Selective Optogenetic Stimulation of Glutamatergic, but not GABAergic, Vestibular Nuclei Neurons Induces Immediate and Reversible Postural Imbalance in Mice

Montardy Q. 1,†, Wei M. 1, 2,†, Yi T. 1,†, Liu X. 1, Zhou Z. 1, 3, Lai J. 1, Besnard S. 4, 5, Tighilet B. 4,*, Chabbert C. 4, * & Wang. L. 1, *

1. 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

2. University of Chinese Academy of Sciences, Beijing, 100049, China

3. McGovern Institute for Brain Research, Department of Brain and Cognitive Sciences, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA

4. Aix Marseille University-CNRS, Laboratory of Sensory and Cognitive Neurosciences, UMR 7260. Team Pathophysiology and Therapy of Vestibular Disorders, Marseille, France. GDR Vertige (GDR#2074).

5. Université de Caen Normandie, CHU de Caen, Caen, France

† These authors contributed equally to this work

* To whom correspondence should be addressed: BT (brahim.tighilet@univ-amu.fr), CC (christian.chabbert@univ-amu.fr) & LW (lp.wang@siat.ac.cn)

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**Highlights**

- For the first time, Vestibular nuclei were optogenetically stimulated in free-moving animals, to assess for glutamatergic and GABAergic neurons functions in posturo-locomotor behaviors.
- Brief optogenetic activation of VNVGluT2+, but not VNGAD2+, induced immediate and strong postural deficit.
- Stimulation of VNVGluT2+ neurons provoked an imbalance with continuous effect on locomotion for a short period of time after stimulation.
- These results are comparable to the classical vestibular deafferentation models during their peak of deficit, and set optogenetic stimulation as a new model to study vestibular deficits.

**Abstract**

Glutamatergic and GABAergic neurons represent the neural components of the medial vestibular nuclei. We assessed the functional role of glutamatergic and GABAergic neuronal pathways arising from the vestibular nuclei (VN) in the maintenance of gait and balance by optogenetically stimulating the VN in VGluT2-cre and GAD2-cre mice. We demonstrate that glutamatergic, but not GABAergic VN neuronal subpopulation is responsible for immediate and strong posturo-locomotor deficits, comparable to unilateral vestibular deafferentation models. During optogenetic stimulation, the support surface dramatically increased in VNVGluT2+ mice, and rapidly fell back to baseline after stimulation, whilst it remained unchanged during similar stimulation of VNGAD2+ mice. This effect persisted when vestibular compensation was removed. Posturo-locomotor alterations evoked in VNVGluT2+ animals were still present immediately after stimulation, while they disappeared 1h later. Overall, these results indicate a fundamental role for VNVGluT2+ neurons in balance and posturo-locomotor functions, but not for VNGAD2+ neurons, in this specific context. This new optogenetic approach will be useful to characterize the role of the different VN neuronal populations involved in vestibular physiology and pathophysiology.
Introduction

The first mentions of vertigo as a specific symptom date back to antiquity with Hippocrates writings (Adams, 1849), and it was not until the beginning of the 19th century that the study of vertigo and balance entered the experimental phases, with the works of Purkinje in humans (Purkinje, 1819), and Flourens in the pigeon (Flourens, 1842). These led to the establishment of a direct link between damage to the inner-ear sensors and the characteristic posturo-locomotor and vestibulo-ocular deficits encountered in dizzy patients. A century later, Lorente de No introduced the first anatomical descriptions of the central pathways that connect the peripheral vestibular sensors to the oculomotor muscles (Lorente De No, 1933).

The anatomical and functional organization of the vestibular system is now better understood, and its contribution to a wide range of functions, from postural and oculomotor reflexes to spatial representation and cognition, has been underscored (Angelaki and Cullen, 2008; Borel et al., 2008). Head accelerations resulting from the interaction with our environment are detected by the mechanosensitive hair cells located in the inner ear sensory organs, and encoded into an electrical signal. This sensory information is then conveyed to the brain stem vestibular nuclei (VN) through the vestibular nerve. Four vestibular nuclei (VN) are located in the brainstem under the floor of the IVth ventricle: the median (MVN), inferior VN (IVN), lateral (LVN) and superior (SVN) vestibular nuclei. At the VN level, this essentially sensory information is integrated and transformed into premotor signals, the transport of which is ensured by vestibulo-spinal and vestibulo-oculomotor pathways. VN Neurons project, through the medial longitudinal fasciculus (MLF), onto the oculomotor nuclei whose motoneurons control gaze stabilization during head movements. The vestibular nuclei influence postural control via two descending pathways to the spinal cord, the lateral (mostly from ipsilateral NVL neurons) and medial (mostly from ipsi and contralateral NVM neurons) vestibulospinal tracts (Wilson and Jones, 1979). Vestibular information is also transferred via vestibulo-thalamo-cortical pathways to different cortical areas providing perceptual and cognitive functions such as perception and orientation of body in space (Lopez, 2016; Lopez et al., 2015). Finally, the vestibular nuclei are interconnected to a multitude of neurovegetative centers in the brainstem (Balaban and Beryozkin, 1994; Balaban and Porter,
1998). Given this anatomo-functional organization of the vestibular system and its projection targets, unilateral alteration of peripheral vestibular inputs induces a quadruple syndrome: posturo-locomo-tor, oculomotor, vegetative and perceptivo-cognitive. Symptoms progressively improve, each with its own kinetics, generally leading to an almost full disappearance of the syndrome. This behavioral recovery phenomenon is referred to as "vestibular compensation" (Lacour et al., 2016).

