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Citation: Science Bulletin 64, 1167 (2019); doi: 10.1016/j.scib.2019.05.005

View online: http://engine.scichina.com/doi/10.1016/j.scib.2019.05.005

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Characterization of glutamatergic VTA neural population responses to aversive and rewarding conditioning in freely-moving mice

Quentin Montardya,1, Zheng Zhoua,b,1, Zhuogui Leia,e,1, Xuemei Liua,b,1, Pengyu Zenga, Chen Chen a, Yuanming Liua, Paula Sanz-Leonc,d, Kang Huanga,b, Liping Wang a,b,*

a Shenzhen Key Lab of Neuropsychiatric Modulation and Collaborative Innovation Center for Brain Science, Guangdong Provincial Key Laboratory of Brain Connectome and Behavior, CAS Center for Excellence in Brain Science and Intelligence Technology, The Brain Cognition and Brain Disease Institute (BCBDI), Shenzhen Institutes of Advanced Technology, Chinese Academy of Sciences, Shenzhen-Hong Kong Institute of Brain Science-Shenzhen Fundamental Research Institutions, Shenzhen 518055, China
b University of Chinese Academy of Sciences, Beijing 100049, China
c School of Physics, the University of Sydney, Sydney 2006, Australia
d Centre for Integrative Brain Function, the University of Sydney, Sydney 2006, Australia
e Department of Biomedical Sciences, City University of Hong Kong, Tat Chee Avenue, Kowloon, Hong Kong, SAR 999077, China

Abstract
The Ventral Tegmental Area (VTA) is a midbrain structure known to integrate aversive and rewarding stimuli, but little is known about the role of VTA glutamatergic (VGluT2) neurons in these functions. Direct activation of VGluT2 soma evokes rewarding behaviors, while activation of their downstream projections evokes aversive behaviors. To facilitate our understanding of these conflicting properties, we recorded calcium signals from VTAVGluT2+ neurons using fiber photometry in VGluT2-cre mice to investigate how this population was recruited by aversive and rewarding stimulation, both during unconditioned and conditioned protocols. Our results revealed that, as a population, VTAVGluT2+ neurons responded similarly to unconditioned-aversive and unconditioned-rewarding stimulation. During aversive and rewarding conditioning, the CS-evoked responses gradually increased across trials whilst the US-evoked response remained stable. Retrieval 24 h after conditioning, during which mice received only CS presentation, resulted in VTAVGluT2+ neurons strongly responding to CS presentation and to the expected-US but only for aversive conditioning. To help understand these differences based on VTAVGluT2+ neuronal networks, the inputs and outputs of VTAVGluT2+ neurons were investigated using Cholera Toxin B (CTB) and rabies virus. Based on our results, we propose that the divergent VTAVGluT2+ neuronal responses to aversion and reward conditioning may be partly due to the existence of VTAVGluT2+ subpopulations that are characterized by their connectivity.

1. Introduction

1.1. The Ventral Tegmental Area

The ventral tegmental area (VTA) is a midbrain structure that has been linked with a variety of behavioral functions including aversion and reward [1–4], and prediction error [2,5], and motivation [2,6,7]. The heterogeneous composition of VTA includes a large proportion of dopaminergic (DA) neurons (60%), and smaller proportions of GABAergic neurons (GABA) (35%) and glutamatergic neurons (2%–5%) [8,9]. Whilst the two first neuronal components (DA and GABA) of VTA have been well studied and characterized [1,10], the glutamatergic population has received less attention and its functional characterization needs to be elucidated to obtain a better understanding of VTA function.

1.2. Physiology, anatomy and function of VTA glutamatergic neurons

While there is a general consensus regarding the role of VTA-DA and VTA-GABA neurons in control of aversive and rewarding behavior, the role of VTA's glutamatergic neurons is not yet understood [10,11]. Glutamatergic neurons, defined by their expression of vesicular glutamate transporter 2 (VGLUT2) [9,12], can form local connections in VTA [13] and send long-range projections to structures such as the Nucleus Accumbens (NAc) or the Lateral Habenula (LHb) [4,14,15].

* Corresponding author.
E-mail address: lp.wang@siat.ac.cn (L. Wang).
1 These authors contributed equally to this work.
Optogenetic stimulation of VTAVGluT2+ neuronal somata promotes rewarding behaviors, such as place preference and appetitive instrumental conditioning [16]; however, at the microcircuit level the function of their interaction with neighboring DA and GABA neurons remains unknown [13]. Conversely, VTA glutamatergic transmission has also been associated with aversive behaviors [17]. Indeed, optogenetic activation of axonal projections from VTAVGluT2+ terminals in the nucleus accumbens (NAC) or to the lateral habenular nucleus (LHb) both promote aversive behaviors, including aversive conditioning [14,16]. Recent electrophysiological recordings of glutamatergic neurons confirmed that individual VTAVGluT2+ neurons can be activated by aversive stimulation, and either excited or inhibited by rewarding stimulation [18], giving an initial insight into this seemingly paradoxical function. However, population response recordings, such as calcium imaging, can potentially help us to understand VTAVGluT2+ function in terms of conditioned aversion and reward. In particular, we use fiber photometry with the genetically-encoded Ca2+ indicators GCaMP6s [19], which is a minimally invasive method that allows in-vivo measurements in freely-moving animals of synchronous neuronal population activity from subcortical structures [20,21], which has proved useful during conditioning [22].

Here, we investigated the VTAVGluT2+ neuronal population response to aversive and rewarding events, both during unconditioned and conditioned stimulation. Conditioning consisted in pairing a tone (CS+) with an aversive (footshock) or rewarding (sucrose) unconditioned stimulus (US). In addition, a retrieval test was performed 24 h after conditioning to see whether VTAVGluT2+ neurons maintain a robust memory of aversive or rewarding conditioning. Finally, to better understand VTAVGluT2+ population responses to aversion and reward, we investigated their connectivity pattern using cell specific monosynaptic retrograde rabies virus tracing, allowing the mapping VTAVGluT2+ inputs.

