Functional analysis reveals differential effects of glutamate and MCH neuropeptide in MCH neurons

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ABSTRACT

Objectives: Melanin-concentrating hormone (MCH) neurons in the lateral hypothalamus (LH) regulate food intake and body weight, glucose metabolism and convey the reward value of sucrose. In this report, we set out to establish the respective roles of MCH and conventional neurotransmitters in these neurons.

Methods: MCH neurons were profiled using Cre-dependent molecular profiling technologies (vTRAP). MCHCre mice crossed to Vglut2fl/fl mice or to DTRfl/fl were used to identify the role of glutamate in MCH neurons. We assessed metabolic parameters such as body composition, glucose tolerance, or sucrose preference.

Results: We found that nearly all MCH neurons in the LH are glutamatergic and that a loss of glutamatergic signaling from MCH neurons from a glutamate transporter (VGlut2) knockout leads to a reduced weight, hypophagia and hyperkinetic behavior with improved glucose tolerance and a loss of sucrose preference. These effects are indistinguishable from those seen after ablation of MCH neurons. These findings are in contrast to those seen in mice with a knockout of the MCH neuropeptide, which show normal glucose preference and do not have improved glucose tolerance.

Conclusions: Overall, these data show that the vast majority of MCH neurons are glutamatergic, and that glutamate and MCH signaling mediate partially overlapping functions by these neurons, presumably by activating partially overlapping postsynaptic populations. The diverse functional effects of MCH neurons are thus mediated by a composite of glutamate and MCH signaling.

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Keywords  MCH; Sucrose preference; Body weight; Glucose metabolism; Neuropeptide; Glutamate

1. INTRODUCTION

Melanin concentrating-hormone expressing (MCH) neurons located in the lateral hypothalamus (LH) play an important role in the regulation of energy balance metabolism, and other functions [1-3]. Both an MCH knockout (KO) and ablation of MCH neurons result in reduced food intake and leanness [2,4]. MCH ablation also enhances glucose disposition in a glucose tolerance test. Recently, MCH neurons have been also implicated in sucrose preference, a reward related process, via projections to dopaminergic neurons in the striatum and midbrain [5]. In previous studies we have noted that there are differences between the effects of MCH neural ablation compared to an MCH knockout, raising the possibility that additional transmitters besides MCH mediate the effects of these neurons. In this report, we set out to establish the molecular phenotypes of MCH neurons by profiling these neurons. A previous study suggested that a subset of MCH neurons are GABAergic while a different study suggested they are nearly exclusively glutamatergic [6,7]. Our data indicated that the great majority of MCH neurons are glutamatergic. We then compared the physiologic effects of ablating glutamate signaling in these neurons to either an MCH knockout or MCH neural ablation. We found that the effects of knocking out the glutamate transporter and MCH were different, suggesting that the function of MCH neurons is a composite of the function of a classic neurotransmitter and a neuropeptide.

2. METHODS

2.1. Mice and diets

C57BL/6 mice were purchased from Jackson (Jaxmice). The generation of MCH-Cre mice has been previously reported [5]. VGlut2<sup>lox/lox</sup> mice were purchased from Jackson (Jaxmice). To generate MCH-
specific VGlut2 knock-out mice (MCHCre;VGlut2fl/fl and MCHCre;VGATfl/fl, respectively), MCH-Cre mice were crossed with VGlut2lox/lox. Colonies were maintained by breeding MCH-Cre, loxp/lox mice with loxp/lox mice. MCHKO mice were purchased from Jackson (Jaxmice), and MCH neuron-ablated mice were generated as described previously [5]. Mice were maintained on a 12:12 h light–dark cycle with free access to water and standard chow or high-fat diet (60% Kcal fat; Research Diets) for 12 weeks (starting at 6 weeks of age). In vivo studies were performed following the National Institutes of Health Guidelines on the Care and Use of Animals and approved by the Rockefeller University Institutional Animal Care and Use Committee (Protocols 15797-H and 15818-H).