**Animal models in vestibular pathologies:**

Different animal models of vestibular disorders have been developed. Researchers set out to determine either the pathogenic conditions likely to induce vestibular dysfunction and its functional consequences on the basis of an epidemiological rationale, or the neurophysiological mechanisms likely involved. The first case included: 1) ototoxic-based vestibular damage, reproducing inner ear-specific toxicity of drugs such as aminoglycosides or cisplatin, and food-borne ototoxicity (Hirvonen et al., 2005; Vignaux et al., 2012; Xia et al., 2012; Zhang et al., 2003); 2) excitotoxically-induced vestibular damage, reproducing the deleterious consequences on the first synapse of sensory cells suffering (Brugeaud et al., 2007; Desmadryl et al., 2012; Dyhrfjeld-Johnsen et al., 2013; Gaboyard-Niay et al., 2016; O’Neill et al., 1999); 3) destruction of peripheral receptors through surgical labyrinthectomy (Hitier et al., 2010; Patk et al., 2003; Smith and Curthoys, 1989); 4) vestibular neurectomy that relates to the full section of the vestibular nerve. This later operation can be achieved either peripheral unilateral vestibular neurectomy (peripheral UVN; nerve section between vestibular endorgans and Scarpa’s ganglion) (Hamann et al., 1998; Li et al., 1995) or central UVN (between the Scarpa’s ganglion and the brainstem). Peripheral unilateral vestibular neurectomy (UVN) leads to the loss of inputs from vestibular endorgans, whilst preserving those arising from vestibular ganglion neurons. Central UVN induces complete loss of inputs arising from both vestibular endorgans and Scarpa ganglion neurons. Several examples of central UVN have been documented previously in different species such as monkeys (Lacour et al., 1976), cats (Xerri and Lacour, 1980) and rats (Li et al., 1995). Along with the lesion models mentioned above, there are models of reversible modulation of the vestibular input. The first to be developed was the so-called "vestibular anesthesia" (Bárány,
1936). Assuming that the episodes of vertigo attacks observed in Menière patients resulted from transient unilateral hyperexcitability of primary vestibular neurons, this approach consisted of counteracting this hyperexcitability through transtympanic administration of neuronal activity, such as lidocaine. A similar modulatory effect was obtained through transtympanic administration of tetrodotoxin (TTX), a selective blocker of voltage-gated calcium channels (Dutheil et al., 2011). Under these conditions, pharmacological modulation of peripheral sensory input either reproduced in animals the vestibular syndrome encountered in patients with acute peripheral vestibulopathy, or led to alleviation of symptoms when administrated to the ear opposite to the damaged one.

Although these models have led to a better understand of the conditions that generate vertigo syndrome, and its temporal evolution, questions related to the anatomical and functional organization of the vestibular central pathways remain. As an example, the target cerebral structures of neurons projecting from the VN have are not all been identified, in particular in the cortex. Likewise, the central pathways projecting from cortical areas towards the VN are poorly documented. This lack of data prevents the development of studies attempting to understand the links between vestibular, cognitive and emotional functions. To answer these questions, we used optogenetics in mice to stimulate glutamatergic and GABAergic VN neuronal subpopulations. The principle of optogenetics is based on the insertion of genes encoding photoactive proteins, such as opsins, into defined cell populations (VGlut2+ and GAD2+). This method makes it possible to specifically stimulate these neurons. Our results demonstrate that it is possible to use optogenetics to generate posturo-locomotor deficits, an essential component of the vestibular syndrome. Here, these posturo-locomotor deficits are compared with those induced by classical chemical labyrinthectomy.

Results

Validation of the optogenetic method in vestibular nuclei
Mice expressing either a cre recombinase driven by VGluT2+ (vesicular glutamate transporter 2) promoter in glutamatergic neurons (VGluT2-ires-cre), or a cre recombinase driven by GAD2 (glutamate decarboxylase 2) promoter in GABAergic neurons (GAD2-ires-Cre) were injected unilaterally with the cre-dependent virus AAV9-DIO-ChR2-mCherry in the vestibular nuclei (VN). In parallel, a control group of mice expressing VGluT2-cre or GAD2-cre received injections of AAV9-DIO-mCherry at the same location. Three weeks after injection, all mice were implanted with an optical fiber above the VN for the purpose of vestibular optical stimulation on awake and free-moving animals (Fig.1A,B). After completion of all experimental tests, brains were extracted and sectioned to verify viral expression and location of optical fibers (VGluT2-cre: Fig.1C,D; GAD2-cre: Fig.1E,F) and data was only used for animals when both parameters were verified. With regard to viral expression, VN neurons that co-expressed mCherry viral fluorescence and DAPI staining indicated that VN neurons were locally infected in both VGluT2-cre (Fig.1C, right) and GAD2-cre (Fig.1E, right) groups. Optical fibers were implanted in the upper parvicellular part of medial VN (Fig.1 C,E,D,F).

Unilateral optogenetic stimulation of VN_{VGluT2+} neurons, but not GAD2 neurons, triggers strong posturo-locomotor deficits

Before starting the optogenetic stimulation protocol, one group underwent a unilateral vestibular deafferentation using chemical labyrinthectomy in order to compare the functional consequences of optogenetic stimulation with characteristic acute vestibular syndrome. This was done by transtympanic administration of sodium arsanilate in WT mice, as previously reported in rats (Vigneaux et al. 2012), and posturo-locomotor behavior was recorded and analyzed in an open field using both subjective (Pericat et al. 2017) and automatized (Rastoldo et al. 2020) analyses. Posturo-locomotor behavioral parameters, included head tilt, bobbing, circling, retropulsion and tumbling (Fig. 2A) were observed at several time points over 3 weeks (Fig. 2B, gray). All mice (n=6) exhibited specific vestibular deficits, such as head tilt and circling, both oriented towards the injured side. In addition, mice exhibited limb extension towards the intact side, accompanied by limb flexion on the opposite side, often leading to falls towards the lesion side. Animals were
strongly and significantly affected on the first few days post-lesion, especially at Day 2 (Baseline = 0 vs. D2 = 8.17, P<0.05, t=2.69, df=8, n=6), then partially recovered from D3 and over the 3 weeks (D2 vs. W3, P<0.05, t=2.94, df=5, Fig. 2.B). A relative heterogeneity of deficits between animals emerged following a qualitative analysis of D2 data, with some showing only head tilt, whilst others displaying all categories of symptoms, including tumbling (Fig. 2.C).