2. Materials and methods

2.1. Animals

All procedures were approved by Animal Care and Use Committee in the Shenzhen Institute of Advanced Technology (SIAT) or Wuhan Institute of Physics and Mathematics (WIPM), Chinese Academy of Sciences (CAS). Adult (6–8 weeks old) male VGluT2-ires-cre (Jax No. 016963, Jackson Laboratory) transgenic mice were used in this study. All mice were maintained on a 12/12-h light/dark cycle at 25 °C. Food and water were available ad libitum.

2.2. Viral preparation

For fiber photometry experiments, AAV2/9-EF1a-DIO-Gcamp6s virus was used. For tracing experiments, we used AAV2/9-EF1a-DIO-BFP. Virus titer were approximately 2–3 × 10^{12} μg/mL. In the rabies tracing experiments, AAV2/9-CAG-DIO-histone-TVA-GFP (4.2 × 10^{12} μg/mL), AAV2/9-CAG-DIO-RV-G (4 × 10^{12} μg/mL) and EnVa-RV-DsRed (1 × 10^{12} pfu/mL) viruses (BrainVTA Co., Ltd., Wuhan) were used.

2.3. Viral injections

VGluT2-ires-cre mice were anesthetized with pentobarbital (i.p., 80 mg/kg) and fixed on stereotaxic apparatus (RWD, Shenzhen, China). During the surgery, mice were kept anesthetized with isoflurane (1%) and placed on a heating pad to keep the body temperature at 35°C. A 10 μL microsyringe with a 33-Ga needle (Neuros; Hamilton, Reno, USA) was connected to a microliter syringe pump (UMP3/Micro4; WPI, USA) and used for virus injection into VTA (coordinates: AP, –3.15 mm; ML, –0.3 mm; DV, –4.4 mm).

2.4. Retrograde tracing

For CTB tracing, VGluT2-ires-cre mice received CTB Alexa Fluor conjugates (CTB 594 or CTB 488, Invitrogen Inc., Grand Island, NY, USA) via injection into NAC and LHb (50 nl per injection) and AAV2/9-EF1a-DIO-BFP into VTA (200 nl per injection). For transynaptic rabies tracing, a total volume of 70 nl mixed viruses AAV-EF1a-DIO-RV-G and AAV-EF1a-FLEX-GFP-TVA (volume ratio 1:1) were injected into VTA of VGluT2-ires-cre mice (coordinates: AP: –3.15 mm; ML: –0.3 mm; DV: –4.4 mm). After 3 weeks, 100 nl of EnvA-RV-DsRed (EnvA-pseudotyped rabies virus) was injected at the same coordinates. Mice were sacrificed one week after this second injection. All rabies tracing experimental procedures were completed in Biosafety level 2 (BSL2) Laboratory.

2.5. Implantation of optical fibers

A 200 μm optical fiber (NA: 0.37; NEWDOON, China) was chronically implanted in the VTA of VGluT2-ires-cre mice 2–3 weeks following virus expression for fiber photometry experiments. The optical fiber was unilaterally implanted in VTA (AP: –3.15 mm, ML: –1.10 mm and DV: –4.2 mm) with a 15° angle in the medial direction of the transverse plane. After surgery all mice were allowed to recover for at least 2 weeks.

2.6. Histology, immunohistochemistry, and microscopy

Mice were sacrificed by overdosing with pentobarbital (1% m/v, 150 mg/kg, i.p.) and transcardially perfused with 1 mol/L cold saline followed by ice-cold 4% paraformaldehyde (PFA; Sigma) in 1 mol/L PBS. Brains were removed and submerged in 4% PFA at 4 °C overnight to post-fix, and then transferred to 30% sucrose to equilibrate. The coronal brains slices (40 μm) were sectioned with a cryostat (CM1950; Leica, Germany). Freely floating sections were washed with PBS and blocked for 1 h at room temperature in blocking solution containing 0.3% Triton X-100 and 10% normal goat serum (NGS). Then the sections were incubated overnight with rabbit monoclonal anti-dsRed (1:500, #632496; Clontech; Japan); GFP (1:500, #ab290, abcam, USA); DAPI (1:50,000, #62248; Thermo Fisher Scientific, USA) diluted in PBS with 3% NGS and 0.1% TritonX-100. The sections were incubated in 1 h at room temperature with Alexa Fluor 488 or 594 goat anti-rabbit secondary antibody (1:200; Jackson Laboratory, USA). Finally, the sections were mounted and photographed using the Zeiss LSM 880 confocal microscope (Zeiss; Germany). The images were acquired using identical gain and offset settings, and analyzed with ImageJ, Image Pro Plus, and Adobe Photoshop software. ROIs were traced with reference to the atlas. CTB and Rabies Virus immunoreactivity was quantified using Image Pro Plus and was verified by comparing with manual counts performed by a trained double-blind observer.

2.7. Unconditioned aversive and conditioned aversive stimulation

VGluT2-ires-cre mice (n = 8) with optical fibers implanted were placed in an unescapable acrylic box (L 25 cm × W 25 cm × H 70 cm) with a metal grid floor that delivered footshock currents (0.6 mA footshock, 0.5 s). Each mouse went through an unconditioned and then a conditioned protocol as below. During unconditioned aversive stimulation, mice were freely moving and footshocks were directly delivered with inter-trial interval durations varying within session randomly set in a range between 60 and
120 s. The session was approximately 10 min long and each mouse received 10 footshocks. The conditioned sessions consisted of 5 trials where an auditory conditioned stimulus (CS; 3 kHz, sine wave, 90 dB, 5 s) was paired with an unconditioned stimulus (US; 0.5 s, 0.6 mA footshock; random inter-trial intervals 60–120 s) that began immediately after tone ended. Mice were presented with 5 CS cues alone, without footshock stimulation, 24 h after conditioning.

