

Physiology of freezing of gait

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Abstract

Freezing of gait (FOG) is a common and debilitating, but largely mysterious, symptom of Parkinson's disease. In this review, we will discuss the cerebral substrate of FOG focusing on brain physiology and animal models. Walking is a combination of automatic movement processes, afferent information processing and intentional adjustments. Thus, normal gait requires a delicate balance between various interacting neuronal systems. To further understand gait control and specifically FOG, we will discuss the basic physiology of gait, animal models of gait disturbance including FOG, alternative etiologies of FOG and functional magnetic resonance studies investigating FOG. The outcome of these studies point to a dynamic network of cortical areas such as the supplementary motor area, as well as subcortical areas such as the striatum and the mesencephalic locomotor region (MLR) including the pedunculopontine nucleus (PPN). Additionally, we will review PPN (area) stimulation as a possible treatment for FOG, and ponder whether PPN stimulation truly is the right step forward.

Abbreviations:

CLR = cerebellar locomotor region; CNF = cuneiform nucleus; DBS = deep brain stimulation; FOG = Freezing of gait; MLR = Mesencephalic locomotor region; MTPT = 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine; PPN = pedunculopontine nucleus; PM = premotor area; SLR = subthalamic locomotor region; SMA = supplementary motor area; SNr = substantia nigra reticulata

Introduction

Patients with Parkinson's disease (PD) often experience debilitating episodes of freezing of gait (FOG), during which their feet feel 'glued to the floor'^{1,2}. FOG is defined "as a brief episode during which patients find it impossible to generate effective forward stepping movements, in the absence a cause other than parkinsonism or higher cortical deficits. It is most commonly experienced during turning and step initiation, but also when faced with special constraint such as a doorway, stress and distraction. Focused attention and external stimuli (cues) can overcome the episode. Because of its sudden and unpredictable nature, FOG often leads to falls and injuries."³ FOG presents itself in different phenomenologies. There are three subtypes: shuffling with small steps, trembling in place without forward movement, or total akinesia⁴. In addition, the effect of special constraints (such as going through doorways or stress) differs between patients. It remains an open question whether these phenomenologies are the result of different severity or represent different pathophysiologies. Most commonly FOG occurs in the OFF state, and may well be levodopa-responsive. However, the relationship with dopaminergic medication is complex, and as the disease progresses levodopa resistant FOG may develop. Paradoxically, sometimes FOG is even induced or worsened by dopaminergic medication.⁵ Freezing of upper limbs and of speech is correlated to FOG and bears remarkable spatiotemporal similarity; however, the added postural component renders FOG a distinct phenomenon. In this review, we will address the physiology of levodopa-responsive and levodopa-resistant FOG. We will discuss the current knowledge of the cerebral substrate of FOG as can be garnered using animal studies, animal and human 'models' of FOG and functional brain imaging. Although FOG can be seen in various diseases such as PD, multiple system atrophy, progressive supranuclear palsy or vascular disease, we will focus on FOG in PD.

Normal walking would seem to demand little attention. However, in actuality it is a complex task requiring a delicate balance between various interacting neuronal systems⁶. It necessitates not only automatic movement processes involving stepping and balance, but also attention, afferent information processing and intentional adjustments⁶⁻⁸.

To further understand gait control and specifically FOG, we will first discuss the normal physiology of gait based on animal studies. Second, we will review animal models of gait disturbance including FOG. Next, the implications of alternative etiologies of FOG are discussed. We will show how functional magnetic resonance studies investigating FOG give insight to its physiology. The outcome of these animal and human studies point to the importance of cortical areas such as the supplementary motor area, as well as subcortical areas such as the striatum and the mesencephalic locomotor centre including the pedunculopontine nucleus (PPN). Last, we will review PPN stimulation as a possible treatment for FOG.

Basic physiology of gait based on animal studies

Lesioning studies have taught us important lessons concerning the cerebral control of locomotion. Multiple cat studies have used decerebration in which input from the brain is eliminated by cutting across different levels of the brainstem⁹. It has been shown that a decerebrate cat can still walk, trot and gallop. When the decerebration is made at precollicular-postmammillary level, the cat initiates locomotion only with electrical or chemical stimulation applied to the mesencephalic or midbrain locomotor region (MLR)¹⁰. However, if the neuraxis is transected slightly higher at the precollicular and premammillary level, cats can spontaneously elicit locomotion with well-coordinated postural control, accompanied by largely appropriate equilibrium control⁹. Therefore a critical region exists between these decerebrate levels. This area is recognized as the *subthalamic locomotor region* (SLR), which mostly corresponds to the lateral hypothalamic area. Stimulation of the SLR evoked locomotion after a large lesion was made in the MLR area¹¹, indicating that the SLR has direct connections with the brainstem locomotor pathway beyond the MLR. However, the walking in the decerebrate preparations is machine-like and is neither goal-directed nor adaptive to the environment. Hence, the SLR connections to the MLR are likely important for normal gait.

Lesions in the *motor cortices* in cats, just like pyramidal tract transection, elicited surprisingly minor deficits in locomotion. However, skilled performance is more severely disturbed by postcruciate than by precruciate lesions. The precruciate area, which corresponds to supplementary motor area (SMA) and premotor area (PM) of the primates, may be involved in movement initiation while the postcruciate cortices (posterior sigmoid gyrus) may utilize specific somatosensory inputs to regulate ongoing movements¹² by anticipatory or feed-forward adjustments¹³. Visuomotor integration is needed when an ambulating subject encounters obstacles, which requires involvement of the posterior parietal cortex¹⁴.

Another method for lesioning studies is to directly inject inhibitory substances into selected brain regions. In bipedal walking monkeys, inactivation of the leg region in primary motor cortex (M1) with injections of muscimol (GABA_A agonist) results in paresis of the contralateral leg¹⁵. On the other hand, injections into trunk or leg regions of the bilateral SMA largely disturb postural control without paralysis¹⁶. When injecting the dorsal PM, spontaneous walking is maintained, but no walking occurs following sensory guidance. While M1 mainly projects to the spinal cord, the SMA and PM have dense projections to the brainstem reticular formation^{17, 18} including the PPN area¹⁹ in addition to projections direct to the spinal cord.

So far three locomotor regions have been identified in animals: the MLR, the SLR and the cerebellar locomotor region (CLR) in the mid-part of the cerebellum²⁰. The MLR appears to be present in all classes of vertebrates²¹. It likely includes the cuneiform nucleus (CNF) and the PPN, although the precise location of the locomotor regulation still remains a matter of debate²². A recent review offers a good overview of history, precise localization and inter-species differences of the MLR and PPN²³. Importantly, the PPN is not just the 'cholinergic part of the MLR', although the cholinergic neurons serve to delineate PPN boundaries (pars compacta and pars dissipata). The PPN is located in the ventrolateral part of the caudal mesencephalic reticular formation, composed of a heterogeneous population of neurons, containing GABA and glutamate in addition to acetylcholine²⁴. Different neuronal types within the PPN area have different functions with their own inter-connections to multiple parts of the

brain. There are connections to cerebral cortex, multiple basal ganglia and limbic areas, the thalamus, the brainstem, the spinal cord and the cerebellum²³. This key location including multiple segregate functions, renders determining the precise function of these regions quite complicated²⁵.