Mice in the VGluT2-cre ChR2 and control mCherry groups were placed in an open field and observed for 5 minutes during which animals did not show any sign of posturo-locomotor deficit. During persistent optogenetic stimulation (20 Hz, 5 ms, 473 nm for 30 s) that followed, the VGluT2-ChR2 group displayed strong posturo-locomotor deficits similar to that observed in the arsanilate group (Fig. 2D, left). However, unlike mice in the arsanilate group, all VGluT2-ChR2 mice exhibited limb extension on the stimulated side, accompanied by limb flexion on the non-stimulated side. Six of nine mice exhibited instantaneous tumbling towards the non-stimulated side (Fig. 2D, 2F, Sup. Video 1). In addition, all mice exhibited circling behavior oriented towards the stimulated side. In comparison, VGluT2-mCherry control group animals did not show deficits during the entire stimulation period (Fig. 2D, right). Quantitatively, the VGluT2-ChR2 group tended to present a higher deficit than the arsanilate group, but not significantly different to D2 post-lesion arsanilate animals (D2 = 8.17 vs. VGluT2-ChR2 = 11.67, P>0.1; Fig. 2B, red). We then followed the same procedure using GAD2-cre animals and stimulated VN GABAergic neurons (60 Hz, 5 ms, 473 nm for 30 s) and did not find any postural deficits in the ChR2 or mCherry animals (Fig. 2E). This is also demonstrated by video examples (Fig. 2G).

In general, the postural behavior in the GAD2-ChR2 mice was comparable to the non-lesion mice (GAD2-ChR2 = 0 vs. VGluT2-ChR2 = 11.67, P<0.001, t=6.48, df=15; Fig. 2B, blue). These data indicate that activation of VN glutamatergic neurons leads to immediate and strong posturo-locomotor deficits that are, on average, comparable to those observed for arsanilate group at day 2. In contrast, activation of GABAergic neurons does not appear to have such a marked effect.

Tail-hanging landing tests confirmed that the VN\(^{\text{VGluT2+}}\) population is responsible for posturo-locomotor deficits, compared with the VN\(^{\text{GAD2+}}\) population
Above, we show that stimulation of VN\textsuperscript{VGluT2+} (but not VN\textsuperscript{GAD2+}) neurons led to posturo-locomotor deficits whilst the animals were standing on a static ground. To verify whether this observation persists when vestibular compensation is removed, we performed a tail-hanging landing test and scored three mains parameters: landing, twirl, and posture after landing (Fig. 3A). First, WT animals following arsanilate injection showed strong and homogeneous deficits. Compared to baseline measures, twirls and posture landing were affected (mean score from all mice=6), whilst posture after landing was only moderately affected (mean score for 4/6 mice=1; Fig. 3C). Across testing, effects were especially marked at D2 post-lesion (Baseline = 0 vs. D2 = 7.33, P<0.0001, t=27.8, df=8, n=6) and did not recover even 3 weeks after surgery (D2 vs. W3, P<0.005, t=4.83, df=5, Fig 3B, gray). Next, mice in the VN\textsuperscript{VGluT2+} and VN\textsuperscript{GAD2+} groups were tested similarly to the WT/arsanilate group and were given 30 s continuous optogenetic stimulation. During stimulation, all VGlut2-ChR2 mice showed the full range of postural deficits, some stronger even than those observed in the WT/arsanilate group, confirming the posturo-locomotor results above, with the exception of one animal that only showed moderate symptoms and no deficit during landing (Fig. 3D, left, Fig. 3F). Control VGluT2-mCherry animals did not show any deficits during light stimulation (Fig. 3D, right), confirming that viral injection surgery did not damage the VN. Overall, stimulation of VN\textsuperscript{VGluT2+} resulted in postural deficits similar to those with arsanilate animals at D2 post-lesion (D2 = 7.33 vs. W3 = 7.78, P>0.1; Fig. 3B, red). Next, GAD2-ChR2 animals were tested and no variation of postural behavior was observed following light stimulation; the GAD2-ChR2 group remained indistinguishable from control GAD2-mCherry group (Fig. 3E, G). Stimulation of GAD2-ChR2 was, overall, similar to arsanilate animals before lesion. Together, these data indicate that brief activation of VN\textsuperscript{VGluT2+} induced strong postural imbalance, whilst transient activation of VN\textsuperscript{GAD2+} neurons do not appear to participate in balance, at least in the short term.

**Optogenetic stimulation of VN\textsuperscript{VGluT2+} induced reversible postural instability**

For a more thorough assessment of static postural instability caused by VN stimulation, we monitored the support surface area (Marouane et al., 2020) before, during and 5 minutes after.
optogenetic stimulation. The support surface area was calculated by measuring the surface delimited by the four legs of each mouse whilst they were static for at least one second. Before stimulation or lesion, VGluT2-ChR2, GAD2-ChR2, and WT-lesion animals had similar scores (Fig. 4A, Pre). During optogenetic stimulation, VGluT2+ mice drastically increased their support surface area compared to pre-stimulation (VGluT2-Pre=0.006% vs. VGluT2-Stim=0.0145%, P<0.0001, t=13.66, df=7), and compared to WT-lesion at D2 (VGluT2-Stim=0.0145% vs. D2=0.0113%, P<0.05, t=2.124, df=12; Fig. 4 A, B, D). In line with previous results, GAD2-ChR2 stimulation did not cause any significant difference in support surface area, and was significantly lower than the support surface area of VGluT2-ChR2 during stimulation (VGluT2-Stim=0.0145% vs. GAD2=0.0055%, P<0.0001, t=10.76, df=14, Fig. 4A, Stim; Fig. 4C). At five minutes post-stimulation, higher support surface area scores were still observed in the VGluT2-ChR2 group than in the GAD2-ChR2 group (VGluT2-Stim=0.0068% vs. GAD2=0.0054 %, P<0.05, t=2.49, df=14; Fig. 4A, C). However, VGluT2-ChR2 group scores post-stimulation were not significantly different from pre-stimulation baseline scores. These results confirm that stimulation of VN\textsubscript{VGluT2+}, but not VN\textsubscript{GAD2+}, artificially induced continuous instability during the entire stimulation duration, comparable to and even stronger than that observed in the unilateral vestibular deafferentation model. Importantly, this effect was reversible and faded some minutes after stimulation ended.

