Trans-inhibition of axon terminals underlies competition in the habenulo-interpeduncular pathway

Graphical abstract

Highlights

- Calcium signals in habenular (Hb) soma do not reflect axon terminal activity
- Synchronized Hb cholinergic activity inhibits non-cholinergic terminals
- Presynaptic inhibition of non-cholinergic terminals is mediated by GABA$_B$ receptors
- GABA$_B$ R activation by retrograde signaling from the interpeduncular nucleus

In brief

Zaupa et al. show that calcium signals in habenular soma are not reflective of activity in axon terminals at the interpeduncular nucleus. They uncover an atypical mode of lateral inhibition between two habenular circuits where the activation of cholinergic neurons inhibits non-cholinergic activity by retrograde signaling from their target site.

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Trans-inhibition of axon terminals underlies competition in the habenulo-interpeduncular pathway

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SUMMARY

Survival of animals is dependent on the correct selection of an appropriate behavioral response to competing external stimuli. Theoretical models have been proposed and underlying mechanisms are emerging to explain how one circuit is selected among competing neural circuits. The evolutionarily conserved forebrain to midbrain habenulo-interpeduncular nucleus (Hb-IPN) pathway consists of cholinergic and non-cholinergic neurons, which mediate different aversive behaviors. Simultaneous calcium imaging of neuronal cell bodies and of the population dynamics of their axon terminals reveals that signals in the cell bodies are not reflective of terminal activity. We find that axon terminals of cholinergic and non-cholinergic habenular neurons exhibit stereotypic patterns of spontaneous activity that are negatively correlated and localize to discrete subregions of the target IPN. Patch-clamp recordings show that calcium bursts in cholinergic terminals at the ventral IPN trigger excitatory currents in IPN neurons, which precede inhibition of non-cholinergic terminals at the adjacent dorsal IPN. Inhibition is mediated through presynaptic GABA_B receptors activated in non-cholinergic habenular neurons upon GABA release from the target IPN. Together, the results reveal a hard-wired mode of competition at the terminals of two excitatory neuronal populations, providing a physiological framework to explore the relationship between different aversive responses.

INTRODUCTION

The bilaterally paired habenular nuclei (Hb) are part of the highly conserved dorsal diencephalic conduction system connecting the limbic forebrain with monoaminergic brainstem areas.1,2 Anatomically, they consist of medial (MHb) and lateral (LHb) nuclei in mammals that are homologous to the dorsal (dHb) and ventral (vHb) nuclei of teleosts. The LHb and vHb send projections through the fasciculus retroflexus (FR) directly to serotonergic raphe neurons and dopaminergic neurons in the ventral tegmental area. Axons from MHb and dHb neurons course through the FR and terminate at an unpaired midbrain target, the interpeduncular nucleus (IPN). The Hb-IPN pathway is critical for understanding addiction and mood disorders due to its involvement in regulating behaviors, such as stress, anxiety, and fear learning.1,3–6

The MHb/dHb glutamatergic neurons are divided into those that co-release acetylcholine (cholinergic) or neuropeptides, including substance P (non-cholinergic) or somatostatin.7–9 In rodents, cholinergic and non-cholinergic neuronal populations project to central and peripheral IPN domains, respectively.7 Gene expression analyses, fluorescent dye tracing, and transgenic reporters have demonstrated left-right differences between the bilateral dHb of larval zebrafish.5,10,11 For example, cholinergic neurons are located mainly in the right dHb and project to the ventral IPN (vIPN), whereas non-cholinergic neurons are more abundant in the left dHb and innervate the dorsal IPN (dIPN) (Figure 1A).9,12 In rodents, cholinergic and non-cholinergic MHB neurons receive afferent input from neurons located in the triangular septum (TS) and the bed of the anterior commissure (BAC), respectively.13 Toxin-induced ablation of TS-cholinergic or BAC-non-cholinergic MHB projections selectively impaired anxiety or fear learning behaviors.3 Conditional genetic deletion of cerebellin (Cbln) genes in the MHB, Cbln4 in cholinergic, or Cbln2 in non-cholinergic neurons affects freezing behavior or spatial learning, respectively.4 The dHb responds differently to external factors, such as light, odor, and aversive stimuli in zebrafish larvae.6,14,15 Inhibition of
Figure 1. Identification of functional domains in dHb-IPN pathway during spontaneous activity

(A) Schematic of dHb-IPN pathway of larval zebrafish. Non-cholinergic (nCh) (red) neurons are mainly in the left dHb and project to the dorsal IPN (d), whereas cholinergic (Ch) (green) neurons are largely in the right dHb and project to the ventral IPN (v) and raphe (bracket). Somatostatinergic neurons are depicted in blue. Anterior (a), posterior (p), left (L), and right (R) are shown.