The PPN is active in suppressing muscle tone via its cholinergic projections to the pontine reticular formation, while the CNF mostly elicits locomotion. Microstimulation of the transition zone between the two areas (ventral CNF, dorsal PPN) induced a mixture of locomotor rhythm with associated muscle tone suppression²⁶⁻³⁰. Moreover, blocking the pontine reticular formation by injecting atropine sulfate, blocked the PPN-induced atonia but facilitated MLR-induced locomotion, indicating that cholinergic PPN neurons not only control the level of muscle tone but also modulate the locomotor pattern, and do this by effects at the pontine level²⁶. Studies in rats show non-cholinergic neurons just medial to the PPN have projections to the spinal cord, while the cholinergic neurons of the PPN do not. This area at the mesopontine junction may be the true MLR. In rats, the dorsal neurons of this MLR area (laterodorsal tegmental nucleus) with spinal projections are active in locomotion, while the ventral neurons are active in standing and do not have spinal projections²².

In the 'locomotor pathway', signals from the MLR also activate medullary reticulospinal neurons, in turn commanding the spinal locomotor network to generate the oscillatory pattern of locomotion^{31, 32}. However, the SLR and the CLR may also activate this reticulospinal locomotor pathway through distinct and direct projections³³. (Figure 1B) Signals from the MLR may also activate monoaminergic descending pathways including the coeruleospinal and raphespinal tracts, acting as a muscle tone excitatory system³⁴⁻³⁶.

An important consideration is how the neurons in these systems are controlled and modulated by neurons in other regions. We will now focus on the inputs of basal ganglia and motor cortices to the MLR area. In decerebrate cats, the basal ganglia control locomotion and posture using different GABAergic output pathways of the substantia nigra reticulata

(SNr); the lateral part of the SNr blocks the PPN-induced muscle tone suppression, whereas the medial part of the SNr suppresses the MLR-induced locomotion^{26, 27, 29}. Recent rat studies confirm that inhibitory input from the SNc (GABAergic and dopaminergic) to ventral MLR regulate posture, while inhibitory projection from the GABAergic SNr to the dorsal MLR regulates locomotion²². In PD, GABAergic outputs of the basal ganglia are abnormally increased³⁷, so excessive SNr-inhibition of the MLR may cause gait disturbance and muscle rigidity^{28, 38}. However, it is unknown whether these mechanisms are the same for bipedal humans and quadrupedal animals. Furthermore, it is unclear what drives or dictates the SNr-induced control of locomotion and posture. There are sub-compartments in the basal ganglia; neostriatum-dorsal pallidal pathway (“dorsal pathway”) and ventral striatum-ventral pallidal pathway (“ventral pathway”). The nucleus accumbens, as a component of the limbic system, may be important in releasing locomotion, via GABAergic projections disinhibiting the MLR via the ventral pallidum^{39, 40} and via fibers to the SNr⁴¹. (Figure 1B) Because the nucleus accumbens also receives inputs from the hippocampus and amygdala, the ventral pathway may be implicated in reward-oriented locomotor behaviors, as it receives inputs from ventral tegmental area, hippocampus and amygdala. On the other hand, the more recently evolved⁴² parts of the basal ganglia make up the dorsal system (Figure 1A). These parts may achieve locomotor control depending on cognitive behavioral context, such as sensory-guided locomotor control.

Studies using neural tracers have demonstrated abundant cortico-fugal projections to the brainstem reticular formation from the premotor cortices (SMA/PM) in quadruped¹⁸ and biped¹⁷ animals. Recent studies have focused onto the importance of cortico-pedunculopontine projection in terms of motor control. The PPN receives partly separate but essentially convergent cortical inputs not only from multiple motor-related areas representing the same body part, but also from multiple regions representing diverse body parts⁴³. Probabilistic diffusion tractography in rhesus monkey as well as humans, shows that the

SMA is strongly connected to the lateral PPN, while the dorsal PM is connected to the medial PPN¹⁹.

The SMA contributes to the anticipatory postural adjustment for step initiation, which is impaired in PD patients⁴⁴. In the higher primate and, in particular, those with bipedal gait, SMA/PM projections to the PPN, in addition to the pontomedullary reticular formation, may play a crucial role of initiation of locomotion with appropriate postural control.

Animal Models of FOG

An important barrier to the discovery of the anatomic and biochemical correlates of FOG has been the lack of an appropriate animal model. Modeling FOG in animals is difficult because of its dependence on intentional motor performance and responses to environment. FOG requires that animals intend to walk during observation and the testing should include responses to obstacles and stressors. In addition, it is necessary to assess motion and associated physiologic parameters by using devices attached to hind legs, which is problematic in animals. Although these problems can be resolved with appropriate training, most animal models of PD or other FOG associated disorders are not suitable for extensive behavioral training due to compromised attention and learning. It is likely that "much of the research into experimental parkinsonism and dyskinesias must be undertaken in primates, for only those animals develop the typical clinical phenomena seen in man"⁴⁵.

Primate studies

In 1983, several cases of parkinsonism in heroin users led to the discovery that the compound 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP) induces parkinsonism in humans⁴⁶. MPP⁺, the toxic metabolite of MPTP, is a mitochondrial complex I inhibitor that selectively kills dopaminergic neurons⁴⁷ via uptake through the dopamine transporter. Thus, a primate model of parkinsonism was created that remains the only mammalian model to

exhibit the whole motor phenotype of PD, including the resting tremor and drug-induced dyskinesias⁴⁸.

In descriptions of humans exposed to MPTP six of seven developed typical FOG⁴⁹. The recent observation of FOG in MPTP intoxicated non-human primates is the first step in overcoming the deficiency of animal models⁵⁰. Twenty-nine macaque monkeys were evaluated for FOG (12 male, mean age 4.3 years) after using a standard protocol of systemic intoxication with MPTP that produced moderate to severe parkinsonism⁵⁰. For this study FOG was defined as episodic inability to generate effective stepping during walking. The monkeys had to be able to walk before and after the FOG episode. Fourteen of 29 (48%) developed FOG in the hind limbs which had many of the characteristics typical of human FOG. The episodes lasted several seconds to a minute and were accompanied by a 4-10 Hz tremor similar to that seen in patients (Figure 2 and Video 1)⁵¹. FOG was most consistently observed in the early morning prior to initiating dopaminergic medications. When the FOG occurred, the monkeys often would sit to break the episode, representing a possible trick. At the time of the publication of that report, FOG occurred in the off state in all but one animal in whom both on and off freezing was observed. However, since the publication of the initial observations L-dopa resistant FOG (during on and off state) has been found to be more prominent in animals with severe parkinsonism (unpublished data). Males were more likely to develop FOG, and the gait problem correlated with severity of parkinsonism and severity of levodopa-induced dyskinesia. Although the differences between humans and primates should of course be kept in mind, the study of both in this monkey model may provide greater insight into both levodopa responsive and unresponsive FOG in the human condition.

Pathological studies of MPTP induced parkinsonism in humans and non-human primates demonstrated changes in the dopaminergic system that recapitulate those seen in PD⁴⁷.

Studies in aged primates also demonstrated non-dopaminergic lesions seen specifically in the locus coeruleus, the primary source of norepinephrine⁵². In primate FOG thus far

preliminary data suggests no significant neuronal loss in the PPN (unpublished data). The PPN is generally unaffected in younger MPTP intoxicated animals, but in some (not all) studies, there has been a loss of 20-30% PPN cholinergic neurons in aged MPTP treated monkeys⁵³.

Attempted modeling in macaque monkeys combining lesioning the dopaminergic system with systemic MPTP intoxication and bilateral PPN injections of UII-Dtx (urotensin II-conjugated diphtheria toxin) to produce specific cholinergic lesions, did not succeed in provoking FOG. However, it did lead to gait disorders, disequilibrium and falls unresponsive to apomorphine⁵⁴. These animals showed mostly postural instability and not FOG. Animal studies based on selective lesioning of the PPN will be discussed further in this review. However, none of the experiments identified have produced freezing. One might conclude to this point that while lesions of the PPN can lead to alterations in cadence of gait, balance and postural changes, it is unclear if there is a direct role in FOG.