**VN\textsubscript{VGluT2+} activation induces short-term disturbances of locomotor function**

To assess the functional consequences of unilateral optogenetic stimulation of VN on locomotor function, we next studied several parameters previously identified as biomarkers of rodent locomotor dysfunction (Rastoldo et al., 2020). These parameters were monitored in mice placed in an open field before, 5 minutes after, and 1 hour after optogenetic stimulation. Before stimulation, VGluT2-ChR2 and VGluT2-mCherry animals traveled a similar distance compared to immediately after stimulation. However, at 5 minutes post-stimulation, the distance traveled by the VGluT2-ChR2 group was significantly lower than that traveled by the VGluT2-mCherry control group (ChR2=2.04 vs. mCherry=5.87, P<0.001, t=4.7, df=11, Fig. 5A). After 1 hour,
VGluT2-ChR2 displacement was not significantly different to the VGluT2-mCherry group. In parallel, the velocity of the VGluT2-ChR2 right after stimulation was significantly lower than the control VGluT2-mCherry group (ChR2=0.012 vs. mCherry=0.019, P<0.001, t=4.6, df=11). This difference disappeared 1 hour after stimulation, where again, the VGluT2-ChR2 group tended to have lower velocity (Fig. 5B). When observing freezing time, the VGluT2-ChR2 group spent significantly more time immobile than the control VGluT2-mCherry group following optogenetic stimulation (ChR2=86.46% vs. mCherry=53.15%, P<0.005, t=3.69, df=11), however, 1 hour later the two groups were not significantly different but there was a trend for higher freezing in the VGluT2-ChR2 group (Fig. 5C). Overall, these results indicate that stimulation of VNVGluT2+ significantly altered locomotion up to 5 minutes after stimulation, but this effect disappeared after 1 hour. In comparison, there was no significant difference or trend between GAD2-ChR2 and GAD2-mCherry groups, either for the distance traveled (Fig. 5D), mean velocity (Fig. 5E) or immobility time (Fig. 5F) at any time point of the experiment. Together, these results corroborate our previous experiment, and indicate that stimulation of VNVGluT2+ neurons, but not VN\textsuperscript{GAD2+}, continued to have an effect on locomotion for a short period of time after stimulation.

**Discussion**

This study demonstrates for the first time that it is possible to induce reversible posturo-locomotor deficits characteristic of vestibular syndrome by optogenetic stimulation of the vestibular nuclei (VN) area in rodents. Optogenetic stimulation of the glutamatergic VN, but not GABAergic, neuronal subpopulation was responsible for immediate and strong posturo-locomotor deficits, comparable to our unilateral vestibular deafferentation model. Optogenetic stimulation reproduced the mirror image of typical posturo-locomotor behavior evoked by unilateral lesion of the vestibular system. By selective and reproducible stimulation of the different neuronal populations in the vestibular nuclei, this approach brings new opportunities to identify downstream pathways of VN neurons, and to better understand their functional relevance under normal or pathological conditions.
Technical considerations

Among the four main vestibular nuclei, we chose to stimulate specifically the medial vestibular nuclei (MVN) for various reasons. The medial vestibular nucleus is the one most extensively studied in animal anatomo-functional studies (for review see: Paterson et al., 2004). It is the largest nuclei form the vestibular nuclear complex (VNC) in most mammals (Alvarez et al., 1998) and is known to receive convergent semicircular and otolithic afferents, and to be involved in both oculomotor and postural functions (Wilson and Jones, 1979). The MVN contains a wide diversity of neuronal classes, which project to the oculomotor nuclei, spinal cord, cerebellum, thalamus, contralateral vestibular nuclei, or function as interneurons (Highstein and Holstein, 2006; Straka et al., 2005). It contains both glutamatergic and GABAergic neurons (de Waele et al., 1995; Gliddon et al., 2005), which can themselves be divided in subgroups depending of their morphology, respective networks, or molecular background. When considered in this context, we would expect AAV virus injections into VGluT2-cre and GAD2-cre mice to stimulate only glutamatergic or GABAergic neurons as if each on them belong to a homogeneous neuronal population. This additional layer of complexity cannot be targeted by using this method and will necessitate future investigation to elucidate.

Functional considerations

It is interesting to note that the posturo-locomotor syndrome induced by unilateral optogenetic stimulation of glutamatergic MVN neurons mirrors that of typical posturo-locomotor behavior evoked by unilateral lesion of the vestibular system with arsanilate. During optogenetic stimulation of MVN glutamatergic neurons, animals rotate towards the side of the simulation, while they circle towards the side ipsilateral to the lesion in the arsanilate model. These typical postural behaviors generated by both optogenetic stimulation and arsanilate lesion result from the electrophysiological asymmetry between ipsi and contralateral MVNs. Indeed, a large number of studies have shown that immediately after unilateral vestibular lesion, resting discharge rates of MVN neurons on the ipsilesional side are abolished, whilst the activity of MVN neurons on the contralesional side is either normal or increased (Newlands and Perachio, 1990; Ris and Godaux, 1998; Smith and Curthoys, 1989). In this situation, the vestibular deficit is always on the side of
the body corresponding to the MVN with reduced electrical activity. Muscle hypotension, falls, head tilt and circling affect the side in which the MVN is electrically deficient. Furthermore, the decline of the posturo-locomotor symptoms coincides with a recovery of the resting activity of ipsilesional MVN (Lacour and Tighilet, 2010). The posturo-locomotor deficits consecutive to optogenetic stimulation of glutamatergic MVN neurons likely result from an electrophysiological imbalance with a highly increased activity on the stimulated side.