(B, F, and J) Representative 6 days post-fertilization (dpf) Tg(gng8:GAL4ff)c426;Tg(UAS:GCaMP7a) brain explants expressing GCaMP in dHb neurons (B), left dHb and their terminals (F), and terminals at the IPN and raphe (J), overlaid by colored circles representing functional domains identified by k-means clustering. Dotted line depicts the fasciculus retroflexus (FR) in (F).

(C, G, and L) Correlation matrix of average ΔF/F signals between different clusters of dHb soma (C), dHb soma and terminals (G), and terminals at the IPN and raphe (L).

(D, H, and M) Average ΔF/F traces from each cluster in 10-min recordings. Colored boxed events in (D) correspond to single-frame images in Figure S1.

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neural transmission in the non-cholinergic (dHbL) versus cholinergic (dHbM) pathways in adult zebrafish is associated with winner or loser states in a social conflict paradigm. Studies in rodents and zebrafish suggest that the cholinergic and non-cholinergic MHy/dHb neuronal populations form functionally distinct, parallel circuits. However, it is unknown whether the two pathways interact or directly influence the activity of one another.

To understand functional connectivity of the dHb-IPN pathway, we took advantage of the small size of the larval zebrafish brain to monitor neural activity in whole explant preparations. Calcium imaging revealed that dHb terminals at the IPN can be resolved into distinct domains based on their signaling dynamics. We discovered an inverse relationship between cholinergic and non-cholinergic neurons only at their axon terminals. Results from pharmacology, imaging, and electrophysiology experiments show that synchronized activity of cholinergic dHb neurons results in GABA release from the IPN, leading to presynaptic inhibition of non-cholinergic axon terminals. We propose a hardwired mode of competition between two neuronal populations, whereby synchronized activation of one group inhibits the activity of the other.

RESULTS

Cholinergic and non-cholinergic dHb terminals exhibit negatively correlated calcium events

The bilateral dHb on the dorsal surface of the diencephalon form connections with the IPN deeply embedded in the ventral midbrain, making it challenging to simultaneously monitor and manipulate both pre- and postsynaptic neuronal populations. To identify the hardwired mode of action of the dHb-IPN pathway, we analyzed spontaneous activity in both regions using brain explants from larval zebrafish. The entire brain was removed from TgBAC(gng8:GAL4ff)c426;Tg(UAS:GCaMP7a)m (B and J) or 50m (F). See also Figures S1–S4.

Calcium signaling data also revealed a local negative interaction between dHb terminals at the IPN. The bilateral dHb on the dorsal surface of the diencephalon form connections with the IPN deeply embedded in the ventral midbrain, making it challenging to simultaneously monitor and manipulate both pre- and postsynaptic neuronal populations. To identify the hardwired mode of action of the dHb-IPN pathway, we analyzed spontaneous activity in both regions using brain explants from larval zebrafish. The entire brain was removed from TgBAC(gng8:GAL4ff)c426;Tg(UAS:GCaMP7a)m (B and J) or 50m (F). See also Figures S1–S4.

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1N, S2A, and S2B). The higher signals were not merely due to greater terminal density at the dIPN relative to the vIPN because we found no difference in the ratio of axon terminals in the two regions when labeled with GFP (Figures S2A’ and S2B).

Quantification of calcium signaling kinetics revealed that the time to reach maximum amplitude (peak time) occurred earlier in terminals at the vvIPN compared to the raphe (1.13 ± 0.42 s versus 1.59 ± 0.72 s), and the duration at half peak time (half duration) of the calcium burst was shorter (2.72 ± 0.56 s versus 3.64 ± 1.55 s; Figures S3A–S3C). Peak inhibition in terminals at the dIPN occurred later (4.33 ± 1.4 s) and lasted longer (9.2 ± 2.9 s) than the calcium bursts (Figures S3D and S3E). Activation at the vIPN was followed by a period of inhibition not observed in terminals at the raphe. Therefore, to resolve the relationship between the negatively correlated events, we measured activity in terminals at the dIPN and at the raphe as a proxy for the vvIPN. We found a 931 ± 50 ms delay between the onset of the calcium burst and inhibition (Figure 2A), suggesting that calcium activity at the vvIPN triggers calcium inhibition in the dIPN.

Figure 2. Synchronized cholinergic terminal activity at vIPN/raphe precedes non-cholinergic terminal inhibition at dIPN
(A) Average ΔF/F trace (black) of negatively correlated events (gray) in dHb terminals at the dIPN (top) and raphe (bottom). Time of increase (green) and inhibition (red) is indicated (dotted lines). Note the delay between the two events (931 ± 50 ms; n = 5 explants, 62 events).
(B) Sagittal sections of anti-GFP (green) and anti-VACHT (magenta) labeling in 6 dpf Tg(nptx2:Gal4-VP16)rw0143a;Tg(UAS:GFP)c426 (left) and Tg(gng8:GAL4ff)c426;Tg(UAS:GFP)c426 (middle) larvae. White arrowheads indicate VACHT labeling in vIPN and raphe. White box (middle) corresponds to magnified panels on right. Scale bar: 20 μm.
(C) Summary of dHb terminal domains. nptx:Gal4;UAS:GFP-positive dIPN domain is indicated (nptx*). The cvIPN is only identified based on calcium signals. Calcium bursts and inhibition are labeled with graph plots.
(D) Electric shock induces negatively correlated events. (Left) Lateral view of Tg(gng8:GAL4ff)c426;Tg(UAS:GCaMP7a) larva expressing GCaMP in dHb terminals is shown. dIPN, vIPN, and raphe are outlined (dotted lines). (Middle) Representative ΔF/F traces upon electric shock (black dotted line) are shown. (Right) Average traces (black) of calcium events (gray) after electric shock are shown (n = 8 larvae, 16 events). Scale bar represents 20 μm.

