Transcranial Magnetic Stimulation in Tourette Syndrome: A Historical Perspective, Its Current Use and the Influence of Comorbidities in Treatment Response

Marco Grados 1,*, Rachel Huselid 2 and Laura Duque-Serrano 3

1 Johns Hopkins University School of Medicine, Baltimore, MD 21287, USA
2 Johns Hopkins University Krieger School of Arts & Sciences, Baltimore, MD 21205, USA; rhuseli1@jhu.edu
3 School of Medicine, Universidad de los Andes, Bogotá 111711, Colombia; l.duque75@uniandes.edu.co

* Correspondence: mgrados1@jhmi.edu; Tel.: +1-443-287-2291

Received: 6 April 2018; Accepted: 28 June 2018; Published: 6 July 2018

Abstract: Background. Tourette syndrome (TS) is a childhood-onset neuropsychiatric disorder consisting of impairing motor and vocal tics which often persists adolescent and adult years. In this older refractory group, standard treatments such as pharmacotherapy and psychotherapeutic interventions may only have limited effects. Based on electrical cortical dysregulation in individuals with TS, a novel approach has employed brain stimulation strategies to modulate the putative aberrant neural electrical activity in pathways that may underlie tics, such as insula-supplementary motor area (SMA) connectivity. Methods. This review will examine all published clinical trials employing transcranial magnetic stimulation (TMS) to ameliorate tics, and discuss a framework for the pathophysiology of TS in relation to electrical brain activity. A framework for future research in tic disorders using TMS and imaging targeting neuroplasticity will be discussed. Results. Therapeutic electrical brain activity modulation with TMS has been carried out in stroke neuro-rehabilitation and neuropsychiatry, including trials in TS. Eleven trials document the use of TMS in TS targeting several brain areas, a positive effect is seen for those trials targeting the SMA. In particular, it appears that younger individuals with concurrent attention-deficit hyperactivity disorder (ADHD) benefit the most. Conclusions. TMS can be used as an effective tool to explore the psychophysiology of TS and potentially provide a therapeutic option. Ultimately, translational research using TMS in TS needs to explore connectivity differences pre- and post-treatment in individuals with TS that are linked to improvement in tic symptoms, with an emphasis on approaches using functional neuroimaging as well as other probes of neuroplasticity.

Keywords: transcranial magnetic stimulation; tourette syndrome; insula; supplementary motor area

1. Introduction

Tourette syndrome (TS) is a developmental neuropsychiatric disorder characterized by a childhood onset, male predominance [1], familial underpinnings [2], and often a life-long course [3]. Clinically, TS is defined by the presence of clinically impairing multiple motor, and at least one vocal tic occurring for over 12 months [4]. TS is frequently comorbid with attention-deficit hyperactivity disorder (ADHD) and obsessive-compulsive disorder (OCD); additionally, an excess of autism spectrum disorder (ASD) traits, learning difficulties, sensory integration disorder are also commonly reported [5]. Etiologically, sharing of genetic vulnerabilities for OCD [6], ADHD [7], disorders related to grooming (trichotillomania, skin picking), and mood dysregulation (bipolar disorder) [8] have been
suggested, while more recent findings point to deficits in neurodevelopmental aspects of synaptic function [9].

The pathophysiology of TS is not fully understood, but increased dopamine release in the striatum (putamen) in conjunction with low serotonergic tone [10], and developmentally immature brain connectivity [11] are posited, among competing theories. Clinically, antecedent to the execution of motor tics, patients often report a premonitory urge, an uncomfortable sensation in a muscle group, which can be alleviated by the tic behavior [12]. The premonitory urge is related to a heightened interoceptive awareness, the sensation of internal bodily states, as opposed to exteroception, the sensation from cutaneous mechanoreception and proprioception [13]. Supporting this theoretical framework, the insula, the brain’s interoceptive center, is observed to be overactive before tic behaviors [14]. Further associating interoceptive abnormalities in TS to the origin of tics, resting-state functional magnetic resonance imaging (fMRI) probes of tic phenomena, show that the dorsal anterior insula is hyperconnected to the supplementary motor area (SMA), a motor planning area [15]. Given the functional properties of insular cortex, such as the integration of sensory and emotional information, and processing of bodily sensations [16], it is plausible that methods that modulate insular activity may decrease internal tic urges and reinforce urge suppression activity.