To better appreciate the behavioral phenotype common to both arsanilate lesion and optogenetic stimulation of VN, it is worth examining VN connectomics. The medial vestibulospinal tract is primarily composed of axons from MVN neurons, but fibers from the lateral and descending vestibular nuclei also contribute to the tract (Carpenter, 1991). Axons of the medial tract travel in the medial longitudinal fasciculus and principally terminate in upper cervical regions of the spinal cord that innervate upper-body musculature, particularly neck musculature. Thus, the MVN and medial vestibulospinal tract provide the circuitry necessary to rapidly control vestibulo-collic reflexes and neck postural changes associated with alterations in head orientation. The lateral vestibulospinal tract is the largest of the two tracts and is primarily composed of axons from neurons in the lateral vestibular nucleus (LVN, also referred as Dieter’s nucleus), with some contribution from the descending (inferior) nucleus. The fibers descend ipsilaterally in the ventrolateral columns and have branches that innervate multiple levels of the spinal cord, thus providing the capacity to modulate spinal motoneuron activity across segments (Abzug et al., 1974). Thus, vestibulospinal pathways are well positioned to modulate the excitability of reflex responses to anticipated or imposed postural displacements. Experiments in animals have shown that electric stimulation of the LVN increases the excitability of extensor motoneurons and decreases those of flexor motoneurons (Grillner et al., 1970; McCall et al., 2017; Wilson and Yoshida, 1969). The facilitatory effect of vestibulospinal inputs on extensor motoneuron excitability suggests that the predominant function of this pathway is to provide appropriate levels of extensor tone to achieve vertical support against gravity. In the present study, the characteristic behaviors observed in optogenetically-stimulated mice, such as tumbling, circling, and retropulsion could be attributed to the specific stimulation of the MVN, whilst the enlargement of the support surface area, and head tilt (observed but not measured) could result
from stimulation of the LVN. Most commissural inhibition is provided by type I excitatory neurons unilaterally acting on type II inhibitory neurons on the opposite side, thus releasing GABA in tonic type I neurons on the opposite side (Kasahara et al., 1968; Mano et al., 1968; Precht et al., 1973). The sole parameter that departs from this rule is the circling sense observed after optogenetic stimulation. As recently reported (Marouane et al., 2020), the circling phenotype is common to rodent models of Parkinson's disease (Smith, 2018). This phenomenon may be explained by the involvement of vestibulo-striatal connections that are essential for the control of postural locomotion (Lopez and Blanke, 2011). Circling results from striatal electrophysiological imbalance common to that induced in the VN by both arsanilate lesion and optogenetic stimulation. Further anatomo-functional study to determine the neurochemical nature of vestibulo-striatal connections should allow determining the bases of circling rotation sense after optogenetic stimulation.

When optogenetic stimulation was interrupted, a restoration of normal postural behavior was observed, likely because electrophysiological balance is restored between opposing MVNs. The advantage of the optogenetic stimulation protocol is that we instantly appreciate the posturo-locomotor recovery phenomenon. In terms of both behavior and neuronal responses, the onset of vestibular compensation occurs later and lasts longer in vestibular lesion models compared to the short recovery times following optogenetic stimulation. The disappearance of behavioral deficits as well as the rebalancing of spontaneous activity in the homologous MVN is observed around one week after unilateral vestibular lesion of the vestibular system in guinea pigs (Ris et al., 1997).

Unlike the glutamatergic neural network, unilateral optogenetic stimulation of GABAergic neurons in the MVN did not produce a posturo-locomotor syndrome. Optogenetic stimulation of the GABAergic neural network should have, in theory, induced ipsilateral inhibition of the MVN, thus generating an ipsilateral postural syndrome similar to that observed after arsanilate injury. One can ask what types of GABAergic MVN neurons were activated by optogenetic stimulation? Answering this question is a challenge considering the complex architectural organization of the MVN GABAergic neural network. Indeed, the GABAergic neurons within the vestibular nuclear
complex (VNC) have been classified into five functionally distinct groups (Highstein and Holstein, 2006). Group one consists of commissural GABAergic neurons. Group two consists of GABAergic neurons that project to the oculomotor motoneurons that are involved in the vestibulo-ocular reflex (VOR) (for review see: Beitz and Anderson, 2000). These two groups of neurons are located predominantly in the superior vestibular nucleus (SVN) and the rostral medial vestibular nucleus (MVN) and send projections through the ipsilateral medial longitudinal fasciculus. Group three consists of GABAergic neurons within the MVN, prepositus hypoglossy (PH), and inferior vestibular nucleus (IVN) that project to the nucleus of the inferior olive. Group four consists of GABAergic neurons within the rostral MVN and lateral vestibular nucleus (LVN) that project to the medial vestibulo-spinal tract. Finally, group five consists of local interneurons. To answer the above question, it would be necessary to know which type of GABAergic neurons were the target of optogenetic stimulation. We could potentially refer to a morphological criterion size. Indeed, it has been suggested that neuronal size can indicate differentiate a local GABAergic neuron from a projection neuron (Tighilet et al., 2007). However, neuronal size alone would very likely not be sufficient to select only the type of GABAergic neurons targeted by optogenetic stimulation, since the injected virus would infect all GAD2 neurons and not specific subgroups. It would be useful to investigate the nature of these neurons using neuroanatomical tracing and electrophysiological experiments. Another possible explanation for the absence of a behavioral effect under optogenetic stimulation of GABAergic neurons is the low ratio of GABAergic neurons in the VNC, which is less than 10% of total VNC neurons (Walberg et al., 1990), making it likely that the optogenetic stimulation did not recruit a sufficient number of neurons to generate inhibition of the NVM. Finally, it is possible that optogenetic stimulation targeted vestibulospinal GABAergic neurons with bilateral projections, thus canceling any effect. The vestibular nuclei influence postural control via the medial (mostly from ipsi and contralateral NVM neurons) vestibulospinal tracts (Wilson and Jones, 1979).