Rodent studies

A study in rodents with cortical cholinergic and caudate dopaminergic lesions described significant propensity to falls pointing to the role of attentional impairment in gait disturbances⁵⁵. Noteworthy, with larger dopaminergic lesion, dopamine loss extending across striatal regions (without the cholinergic lesion), gait abnormalities increase and rats also tend to slow or stop when traversing changing surfaces, leading to falls⁵⁶. This tendency to stop and fall in rats with extensive dopamine depletion is congruent with the observed correlation between FOG and increased severity of parkinsonism in the primate⁵⁰ as well as patients with PD. These studies did not address the impact of MLR involvement. However, a role of the PPN in FOG has been supported by a recent rodent study where bilateral dopaminergic lesions were combined with partial PPN destruction and the effects of stimulation on the anterior and posterior PPN areas were analyzed⁵⁷. In this study, stimulation of the anterior PPN resulted in apparent freezing with start hesitation, while stimulation of the posterior region improved

their gait. The freezing episodes in rats that were observed only in certain tasks (contextual gait), were not fully characterized and may not express the same parameters (tremor, timing, etc.) as of the FOG in PD. Still, rodent studies are important to assess the contribution of different brain regions in the control of gait. For example, another study in rodents implied that the more dorsal MLR may be more important for FOG. While lesioning the ventral MLR (pontine reticular formation) with OX-SAP toxin in rats induced typical cataplexy attacks, lesioning the dorsal MLR (laterodorsal tegmental nucleus) produced prolonged periods of immobility²². In contrast, stimulating this 'dorsal MLR' bilaterally with ibotenic acid caused continuous rhythmic walking and vocalizations²².

A recent study using optogenetics in mice showed that glutamatergic neurons in the MLR area are essential to initiate locomotion⁵⁸. This experiment supports that cholinergic PPN neurons are not a critical element to initiate locomotion, but rather have a modulatory role.

Another recent study in mice showed that the medullary reticulospinal system halted locomotion⁵⁹. Together with the studies previously mentioned, we corroborate that basal ganglia output controls the excitability of the reticulospinal system to control locomotion via non-cholinergic (GABAergic and glutamatergic) and cholinergic neurons in the PPN/MLR area.

In addition, there is increasing evidence concerning the role of attentional performance through cholinergic projections of the forebrain in falls⁶⁰. The basal forebrain has projections to the PPN⁶¹. In addition, it is often degenerated in PD patients⁶², which correlates with reduced walking speed⁶³. Loss of cholinergic neurons in the basal forebrain and PPN in addition to the damage of midbrain dopamine neurons was associated with falls in PD⁶⁴.

Deciphering which of these is the primary source of falls, recent challenging experiments in rats point to the importance of the combination of basal forebrain cholinergic cell loss with striatal dopamine loss. Adding a selective cholinergic PPN lesion did not further deteriorate performance of the rats.⁵⁶ Although these experiments point to the importance of the basal

forebrain cholinergic – striatal disruption for falls, its role for freezing remains to be established.

Another step forward in animal modeling of FOG will be regional specific MLR lesions in the MPTP-treated monkey, to examine if they actually generate or enhance primate FOG. It would be critical to determine and quantify the changes in MLR cholinergic, glutaminergic and other neuronal subpopulations that may be associated with the development of FOG in primates as a next step toward mechanistic studies of the FOG pathophysiology.

Functional MRI studies in humans with freezing of gait

Imaging studies in subjects with FOG are also important tools in unraveling the physiology of this gait disorder. We will focus on what functional MRI (fMRI) studies during action teach us about the cerebral substrate of FOG, with a special focus on the PPN region / the MLR. In addition, we will discuss how to interpret the outcomes of different fMRI paradigms.

Functional MRI measures changes in cerebral blood flow related to neural activity, while the subject is lying in an fMRI scanner. Creative solutions have been found to solve the limitation that walking is not possible during scanning. These solutions include the use of motor imagery of gait, virtual reality in combination with alternating leg movements and freezing of finger movements as alternatives to freezing while walking.

The first fMRI study in PD subjects with FOG investigated gait planning, using motor imagery of walking⁶⁵. This approach exploits the large neural overlap that exists between planning and imagining a movement. In addition, it avoids the confounds introduced by brain responses to altered motor performance and somatosensory feedback during actual freezing episodes. Twelve PD-freezers, twelve matched PD non-freezers and 21 matched healthy controls were studied. Subjects performed two validated tasks: motor imagery of gait, and a visual imagery control task. Imagery performance was quantified by measuring the time

required to imagine walking on paths of different width and length, and showed matched task performance. During motor imagery of gait, freezers showed more activity than non-freezers in the MLR. Freezers also tended to have decreased responses in mesial frontal and posterior parietal regions. The gait-related hyperactivity of the MLR correlated with clinical parameters (FOG severity and disease duration). The hyperactivity was located dorsally in the MLR, including the PPN but with a local maximum in the cuneiform nucleus and reaching the periaqueductal grey. Another study using motor imagery of gait on 9 PD freezers versus 9 PD non-freezers showed a decrease in right globus pallidus and trends to a similar decrease in SMA activity in those with FOG⁶⁶. However, this study did not show PPN hyperactivity (they actually showed a trend to hypo-activity), while using a stereotactically defined region of interest analysis of the PPN with the local maximum slightly more caudal to the area found in the first study (Talairach coordinates ⁶⁵ X-0, Y -28, Z -15, ⁶⁶ X ±6, Y -26, Z -11).

When considering motor imagery studies, it should be taken into account that they may investigate motor planning and not motor execution. In addition, patients imagined they were walking, but not having FOG episodes. Another approach that looked at actual motor execution and not planning, investigated repetitive hand movements with upper limb freezing⁶⁷. Upper limb freezing has been shown to be similar to leg freezing and correlated to FOG^{68, 69}. In an fMRI study, 16 PD freezers, 16 PD non-freezers and 16 controls performed a bimanual flexion-extension task of the index fingers. During regular movement, freezers showed decreased activity in cortical regions in the left dorsal premotor and primary motor cortex, as well as the right dorsolateral prefrontal cortex. In addition, those with FOG showed subcortical bilateral activation of STN, putamen and pallidum during regular movement. Comparing regular movement with freezing of upper limb episodes, upper limb motor blocks were associated with increased activation in the right motor cortex, dorsal premotor cortex, SMA and left prefrontal cortex, whereas bilateral pallidum and putamen activity was decreased. MLR involvement would not be expected in these studies.

A third approach also investigated non-gait repetitive movements, specifically alternate leg movements in a virtual reality walking environment. This approach showed by increasing cognitive load during continuous movement, a decreased activity in the ventral striatum, preSMA and left STN⁷⁰. During motor blocks, a decrease in dorsolateral prefrontal cortex, increase in frontoparietal and decrease in bilateral caudate, GPi and thalamus were found. In addition, a decrease in MLR activity (almost same maximum as ⁶⁵) was found which correlated to the severity of FOG⁷¹. The outcome of these studies point to the importance of cortical areas, particularly the SMA, as well as subcortical areas including the striatum and the MLR.