2.8. Unconditioned reward and conditioned reward

Following viral injections and optical fiber implantations, VGluT2-ires-cre mice (n = 8) underwent a third surgery to implant a steel headplate for head-fixing purposes. The mice were habituated (~30 min/d) to the head-fix system over two-three days. During the experiment, each mouse was head-fixed and a tube delivering liquid reward was directly aimed at their mouth, through which single drops of sucrose (5% w/v) could be delivered as reward. Each mouse went through an unconditioned and then a conditioned protocol as described below. Unconditioned reward sessions were conducted during which 30 reward trials were presented with inter-trial interval durations varying within session, randomly set in a range between 25 and 40 s. Conditioned reward sessions consisted of one session of 30 trials in which an auditory conditioned stimulus (CS; 10 kHz, sine wave, 80 dB, 5 s) was paired with one sucrose delivery, which was delivered immediately after the tone ended. Mice were presented with 30 CS cues alone, without sucrose reward, 24 h after conditioning.

2.9. Fiber photometry

Ca2+ signals were recorded using a fiber photometry system (Thinker Tech, Nanjing). Two weeks post AAV2/9-DIO-GCaMP6s virus injection, an optical fiber (NA: 0.37; NEWDOON, China) was implanted into VTA as described above.

The fiber photometry system included a 502–730 nm transmission band (Edmund, Inc.), a 480 nm excitation light from LEDs (CREE XPE), reflected off a dichroic mirror with a 435–488 nm reflection band and coupled into a 200 μm 0.37 NA optical fiber (Thorlabs, Inc.) by an objective lens. At the fiber tip, the laser intensity was about 20 μW. The collection of GCaMPs6s fluorescence used the same objective, transmitted by the dichroic mirror filtered through a green fluorescence protein (GFP) bandpass emission filter (Thorlabs, Inc. Filter 525/39), and detected by a CMOS camera sensor (Thorlabs, Inc. DCC3240M). The calcium signals were recorded by CMOS camera at 50 Hz. A LabVIEW (National Instruments, US) program was developed to control the CMOS camera. Behavioral event signals were recorded by a DAQ card (NI, usb-6001) at 1000 Hz using the same LabVIEW program.

2.10. Photometry data analysis

Calcium Imaging signals were first extracted using Blackrock NPKM (Neural Processing MATLAB Kit), using provider instructions (Thinker Tech, Nanjing). Custom MATLAB (The MathWorks Inc. ©) scripts were developed for further analysis using R2012a. Signals were analyzed as dF/F = (F – Fb)/Fb, where Fb was defined as the baseline fluorescence before stimulation. Data were then smoothed using a 10 ms sliding windows. Time courses were calculated by aligning the time of stimulation across all individual trials and then calculating the mean change in calcium at each time window. To compare calcium activity between conditions, mean calcium activity was calculated for 0.5 s time windows centered around the time of the activity peak (2 s before stimulation vs. CS vs. US); A multivariate permutation (1000 permutations, α level of 0.05) test was used to test the statistical significance of the difference between conditions over the time course, and a threshold indicating a statistically significant difference from the baseline was applied (P < 0.005). Area under Curve index is the sum of transient Ca2+ activity [22] over a period of 0.5 s centered around the peak of activity.

3. Results

3.1. VTAVGluT2+ population increases activity to unconditioned aversive stimulation and this response remains constant over successive trials

We first investigated whether the VTAVGluT2+ neuronal population responds to unconditioned aversive stimulation. To do that we infected VGluT2-cre animals with adeno-associated virus (AAV) expressing GCaMP indicator by injecting the AAV9-EF1α-DIO-GCaMP6s virus in VTA (Fig. 1a). Three weeks later an optical fiber was implanted above VTA, allowing in-vivo recording of VTAVGluT2+ calcium signals during freely-moving behavior (Fig. 1a, b). At the end of experiments, GCaMP6s virus expression in VTA (Fig. 1c) and fiber positioning were systematically checked in every mouse. (Fig. 1d). During the unconditioned aversive experience, mice received footshocks (0.5 s at 0.6 mA) whilst the activity of VTAVGluT2+ neurons was recorded. Immediately after the beginning of the footshock, the calcium signal of VTAVGluT2+ neurons strongly increased for each individual mouse (Fig. 1e, top), which was a stereotypical effect well aligned with the onset of stimulation (Fig. 1e, bottom). All mice expressed a similar increase of activity (4.26% DF/F, n = 8) directly after aversive stimulation, which was significantly different from baseline expression for 1.54 s before returning to baseline level (Fig. 1f, red part of the curve indicates P < 0.05 using the multivariate permutation test). The mean signal values for all mice for a period of 0.5 s around the peak response amplitude (T = 0.68 s) revealed that activity was significantly higher than baseline (BL = 0.002% DF/F vs. Footshock = 3.96% DF/F, P < 0.0001; Fig. 1g). To observe the effect of repeated unconditioned stimulation on VTAVGluT2+ neurons, we analyzed response trends on a trial-by-trial basis and across animals (Fig. 1h). The peak responses in successive trials remained at a similar level (Fig. 1h), which was confirmed by computation of the Area under Curve index (AUC) (Fig. 1i).

In summary, this experiment demonstrated that VTAVGluT2+ neurons were strongly activated by unconditioned aversive stimulation and the amplitude of the peak response remained constant across trials.

3.2. The VTAVGluT2+ neuronal population responds to aversive conditioning

To characterize the responses of VTAVGluT2+ to conditioned aversive stimulation (Fig. 2a) we applied the following protocol: (1) Habituation Day, during which animals received tone stimulation only; (2) Conditioning Day, during which a tone (CS) was paired with an unconditioned stimulation (US); (3) Retrieval Day, during which only the CS was presented.