Compared with recently documented rat UVN model (Rastoldo et al., 2020), optogenetic stimulation seems to similarly affect parameters such as mean distance moved, velocity and immobility time. In addition, the support surface area is also similarly affected in these three models (Liberge et al., 2010; Marouane et al., 2020). These observations strengthen the
optogenetic model as a new vestibular pathological model. Given that efferences arising from the MVePC also project onto oculomotor muscles, in addition to multiple cortical areas, (for review see: Lopez, 2016; Lopez and Blanke, 2011), it is likely that VN unilateral optogenetics stimulation also triggers other functional alterations through modulating the vestibulo-ocular reflex (nystagmus and oscillopsia), vestibulo-sympathetic reflex (increase heart rhythm, drop of body temperature) (Balaban, 1999; Yates et al., 2014), and vestibulo-cortical pathways (spatial disorientation). Further studies are necessary to determine whether these functional consequences reported upon unilateral labyrinthectomy and UVN also take place after optogenetics VN stimulation. The optogenetic approach, which makes it possible to specifically stimulate neurons in different areas of the brain, is suitable for identifying downstream networks originating from the VN, and also the upstream pathways which can regulate function.

Clinical relevance

The development of an optogenetic approach that selectively stimulates neuronal bundles arising from different zones of the vestibular central pathway is a new step, which will be vital in determining the downstream pathways at the origin of the various symptoms of vestibular syndrome. The transfer of this technology to humans is not yet a reality. However, electrical stimulation of the deep nuclei is already being used clinically for Parkinson's disease (Gradinaru et al., 2009; Kravitz et al., 2010), and recent advances in the use of optogenetics in humans for therapy raises the possibility of stimulating patients with unilateral or bilateral vestibular areflexia or hyporeflexia, a restoration of the peripheral inputs, through vestibular optic stimulation or implant interface (Guyot and Perez Fornos, 2019; Sluydts et al., 2020). The use of such technological advances is quickly becoming a reality.

Materials & Methods

Animals
All procedures were approved by Animal Care and Use Committees in the Shenzhen Institute of Advanced Technology (SIAT), Chinese Academy of Sciences (CAS). Wild type C57BL/6 (6–8 weeks) mice (Beijing Vital River Laboratory Animal Technology Co. Ltd, Beijing, China) were used in this study. In addition, adult (6-8 weeks) male VGluT2-ires-cre (Jax No. 016963) and male GAD2-ires-cre (Jax No. 010802) transgenic mice were used. All mice were maintained on a 12/12-h light/dark cycle at 25 °C. Food and water were available ad libitum.

**Viral preparation**

For optogenetic experiments, we used one of the following viruses depending on the protocol: AAV9-EF1a-DIO–hChR2(H134R)–mCherry or AAV9-EF1a-DIO–mCherry. Virus titers were approximately 2-6x10^{12} vg/ml. Viruses were purchased from Brain VTA Co., Ltd., Wuhan and Shanghai Taitool Bioscience Co.Ltd.

**Viral injections**

VGluT2-ires-cre and GAD2-ires-cre mice were anesthetized with pentobarbital (i.p., 80 mg/kg) and fixed on a stereotaxic apparatus (RWD, Shenzhen, China). During surgeries, mice were kept anesthetized with isoflurane (1%) and placed on a heating pad to maintain a body temperature of 35 °C. A 10-µL micro-syringe with a 33-Ga needle (Neuros; Hamilton, Reno, USA) was connected to a microliter syringe pump (UMP3/Micro4; WPI, USA) and slowly placed for virus injection into the VN (coordinates: AP: –6.0 mm, ML: –1.2 mm and DV: –4.5 mm). Virus was then injected (120 nl at a rate of 50 nl/min). After injection, the needle was kept in the place for an additional 5-10 minutes to facilitate the diffusion of the virus, after which it was slowly withdrawn.

**Implantation of optical fibers**

Optical fibers (200 um diameter, NA: 0.37) were chronically implanted in the VN 3 weeks after virus infusion. For optogenetic experiments, an optical fiber was unilaterally implanted above the VN (AP: –6.0 mm, ML: –1.2 mm and DV: –3.8 mm). After surgery all animals were given at least one week to recover before experiments began.
Histology, immunohistochemistry, and microscopy

Mice were euthanized with an overdose of chloral hydrate (300 mg/kg, i.p) and transcardially perfused with ice-cold 1 x PBS and then with ice-cold 4% paraformaldehyde (PFA, sigma) in 1 x PBS. Brains were extracted and submerged overnight in 4% PFA at 4 °C before being switched to 30% sucrose in 1 x PBS to equilibrate. Brains were cut into 40-µm thick coronal sections using a cryostat microtome (Lecia CM1950, Germany). Finally, slices were mounted and cover-slipped in anti-fade aqueous mounting reagent with DAPI (ProLong Gold Antifade Reagent with DAPI, life technologies). Brain sections were then photographed and analyzed with an Olympus VS120 virtual microscopy slide scanning system and ImageJ and Photoshop software.