Beyond the individual role of these separate regions, it shows the importance of their interplay and communication through different neural networks. It appears to be a dynamic process (Figure 3): during continuous movement cortical activity is decreased and (compensatory to this decreased cortical activity?) subcortical activity is increased (MLR for the legs, STN for the hands). During subsequent episodes of freezing (of upper or lower limb) the STN and MLR hyperactivity breaks down to hypoactivity. Subsequent increased activity in dorsolateral prefrontal cortex and insula may be called upon to overcome the block but may play a role in subsequent emotional response as well. This failing dynamic process leading to freezing might become particularly evident during challenging events that require precise regulation of step length and gait timing, such as turning or initiating walking, which are well known triggers for FOG.⁷² In this regard, greater resting state functional connectivity between the SMA and the MLR positively correlates to FOG severity, possibly reflecting a maladaptive neural compensation⁷³. Indeed, the changes found in cerebral blood flow may well be compensatory and not reflect the primary pathological process. FOG may be caused by a combination of altered brain activity with impaired ability to compensate for that alteration. This may explain the manifestation of FOG during changes in motor behaviour, such as turning or initiating walking. These gait adaptations not only require a switch of motor program, but also more precise regulation of step length, gait timing and postural

adjustments⁶⁵. It should be taken into account that all fMRI studies investigated levodopa-responsive freezers. Hence, in levodopa-resistant FOG, compensation may be further impaired.

Beyond fMRI, intrasurgical single unit recordings of (supposed) PPN and STN in PD patients give more insight to this complicated dynamic process. Recent investigations reveal the presence of both alpha and beta oscillations within the PPN⁷⁴⁻⁸⁰. Beta oscillations were identified within rostral PPN while alpha oscillations are more prominent in caudal PPN^{76, 79, 80}. In general alpha oscillations were facilitatory for movements while beta oscillations were antikinetic⁷⁶⁻⁷⁸. Task-related modulation of PPN activity is shown to be spatially-specific: during motor imagery of gait, alpha and beta band power is increased mostly in the intermediate and dorsal part of the rostral PPN (around the inferior colliculus)⁸⁰. In contrast, the STN mostly changes activity on real executed hand movements⁸⁰. Another study showed more caudal PPN alpha activity to correlate with gait function.⁷⁶ It should be noted that these studies were performed in the diseased and not in the healthy state. In addition, even within the PPN, different cells on the same spot are connected to different networks⁷⁹. To reliably intervene in this complicated process will be a tremendous challenge.

Primate research and interfering with the PPN

The MLR is clearly important for gait, and its PPN component has generated much attention in relation to motor disorders⁸¹. It also has involvement in other functions including the regulation of cortical activity, sleep and diurnal rhythms, attention, learning and reward. In recent years, studies in the non-human primate have suggested that dysfunction of the PPN is at least partly responsible for axial motor symptoms of PD and that PPN stimulation may improve gait and postural control in patients with PD.

Early studies revealed that there is significant degeneration of PPN neurons in patients with advanced PD and in patients with multiple system atrophy with severe akinesia^{82, 83}. The

extent of neuronal loss in the PPN corresponds to the degree of loss of dopaminergic neurons in the substantia nigra compacta in PD. There is also some correlation between the degree of degeneration in the PPN and the severity of PD symptoms⁸³.

In non-human primates, unilateral lesioning of the PPN leads to a hemi-Parkinsonian state that resolves over time, whereas bilateral lesions in the PPN lead to a much more severe and permanent Parkinsonian state^{84, 85}. However, the gait disorder seen with PPN lesions is mostly postural instability and not FOG⁸⁰.

Another step forward was to electrically (macro) stimulate the PPN in normal primates⁸⁶. After stereotactic implantation of a DBS electrode unilaterally in the PPN in a normal macaque, no adverse effects on the motor performance of the animal were demonstrated. When subsequent trials of stimulation were performed (multiple sessions at several different frequencies ranging from 5 to 100 Hz), there was a significant decrease in motor activity with higher frequency DBS (45 to 100 Hz; Figure 4). There was also loss of postural control with 100 Hz stimulation. Lower frequency (5 to 10 Hz) stimulation caused contralateral arm tremor⁸⁷. This could be confounded by the close proximity of the superior cerebellar peduncle. This series of experiments also established the practical feasibility of chronic DBS of the PPN by a fully implanted system with no significant adverse effects—an important consideration for further primate studies and possible clinical application.

Activating the PPN by injection of GABA antagonists leads to improvement of motor activity in post-MPTP animals. In a series of experiments a stereotactically implanted stainless steel cannula was placed into the unilateral PPN of a macaque (n = 2) was held in place with dental acrylic (Figure 4). The activity of the PPN was then modulated with sequential injections of GABA agonist muscimol, GABA antagonist bicuculline and saline (control). The injections were then repeated after the monkey was rendered parkinsonian with intravenous MPTP. Motor activity was recorded using automated activity counts, clinical scoring and blinded videotape assessments⁸⁸. In the pre-MPTP state, inhibition of PPN using muscimol significantly reduced the activity of the monkey. In the post-MPTP animal, PPN stimulation using bicuculline significantly increased the overall motor activity. Moreover, it also improved

balance and posture. These experiments again demonstrated that inhibition of the PPN leads to akinesia. They further established that stimulation of the PPN in the parkinsonian state can lead to increased activity and reversal of other axial symptoms like postural control.

Moreover, both sets of experiments demonstrated marked bilateral clinical effect from unilateral manipulation of the PPN.

Axial symptoms in advanced PD generally respond poorly to current medical and surgical treatments, including STN and GPi DBS. To establish the feasibility of treating advanced PD with marked axial symptoms with PPN DBS, a DBS lead was stereotactically inserted into the PPN of monkey's (n = 2) and connected to the fully implanted internal pulse generator. Serial stimulation experiments were performed. The animals were then made parkinsonian with intravenous MPTP and DBS repeated⁸⁹⁻⁹¹. The results showed that low frequency PPN DBS (5 to 10 Hz) improved the activity of the MPTP-monkeys. Moreover, when PPN DBS and L-Dopa treatment were combined, the activity of the monkeys was significantly greater than either treatment given alone. Based upon these observations clinical translation rapidly followed.

PPN: Potential for treating FOG

World-wide, approximately 100 patients have received DBS of the PPN area⁹². The data referred to here should be considered PPN area rather than PPN since precise localization is difficult (See figure 5 for localization). Most human studies reported gait improvement with low frequency (10-25 Hz) stimulation although with a significant heterogeneity within and across studies (Table 1). Some studies show improvement of axial symptoms during ON⁹³ or OFF⁹⁴ with reduction of falls^{94, 95}. Low frequency stimulation (10-25 Hz) seems most effective⁹⁶. Interestingly, no human study found that high frequency (>100 Hz) stimulation had a detrimental effect on axial motor function, in contrast with the animal studies⁸⁶. Similar improvements are reported with PPN DBS in combination with other targets (caudal Zona incerta, STN, GPi)⁹⁷⁻⁹⁹. The effects of PPN implanted into the right side may have slightly

better outcome, possibly suggesting the presence of a 'dominant PPN'¹⁰⁰. However, the comparative efficacy of unilateral versus bilateral stimulation is far from resolved.

After bilateral PPN stimulation compared with unilateral stimulation, the time spent on a turn task was less, without changes in spatio-temporal measures of gait¹⁰¹. Thus, there seems to be a specific effect of PPN DBS on turning and body weight shifting between lower limbs.

This notion is consistent with animal studies showing that PPN stimulation mainly affects postural tone while cuneiform nucleus (CNF) stimulation promotes stepping. Accordingly, two case reports found a positive effect of PPN DBS on Pisa syndrome (i.e. a lateral trunk flexion)^{102, 103}. A clinicopathological study in two patients with progressive supranuclear palsy (hence not idiopathic PD) found that a placement in the PPN adjacent to the CNF produced some clinical benefit on gait initiation¹⁰⁴.