We confirmed that GFP expression does not colocalize with VACHT protein (Figure 2B), whereas dHb terminals at the vIPN and raphe are positive for VACHT staining in TgBAC(gng8:GAL4ff)c426;Tg(UAS:GFP)c554 larvae (Figure 2B). We therefore conclude that calcium bursts at these regions are due to the synchronized activation of dHb cholinergic neurons (Figure 2C). Negatively correlated events are already apparent in 3-day-old larvae (Figure S4) upon emergence of spontaneous activity in dHb neurons (data not shown), suggesting that it is a hardwired mode of activity, coincident with establishment of the dHb-IPN pathway.

To determine whether external stimuli provoke negatively correlated events in vivo, we applied a mild electric shock to larvae. Indeed, we observed negatively correlated events between dHb terminals at the dIPN and vIPN/raphe (Figure 2D). However, due to tissues surrounding the brain, the resolution of calcium activity was inferior in larvae compared with the brain explants. Nevertheless, this result indicates that negatively correlated events can be induced in vivo, suggesting it is a physiologically relevant mode of neuronal activation.

Nicotinic acetylcholine receptors do not mediate negatively correlated events
The MHB/dHb-IPN pathway shows the highest expression and greatest variety of nicotinic acetylcholine receptor...
Because calcium inhibition in non-cholinergic dHb terminals is preceded by a calcium burst in cholinergic terminals, we investigated whether inhibition depends on the activation of nAChRs.26 Perfusion of nicotine resulted in prolonged calcium inhibition of dHb terminals at the dIPN, which corresponded with an extended series of calcium bursts in dHb terminals at the raphe (Figures 3 A and 3A'); 260 ± 176 s. To determine the contribution of nAChRs in the generation of negatively correlated events, we administered mecamylamine and d-tubocurarine, effective blockers of cholinergic transmission in the larval IPN.26 nAChR antagonists did not attenuate but rather increased the frequency of negatively correlated events in dHb terminals at the raphe (green, bursts) and dIPN (red, inhibition). One-way ANOVA followed by Tukey's test; **p < 0.005. b, 15 min of baseline; I, 0–15 min after application; II, 15–30 min after application; III, 30–45 min after application.

**GABAergic inhibitory postsynaptic currents.** The cholinergic/glutamatergic postsynaptic barrage was relatively short (half-duration = 262 ± 124 ms) and synchronized with the onset of the calcium burst (delay = 67 ± 69 ms). In contrast, the barrage of GABAergic synaptic currents had a delayed onset (delay = 162 ± 80 ms) and lasted longer (half-duration = 387 ± 143 ms). These results suggest that calcium bursts correspond to the release of glutamate and acetylcholine from dHb terminals at the IPN, which is followed by local release of GABA from IPN neurons.

**Presynaptic GABA<sub>A</sub> receptors in dHb axon terminals mediate calcium inhibition**

GABA has an inhibitory presynaptic action on transmitter release through metabotropic GABA<sub>A</sub> receptors.27 We asked whether GABA released by the IPN could activate presynaptic metabotropic GABA<sub>A</sub> receptors, thereby repressing dHb terminal activity. Application of cgp55845, a GABA<sub>A</sub> receptor antagonist, resulted in a 74% ± 9% reduction in the amplitude of calcium inhibition (0.16 ± 0.03 versus 0.04 ± 0.01) without affecting the amplitude of calcium bursts (0.22 ± 0.09 versus 0.26 ± 0.13; Figures 5A and 5A'). To test the role of GABA<sub>A</sub> receptors directly, we applied tetrodotoxin to inhibit the propagation of action potentials and eliminate all negatively correlated events. We...
then administered baclofen, a GABA\(_B\) receptor agonist, and observed a prolonged decrease in the calcium baseline, similar in amplitude to that observed during inhibition of dHb terminals at the dIPN (Figures 5B and 5B'). Calcium imaging of dHb terminals together with patch-clamp recordings of IPN neurons confirmed that prolonged inhibition corresponds with a decrease in the frequency of fast postsynaptic inward currents recorded in the IPN neurons (Figures 5C and 5C').