On Habituation Day VTAVGluT2+ neurons were insensitive to the CS as shown in the example (Fig. 2b, c, left), demonstrating that a neutral stimulus was insufficient to evoke a significant response in this neuronal population. During the Conditioning Day, the response of individual mice to CS was larger compared to Habituation Day. All mice had a strong evoked response to the US, similar to the unconditioned footshock experiment (Fig. 2b, c, middle). Following conditioning, increased sensitivity to CS was evident in the mean responses of all mice (Fig. 2d; n = 8), where CS-evoked and US-evoked response amplitudes were significantly higher than the baseline (BL-evoked = 0.12% DF/F; CS-evoked = 3.8% DF/F, P < 0.0001; US-evoked = 8.86% DF/F, P < 0.0001; (Fig. 2e), and
US-evoked signal was still significantly stronger than CS-evoked signal ($P < 0.01$). As a control, another group of mice infected with a GFP virus followed the same aversive conditioning protocol ($P < 0.005$; Fig. 2j), indicating that VTAVGluT2+ neuronal populations of individual mice, as shown in Fig. 3c. An increase of calcium activity just after reward delivery was observed across all mice that was significantly different from baseline measurement (Fig. 2k). Velocity signal evoked activity remained constant for at least a period of 24 h. Together, these results show that during Conditioning Day, the amplitude of the CS-evoked responses of VTAVGluT2+ neurons gradually increased over repetitions, while US-evoked activity remains relatively constant. During Retrieval Day CS-evoked responses are similar to those observed during Conditioning Day and there is an Expected-US-evoked response.

3.3. VTA VGluT2 neurons respond to successive repetition of unconditioned rewarding stimulation

Since direct stimulation of VTAVGluT2+ neurons promote reward [16], and that unitary electrophysiological recording has shown that some VTAVGluT2+ neurons are sensitive to rewarding stimulation [18], we next wanted to investigate glutamatergic neuron population responses to rewarding stimulation. To answer this question, calcium signals were recorded whilst mice were head-fixed and a tube directly delivered a liquid reward in their mouth (5% sucrose water, ITI = 60–120 s, 30 trials) (Fig. 3a). Fiber position was carefully verified in brain slices for each individual mouse at the end of experiments (Fig. 3b). Following rewarding stimulation, a rapid increase of calcium activity was observed in VGluT2 neuronal populations of individual mice, as shown in Fig. 3c. An increase of calcium activity just after reward delivery was observed across all mice that was significantly different from baseline mea-

**Fig. 1.** VTA response to unpredicted aversive stimulation. (a) Schematic representation of AAV9-DIO-GCaMP6s injection in VTA. (b) Schematic representation of Fiber Photometry setup. (c) Representative image of GCaMP6s virus expression in VTA VGluT2-cre mice (Green, GCaMP6s; scale bar, 200 and 20 μm respectively). (d) Location of optical fiber position for every mouse. (e) Example showing individual trials Ca$^{2+}$ signal in VTA VGluT2 for one mouse (top) plotted as an individual time courses in black, and averaged in red; (bottom) plotted as heatmap. Footshock stimulation at T = 0 ms; Lightning symbol represents footshock time (T = 0). (f) Mean time course of VTA VGluT2 $+$ calcium signal (n = 8 mice), increasing during unconditioned stimulation. Data points in red represent a section of the time course where differences compared to base line (BL) are statistically significant, using a Multivariate Permutation Test. (g) Mean Ca$^{2+}$ signal during the baseline vs. Shock (BL = 0.002% DF/F vs. Shock = 3.96% DF/F, $P < 0.0001$). (h) Mean trial-by-trial responses (trial 1 to trial 5) (i) Trial-by-trial Area Under Curve (AUC) centered on shock stimulation, computed for all animals.
measurements for 8.16 s (Fig. 3d). Analysis of the amplitude of the peak of activity revealed that it was significantly higher than the baseline (BL = 0.07% DF/F vs. Reward = 4.72% DF/F, P < 0.0001). To check for a potential gradual change of signal amplitude across stimulation, we calculated the mean in groups of 5 consecutive trials and found stable activity across trials (Fig. 3f, g), similar to responses to unconditioned aversive stimulation.

In summary, we show that VTAVGluT2+ neurons were excited by rewarding stimulation and the amplitude of their activation remained stable across stimulation trials.

Fig. 2. VTA response to aversive conditioning. (a) Schematic representation of CS-US conditioning, where CS is a 5 s tone and US a 0.5 s footshock starting immediately after CS; Symbols represent tone CS and US footshock time (respectively T = 0 and 5 s). (b) Single animal example signal plotted as individual time courses in black, and averaged in red, for Day 0 Habituation (tone only), Day 1 Conditioning (CS-tone + US Shock), and Day 2 Retrieval (CS only). (c) Same as panel B, plotted as heatmap. (d) Day 1 Mean Ca2+ signal over the conditioning time course (n = 8 GCaMP6s mice + 2 GFP), showing an increase during CS and US stimulation. (e) Mean Ca2+ signal comparing Baseline (BL), CS and US (BL-evoked = 0.12% DF/F; CS-evoked = 3.8% DF/F, P < 0.0001; US-evoked = 8.68% DF/F, P < 0.0001). (f) Day 1 Mean conditioning trial-by-trial responses (bottom) zoom on CS-evoked responses and US-evoked responses. (g) CS-evoked and (h) US-evoked trial-by-trial Area Under Curve computed among for mice. (i) Day 2 Mean Ca2+ signal retrieval time course. (j) Mean Ca2+ signal during BL vs. CS vs. expected-US stimulation. (k) Day 2 Mean retrieval trial-by-trial responses (l) CS- and US-evoked trial-by-trial AUC during Retrieval.

