morphine dosing and CPP behavior experiments and analyzed CPP behavioral data. J.B. performed all viral surgeries for CPP, conducted molecular assays, and analyzed and interpreted all molecular data. B.J. performed IHC and provided representative histological image for viral injections. S.A., E.J.N., and V.Z. contributed design of experiments and interpretation of data. J.B., C.J.B., and S.A. wrote the manuscript with input from all authors.

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COMPETING INTERESTS

The authors declare no competing interests.

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