In zebrafish, there are two GABA\(_B\) receptor genes, GABA\(_B\) receptor 1a (gabbr1a) and 1b (gabbr1b).\(^{19}\) We found that gabbr1a, but not gabbr1b, is prominently expressed in the dHb in a larger domain of the left nucleus compared with the right, in the location of non-cholinergic neurons (Figures 5D and 5E). Strikingly, gabbr1a transcripts localized to the dHb nuclei and were not detected in other brain regions, including the IPN (Figure 5D). Together, these results suggest that calcium inhibition is due to the activation of presynaptic GABA\(_B\) receptors in non-cholinergic terminals at the dIPN.

Retrograde signaling from GABAergic IPN neurons to non-cholinergic habenular inputs

To determine that inhibition of dHb terminals at the dIPN results from GABA release upon IPN activation, we carried out electrical stimulation of the IPN with calcium imaging (Figure 6A; Video S4). Activation of the IPN effectively triggered negatively correlated events consisting of the vIPN/raphe calcium burst (likely due to the activation of the dHb terminals at the vIPN) and dIPN calcium inhibition (Figure 6B). We next perfused a GABA\(_B\) antagonist, cgpg55845, during negatively correlated events induced by electrical stimulation and found a 90% ± 13% decrease in calcium inhibition at the dIPN (−0.15 ± 0.07 versus −0.02 ± 0.02) without any change in calcium bursts at the raphe (0.22 ± 0.08 versus 0.2 ± 0.06; Figures 6B–6E).

To verify the presence of GABAergic neurons in the IPN, we imaged the TgBAC(gng8:GAL4iff)\(^{24,26}\), Tg(UAS:GFP)\(^{24,26}\), Tg(gad1b:|R|-GFP) larvae in which dHb terminals are labeled with GFP and GABAergic neurons by RFP.\(^{20}\) We found that nearly half of IPN neurons were labeled by gad1b:RFP (48% ± 2%; n = 3 larvae; N = 479 cells; Figure 6F), with more located in the dIPN than vIPN (79% ± 11% versus 21% ± 11%; n = 3 larvae; N = 232 gad1b:RFP\(^{+}\) cells). These results suggest that a higher concentration of GABA is released from neurons in the dIPN.

DISCUSSION

The dHb of zebrafish larvae respond to external factors, including light, odor, and aversive stimuli.\(^{5,14,15}\) A prerequisite to investigating the innate hardwired activity of the dHb-IPN pathway was eliminating the influence of external factors. We achieved this by performing calcium imaging using a whole brain explant preparation, in which central neural circuits, including the dHb-IPN pathway, remain intact. We found a stereotyped mode of spontaneous activity in dHb soma and axon terminals at the IPN.

Through pharmacological and electrophysiological manipulations, we determined that synchronized activation of dHb cholinergic neurons inhibit non-cholinergic neurons by an atypical mechanism of target-mediated, trans-inhibition at the level of axon terminals.

Negatively correlated events at axon terminals of cholinergic and non-cholinergic dHb neurons

Clustering algorithms are commonly used to segregate neurons into different functional groups during calcium imaging of neuronal soma.\(^{21–23}\) The k value and threshold for hierarchical clustering were determined from a previous study of the zebrafish dHb\(^{23}\) and manual inspection of the activity of individual neurons assigned to different clusters. We obtained similar, but not identical, results from k-means and hierarchical clustering methods. Nevertheless, the minor difference resulting from the clustering algorithms does not alter the conclusion showing that most dHb neurons on the right and a small cluster on the left show high levels of synchronized spontaneous activity. The location of these neurons corresponds to different neurotransmitter phenotypes: cholinergic and somatostatin,\(^{12}\) suggesting that dHb neurons expressing the same neurotransmitter function together.

In analyzing group dynamics of axon terminals, we found it important to also carry out manual inspection of raw data. For example, only calcium peaks at the vvIPN and not the dvIPN were determined from a previous study of the zebrafish dHb\(^{23}\) and manual inspection of the activity of individual neurons assigned to different clusters. We obtained similar, but not identical, results from k-means and hierarchical clustering methods. Nevertheless, the minor difference resulting from the clustering algorithms does not alter the conclusion showing that most dHb neurons on the right and a small cluster on the left show high levels of synchronized spontaneous activity. The location of these neurons corresponds to different neurotransmitter phenotypes: cholinergic and somatostatin,\(^{12}\) suggesting that dHb neurons expressing the same neurotransmitter function together.

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Figure 5. Calcium inhibition occurs via presynaptic GABA<sub>B</sub> receptors

(A and B) Representative ΔF/F traces in terminals at dIPN (top) and raphe (bottom) after application of (A) 1 μM cgp55845 or (B) 3 μM tetrodotoxin (TTX) + 2 μM baclofen. Time of drug application is indicated (vertical lines).