Among the methods to induce or suppress targeted neural activity, inducing neuroplastic changes in cortical neurocircuitry, is transcranial magnetic stimulation (TMS). TMS is a noninvasive procedure which employs extracranial magnetic fields to create cortical electrical activity, which creates a stimulatory impulse to cortical neurons. Depending on TMS parameters, either induction, suppression or neural activity can be attained. In TS, natural targets for TMS have historically included motor and premotor cortex [17], but more recently, the SMA [18]. The aim of this narrative review is to examine the historical context of the use of TMS in TS, and to highlight published studies which use TMS to ameliorate TS symptoms both in adults and children. A framework for understanding predictive factors for the efficacy of TMS in TS is also offered.

2. Transcranial Magnetic Stimulation

TMS is based on the application of a magnetic field generated by pulsating electrical currents in a coil applied to cranial structures, which itself induces electrical activity intracranially in discrete brain regions. Magnetic fields used in TMS are based on the Biot–Savart law, discovered by Jean-Baptiste Biot and Felix Savart in 1820, which explains how electrical current generate magnetic fields [19], and Faraday’s law which postulates that fluctuating magnetic fields induce a “wave of electricity” [20]. These mechanisms generate, electricit in neuronal spaces and depolarize columns of neurons [21]. Technical considerations coil shape, brain region landmarks, and field strength (1.5–2 tesla). In repetitive TMS (rTMS), the magnetic stimulators that produce the electrical current are able to cycle at high rates to produce high pulse frequencies. High pulse frequencies generate a high degree of synchrony firing in neurons given that the period of inhibition after a TMS pulse lasts about 100 ms. The effect of TMS is able to establish hemispheric dominance [22]. And affect learning [23]. More recent data support rTMS reduction of inhibitory neurotransmission to facilitate associative plasticity in hippocampal slice cultures [24]. Enhancing associative synaptic plasticity [25]. The first therapeutic use of TMS was foreshadowed by Barker in 1985, reporting on the “novel method” of contactless stimulation of the human motor cortex [26]. By 2014, a targeted review of TMS in psychiatric disorders noted more than 600 papers mentioned TMS in relation with psychiatric disorders [27].

3. Neurocircuitry and Neurotransmitters Glutamate/GABA Suggest Abnormal Excitatory Activity in Tourette Syndrome

In the pathophysiology of TS, cortico-striatal-thalamo-cortical circuits (CSTC) connecting brain cortical regions to basal ganglia, are critical to its presumed pathophysiology [28]. These loops are active in parallel, but also harbor interconnections to support the interplay of the motor, emotional, and cognitive domains [29]. A direct excitatory pathway from striatal medium spiny neurons (MSN) to
globus pallidus, and an indirect inhibitory pathway from the globus pallidus through the subthalamic nucleus [30] set the stage for the basal ganglia to function as a modulating center for planned and executed movements, inhibiting motor mechanisms which compete with planned movements [31]. At the neurotransmitter level, dopamine and glutamate—via the glutamatergic subthalamic nucleus STN in the indirect pathway—are the key neurotransmitters implicated. Single photon emission tomography (SPECT) and positron emission tomography (PET) studies, which tag dopamine ligands with radioactive traceable isotopes confirm increased density of striatal dopamine transporter density [32,33], increased 18F-fluoro-dopa uptake in the left caudate [34], greater dopamine release in putamen after amphetamine challenge [35], and increased 11C-dihydro-tetrabenazine binding in the right ventral striatum [36]. Mesocortical dopaminergic inputs from the ventral tegmental area, in turn, can alter glutamatergic pyramidal neuronal excitability both directly and indirectly; the indirect pathway using modulation from gamma-amino-butyric acide GABAergic interneurons [37]. In addition, a crucial role is played by glutamate in tonic/phasic ventral tegmental dopaminergic signaling via excitatory ventral subiculum hippocampal glutamate neurons [38]. A transgenic mouse has modeled chronic glutamate excitation of striatal motor output biased towards overactivity of the motoric direct pathway and inactivity of the inhibitory indirect pathway. In this model, a hyperglutamatergic CSTC “tic circuit” is proposed [39]. The only human post-mortem study assessing brain glutamate levels found reduced amounts in globus pallidus interna, globus pallidus externa, and substantia nigra pars reticulata [40]. More recently, a genomic study implicates astrocytic function and glutamate and glucose metabolism with TS [41], and a case control study showed that a gamma-amino butyric acide (SLC1A3; also known as excitatory amino acid transporter 1 or EAAT-1) variant (E219D) resulted in increased uptake of glutamate by astrocytes [42]. Despite pathophysiological studies implicating glutamate in TS, clinical trials have not provided robust support for use of glutamate-modulators in this condition [43]. However, resting-state fMRI and magnetic resonance spectroscopy (MRS) studies have continued to provide support for “immature connectivity” [11] and reductions in striatal concentrations of glutamine, glutamate + glutamine (Glx) in TS. Glutamine levels in the striatum were inversely associated with tic severity, while thalamic glutamate was inversely associated with tic urges [44]. In summary, dopamine and glutamate influences are widely recognized in TS, however, the counterbalance to glutamate transmission in the CNS, inhibitory GABA loads, may also play an influential role in the expression of TS.