2.2. Physiological measurements

Body weights were determined weekly. Blood samples were collected via tail vein or trunk bleeds using a capillary collection system (Sarstedt). Blood glucose was measured using an Ascensia Elite XL glucometer (Bayer HealthCare, Tarrytown, NY). Plasma leptin levels were analyzed by ELISA techniques (Alpco immunosays). Glucose tolerance tests were performed on overnight fasted mice. c-glucose (2 g/kg) was injected intraperitoneally (i.p.) and blood glucose determined at the indicated time points. Locomotion activity was measured using beam breaks in metabolic cages from TSE systems.

2.3. Construction of AAV-IV-GFPL10

For construction of AAV-IV-GFPL10, see [8]. Briefly, we generated the plasmid for AAV-IV-GFPL10 by synthesizing an inverted EGFP sequence flanked by synthetic introns, lox sites, and forward Rpl10a. This fragment was then subcloned into pAAV-DIO-Chr2-mCherry to establish Cre-dependent expression, and it was then packaged into AAV pseudotype 5 (JNC Vector Core).

2.4. Immunoprecipitation and RNA-seq analyses

Details on GFP immunoprecipitations can be found in [9].

2.5. Stereotaxic surgeries

MCH Cre transgenic mice of 12–16 weeks of age were induced and maintained on isoflurane before being bilaterally injected with 0.5 μL of AAV IV-GFPL10 in the LHA (LHA coordinates: −1.4 AP, −1.0 −/+ 0.7 L and −5.2 DV). After viral injection, the needle was left in place for 10 min before slowly retracting. The skin was closed with a surgical clip.

2.6. Sucrose preference

Animals were acclimated to the chambers until side preference for either bottle was even. When preference was even, mice were studied as follows: first, sucrose in one bottle and water on the other, then, sucralose in one bottle and water in the other, and, lastly, sucrose in one bottle and sucralose in the other. During the acclimation and exposure periods, mice were water-deprived for 16–23 h and then given water through the bottles inside the chamber for half an hour. Two-bottle preference was calculated as the ratio: preference for 1 = number of licks on bottle 1/(number of licks on bottle 1 + number of licks on bottle 2) and expressed as percentage values, with 50% representing the indifference ratio. Behavioral data was analyzed with Excel and Prism and is expressed, as mean ± SEM. Significance tests comparing groups were t tests. The size of each animal group is given in Table 2. The investigator was blind to the genotype. In all cases, concentration of sucrose was 0.4 M, and concentration of sucralose was 1.5 mM. These concentrations were based on previous literature [10]. Briefly, the differences in molarity of sucrose and sucralose reflect differences in ligand-binding affinity of either sweeter than to taste receptors and were chosen among the plateau values of behavioral dose–response curves (preference for either sweeter versus water in [10]). Volume dispensed by the lickometers averages 2 μL/vick [10].

2.7. Immunohistochemistry

Mice were first anesthetized with isoflurane followed by transcardial perfusion with PBS and then 10% formalin. Brains were dissected, incubated in 10% formalin overnight at 4 °C, and 40 μm sections were prepared using a vibratome. Free floating sections were blocked for 1 h at room temperature in blocking buffer (PBS, 0.1% Triton-X, 2% goat serum, 3% BSA) and then incubated overnight at 4 °C with primary antibody for rabbit anti-MCH (1:5000; Phoenix Pharmaceuticals). The next day, sections were washed in washing buffer (PBS, 0.1% Triton-X) 3 times for 20 min, incubated with dye-conjugated secondary antibody (1:1000; Alexa Fluor 488 or 594 goat anti-rabbit) for 1 h at room temperature, washed in washing buffer, and then mounted.

2.8. Microscopy and quantification

Stained brain sections were imaged using Zeiss LSM 510 laser scanning confocal microscope. All parameters were kept constant for quantification purposes and were conducted with ImageJ.

2.9. Statistics

Results are expressed as mean ± SEM. Student t test, one-way ANOVA, or two-way ANOVA was used to assess the significance of difference of the mean or the comparative analyses. Sidak post-hoc tests were used for multiple comparisons. Differences were considered significant at p < 0.05.