(A') Average peak amplitude before and after cgp55845 application in terminals at the dIPN (0.16 ± 0.03 versus 0.04 ± 0.01) and raphe (0.22 ± 0.09 versus 0.26 ± 0.13) during inhibition (top) and burst (bottom). n = 6 explants, 50 events. Red asterisks indicate attenuated calcium inhibition at the dIPN after cgp55845 application.

(B') Paired plots comparing average inhibition amplitude in control conditions (0.22 ± 0.07) to the average baseline decrease after baclofen application (0.24 ± 0.08), n = 7 explants, 46 events. Note abrupt decrease (red arrowhead) in baseline at the dIPN following baclofen.

(C) ΔF/F traces of INP (top) aligned with simultaneous whole-cell, patch-clamp recording trace (middle) upon baclofen perfusion. Time of baclofen application is indicated (black bar). Magnified traces of synaptic activity before (left) and after (right) baclofen (bottom panels) are shown.

(C') Paired plot of synaptic activity frequency before (4.29 ± 1.05 Hz) and after (0.47 ± 0.15 Hz) baclofen. n = 6 explants. Mean ± SD is indicated in (A'), (B'), and (C'). Wilcoxon signed-rank test. *p < 0.05.

(D and E) Dorsal (left) and sagittal (right) views of gabbr1a (D) and gabbr1b (E) expression at 6 dpf. dHb nuclei (red arrowheads) and absence of expression at INP (white arrowhead; D) are shown. Scale bars: 50 μm.
they are targeted by different dHb neurons. In conclusion, cholinergic neurons exhibit synchronized activity and project to the vviPN and raphe, which display negatively correlated events with axon terminals at the dIPN (Figure 2C).

Nicotine promotes negatively correlated events in the dHb–IPN pathway

We found that nicotine induces prolonged duration of the negatively correlated event. We previously showed that a cocktail of nAChR antagonists inhibits approximately 80% of cholinergic currents in IPN neurons. However, application of antagonists failed to attenuate but rather increased the frequency of negatively correlated events. The cholinergic system can increase the excitability of GABAergic interneurons in the neocortex and the lateral habenula. Likewise, our data suggest that GABAergic neurons are activated by nACHRs in the IPN. We hypothesize that blocking nACHRs throughout the whole brain inhibits GABAergic signaling, resulting in a global increase in synchronized network activity.

Although nACHR activation can generate a negatively correlated event, the release of GABA from IPN neurons to induce inhibition is predominantly triggered by glutamatergic rather than cholinergic transmission from the dHb terminals at the IPN during spontaneous activity.

Localization of the GABA<sub>β</sub> receptor to non-cholinergic terminals at the dIPN mediates presynaptic inhibition

We found that gabbr1a is expressed asymmetrically in a large area in the left and smaller area in the right dHb, indicating that the GABA<sub>β</sub> receptor is present in non-cholinergic neurons that innervate the dIPN in larvae. Moreover, gad1b-positive neurons predominate in the dIPN, suggesting that GABA released from dIPN neurons activates presynaptic GABA<sub>β</sub> receptors on non-cholinergic terminals at the dIPN. Based on calcium imaging, pharmacology, electrical stimulation, and molecular studies, we propose a model whereby (1) activation of cholinergic dHb neurons (2) activates GABAergic neurons in the IPN, resulting in the (3) release of GABA that through retrograde signaling (4) activates inhibitory presynaptic GABA<sub>β</sub> receptors on non-cholinergic dHb terminals at the dIPN, leading to (5) inhibition of glutamatergic transmission (Figure 6G).

How does activation of vviPN neurons induce GABA release by dIPN neurons? Single cell labeling experiments showed the diverse morphology of IPN neurons, including those that appear to be interneurons extending their processes between the dorsal and vviPN. We hypothesize that communication between vviPN and dIPN neurons could be mediated by such interneurons connecting the dorsal and ventral IPN. Another possibility is that more vviPN neurons are GABAergic than visualized by the transgenic line used in this study, akin to what has been shown for the adult IPN. Thus, GABAergic vviPN neurons might form direct synaptic connections or, through non-synaptic volume transmission, release GABA to inhibit non-cholinergic terminal activity at the dIPN.

As the MHB/dHb-IPN pathway is evolutionarily conserved in all vertebrates, it is likely that this mode of hardwired activity is not unique to zebrafish. Indeed, a recent study on mice reports that presynaptic GABA<sub>β</sub> receptors at non-cholinergic terminals at the IPN play an inhibitory role on plasticity, suggesting a conserved role for these receptors.

Model for competitive selection between dHb circuits on behavior

It is crucial for an animal to identify the most salient stimulus among many concurrent external stimuli and select a single behavioral response. One example is the flight, freeze, or fight response. During early stages of development, hardwired circuits to execute specific behavioral repertoires, such as escape, are essential for survival. The process of transforming multiple sensory inputs into a single behavioral outcome is termed competitive selection and includes selective attention and decision making. Computational formulations, such as winner-take-all models, have been proposed and recent studies have begun to elucidate the mechanisms for behavioral selection in competing circuits.