The mechanism of action of PPN DBS is still far from being elucidated and may be related to an effect on the cholinergic pathways connecting PPN and thalamus and involved in vigilance and sleep/awake cycle. In keeping with this hypothesis, PPN DBS improves reaction times¹⁰⁵ and attention in PD^{103, 106}. Alternative mechanisms engaging local locomotor networks, descending pathways and disrupting pathological cortical and subcortical oscillations with, for example, stimulation spread to the adjacent medial lemniscus or reticular formation also need to be considered. A study using [¹⁵O] H₂O PET in patients with advanced PD showed unilateral PPN DBS increased blood flow in subcortical regions such as the thalamus, cerebellum and midbrain region (including the MLR) as well as different cortical areas involving medial sensorimotor cortex extending into caudal SMA.¹⁰⁷ Interestingly, improved imagined walking distance after DBS of the subthalamic nucleus in PD patients is related to activity changes in the MLR, as shown by a recent PET study¹⁰⁸.

Problems with PPN area stimulation

What is the actual outcome of PPN area stimulation? Results to date are unfortunately inconclusive, both for theoretical and for methodological reasons.

Theoretical issues

It is not clear whether it is actually the PPN that is being stimulated, and, if yes, what area within the PPN. Indeed, the actual location of the PPN in humans remains a matter of debate¹⁰⁹⁻¹¹¹ and its precise delineation is unknown. Consequently, surgical teams have set, and aimed at, different targets^{94, 101, 112}, making it difficult, if not impossible, to conclude on the role of the PPN and to compare outcomes. Inter-individual variability of the brainstem further complicates the matter. As noted above, the PPN has degenerated in PD^{113, 114} with a cell loss correlated with gait disability¹¹⁵. Such degeneration of the targeted structure, together with the heterogeneity of its cell population, questions the mechanisms underlying the efficacy of stimulation, or lack thereof.

Practical issues

Patients under study underwent different surgical procedures from one center to the next, ranging from bilateral PPN area implantation together with STN or caudal Zona Incerta implantation^{93, 97, 98, 112}, bilateral PPN area implantation in patients already benefiting from STN stimulation⁹⁴, lone unilateral or bilateral PPN area implantation in patients without previous brain surgery^{95, 105}. This, together with the uncertainty regarding the structure actually stimulated, makes it difficult to interpret the reported results regarding specific PPN area stimulation effects.

Most studies failed to meet high quality standards, being open label and sometimes missing pre-surgery baseline data altogether. Other confounding factors include small sample sizes, ill-defined inclusion criteria, varied gait disorders, lack of specific gait and FOG assessments, differences in assessment procedures and outcome criteria, and the likely withholding of negative outcomes. Furthermore, unlike that to STN stimulation, the response to PPN area stimulation is slow, taking days to set in. It also wanes very slowly following stimulation

arrest. The effects also wane under stable stimulation conditions, making it mandatory to frequently adjust parameters settings. Because of these characteristics the widely used acute assessment protocols are inadequate to assess the effects of PPN area stimulation.

Quality of data: Close to one hundred papers have reported on PPN area stimulation in humans, but only 6 studies involved double-blind assessments^{94-96, 101, 116, 117}, and one was single-blinded⁹³. These studies involved small sample sizes, ranging from one¹¹⁷ to 7 patients¹⁰¹, as well as variable stimulation periods before assessment (one hour to six weeks). Objective assessment protocols also differed.

Assessment protocols: Methodological flaws and differences also make it difficult to pinpoint the specific effects of PPN area stimulation. Regarding motor assessments, many studies fail to report specific gait evaluation or objective measures of FOG before and after surgery.

Often, the UPDRS Activities of Daily Living (II) or Motor section (III) total scores and various combinations of axial and gait scores were the only outcomes. These are not sensitive or specific enough to clarify the effects of PPN area stimulation.

Furthermore, specific gait and FOG assessments lead to inconsistent or even contradictory results. The same is true regarding quantitative gait analysis^{101, 116, 118}. Thus, it is not clear whether PPN area stimulation, when effective, improves gait, FOG^{94, 96} or falls, related^{94, 96} or unrelated to FOG⁹⁵.

Inclusion criteria: Finally, which patients could benefit from PPN area surgery, if any, is not clear either. Well controlled studies showed that chronic outcome is highly variable, ranging from no effect (or even worsening), to mild or major improvement^{94, 95, 117} in FOG or frequency of falls. Yet, available data provide distressingly little information on the clinical profile of the patients and some studies failed to perform thorough pre-surgery assessments^{93, 105, 118}. In all studies, the phenotype of the patients appears quite variable, especially regarding quality of gait pattern, frequency and characteristics of falls, response to levodopa (both overall and for FOG specifically) and, as already mentioned, the prior brain

surgery status. Such variability in the characteristics of the patients is likely to influence the outcome of the surgery.

The distinction between levodopa responsive and resistant FOG is an important consideration when evaluating the effect of PPN stimulation. Levodopa responsive FOG generally responds well to treatment of the STN^{119, 120}. With advancing disease, FOG becomes levodopa-resistant, probably by progression of non-dopaminergic disease. Whether this is due to cholinergic failure and can be addressed by stimulating the cholinergic part of the PPN, or whether other neurotransmitters such as glutamate play their part, remains to be determined. Future studies need to be clear concerning what phenotype of FOG is included and which phenotype reacts most.

In summary, objective assessments suggest that PPN area stimulation might improve FOG in some patients, without significant effects on other gait parameters. FOG reduction may contribute to explaining a positive outcome on falls. Whether stimulation might also improve postural stability remains to be clarified. Altogether, it will not be possible to conclude on the best indication, if any, of PPN area stimulation until we know better where and how to stimulate, and until controlled studies with thorough inclusion criteria and consistent evaluation protocols are conducted.

Conclusion

While FOG remains a mysterious phenomenon, animal and imaging studies have started to unravel the mystery. The outcome of these studies point to a dynamic process of hypo and hyperactivation in cortical areas such as the SMA, as well as subcortical areas such as the striatum and the MLR including the PPN. The specific role of the PPN remains unclear.

Control of locomotion is very complex. It is a network operation of multiple interacting brain areas (or even brain networks) using multiple neurotransmitters. One way forward may be to look at the entire network instead of one detailed cell group in the network. When doing experiments, one has to be highly specific on the part of locomotion (freezing, posture,

slowness of gait) that is being studied. Whether different phenomenological subtypes of freezing (akinetetic, trembling, responsiveness to environment) have truly a different pathology or are a reflection of a different (extent of) defective node of the network, remains to be determined.

The dopaminergic component is a critical element. If only dopamine depletion is the problem, this will lead to OFF freezing. If then another element (cholinergic, glutamatergic) breaks down, the system breaks down leading to more complex freezing including ON freezing. Too much dopamine will also cause the system to fail because of its U-shaped function¹²¹.

For levodopa responsive FOG, STN stimulation offers opportunities¹²² but for levodopa resistant FOG in PD and other disorders, we are still left to ponder. Although the PPN is embraced as a possible target for FOG, especially for levodopa resistant FOG leading to falls, to date evidence is inconclusive. Further work is needed to investigate where, how and who to stimulate to get good results. Clear patient selection, thorough gait and falls assessment and strict definition of the surgical target will be fundamental for future research. Even then, the compensatory role of PPN area stimulation may be limited by disease progression.

Basic science studies suggest the PPN is most involved in tone and balance, more than in stepping. Consequently and fitting, patient studies hitherto suggest PPN stimulation may improve falls, more than freezing of gait. Giving attention to other possibly relevant areas in the dynamic pathophysiological process of FOG, such as the SMA or the MLR area dorsomedial to the PPN (including the laterodorsal tegmental nucleus) may provide further direction in the development of effective therapies.

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The authors have no relevant conflict of interest to report.