While the abnormal glutamatergic drive theory in TS favors an excess of glutamate in the origin of tics, a lack of neuronal inhibitory capacity could equally account for tics through an excitation-inhibition (E/I) imbalance of the GABA-glutamate-glutamine cycle [44]. For TS, a developmental dysfunction of GABAergic striatal interneurons is a plausible component of an E/I imbalance. GABA neurons, in fact, temporally antecede glutamate neurons in development and act as the first excitatory neurons, with GABA stimulation being the sole excitatory transmitter early in fetal development [45]. While a GABA-driven hyperexcitable state is developmentally present early in life, over time, inhibitory neurocircuitry achieves maturity through the “GABA switch” from excitatory to inhibitory functions. This process in GABA neuron development may be facilitated by oxytocin [46] and is evolutionarily preserved: early in C. elegans development GABA neurons produce muscle contraction through depolarization, while at the mid-larval stage the same neurons are hyperpolarizing and relax muscle [47]. If this switch is incomplete it can lead to epileptogenesis [48] and possibly other neurodevelopmental conditions [49]. In humans, the E → I switch for GABA neurons occurs postnatally at weeks 1–2 [50], with GABA load in neonates being lower than that of children [51]. The lack of developmental maturity in TS [11], specifically in basal ganglia interneurons, may be one factor predisposing children to tics. One post-mortem study has found that the tangential migration of GABA parvalbumin-positive interneurons may be abnormal in TS, with lower densities of interneurons in caudate-putamen (striate nuclei) and a higher proportion in the internal globus pallidus [52]. In sum, the delayed maturation of striatal GABAergic interneurons is an attractive emerging model for TS, a model which has been recently shown to cause dystonia, a related movement disorder [53].
The influence of cortical GABA in the clinical expression of TS has been examined, suggesting decreased cortical GABA in TS, although with developmental differences. In children 5–12 years with TS, one study used 1H magnetic resonance spectroscopy (MRS) at 7T and found no cortical GABA perturbations, but confirmed abnormal glutamate function in the premotor cortex [54]. In another study of 8–12-year-old children with TS and healthy controls, a reduced GABA concentration in primary sensorimotor cortex was detected using MRS [55]. At older ages, GABA imbalances may be more evident in TS. In 15 adolescents with TS, similarly using 1H MRS at ultra-high field (7 T), GABA was increased in SMA and was negatively correlated with cortical excitability [56]. In brain regions dependent on GABA interneurons for E/I homeostasis, inhibitory synaptic potentials coming from even single striatal interneurons can easily modulate the origin of action potentials in a large number of projection neurons [57], implying that maturational or other dysfunction affecting interneurons can readily dysregulate cortical or striatal neurocircuitry. However, complicating the clinical argument for an E/I imbalance in TS, compensatory mechanisms in individuals with TS may develop over time due to chronic tic suppression, resulting in an acquired “tonic” inhibition state [58].