Fig. 3. VTA response to unconditioned reward. (a) Schematic representation of Fiber Photometry setup during reward delivery. Head-fixed mice directly receive 5% sucrose in their mouth. (b) Location of optical fiber tip for all animals. (c) Single animal example Ca2+ signal (left) plotted as individual time courses in black (30 trials), mean Ca2+ signal in red; (right) and plotted as heatmap for 30 trials. Water drop represents rewarding stimulation time (T = 0). (d) Mean Ca2+ signal time course (n = 10 mice), increasing during rewarding stimulation. (e) Averaged Ca2+ signal during the baseline (BL = 0.07% DF/F) vs. rewarding stimulation (Reward = 4.72% DF/F, P < 0.0001). (f) Mean reward-evoked 5-trial block Ca2+ signal responses. (g) Mean reward-evoked AUC of 5-trial blocks.
3.4. VTA VGluT2 neurons respond to reward conditioning

We have demonstrated that the VTA\textsuperscript{VGluT2+} population gradually learned to respond to a CS tone preceding aversive stimulation. To determine if rewarding stimulation can evoke a similar response pattern, mice were conditioned to a reward; the CS was a 5 s tone and the US was 5% sucrose water (random ITI 60–120 s). The experiment was conducted over three consecutive days, similarly to aversive stimulation described above: Habituation Day, Conditioning Day, and Retrieval Day (Fig. 4a). During the Conditioning Day, an increase in calcium signal evoked by CS was observed, followed by a higher amplitude increase evoked by the US (Fig. 4b). Group data confirmed this pattern (Fig. 4d), revealing that the difference between baseline and CS stimulation-evoked VTA\textsuperscript{VGluT2+} population response was statistically significant and sustained for 3.88 s, and that the US stimulation evoked a larger activity increase that slowly returned to baseline level over 13.88 s. The CS-evoked and US-evoked activity peaks were significantly higher than the baseline (BL = 0.02% DF/F, CS = 2.53% DF/F, \( P < 0.0001 \); US = 11.88% DF/F, \( P < 0.0001 \), Fig. 4e), and the US-evoked activity significantly higher than the CS activity (\( P < 0.0001 \)). To investigate trial-by-trial changes, we plotted groups of 5 trials (Fig. 4f); this shows that CS-evoked activity slowly increased across trials, which was confirmed by calculating the area under curve (Fig. 4g), whereas US-evoked responses remained stable across trials (Fig. 4f, g). In addition, it appeared that across conditioning, CS-evoked activity remained sustained until the beginning of the rewarding stimulation, which can be seen by increasing the time window to take into account the long-sustained activity during CS (Trial 1–5 = 0.84 AUC vs. Trial 26–30 = 3.81, \( P < 0.05 \), Fig. 4g). During the Retrieval day, a calcium signal peak was visible during CS and US as shown in Fig. 4c, but of low amplitude compared to baseline. Group mean calcium signal changes show that the CS did not evoke any clear change, whereas expected-US evoked very brief activity trend (Fig. 4h); however, this was not statistically significant when looking at a 0.5 s time window around activity peak (Fig. 4i). In line with these results, no pattern was found across trials (Fig. 4j).

Together, these data demonstrate that the VTA\textsuperscript{VGluT2+} population responds to rewarding conditioning with increasing CS-evoked activity amplitude and duration, whilst US-evoked activity remained stable. But during retrieval, VTA\textsuperscript{VGluT2+} neurons did not respond.

3.5. VTA VGluT2 neurons may be characterized by their specific network

The VTA\textsuperscript{VGluT2+} neuronal population can respond differently to both aversive and rewarding stimulation whilst having a similar molecular background. To understand better how such differences in a homogeneous population may occur, we proposed to determine to which broader network they belong to as an alternative strategy to characterize them [10]. First, to determine whether VTA\textsuperscript{VGluT2+} neurons send collaterals to structures serving a similar function, we injected in the same VGLuT2-cre mice both the retrograde tracer Cholera Toxin-B conjugated with Alexa 594 (CTB-594) in NAc and Cholera Toxin-B conjugated with Alexa 488 (CTB-488) in LHb, two structures downstream of VTA\textsuperscript{VGluT2+} neurons that are associated with aversive responses. Mice were then infected with AAV-DIO-BFP virus in VTA (Fig. 5a). After expression, CTB was...
Fig. 5. LHb and NAc projections to VTA VGluT2 neurons. (a) Schematic representation of NAc CTB Alexa 594 tracer (CTB 594) injection (in red) + LHb CTB Alexa 488 tracer (CTB 488) injection (in green) + AAV9-DIO-BFP injection into VTA of VGluT2-cre animals. (b) Representative images showing the injection site of CTB 488 (top left, in green) in LHb, CTB 594 (top right, in red) in NAc, AAV-DIO-BFP virus expression in VTA (bottom, in blue) and retrogradely labelling CTB cells in VTA (scale bar, 200 μm). (c) Magnified pictures of VTA neurons expressing each fluorescence, merged on the bottom right. No co-expression of CTB 488 and CTB 594. (d) Representative pictures of VGluT2 positive neurons (in blue) VGluT2-VTA-NAc neurons (in red) and VGluT2-VTA-LHb neurons (in green) along anteroposterior VTA axis. (e) Quantification of VGluT2-VTA-NAc and VGluT2-VTA-LHb cells along anteroposterior VTA axis.
Fig. 6. VTA VGluT2 neuronal inputs. (a) Schematic representation of RV-virus injection protocol. (b) Representative picture of VTA neurons expressing RV virus after injection in VTA of VGluT2-cre animals (Red, rabies-dsRed; green, TVA-GFP; blue, DAPI, scale bar, 200, 50 and 20 μm, respectively). Boundaries of VTA are drawn in white dashed lines. (c) Quantification of neurons expressing RV virus after injection in VTA of VGluT2 animals. (d) Representative pictures showing retrograde labeling in the LHb & MHB; BNST and NAc with inputs to VTA VGluT2 + neurons (Red, rabies-dsRed; blue, DAPI, scale bar, 200 μm). ACC, Anterior Cingulate Cortex; BnST, Bed nucleus of the Stria Terminalis; CEA, Central nucleus of the Amygdala; CNF, Cuneiform Nucleus; DMH, Dorsomedial Nucleus of the Hypothalamus; DRN, Dorsal Raphe Nucleus; IL, Infralimbic Cortex; LC, Locus Coeruleus; LDTg, Laterodorsal Tegmental Nucleus; LH, Lateral Hypothalamus; LHb, Lateral Habenula; LS, Lateral Septum; MHb, Medial Habenula; NAC, Nucleus Accumbens; dPAG, Periaqueductal Gray; iPAG, lateral Periaqueductal Gray; vPAG, ventrolateral Periaqueductal Gray; PBN, Parabrachial nucleus; PL, Prelimbic Cortex; PPTg, Pedunculopontine tegmental nucleus; PVN, Paraventricular Nucleus; RMTg, Rostromedial Tegmental Nucleus; VMH, Ventromedial Hypothalamus; VP, Ventral Pallidum.
retrogradely transported from NAc and LHb terminals to VTA cell bodies (Fig. 5b, c), confirming previous studies in the literature [15,23]. We found 28.99 CTB-594-positive neurons in VTA (Anterior = 10.33 cells; Middle = 9.66 cells; Posterior = 9 cells; Fig. 5d), and 2.66 CTB-488-positive neurons (Anterior = 2.66 cells; Middle = 2 cells; Posterior = 0.66 cells), revealing that NAc and VTA receive VTA projections. The distribution of CTB along anteroposterior axis of VTA revealed both NAc and LHb received homogeneous projection from VTAVGluT2+ neurons (Fig. 5d, e). Only a minority of labeled neurons coexpressed VGluT2 and CTB marker (0.11% for NAc, and 0.04% for LHb), revealing that glutamatergic neurons account for a minority of VTA projections to NAc and LHb. Importantly, we did not observe VTAVGluT2+ neurons expressing both CTB-494 and CTB-488 (Fig. 5b), suggesting that VTAVGluT2+ neurons do not send collaterals to these two aversive-response associated downstream VTA targets. But nine cells expressing simultaneously CTB-594 and CTB-488 were found, demonstrating that VTA non-glutamatergic neurons can send collateral to NAc and LHb.