Figure Legends

Figure 1:

Neuronal mechanisms of controlling locomotion

- A. Dorsal system for cognitive locomotor control. A visuo-motor pathway from the visual cortex to motor cortex via the parietal cortex contributes to this control. Corticofugal projections act on to the basal ganglia nuclei, brainstem and spinal cord. Dopaminergic (DA) projection from the substantia nigral pars compacta (SNc) to the caudate and putamen (CPu) may be involved in learning the locomotor behaviors. GABAergic output from the basal ganglia nuclei (internal segment of the globus pallidus and substantia nigra pars reticulata; GPi/SNr) acts on MLR/PPN area may control locomotion and posture. Efferents from the midbrain locomotor region (MLR) activate both excitatory system and locomotor pathway. The excitatory system arises from the locus coeruleus (LC) and the raphe nuclei (RN). The locomotor pathway consists of reticulospinal neurons arising from the ventromedial medulla corresponding to the nucleus reticularis magnocellularis (NRMc). Cholinergic and glutamatergic projections from the PPN excite SNc-DA neurons. Cholinergic PPN neurons also excite pontine reticular formation neurons, which, in turn, activate reticulospinal neurons in the medial medulla corresponding to the nucleus reticularis gigantocellularis (NRGc). These descending tracts act on central pattern generator (CPG) in spinal cord to evoke locomotion. Efferents from the cerebellar locomotor region (CLR) may excite locomotor pathway.
- B. Ventral system for emotional locomotor control. Efferents from the amygdala (AMD) and hippocampus (Hipp) project to the nucleus accumbens (NAc). GABAergic NAc neurons project to ventral pallidus (VP) and the SNr, which control activity of the MLR/PPN neurons. Efferents from the amygdala and the hippocampus also act on lateral

hypothalamic area, which corresponds to the subthalamic locomotor region (SLR). DA projections from the ventral tegmental area (VTA) may contribute to the reward-oriented locomotor behaviors. See text for further details.

Abbreviations: E; extensor motoneurons, F; flexor motoneurons.

Figure 2:

Analysis of the evolution of FOG episodes in the MPTP-treated monkey.

A and B, recordings of leg movements with an accelerometer placed on the back of the leg during the whole duration of FOG episodes (~15 s). Each episode (A and B) has regular oscillations corresponding to tremor before the end of freezing. The traces show raw accelerometry data. C and D, rate meters for the whole duration of the FOG episodes corresponding to A and B, respectively. The peaks correspond to the tremor periods towards the end of the freezing episode when walk restarts and initial oscillatory movements. Rate meters used the data produced after detection of full phase oscillations above the threshold. The graphs were constructed with a bin width of 500 ms, and smoothed using a Gaussian filter. E, distribution of FOG tremor frequencies. The graph shows the frequencies (Hz) found across 20 recordings of tremor, each in a separate FOG episode. The average rate in the recorded tremors was 7.07 Hz (± 1.47 STD). Taken from ⁵⁰ with permission.

Figure 3:

Dynamic cerebral substrate of FOG, based on data from fMRI studies Snijders et al 2011, Vercruyssen et al 2014, Shine et al 2013a and 2013b

During repetitive continuous movement of hands or legs, or motor imagery of gait, cortical activity in areas as the SMA and PFC is reduced, while subcortical activity is increased; for the STN in the hands, in the MLR for leg movements or imagery of gait. During a freezing

episode, SMA activity is still reduced, but also subcortical areas STN and MLR show reduced activity. In contrast, frontoparietal areas and the insula show increased activity.

Abbreviations: SMA = supplementary motor area; PFC = prefrontal cortex; STN = subthalamic nucleus; MLR = Mesencephalic locomotor region; DLPFC = dorsolateral prefrontal cortex

Figure 4:

In normal macaque, multiple sessions of DBS unilateral PPN stimulation at several different frequencies: significant decrease in motor activity with higher frequency DBS (45 to 100 Hz).

Figure 5:

(A) (A) Location of the PPN identified on the ICBM 2009b NLIN T1 asymmetric template by Andreas Horn, Michael D. Fox and Clifford B. Saper, by transferring the location of ChAT-staining of neurons in the PPN in human brain¹²³ to standard atlas space using neuroanatomical landmarks.

(B) MRI with bilateral electrodes for stimulating the PPN. Note that to implant the PPN region the surgeon has to avoid traversing the lateral ventricles which would increase the risk of hemorrhage. To remain intra-parenchymal the surgeon approaches the target calculating the azimuth and declination to place as many contacts in the PPN area as possible. (Part B courtesy of Dr. Peter Silburn, Brisbane Australia)

Supplementary Video 1:

Monkey with FOG in the hind limbs with many of the characteristics typical of human FOG, including a 4-10 Hz tremor similar to that seen in patients⁵¹.

Table 1. Summary of published clinical studies of PPN DBS for freezing of gait in Parkinson's disease

Study design	Author-year	N	Type patient	Target	Stimulation parameter	Follow-up in months	Outcomes
Prospective	Stefani-2007 ⁹³	6	Disabling axial signs	Bilateral STN and PPN	2V 60 μ s 25 Hz	6	Combined therapy improved axial motor symptoms. Efficacy declined over time.
	Ferraye-2010 ⁹⁴	6	Severe freezing unresponsive to levodopa and STN DBS	Bilateral STN and PPN	1.2-3.8V 60-90 μ s 15-25 Hz	12	Double-blind: no change. Improvement in OFF freezing and falls. Significant inter-subjects variability.
	Moro-2010 ⁹⁵	6	Severe freezing & falls unresponsive to medical treatment	Unilateral PPN	2V 70 μ s 67 Hz	12	No overall improvement. Improvement in ON & OFF falls. Post-hoc: slightly better right PPN ¹⁰⁰
	Khan-2011 ⁹⁸	7	Freezing & falls unresponsive to levodopa	Bilateral PPN & Zona Incerta	2.6V 60 μ s 60 Hz	13	Improved axial UPDRS-III subscore in OFF state. Synergic effect between Zi and PPN stimulation.
	Khan-2012 ⁹⁷	5	Freezing & falls unresponsive to levodopa	Unilateral and Bilateral PPN & Zona Incerta		27.0 \pm 19.6	Improved axial UPDRS-III subscore in ON state with bilateral

							PPN stimulation. Synergic effect of PPN with Zi stimulation.
Nosko-2015 ⁹⁶	9	Severe gait disorders & freezing	Bilateral PPN & STN	0.8-2.6V 60 μs 10-25 Hz	12		Lower frequency more efficacious than high frequency.
Thevathasan-2010 ¹⁰⁵	11	Severe freezing & falls unresponsive to medical treatment	Bilateral PPN, also Zona Incerta in 3 patients	2-3.7V 60 μs 20-35 Hz	12.7±12.1		Speed of reaction improved by stimulation.
Thevathasan-2011 ¹²⁴	8	Severe freezing	Bilateral PPN	2.5-4.3V 60 μs 30-35 Hz	19.7±9.2		PPN DBS restores proximal StatReact and reaction time
Thevathasan-2012 ⁷⁶	7	Severe freezing	Bilateral PPN	2.2-4.3V 60 μs 35 Hz	<2.5		Freezing improved with mere electrode insertion. Alpha oscillations caudal PPN correlate with gait function
Mazzone-2011 ¹¹⁶	23	Freezing	Unilateral PPN		21.3±10.0		DBS improved both UPDRS-III and the axial subscores
Mazzone-2014 ¹²⁵	10	Drug resistant gait disabilities	Unilateral PPN	2.5-3V 60 μs 40 Hz	12		DBS improved some parameters of gait analysis