3. RESULTS

3.1. MCH neuron profiling confirms glutamatergic identity of these neurons

We first set out to establish the phenotypes of MCH neurons in an unbiased manner by molecular profiling using viral TRAP (vTRAP) (short for viral Translating Ribosome Affinity Purification). We injected an adeno-associated virus (AAV) expressing the ribosomal fusion protein L10 flanked by lox sites (AAV-GFPL10) into the LH of MCH Cre mice. After three weeks, polysomes from hypothalamic blocks were precipitated using an anti-GFP antibody followed by high-throughput RNA sequencing (RNA-seq) of both the precipitated polysomal RNA (IP RNA) and total Input RNA (Figure 1A). We plotted FPKM (fragment per kilobase of transcript per million mapped reads) values for the IP RNA vs Input RNA on a log–log scale (Figure 1B) and found that Pmch itself was 121 Fold differentially enriched (DE is IP/Input) and that Tacr3 was 200 Fold differentially enriched (DE is IP/Input). Additionally, we did not see enrichment of Mup6 (78.3 fold), Lrrn7g (53.91 fold), and lapp (30.8 fold). The fold enrichment for all of the enriched genes is listed in Supplemental Table 1. We next replicated the RNA-seq data for a subset of genes by performing qPCR on the IP and total RNA and reconstructed highly significant enrichment of Pmch (102.5 Fold), Tacr3 (18.4 Fold), and Carpt (29.9 Fold) (Figure 1D). As expected, we did not see enrichment of the arcuate markers Agrp and Pomc or glial markers (Glap, Mal and Mbp) (Figure 1B–D).
Next, we analyzed the fold enrichment of markers for glutamate and GABA and found significant enrichment of the vesicular glutamate transporter 2 (vGlut2) RNA (2.1 Fold, \( p < 0.01 \)) (Figure 1H) and vGlut3 (2.88 Fold, \( p < 0.01 \)) but not for vGlut1 (Supplemental Table 1). For GABAergic markers, there was a 1.8-fold enrichment for the 67 kD isoform of glutamic acid decarboxylase GAD67 (\( \text{Gad}1 \), 1.8-fold, \( p < 0.01 \)) and 1.6-fold enrichment for the 65 KDa isoform, GAD65 (\( \text{Gad}2 \), 1.6-fold, \( p < 0.01 \)) (Figure 1F–G) raising the possibility that the MCH neurons could be glutamatergic or GABAergic. However, VGAT, the GABA transporter, was not enriched (\( \text{VGAT} \), 1.01-fold, \( p = 0.95 \)). We further characterized these neurons by performing dual ISH/IHC for MCH (red) in a vGlut2GFP mouse. Colocalization is shown in yellow and with white arrows pointing to the colocalizing cells. \( n = 3 \) in each group. Data are expressed as mean ± SEM. **\( P < 0.01 \). Scale bar = 100 nm. See also Supplemental Table 1.

3.2. Glutamatergic MCH signaling and energy balance

We evaluated the function of glutamate signaling by MCH neurons by generating MCH neuron-specific vGlut2 knockout mice by mating previously validated MCH-Cre mice [2] to animals with a floxed allele of vGlut2 (hereafter referred as \( \text{MCHCre};\text{vGlut2Fl/Fl} \) [2]. Immunohistochemical staining for MCH revealed that the vGlut2 knockout in MCH neurons did not alter the MCH population size or somatic area (Figure S1A), indicating that a vGlut2 knockout in these neurons does not alter their differentiation and/or survival. We next compared the