Higher baseline GCaMP fluorescence in the terminals at the dIPN relative to the vviPN suggests that non-cholinergic neurons display high-frequency oscillation. We propose that the synchronized burst in cholinergic terminals at the vviPN transiently inhibits this constitutive activity that occurs in non-cholinergic neurons, ensuring only one pathway is active at any given time.

What could be the advantage of having local inhibition at the terminal level? Different external stimuli preferentially activate left or right dHb neurons. However, the ratio of asymmetric dHb activation depends on the intensity of the stimulus for both light and electric shock (personal observations), suggesting that the left and right dHb can be co-activated by the same stimulus.

Susceptible behavior refers to emotional states with negative valence, such as fear or anxiety. Fear is defined as a response to a factual known threat while anxiety is an unknown, poorly defined threat. Brain areas that contribute to fear and anxiety...
exhibit great overlap, including the amygdala, medial prefrontal cortex, and the hippocampus. Although recent optogenetic and in vivo studies have begun unraveling the microcircuitry that participate in these behaviors, how they interact with other areas of the brain is unclear. The highly conserved MhB/dHb-IPN pathway consists of two circuits composed of different neuronal populations, which mediate fear- or anxiety-related behaviors. We propose that the trans-inhibition mechanism between cholinergic and non-cholinergic dHb-IPN circuits underlies competitive selection of aversive behavioral responses.

A brief electric shock induces fast swim followed by freezing behavior before returning to baseline locomotion in larval zebrafish. Here, we demonstrate that it can also promote negatively related behaviors. We hypothesize that activation of the cholinergic pathway by electric shock results in a stereotypical fast swim (flight) response, which is followed by inhibition of the non-cholinergic pathway, allowing the animal to freeze. Prior to puberty, a flight or freeze response would be more beneficial than a flight response. In juveniles and adults, this dual circuit likely becomes more complex as different neuropeptides become expressed in the dHb-IPN pathway and with the maturation of behavioral repertoires from prior experience. For example, adult zebrafish exhibit a fight or flight response, manifested by a “winner” or “loser” phenotype in a social conflict paradigm, which is correlated with potentiation in non-cholinergic and cholinergic neuronal terminals, respectively. We hypothesize that the “loser”/flight behavior corresponds to the default response to an aversive stimulus upon activation of the cholinergic pathway. The “winner”/fight behavior could result from preferential hyper-activation of the non-cholinergic pathway, corresponding to a heightened level of anxiety.

In conclusion, investigating neuronal population dynamics within an intact pathway has revealed an atypical mode of trans-inhibition between two excitatory neuronal populations at their axon terminals. This hardwired mode of competition could underlie the mechanism for competitive selection between these neural circuits and provide a physiological framework to explore the relationship between anxiety and fear.

STAR+METHODS

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AUTHOR CONTRIBUTIONS


DECLARATION OF INTERESTS

The authors declare no competing interests.

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REFERENCES

### KEY RESOURCES TABLE

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### Software and algorithms

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RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Elim Hong (elim.hong@inserm.fr).

Material availability
All plasmids generated in this study are available upon request to the Lead Contact.

Data and code availability
- All data reported in this paper will be shared by the Lead Contact upon request.
- This paper does not report original code.
- Any additional information required to reanalyze the data reported in this paper is available from the Lead Contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

AB wild-type and transgenic zebrafish lines TgBAC(gng8:GAL4ff)c426, Tg(UAS:GCaMP7a), Tg(nptx2:Gal4-VP16)rw0143a, Tg(gad1b:|-GFP), Tg(UAS:GFP)c354 5,9,20,42,43 were used. Fish were maintained at 28°C/14°C on a 14:10h light: dark cycle in a recirculating system. All experiments were carried out in agreement with the European Directive 210/63/EU on the protection of animals used for scientific purposes and the French application decree ‘D/C19ecret 2013-118’. The projects of our research group have been approved by the ethical committee ‘Comité d’éthique Charles Darwin’ (APAFIS#15909-2018070912072530 v5). The fish facility has been approved by the French ‘Service for animal protection and health’ (A-75-05-25).

METHOD DETAILS

Explant Dissection
TgBAC(gng8:GAL4ff)c426; Tg(UAS:GCaMP7a) 3, 6 and 7 day-post-fertilization (dpf) larvae were used for experiments. Larvae were anesthetized in Tricaine (0.01%) and brains dissected using fine forceps in ice cold Ringer’s solution (134 mM NaCl, 2.1 KCl, 1.2 mM MgCl2, 10 mM HEPES, 10 mM Glucose, 2.1 mM CaCl2, pH = 7.6) and mounted in 1.2% low melting agarose for calcium imaging and electrophysiology experiments.