Transactional or retrospective

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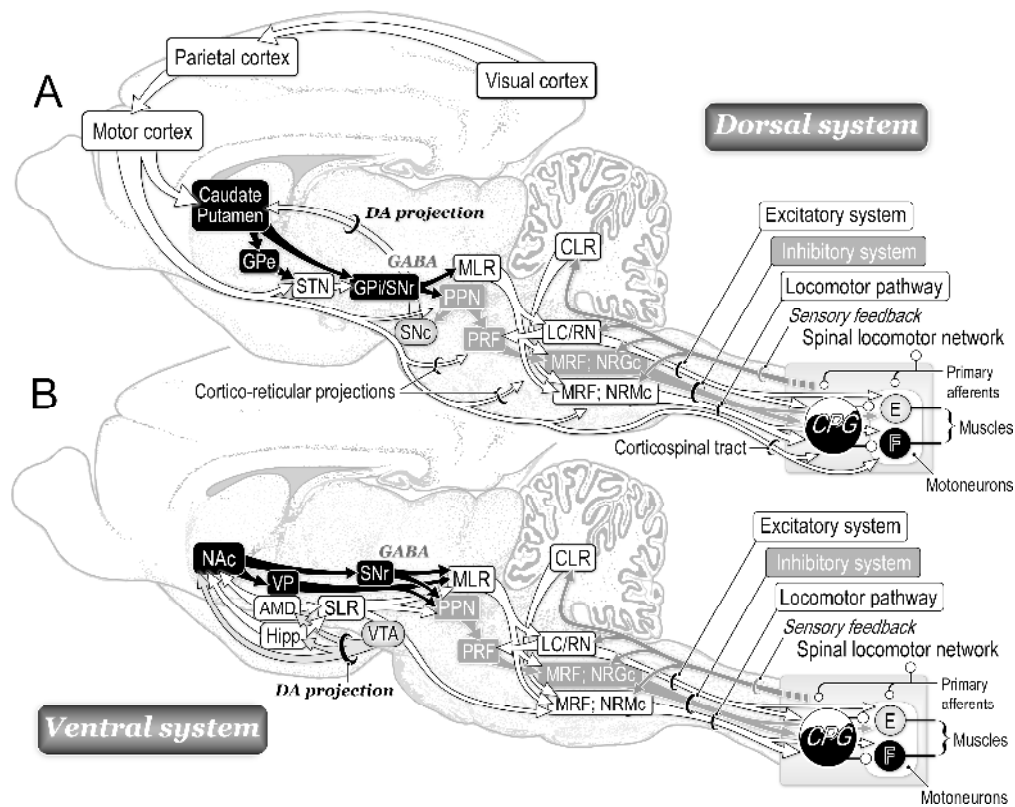


Figure 1:

Neuronal mechanisms of controlling locomotion

A. Dorsal system for cognitive locomotor control. A visuo-motor pathway from the visual cortex to motor cortex via the parietal cortex contributes to this control. Corticofugal projections act on to the basal ganglia nuclei, brainstem and spinal cord. Dopaminergic (DA) projection from the substantia nigral pars compacta (SNc) to the caudate and putamen (CPu) may be involved in learning the locomotor behaviors. GABAergic output from the basal ganglia nuclei (internal segment of the globus pallidus and substantia nigra pars reticulata; GPi/SNr) acts on MLR/PPN area may control locomotion and posture. Efferents from the midbrain locomotor region (MLR) activate both excitatory system and locomotor pathway. The excitatory system arises from the locus coeruleus (LC) and the raphe nuclei (RN). The locomotor pathway consists of reticulospinal neurons arising from the ventromedial medulla corresponding to the nucleus reticularis magnocellularis (NRMc). Cholinergic and glutamatergic projections from the PPN excite SNc-DA neurons. Cholinergic PPN neurons also excite pontine reticular formation neurons, which, in turn, activate reticulospinal neurons in the medial medulla corresponding to the nucleus reticularis gigantocellularis (NRGc). These descending tracts act on central pattern generator (CPG) in spinal cord to evoke locomotion. Efferents from the cerebellar locomotor region (CLR) may excite locomotor pathway.

B. Ventral system for emotional locomotor control. Efferents from the amygdala (AMD) and hippocampus (Hipp) project to the nucleus accumbens (Nac). GABAergic Nac neurons project to ventral pallidum (VP) and the SNr, which control activity of the MLR/PPN neurons. Efferents from the amygdala and the hippocampus also act on lateral hypothalamic area, which corresponds to the subthalamic locomotor region (SLR). DA projections from the ventral tegmental area (VTA) may contribute to the reward-oriented locomotor behaviors. See text for further details.

Abbreviations: E; extensor motoneurons, F; flexor motoneurons.

Figure 1
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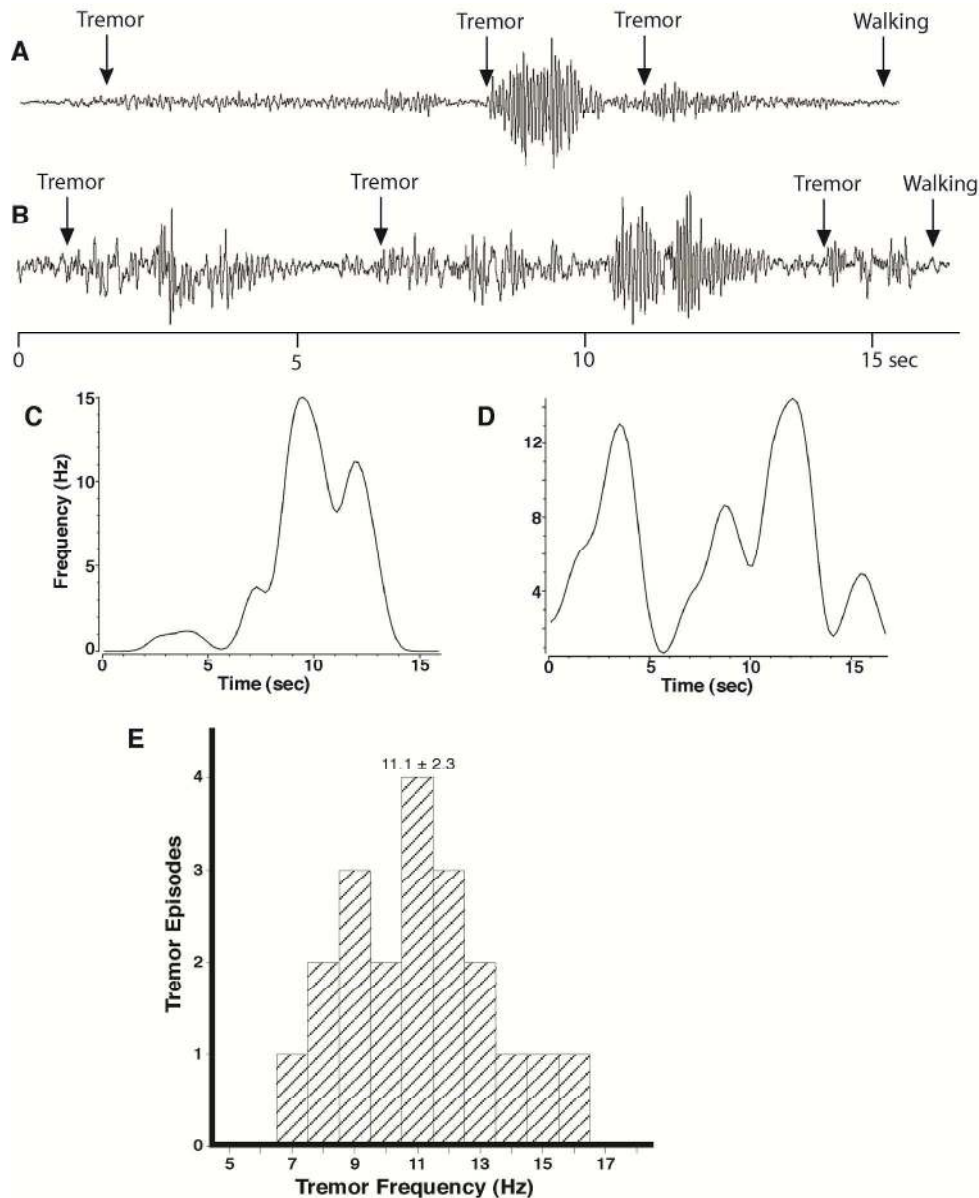