Figure 1: MCH neurons express are uniquely glutamatergic. (A) Mixed illustration. Schema of the adeno-associated virus (AAV) design for capturing transcribing RNA by immunoprecipitation and representation of stereotactic injection of AAVIVGFPL10 into the lateral hypothalamus of the brain. (B) FPKM GFP IP plotted against FPKM input on a Log–Log scale. Red dots highlight differentially enriched markers of MCH cells (\( \text{Pmch} \), Tacr3 and Carpt). (C–D) RNAseq data and Taqman analysis of MCH neuron markers (\( \text{Pmch} \), Tacr3 and Carpt), glial markers, \( \text{App} \) and \( \text{Pomc} \) (Non MCH-expressed genes), (E) Histogram display of number of differentially enriched genes (IP/Input) (F–H) RNAseq enrichment of glutamatergic vGlut2 gene and GABAergic Gad1 and Gad2 genes. (I) RNAscope for VGAT paired to an IHC for MCH showing no colocalization between MCH and VGAT (J–K) IHC of MCH (red) in a vGlut2GFP mouse. Colocalization is shown in yellow and with white arrows pointing to the colocalizing cells. \( n = 3 \) in each group. Data are expressed as mean ± SEM. **\( P < 0.01 \). Scale bar = 100 nm. See also Supplemental Table 1.
metabolic phenotype of these mice with animals to an ablation of MCH neuron using diphtheria toxin (hereafter referred as MCHDTA mice) or a germline knockout of the MCH neuropeptide (MCHKO). VGlut2Fl/Fl mice showed a significant reduction in body weight that was similar in magnitude to those of MCH ablated and MCHKO mice (Figure 2A). Also similar to both, the MCH vGlut2 knockout mice were hyperactive (Figure S1B) and also showed a late onset hypophagia (Figure 2B) resulting in a significant reduction in adiposity (Figure 2C). When fed a HFD, also similar to the MCH KO and MCH ablation, MCHCre; VGlut2Fl/Fl mice gained less weight than controls (Figure S1C) with significantly reduced adiposity (Figure S1D) and food intake (Figure S1E) [13]. Consistent with this, MCH VGLUT2 KO mice had significantly lower leptin levels than control animals (without a knockout) fed either normal chow diet or a high fat diet (HFD) (Figure S1F).

3.3. Glutamatergic signaling by MCH neurons, but not MCH, regulates glucose metabolism and sucrose preference

Previous studies have shown that MCH neurons are glucose sensitive and that ablation of these neurons alters glucose homeostasis [3,14,15]. At baseline, MCHKO, MCHDTA, and MCHCre; VGlut2Fl/Fl mice were normoglycemic (Figure 2D). However, similar to mice with an MCH ablation, MCHCre; VGlut2Fl/Fl mice showed a significant improvement of glucose tolerance with a 21.16% lower area under the curve compared to control littermate mice after a glucose tolerance test with an injection of a 2 g/kg bolus of glucose. In contrast, mice with a knockout of the MCH peptide did not show an altered glucose tolerance test (Figure 2E–F and [4]). These data demonstrate that, similar to MCH ablation but in contrast to an MCHKO, a deletion of vGlut2 in MCH neurons alters glucose metabolism. MCH neurons convey the post-ingestive value of sucrose and project to midbrain regions regulating reward [5]. For example, we previously showed that animals with an ablation of MCH neurons no longer prefer sucrose to a sucralose, non-nutritive sweetener. In contrast, MCHKO mice show a normal preference for sucrose. To further assess the roles of glutamate signaling by these neurons, we evaluated the sucrose preference of MCHCre; VGlut2Fl/Fl relative to MCHKO and mice with an MCH neural ablation [10]. The mice were given a sequential set of choices using a two-bottle test as follows: sucrose vs. sucralose, sucrose vs. water and sucralose vs. water, and the preference ratios were computed. Normal mice show a strong preference for sweet taste (either sucralose or sucrose) compared to water and also prefer sucrose to sucralose.

As previously reported, an ablation of MCH neurons reduced an animal’s preference for sucrose relative to sucralose while sucrose preference in an MCHKO was unchanged relative to controls (Figure 3A–C). However similar to mice with an MCH ablation, MCHCre; VGlut2Fl/Fl mice did not show a preference for sucrose relative to sucralose (Figure 3A). Thus, the effect of an ablation of MCH neurons on sucrose preference is a result of a loss of glutamate signaling, not MCH itself. A knockout of glutamate signaling in MCH neurons was specific for nutrient preference as the knockout mice still showed a normal preference for either sucrose or sucralose relative to water. (Figure 3B–C).