Finally, to check and detail the VTAVGluT2+ network, we mapped the structures projecting to VTAVGluT2+ neurons using Cre-dependent monosynaptic retrograde tracing. VGluT2-ires-Cre transgenic mice received AAV-CAG-HISTO-GFP (AAV2/9) and AAV-CAG-DIO-RC (AAV2/9) virus injection into VTA. Three weeks later, VTA was infected with RV-EvNA-DsRed (EnvA pseudotyped, G-deleted and DsRed-expressing rabies virus) using the same coordinates (Fig. 6a). Mice were sacrificed one week after this second injection and injection sites were verified as VTA (Fig. 6b). Neurons projecting to VTAVGluT2+ neurons were defined as expressing red retrogradely label virus only (Fig. 6d), allowing a precise count of cells in each upstream target (Fig. 6c). VTAVGluT2+ upstream projections were particularly strong from the Dorsal Raphe Nucleus (DRN, 293.3%), Lateral Hypothalamus (LH, 106.4%), and the Medial Habenula (MHB, 166.5%). The Lateral Habenula (LHB, 129.7%), Rostromedial Tegmental Nucleus (RTmG, 107.2%), Laterodorsal Tegmental Nucleus (LDTg, 101.9%), and Periaqueductal Gray (PAG, 181.93%, including vPAG, pPAG and dPAG) were also particularly strong, as well as NAc (122.6%, including NACshell and NACcore).

Together, these data show that, on one side, a majority of projections to VTAVGluT2+ arise from DRN, LH and MHB. On the other side, VTAVGluT2+ neurons project to NAc and LHb, which represents only a minority of VTA projections to these structures. While VTA send collateral to NAc and LHb, they do not originate from VTAVGluT2+ neurons.

4. Summary of results

We used fiber photometry to investigate how the Ventral Tegmental Area glutamatergic neuronal population was recruited by unconditioned and conditioned aversive and rewarding stimulation. We demonstrated that VTAVGluT2+ population was activated by both aversive and rewarding unconditioned stimulation, with a response amplitude remaining stable across trials. During the conditioning protocol, CS-evoked responses gradually increased over trials, briefly for aversive conditioning and in a sustained manner for rewarding conditioning; in parallel, US-evoked activity remained stable. During a retrieval test, CS-evoked and expected-US neuronal activities remained strong only for the aversive conditioning protocol, but not for the rewarding protocol. This suggests that aversive and rewarding conditioning signals are integrated by VTAVGluT2+ neurons through different mechanisms. Finally, to help better characterize VTAVGluT2+ neurons based on their connectivity pattern, we injected a CTB retrograde tracer in LHB and NAc nuclei, which revealed that only VTA non-glutamatergic neurons send collaterals to NAc and LHb. In parallel, by injecting rhabies retrograde tracer in VGluT2+ animals, we identified that VTAVGluT2+ neurons received inputs from variety of brain structures, with especially strong inputs from DRN, LH and MHB.

5. Discussion

5.1. Fiber photometry and neuronal response

We used fiber photometry to perform a systematic exploration of VTAVGluT2+ neurons at the population level. Photometry is a recording method used to index synchronous neuronal activity, through measures of intracellular variations of calcium concentration [21]. Although the GABAPs signal mainly corresponds to soma response [19], changes at the level of the terminal fields can also be detected [24]. Consequently, we cannot rule out that small proportion of the signal we recorded may originate from terminals of locally infected VGluT2 VTA neurons, forming synapses with neighboring dopamine or GABA neurons. Finally, unlike single neuronal recordings, variation in calcium signal can also be interpreted as a function of number of active neurons. An increase of fluorescence may reflect that more neurons are recruited by a neuronal process.

However, use of both single neuron and population recording have disadvantages because VTAVGluT2+ is a heterogenous population [10]. Indeed with photometry alone, subgroups of VTAVGluT2+ neurons that have a weak contribution to the signal are difficult to distinguish and isolate. Looking ahead, it will become very useful to combine photometry with technologies such as electrophysiology [25] to allow simultaneous investigation of single unit and population responses. This strategy could become a key method used to understand neural network function.