Pharmacology
Drugs were applied to the explants either directly using a pipette or by using a perfusion peristaltic pump (Ismatec) at a rate of 5-6 ml/min. Mecamylamine hydrochloride (100 μM), (+)-D-tubocurarine chloride (10 μM), nicotine ditartrate (10 μM, Acros Organics), tetrodotoxin (3 μM), baclofen (2 μM) and cgp55845 (1 μM) were used. The drugs were purchased from Tocris Bioscience unless otherwise noted.

Calcium Imaging
Calcium imaging was performed on a spinning disk microscope (Zeiss Axio Examiner.Z1) using a 40X water immersion objective (NA = 0.95) or Leica SP5 laser scanning confocal microscope using a 25X water immersion objective (NA = 0.95). Images were acquired at a rate between 2.2-6.7 Hz, depending on the experiment.

Nicotine was perfused for 5 minutes followed by E3 medium to wash out the drug. For experiments with mecamylamine hydrochloride and D-tubocurarine chloride, as a short-term application did not show any changes in calcium signals, we carried out the following experiment to assay for long-term change: baseline calcium signals were recorded for 15 minutes before pipetting the drug directly to the explant. The calcium signals were recorded for an additional 45 minutes after drug application. For the cgp55845 and baclofen experiments, 10-15 minutes baseline activity was recorded followed by drug application using the perfusion system and imaged for another 15 minutes.
Patch-clamp recordings and simultaneous Ca$^{2+}$ imaging

Explants from TgBAC(gng8:GAL4ff)C426; Tg(UAS:GCaMP7a) larvae were placed in a recording chamber and constantly perfused with Ringer’s solution at a rate of 1–2 mL/min. The IPN was made accessible for patch-clamp recordings and Ca$^{2+}$ imaging by orienting the explants with the ventral surface upward. dHb terminals and the IPN were located and imaged using a 63x water immersion objective mounted on an epifluorescence microscope BX51W1 (Olympus) equipped with a 470 nm LED fluorescent excitation light. Patch electrodes were filled with an intracellular solution containing 130 mM CsMeSO4, 4 mM MgCl2, 4 mM Na-ATP, 0.3 mM Na-GTP, 10 mM HEPES, and 10 mM EGTA, 5 mM QX-314 adjusted to pH 7.2, 290 mosm, for resistances between 5 and 8 MΩ. This intracellular solution allowed discrimination of cationic excitatory currents and chloride inhibitory currents based on their distinct reversal potentials (E_{cation} = 0 mV; E_{Cl} = -60 mV). By recording the cells at -60 mV, EPSCs appeared as inward currents, whereas IPSCs were barely detectable. Conversely, IPSCs appeared as outward currents when recording the cells at a holding potential of -0 mV, while EPSCs became undetectable. Whole-cell recordings were obtained using a Multiclamp 700B amplifier connected to a digidata 1440 acquisition system monitored using pClamp 10 software (Molecular Devices). Analyses were performed using Clampfit 10 (Molecular Devices). For all experiments, data were filtered at 4 kHz during recording and the traces were digitized at 20 kHz.

Simultaneous time-lapse imaging of GCaMP fluorescence in dHb terminals was performed using either an Orca Flash4.0 (Hamamatsu, Japan) or Orca 03G (Hamamatsu, Japan) camera and recorded using the HC Image Live software (Hamamatsu, Japan) at an acquisition frequency of 5 Hz or 10Hz with an exposure time of 200 ms and 100 ms, respectively. Time-lapse and patch-clamp acquisition were synchronized via a TTL signal sent to the camera via the pClamp 10 software. Time-lapse frames were encoded at a 256x256 pixel resolution using a 16-bit gray scale. Time-lapse images were analyzed using the ImageJ software (National Institute of Health, USA) and fluorescence signals were plotted using Clampfit 10.

Immunohistochemistry

TgBAC(gng8:GAL4ff)C426; Tg(UAS:GFP)C354 6 dpf larvae were fixed in BT fix (4% PFA, 0.15mM CaCl$_2$, 4% sucrose in 1x PBS) overnight in 4°C for rabbit anti-VAChT (1/200, Synaptic systems) labeling. Larvae were embedded in 4% low melt agarose and sectioned to 50 μm using a vibratome (Leica, Inc). Floating sections were incubated in primary antibody in incubating solution (0.8% Triton-X in 1x PBS) for 3 days on a shaker in 4°C. The sections were washed and incubated in secondary anti-rabbit Alexa Fluor 594 (1:10000) with Hoechst 33342 (1:2000) overnight at 4°C on a shaker. TgBAC(gng8:GAL4ff)C426; Tg(UAS:GFP)C354, Tg(gad1b::R)-GFP larvae were fixed overnight in 4% PFA, sectioned to 50 μm and incubated with Hoechst 33342 (1:2000) overnight at 4°C on a shaker. All sections were then mounted on slides using Mowiol (Sigma). They were imaged using the Leica TCS SP5 A0BS upright microscope using a 63x (N.A. = 1.4) objective.