Figure 2: MCHCre;Vglut2Fl/Fl mice are lean and show improved glucose tolerance similar to diphtheria toxin ablated MCH mice. (A) Weekly body weight assessment in MCHKO, MCHDTA, and MCHCre; VGlut2 Fl/Fl mice. (B) Food intake assessment at 12 weeks of age in MCHKO, MCHDTA, and MCHCre; VGlut2 Fl/Fl mice. (C) Adiposity shown by means of an EchoMRI scan of MCHKO, MCHDTA, and MCHCre; VGlut2 Fl/Fl, (D) Basal blood glucose levels and (E–F) glucose tolerance test of MCHKO, MCHDTA and MCHCre; VGlut2 Fl/Fl mice. n = 5–8 in each group. Data are expressed as mean ± SEM. *p < 0.05 **p < 0.01 ***p < 0.001 ****p < 0.0001. See also Figure S1.
4. DISCUSSION

Numerous studies have shown that MCH neurons in the LH regulate feeding, metabolism, reward, and other biologic processes [1–3,5,16], and many of these effects have been ascribed to MCH itself [1]. However, MCH neurons also express classical neurotransmitter(s) and their functional role compared to the neuropeptide has not been evaluated.

The neurotransmitters co-expressed in MCH neurons have been controversial. While some studies of MCH neurons have only found co-expression of markers for glutamatergic neurotransmission, others have suggested that there is a separate, possibly overlapping, GABAergic subpopulation. For example, a recent report demonstrated expression of glutamate transporter vGlut2 in MCH neurons in 68/69 cells with no co-expression of VGAT [17]. In contrast, another study has shown that MCH neurons co-express the GABA marker GAD67, suggesting that these same neurons may co-release GABA [6]. In yet another publication, the GABAergic LH neurons were reported to project to different VTA regions than the glutamatergic subpopulation and mediate positive reinforcement while LH glutamatergic inputs to the VTA were reported to cause avoidance and suppress DA release in the NAc [18].

To resolve these discrepancies, we used molecular profiling to characterize MCH neurons and found that vGlut2 is expressed in 97% of MCH neurons. In contrast, we failed to find expression of the GABA transporter VGAT in these neurons. This suggests that the vast majority of MCH neurons are glutamatergic. However, consistent with other reports, we found a small but significant level of expression of GAD65 and GAD67, which generates GABA by decarboxylation of glutamic acid, in MCH cells (see Figure 1E–G). However, it is unclear what the expression of GAD in these cells means since, in the absence of a reuptake mechanism, GABA release would likely deplete the intracellular pool. Of note, we did not find enrichment for other monoamine transporters, and moreover, any GABA production by these cells would be co-expressed with vGlut2 in the vast majority of MCH cells. Thus while our anatomic data strongly suggest a glutamatergic identity of most MCH cells, we cannot rule out the existence of other MCH cell subtypes or their co-release or reuptake with other neurotransmitters such as GABA through non-canonical mechanisms [19,20].

Consistent with a more prominent role for glutamate and not GABA signaling by
these neurons, our functional studies show that the aggregate effects of the MCH peptide and glutamate signaling by MCH neurons can fully explain the known functions of these neurons. Indeed, in each case where the phenotype of a mouse with an MCH knockout differs from that of mice with ablation of MCH neurons, the discrepancy can be explained by invoking glutamate signaling. For example, while the effect to reduce the body weight of a MChVglut2KO is similar to that of an MCHKO, the function of these neurons to control glucose metabolism requires glutamate but not MCH signaling (Figure 2). Similarly while it was hypothesized that the effect of MCH ablation in adult mice to alter glucose tolerance might be a result of disruption of CART, nestin-1 or GABA [4], our results show that glutamate signaling is responsible for this effect. Additional reports have established that MCH neurons communicate the reward value of sucrose at synapses while MCH can signal in a volumetric manner which also might result in signaling at partially overlapping cell types. Consistent with a prominent function of glutamate signaling by these neurons, comprehensive genetic analyses indicate that either MCH and/or glutamate are sufficient to mediate all of the known functions of these neurons. These data suggest that the effects of MCH neurons are a composite of the effects of these two neurotransmitters.

**AUTHOR CONTRIBUTIONS**

MS, KT, LP, and AN designed and performed experiments and analyzed data. MS and JF wrote the manuscript. EA and AD performed sucrose preference experiments. ZL and CC provided experimental support and intellectual input. JF conceived the study, supervised research, and wrote the manuscript with input from all authors.

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**CONFLICTS OF INTEREST**

Authors declare no conflicts of interest.

**APPENDIX A. SUPPLEMENTARY DATA**

Supplementary data related to this article can be found at https://doi.org/10.1016/j.molmet.2018.05.001.

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