5.2. VTA glutamatergic neurons response to aversive and rewarding conditioning

This study focused on the population response of VTAVGluT2+ neurons during unconditioned and conditioned aversive or rewarding stimulation. We have demonstrated that: (1) VTAVGluT2+ exhibits a similar response to unconditioned aversive and rewarding stimulation and (2) for conditioned aversive and rewarding stimulation, during Training Day, CS-evoked activity gradually increased over trials, and US-evoked activity remained constant. The key finding here is the difference in terms of conditioned evoked responses during the Retrieval Day. For aversive conditioning, the amplitude of the CS-evoked responses increased gradually over trials, and there was a significant response at the expected time of footshock delivery. This behavior was not observed for reward conditioning.

By demonstrating that the VTAVGluT2+ population responds to both rewarding and aversive stimulation, our results are in line with optogenetic studies showing VTAVGluT2+ neurons can promote rewarding behaviors when directly stimulating VTA [16], and aversive behaviors when stimulating VTAVGluT2+ downstream targets [4,14]. These opposing functions may be explained if, for example, on one hand, VTAVGluT2+ projecting neurons encode aversion, whilst on the other hand, VTAVGluT2+ local neurons encode reward. A second hypothesis is that VTAVGluT2+ population is not a pure homogeneous population at the molecular level: indeed a small subpopulation of VTAVGluT2+ neurons coexpress VGluT2 and TH [10,26,27], and these potentially contribute to the responses we recorded. A last hypothesis is that VTAVGluT2+ neurons encode perceived saliency, like recently demonstrated in the paraventricular thalamus [28], but further investigation is required to account for the scalability of response with stimulus intensity.
Our data also complements another recent electrophysiological study showing that VTAVGlut2+ neurons are sensitive to both aversive and rewarding stimulation [18]. This provided a precise characterization of individual VGlut2 neurons based on their individual response pattern: they revealed that most VGlut2 neurons increased firing rate during aversive stimulation and decreased during rewarding stimulation [18]. In addition, they also showed that the majority of VTAVGlut2+ neurons decreased their activity during reward, and only a small fraction was activated by both reward and aversion. Our results are in accordance with and support their findings that VTAVGlut2+ neurons increase activity during unconditioned aversive stimulation. Another of their findings, that most of VTAVGlut2+ neurons decreased activity during reward, appears at odds with our results. However, this can perhaps be explained by the fact that the GCaMP6s signal is mainly correlated with neuronal activity, and it is virtually insensitive to inhibition [19]. Consequently, the increase of calcium signal we recorded during reward was likely driven mainly by the small VTAVGlut2+ subpopulation described as responsive to both aversion and reward [18], and thus may not reflect the other subpopulation that is inhibited by reward. However, further investigation combining population and single neuron recording is required to test this hypothesis.

We next asked what the VTAVGlut2+ neuronal response to aversive conditioning and its retrieval are. We demonstrated that CS-evoked population response gradually increased across trials, whilst in parallel, the US-evoked response remained stable. We also demonstrated that VTAVGlut2+ neurons strongly respond to CS and expected-US 24 h after aversive conditioning. Some of these VTAVGlut2+ features, in particular the gradual increase of response to CS over trials, and response to an expected stimulation, may resemble VTA dopamine neurons [2,29–31]. One possible explanation for our results arises from the finding that a subpopulation of VTA glutamatergic neurons, VGlut2 neurons, can coexpress VGlut2 and TH [10,26,27]. Indeed, we cannot exclude that these specific VGlut2+ TH neurons contributed to the pattern of response we recorded and subsequent studies will be necessary to give a definitive answer to this question. Another non-exclusive explanation may be that DA and VGlut2 neurons form connections at the microcircuit level [13], especially modulating the VGlut2 population response during CS.

Finally, CS preceding-reward or CS preceding-aversive responses are slightly different, the former being sustained and the later brief. In addition, and contrary to the aversive experiment result (Fig. 2I), the retrieval-evoked response to the CS, following rewarding conditioning, remained extremely weak, if not absent (Fig. 4H). This suggest that VGlut2 neurons responding to reward and aversive conditioning may belong to segregated subpopulations, probably in part corresponding to the different types of VTAVGlut2+ neurons recently characterized [18]. These divergences may be in part explained by specific connectivities of VTAVGlut2+ neurons sensitive to reward and aversion, in particular at the microcircuit level where interactions with dopamine and GABA neurons are known to exist [13]. For example, we can hypothesize that local VTAVGlut2+ neurons encode rewarding functions, while VTAVGlut2+ projections encode aversive ones. In parallel, it is important to note that fiber photometry methods alone do not allow observation of the contribution of a minority activity within a population signal. Consequently, we cannot exclude that a small group of VTAVGlut2+ neurons may similarly respond to both reward and aversive conditioning, whilst remaining invisible at the population signal level. Future experiment involving a combination of electrophysiology and calcium imaging should compare the functional and response profiles of these VTAVGlut2+ neurons, especially focusing on neurons that have diverging connectivity patterns.

Together, we showed that across trials, VTAVGlut2+ neurons are similarly recruited during rewarding and aversive unconditioned stimulation, but differences emerge during conditioning. In particular, retrieval responses diverged, which could suggest that a majority of neurons responding to reward and aversion conditioning may belong to different subpopulations. Investigating the response of VTAVGlut2+ neurons during conditioning could help understand VTA function, including mechanisms that sustains learning and expectation in DA and GABA neurons.

5.3. Understanding VTAVGlut2+ neurons based on their network

We used CTB retrograde tracing to investigate VTAVGlut2+ projections to NAc and LHb and found that VTAVGlut2+ represent a minority of projections to these structures. Of particular importance, we showed that, whilst VTA sends collaterals to NAc and LHb, these collaterals do not arise from VTA glutamatergic populations. Knowing that both VTAVGlut2+–to-NAc and VTAVGlut2+–to-LHb pathways are known to serve aversive function [16,26], our results raise the question of their individual functional characteristics. In particular, it would be important to compare VTAVGlut2+–to-NAc with VTAVGlut2+–to-LHb activity patterns during aversive stimulation.