RNA in situ Hybridization

PCR fragments for gabbr1a (ENSDARG00000018967) (553-bp) and gabbr1b (ENSDARG00000016667) (539-bp) were subcloned into pCRII-TOPO vector using the TOPO®TA cloning Kit (Invitrogen). Reverse primer sequences were from Cocco et al. The following forward primers were used: gabbr1a (AGCTTGGTCTCTACGCG) gabbr1b (AGACAGAAGAAGACCTGGA). Restriction enzymes and RNA polymerase used to synthesize the antisense RNA probes are as follows: gabbr1a (KpnI/T7), gabbr1b (KpnI/T7). Colorimetric in situ hybridization (ISH) assay was performed as described in Thise and Thise Following ISH, the larvae were mounted in 100% glycerol and imaged on a Nikon Eclipse E800 microscope.

Electrical shock assay

Custom built electrical stimuli setup consisted of copper mesh electrodes that were mounted 6 cm apart on opposing sides and connected to a DS3 isolated constant current stimulator (Digitimer, Ltd.). Three electrical field stimulations (6mA, 20V, 500msec) were triggered every five minutes by MetaMorph (Molecular Devices). The larvae were paralyzed in 2% pancurium bromide and mounted in the middle of 90mm diameter Petri dishes filled with 30 mL E3 medium to ensure consistent stimulation. Shock assay was performed in 6-7 dpf agarose-restricted larvae, paralyzed with 2% pancurium bromide.

Electrical stimulation

The IPN neurons were stimulated using a low-resistance (< 1MΩ) glass pipette filled with Ringer solution containing an AgCl electrode. Train stimulations (20μs duration 50Hz stimulations lasting 400ms) were triggered every minute with an isolated current stimulator (DS3; Digitimer Ltd.) using a Master-8 pulse generator (A.M.P.I). Prior to each experiment, a dose-dependent curve was performed to select the optimal setting that induced negatively correlated events in the terminals at the IPN. Stimuli ranged between 100 µA - 1.2 mA.

QUANTIFICATION AND STATISTICAL ANALYSIS

Calcium data analysis

Average fluorescence intensity of habenular soma was calculated by manually drawing regions of interest (ROI) in FIJI as described in Duboué et al. To carry out a non-biased analysis of the dHb-IPN pathway and dHb terminals at the IPN, 2x2 μm$^2$ grid ROIs were drawn over the entire recording using FIJI macro. Average fluorescence intensity for each ROI was then extracted per frame and analyzed on MATLAB using custom written scripts.
Amplitude of calcium signal was calculated according to the formula:

\[ \Delta F/F = (F_i - F_{\text{mean}})/F_{\text{mean}}, \]

whereby, \( F_i \) is the mean intensity in a single ROI at a single time point while \( F_{\text{mean}} \) is the mean intensity in a single ROI throughout the entire recording time. Multiple clustering algorithms were tested including functional clustering algorithm\(^{45}\) and spectral clustering on calcium traces and correlation coefficient values. After comparing the clusters generated using various algorithms, we found \( k \)-means and hierarchical clustering based on pairwise linear correlation matrix were the most representative of calcium signals in the spontaneous activity recordings. \( k \)-means and hierarchical clustering based on pairwise linear correlation matrix were carried out on MATLAB using \textit{kmeans} and \textit{linkage} functions (‘euclidean’ distance and ‘complete’ method), respectively. Different \( k \) values were tested and \( k = 5 \) was selected based on manual inspection of calcium signals in the time-lapse recordings and also using a heatmap. The correlation matrix displayed in figures was generated by calculating the correlation coefficients between the average calcium signal traces for each cluster.

The threshold for hierarchical clustering was determined by manual inspection of the calcium signals in the recording and heatmap. Once specific IPN domains were identified by \( k \)-means, larger ROIs were manually drawn containing the different IPN domains. Analysis of calcium burst frequency using ‘findpeaks’ function was performed in MATLAB (Mathworks). Peak of calcium burst, calcium inhibition and half duration were identified using \textit{findpeaks} and \textit{half prominence} functions in MATLAB and verified manually. Calcium burst peak was defined by the highest value during a calcium burst. Peak of calcium inhibition was defined by the lowest value during a calcium inhibition event. Half duration was calculated as the width of the calcium signal at the half value of a peak during a calcium burst or inhibition. All graphs and plots were made on MATLAB using custom written scripts.

**Statistical analysis**

Statistical tests were carried out using a Wilcoxon signed rank test for averaged paired samples, Kruskal-Wallis test followed by a Dunn’s test for comparison between more than 2 groups, Mann-Whitney test was used with independent samples and a linear mixed model with fixed effect was used to compare multiple values from repeated events. All statistical tests were performed on R or GraphPad Prism (GraphPad Software, Inc.). \( p < 0.05 \) was considered as statistically significant. All statistical details including statistical tests, exact values of \( n \), value representation (mean ± SD) are indicated in the figure legends.