Arsanilate injection

Once mice were deeply anesthetized using pentobarbital (i.p. 80 mg/kg), a micro needle (30 gauge) was used to create a hole in the eardrum and infuse 0.150 ml of a 10% p-arsanilic acid solution (A9258, dissolved in PBS). Following transtympanic instillation, each mouse was laid on its side for 30 minutes with the lesion side was on top.

Behavior

The behavioral capacities of mice in the arsanilate group was assessed at 7 times points relative to day of surgery: baseline (before surgery), then 24, 48 and 72 hours after, and at 1, 2 and 3 weeks. Animals performing optogenetic stimulations were exposed to the same apparatus as the arsanilate group, and were habituated to the open field one day before the experiment. During the stimulation day, animals were first placed in the field for a 5-minute phase without stimulation, and then optogenetic stimulation (473-nm blue laser; Aurora-220-473, NEWDOON, Hangzhou) was delivered for 30 seconds (VGluT2: 20 Hz, 5 ms, 473 nm; GAD2: 60 Hz, 5 ms, 473 nm), followed by a final 5 minute without stimulation. The locomotion test was performed in an open field (40 cm x 40 cm x 40 cm) to which mice were habituated to the day before the test. During the test, mice were lifted by their tails for 5 quick vertical movements, then were immediately placed in the open field. Behavioral recordings were made by 2 cameras (one top-view, one side-
view) for a duration of 1 minute, to assess vestibular symptoms, which include (i) tumbling: spontaneous or evoked rotations of the animals along their body axis, rated 5 in our evaluation scale, (ii) Retropulsion: characterizes backwards movements of the animals, rated 4; (iii) Circling: circular movements of the rats in the horizontal plane, rated 3, (iv) Bobbing: relates to rapid head tilts to the rear and is assimilated to cephalic nystagmus, rated 2, and (v) Head tilt, rated 1. When none of these behaviors were observed, animals received a score of 0.

For the tail-hanging landing paradigm (THL), mice were lifted by their tails for 5 quick vertical movements. This test normally induced forelimb extension as the animals reach the ground. Unilateral vestibular deficit resulted in trouble during the landing process. Such deficits were quoted from 0 (perfect preparation of the two front paws before reaching the ground) to 3 (absence of preparation for landing). The landing process was accompanied by axial rotation of the body, i.e. a twirl movement that we quantified from 0 (no rotation) to 3 (continuous twisting). We also monitored the intensity of syndrome reactivation once landed, from 0 (no sign) to 3 (max expression/accentuation of circling, tumbling, muscle dystonia, bobbing or head tilt). Mice receiving optogenetic stimulation during THL tests were first exposed to the same protocol as arsanilate group (lifted by the tail without additional stimulation). Then they received optogenetic stimulation continuously for 30 s, whilst being tested for lifting and landing. Two cameras recorded, 1 from the side, 1 from the bottom.

**Data analysis**

Speed, immobility, and distance data were analyzed using Anymaze software (Stoelting Co.). Support surface area and video frames were first analyzed using Kinovea software (https://www.kinovea.org/), and then MATLAB was used to compute the size of the surface. Statistical significance was determined using two-tailed Student's t-tests with a significance level of p<0.05.

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Author contributions

Quentin Montardy: Conceptualization, methodology, validation, formal analysis, data curation, writing – original draft, Writing - Review & Editing, visualization, project administration

Mengxia Wei: Investigation (virus injection, fiber implantation, arsanilate injection and behavior, immunohistochemistry and microscopy), data curation and analyzes, writing – original draft, Writing

Tian Yi: Investigation (virus injection and fiber implantation, arsanilate injection and behavior), data curation and analyzes, writing – original draft

Xuemei Liu: Investigation (performed immunohistochemistry and quantitative analyzed of the tracing data), manuscript review and editing

Zhou Zheng: Methodology (setup injection protocol), conceptualization, manuscript review and editing

Lai Juan: Investigation (immunohistochemistry)

Stephane Besnard: Conceptualization, manuscript review and editing

Brahim Tighilet: Conceptualization, original draft writing, manuscript review and editing

Christian Chabbert: Conceptualization, supervision, original draft writing, manuscript review and editing

Liping Wang: Conceptualization, supervision & Project administration, manuscript review and editing

Figure Legends

Figure 1 - VN virus injection and optical fiber implantation. A. Schematic showing the experimental protocol. AAV9-DIO-ChR2-mCherry was injected int the VN of VGlut2-ires-cre or
GAD2-ires-cre. Three weeks later, an optical fiber was positioned in the vicinity of the VN, and behavioral tests were performed 1 week later. B. Schematic representation of virus injection and optical fiber location in the VN. C. Representative photograph of virus expression in the VN of VGlut2+ mice. Red, ChR2-mCherry virus; Blue, DAPI. Scale bar, 100 µm (left), and 50 µm (right). MVePC (parvcellular medial vestibular nucleus) MVeMC (magnoellular medial vestibular nucleus), SPVe (superior vestibular nucleus), LVe (lateral vestibular nucleus) D. Representation of fiber tip position for individual VGlut2+ animals (in red) E. Representative picture of virus expression in VN of GAD2+ animals. F. Representation of fiber tip position for individual VGlut2+ animals (in blue).

**Figure 2 - Optogenetic stimulation of VN^VGlut2+ neurons, but not GAD2 neurons, induced strong posturo-locomotor deficits.** A. Evaluation grid used to score behavior in the open field. If all symptoms are expressed, then a rating of 15 is scored, if all except tumbling are expressed, the rating 10 is scored, and so on. B. Illustration of the time course of the behavioral evaluation score for WT animals with arsanilate lesions (gray), compared with VN^VGlut2+ (red) and VN^GAD2+ mice during optogenetic stimulation. C. Behavioral evaluation score of WT-lesion animals, at D2 post-lesion, whilst animals were placed on a static ground. D. Behavioral evaluation score for the VGlut2-ChR2 (left) and VGlut2-mCherry (right) groups. E. GAD-ChR2 (left) and GAD2-mCherry (right) groups during a 30 s stimulation whilst animals were on the ground. GAD-ChR2 and mCherry do not show a deficit. F. Example of posturo-locomotor behavior from a VGlut2-ChR2 mouse (essentially tumbling) G. Illustration of a GAD-ChR2 mouse during stimulation.