We next used RV tracing to map the inputs of VTAVGlut2+ neurons and observed that particularly strong projections to VTA glutamatergic neurons were coming from DRN, LH and MHb. Our data are consistent with previous studies [15,23], and confirm that structures such as LHb or NAc also sends projections specifically to VTAVGlut2+ populations, which may supply feedback that promotes aversive behaviors. It is known that DRN and PAG send projections to VTA-DA and VTA-GABA neurons linked to aversive and rewarding behaviors [32,33]; however, we observed that these structures also send parallel projections to VTAVGlut2+, whose function remains unknown.

The diversity of VTAVGlut2+ neurons inputs and outputs support the idea that VTAVGlut2+ function is not only based on their molecular background, but also on the network the belong to [10]. For example, functions and response patterns of dopamine neurons are highly heterogeneous and could depend of their specific projecting pattern [3,10,34]. In particular, knowing VTAVGlut2+ neurons stimulation can be either aversive or rewarding, we can posit that VTAVGlut2+ projections promote aversion, while local VTAVGlut2+ neurons promote reward, likely via neighboring connections to DA and GABA neurons. Alternatively, it has been demonstrated that, depending of the pattern of optogenetic activation, VTA glutamatergic projections to NAc, LHb and ventral pallidum can drive both place avoidance and promote appetitive self-stimulation [35]. Together with the possibility of distinct subpopulations encoding reward and aversion, the question arises of how, and under which condition, these subpopulations are recruited. One may think that the VTAVGlut2+ pattern of activity could act at VTA microcircuit level by participating in the preferential activation of a subpopulation of glutamatergic neurons. An interesting future direction would be to specifically target these potential subpopulations based on their connection patterns to compare their activity profile and their molecular background.

In summary, we used fiber photometry to demonstrate that VTAVGlut2+ neuronal population response to aversive and rewarding conditioning are divergent, especially during retrieval of conditioning. This suggests that VTAVGlut2+ populations responding to reward or aversive conditioning may belong to different subpopulations. We posited that these subpopulations may be better characterized by their associated network rather than by their molecular background. In the future, investigating VTAVGlut2+ neurons based on their local or long-range connectivity pattern may be
important to better understand VTA function. In particular, deciphering the function of VTAVGluT2+ at the microcircuit level would shed new light on our understanding of how local VTA DA and GABA neurons process reward and aversive conditioning.

**Conflict of interest**

The authors declare that they have no conflict of interest.

**Acknowledgments**

We thank Ji Hu for providing the VGluT2-Cre mice. This work was supported by the National Natural Science Foundation of China, China (31630031, 81425010, 31471109, 31671116, and 31500861), International Partnership Program of Chinese Academy of Sciences, China (172644KYSB20170004), Helmholtz-CAS Joint Research Grant (GJHZ2015B), Guangdong Provincial Key Laboratory of Brain Connectome and Behavior, China (2017B030301017), Shanmen Governmental Grants, China (JCY20160424919027063, KJQXSC20160301144002, JCY20170413164535041, JCY2015041150223647, JGCG201604249190521240), Research Instrument Development Project of the Chinese Academy of Sciences, China (YJKYYQ2017064); Youth Innovation Promotion Association of Chinese Academy of Sciences (2017413), Shanmen Municipal Funding, China (GJHZ20160222900136090), Shanmen Discipline Construction Project for Neurobiology, China (DRSCM [2016]1379), Ten Thousand Talent Project, Guangdong Special Support Program, China and Science and Technology Planning Project of Guangdong Province, China (2018B030331001).

**Author Contributions**

M.Q., Z.Z., Z.L. and X.L. contributed equally to this work. M.Q. and Z.Z. designed and initiated the project. Z.Z. performed virus injections and fiber implantation. Z.L., P.Z., and K.H. set up the behavior protocol. Z.L., P.Z., and C.C. performed photometry experiments. Y.L., C.C. and K.H. helped to collect the data. M.Q. and S.L.P. processed and analyzed photometry data. X.L., Z.L. P.Z. and Y.L. performed immunohistochemistry and quantitative analysis of the tracing data. M.Q. and Z.Z. interpreted the results. M.Q, Z.Z., Z.L. and X.L. SL.P. and L.W commented the manuscript. M.Q., Z.Z. and L.W. wrote the manuscript. L.W. supervised all aspects of the project.

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Zheng Zhou is a Ph.D. candidate student in the Brain Cognition and Brain Disease Institute of SIAT, Chinese Academy of Sciences (CAS). He received his Ph.D. degree in 2019. He worked as research assistant at Institute of Neuroscience, CAS from 2008 to 2011. His major research interests are focused on the neural circuitry basis of negative emotion and mental disorders.

Zhuogui Lei was a research assistant in the Brain Cognition and Brain Disease Institute of Shenzhen Institutes of Advanced Technology (SIAT), Chinese Academy of Sciences (CAS) from 2016 to 2018. His major research interest focused on the neural circuitry basis of fear emotion. Currently, he is a Ph.D. student in the Department of Biomedical Science, City University of Hong Kong.

Xuemei Liu is a Ph.D. candidate student and an assistant investigator in the Brain Cognition and Brain Disease Institute of SIAT, Chinese Academy of Sciences (CAS). Her major research interest are: (1) the neural circuits of the innate defensive behaviors evoked by visual threat signal; (2) the distinct neural mechanism of the adaptive and non-adaptive innate defensive behaviors.

Liping Wang is a director of the Brain Cognition and Brain Disease Institute of SIAT, Chinese Academy of Sciences (CAS). He received his Ph.D. degree in medical neuroscience from Charité-University Medicine, Berlin, Germany (2005). Then he started his postdoctoral training at the Department of Bioengineering at Stanford University with Prof. Dr. Karl Deisseroth (2005-2008). His research interests is to understand the neural circuitry basis of emotion and mental disorders, which is focused on: (1) determining the structure and function of brain specific neuronal subtypes control specific emotional behavior, including innate fear, learned fear and anxiety; (2) elucidating the modified neural circuits properties in mental disorders, such as schizophrenia and post-traumatic stress disorder.