Figure 2: Analysis of the evolution of FOG episodes in the MPTP-treated monkey. A and B, recordings of leg movements with an accelerometer placed on the back of the leg during the whole duration of FOG episodes (~15 s). Each episode (A and B) has regular oscillations corresponding to tremor before the end of freezing. The traces show raw accelerometry data. C and D, rate meters for the whole duration of the FOG episodes corresponding to A and B, respectively. The peaks correspond to the tremor periods towards the end of the freezing episode when walk restarts and initial oscillatory movements. Rate meters used the data produced after detection of full phase oscillations above the threshold. The graphs were constructed with a bin width of 500 ms, and smoothed using a Gaussian filter. E, distribution of FOG tremor frequencies. The graph shows the frequencies (Hz) found across 20 recordings of tremor, each in a separate FOG episode. The average rate in the recorded tremors was 7.07 Hz (± 1.47 STD). Taken from (Revuelta et al) with permission.

Figure 2

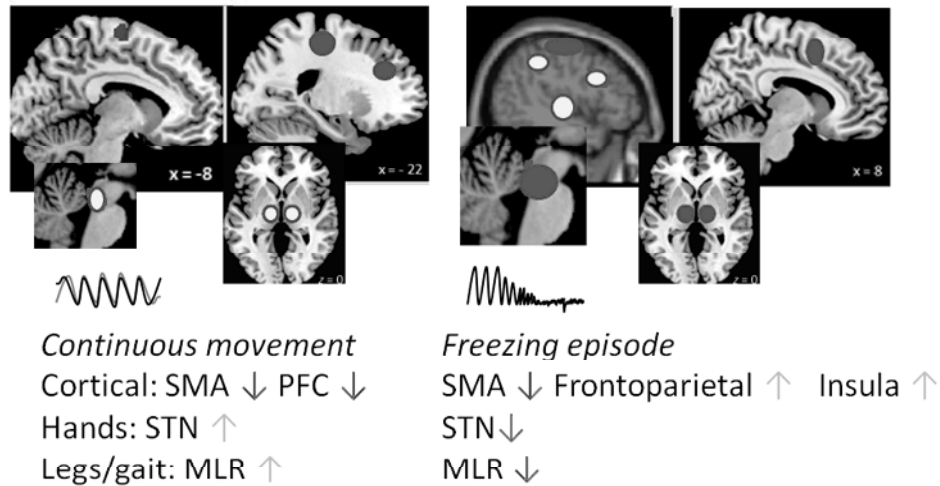


Figure 3: Schematic cerebral substrate FOG following from fMRI studies

Figure 3:

Dynamic cerebral substrate of FOG, based on data from fMRI studies Snijders et al 2011, Vercruyse et al 2014, Shine et al 2013a and 2013b

During repetitive continuous movement of hands or legs, or motor imagery of gait, cortical activity in areas as the SMA and PFC is reduced, while subcortical activity is increased; for the STN in the hands, in the MLR for leg movements or imagery of gait. During a freezing episode, SMA activity is still reduced, but also subcortical areas STN and MLR show reduced activity. In contrast, frontoparietal areas and the insula show increased activity.

Abbreviations: SMA = supplementary motor area; PFC = prefrontal cortex; STN = subthalamic nucleus; MLR = Mesencephalic locomotor region; DLPFC = dorsolateral prefrontal cortex

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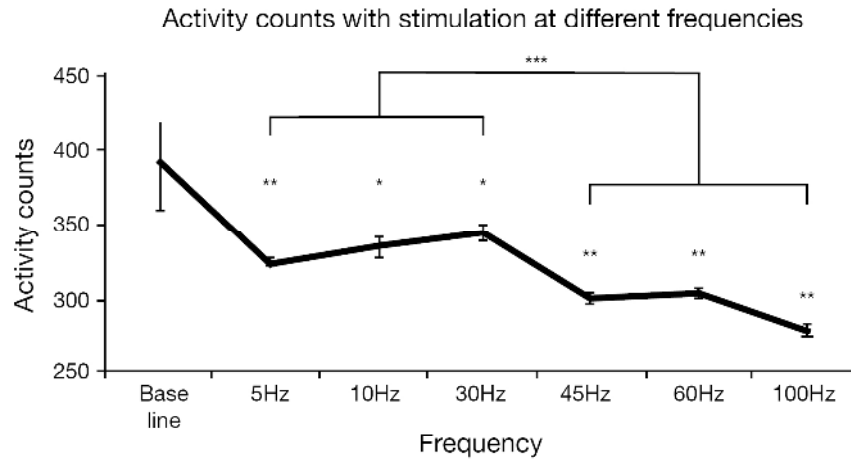


Figure 4: In normal macaque, multiple sessions of DBS unilateral PPN stimulation at several different frequencies: significant decrease in motor activity with higher frequency DBS (45 to 100 Hz).

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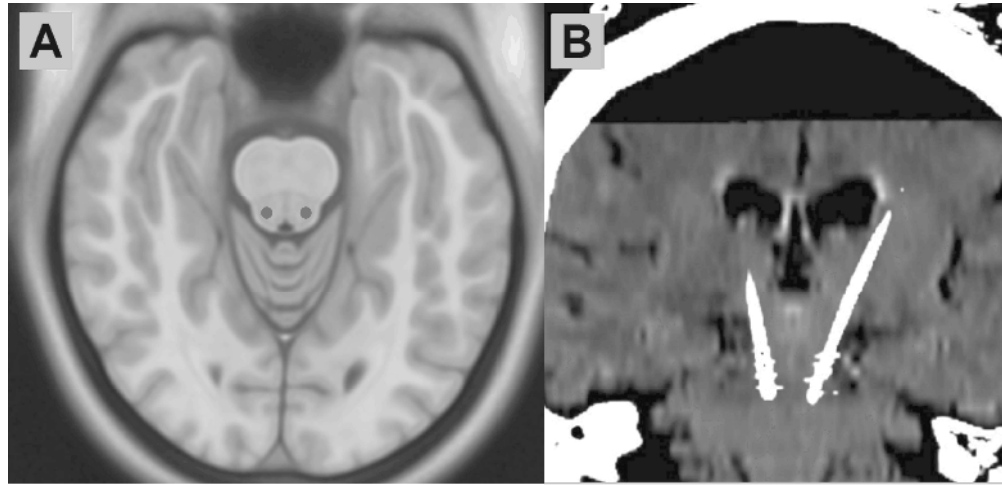


Figure 5:

(A) Location of the PPN identified on the ICBM 2009b NLIN T1 asymmetric template by Andreas Horn, Michael D. Fox and Clifford B. Saper, by transferring the location of ChAT-staining of neurons in the PPN in human brain 123 to standard atlas space using neuroanatomical landmarks.

(B) MRI with bilateral electrodes for stimulating the PPN. Note that to implant the PPN region the surgeon has to avoid traversing the lateral ventricles which would increase the risk of hemorrhage. To remain intraparenchymal the surgeon approaches the target calculating the azimuth and declination to place as many contacts in the PPN area as possible. (Part B courtesy of Dr. Peter Silburn, Brisbane Australia) !! †

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