**Figure 3 - Tail-hanging landing test confirms that the VN^VGlut2+ population is mainly responsible of posturo-locomotor deficits, compared to the VN^GAD2+ population.** A. Evaluation grid used for tail-hanging landing tests. B. Time course of the behavioral evaluation score for WT animals with arsanilate lesions (gray), compared to VN^VGlut2+ (red) and VN^GAD2+ mice during optogenetic stimulation. C. Behavioral evaluation score of the WT-lesion group at D2 post-lesion, whilst undergoing tail-hanging landing tests. D. Behavioral evaluation scores of the VGlut2-ChR2 (left), VGlut2-mCherry (right) groups. E. GAD-ChR2 (left) and GAD2-mCherry (right) groups during a 30 s stimulation during tail-hanging landing test. The GAD-
ChR2 and mCherry groups did not show a deficit. F. Example of posturo-locomotor behavior in a VGlut2-ChR2 mouse. G. Representative GAD-ChR2 mouse during stimulation. Time in frame: left, before stimulation; right, during stimulation.

Figure 4 - Reversible postural instability upon optogenetic stimulation of VN VGlut2+ neurons. A. Support surface area as a percentage, averaged across all VN VGlut2+ (red), VNGAD2+ (blue) and WT-lesion animals (gray). Three 5-min time points were investigated: before stimulation, 5 min after stimulation, and 1 h after stimulation. B. Example of the support surface area computed on one VN VGlut2+ mouse and C. a VNGAD2+ mouse, before, right after, and 1 h after stimulation, as well as D. WT-lesion animal before and after lesion.

Figure 5 - VN VGlut2+ activation induces short-term disturbances of locomotor function. Averaged locomotor behaviors during free movement in an open field of mice that received either VGlut2-ChR2 (dark red) or VGlut2-mCherry injection (light red) in the VN. Three 5-min time points were investigated: before stimulation, 5 min after stimulation, and 1 h after stimulation. A. Distance moved in meters. B. Mean velocity during movement in m/s. C. Time spent immobile whilst in the apparatus expressed as a percentage of total experiment time. Averaged locomotor behaviors for GAD2-ChR2 (dark blue) or GAD2-mCherry injection (light blue), computed as previously described D. Distance moved. E. Mean velocity. F. Percentage of time spent immobile.
References


Lorente De No, R., 1933. The central projection of the nerve endings of the internal ear. The Laryngoscope.


Ris, L., Godaux, E., 1998. Neuronal Activity in the Vestibular Nuclei After Contralateral or Bilateral Labyrinthectomy in the Alert Guinea Pig. Journal of Neurophysiology 80, 2352–2367. https://doi.org/10.1152/jn.1998.80.5.2352


Fig. 1

A

Virus injection in VN
VGlut2-cre or GAD2-cre
Fiber implantation
Above VN
Behavior

3 weeks
1 week

B

Virus
AAV-DIO-ChR2-mCherry

Optical fiber

VGlut2-cre or GAD2-cre

C

DIO-ChR2-mCherry
VGlut2-cre

Optical fiber

4V
MVePC
MVeMC
SpVe
LVe

D

E

Optical fiber
DIO-ChR2-mCherry
GAD2-cre

4V
MVePC
MVeMC
SpVe
LVe

F
Fig. 3

A

<table>
<thead>
<tr>
<th>Action</th>
<th>After landing</th>
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</thead>
<tbody>
<tr>
<td>Landing</td>
<td>3 3 3 3 3</td>
</tr>
<tr>
<td>Twirl</td>
<td>3 3 3 3 2</td>
</tr>
<tr>
<td>Total score</td>
<td>9 8 5 3 3</td>
</tr>
</tbody>
</table>

B

Tail-Hanging landing (averaged)

<table>
<thead>
<tr>
<th>Time</th>
<th>Baseline</th>
<th>Day 1</th>
<th>Day 2</th>
<th>Day 3</th>
<th>Week 1</th>
<th>Week 3</th>
<th>VGluT2-ChR2</th>
<th>GAD2-ChR2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>9</td>
<td>9</td>
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<td>9</td>
<td>9</td>
<td>5</td>
<td>9</td>
</tr>
</tbody>
</table>

C

Tail-Hanging landing

Day 2

WT lesioned

D

VGluT2-ChR2

Stim 'OFF'

Stimulation 'ON'

E

GAD2-ChR2

Stim 'OFF'

Stimulation 'ON'

F

VGluT2-ChR2

Stim 'OFF'

Stimulation 'ON'

G

GAD2-ChR2

Stim 'OFF'

Stimulation 'ON'
Fig. 4

A

![Graph showing Area (%) for different conditions.](https://doi.org/10.1101/2020.09.03.281980)

B

**VGluT2-ChR2**

Pre-stim  | Stim  | 3 min post-stim

C

**GAD2-ChR2**

Pre-stim  | Stim  | 3 min post-stim

D

**WT-lesioned**

Pre-stim  | Day 2
**Fig. 5**

- **A**: Graph showing distance traveled (m) with error bars for pre-stim, 5 min post-stim, and 1 hour post-stim. The legend indicates significant differences between GAD-ChR2 and GAD-Mcherry.
- **B**: Graph showing mean velocity (m/s) with error bars for pre-stim, 5 min post-stim, and 1 hour post-stim. The legend indicates significant differences between post-stim and pre-stim.
- **C**: Graph showing time immobile (%) with error bars for pre-stim, 5 min post-stim, and 1 hour post-stim. The legend indicates significant differences between GAD2-Chr2 and GAD2-mCherry.

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