Cortical probes of GABA function with TMS include use of a paired-pulse paradigm with TMS to measure the GABA-mediated short-interval intracortical inhibition (SICI). In SICI, subthreshold stimulation, not sufficient to produce a corticospinal output, is applied as a condition pulse 1–5 msec before the test suprathreshold pulse. This paired stimulation significantly suppresses the amplitude of the motor evoked potential (MEP) of the second pulse [59]. Several studies have explored the association of SICI with TS. Interestingly, tic severity and ADHD hyperactivity-impulsivity scores in children and adults are correlated with less intracortical inhibition (lower SICI), while there is no relationship with obsessive-compulsive disorder (OCD) severity scores. Also, intracortical facilitation, the production of larger MEPs through more widely spaced TMS paired pulses that facilitate motor output, is not related to ADHD, tic or OCD clinical severity [60]. The relationship between lower SICI and ADHD symptoms is attenuated significantly in the presence of dopamine receptor blockers, possibly based on the regulation of cortical inhibition by D1 and D4 receptors involved in glutamate pathways. Speculatively, the same mechanisms that regulate subcortical inhibition, which when dysregulated could lead to tics, may be those that influence cortical inhibition [61].

4. Brain Electrical Activity and Tourette Syndrome

Having established that glutamatergic and GABAergic transmission may have a role in the development and maintenance of tics in TS, electrical activity in TS has been explored. An examination of surface electromyography recorded from the right abductor digiti minimi muscle was completed after TMS application to the left motor cortex. While motor threshold and peripheral motor excitability were normal in TS individuals, the cortical silent period was shortened and intracortical inhibition reduced. A subgroup analysis revealed that these differences were prominent in the presence of tics [62]. A more recent study used single-pulse TMS in conjunction with a manual Go/NoGo task to investigate alterations in corticospinal excitability ahead of volitional movements. In ten adolescents with TS, corticospinal excitability was significantly reduced in TS in the period immediately preceding a finger movement. TMS-induced motor evoked potentials were also abnormal only in the TS group, suggesting an inability to modulate motor cortical excitability prior to tics [63]. Motor evoked potentials have also been studied in relation to paired associated learning in TS, in order to explore neuroplasticity potential in the disorder. Paired associative stimulation in healthy controls produces long-term improvement, showing long-term potentiation due to neuroplasticity; however, individuals with TS did not show the improvement when tested nine months after paired associative stimulation. There was a statistical signal for the association of less synaptic plasticity with more severe tics [64]. Others studies have also shown aberrant motor cortical plasticity in TS using the paired associative stimulation paradigm [65].

While these studies establish that cortical electrical activity is abnormal in TS, the influence of OCD and ADHD on brain electrical activity needs to be elucidated. A study comparing cortical and brainstem plasticity in individuals with TS versus OCD alone, found that abnormal plasticity is consistently present...
in TS, but is normal in those individuals with OCD alone [66]. Along these lines, the main comorbid disorder with TS, ADHD has also been found to be associated with increased cortical excitability. In one study, motor thresholds, GABA-determined SICI, intracortical facilitation, and short latency afferent inhibition were measured with TMS in 18 TS only, 6TS + ADHD, and 5TS + OCD individuals, in comparison to 24 healthy subjects. In one notable finding in this report, individuals with TS + ADHD had more intracortical facilitation than controls, compared to individuals with uncomplicated TS and TS + OCD. It appears that ADHD cortical instability confers greater brain electrical imbalance in addition to that already present in TS [67]. These findings underscore the notion that neuropsychiatric disorders, such as ADHD and TS, with excess movement may involve increased glutamatergic excitatory output from thalamus to cortical centers, resulting in either excessive motor dyscontrol (hyperactivity/impulsivity) or tic behaviors (suppressible but difficult to contain). In another study which used TMS to measure motor cortex inhibition in 36 children and adults with TS, severity of ADHD symptoms, and motor tics were independently associated with SICI (r(2) = 0.50; F[2,27] = 13.7; p < 0.001), particularly in subjects not taking neuroleptics (r(2) = 0.68; F[2,17] = 17.8; p < 0.0001). The correlation with cortical disinhibition was mostly dependent on ADHD symptom severity (r = 0.53; p = 0.003) compared to tic severity (r = 0.42; p = 0.02). A continuum of electrical brain overactivity may thus exist in a gradient from TS + ADHD, TS + OCD to OCD only. Critically, the findings suggest a potential preference for using comorbid groups as targeted subpopulations in TMS [60].

5. Transcranial Magnetic Stimulation as a Measure of the Electrophysiology in Tourette Syndrome

As TMS can measure cortical brain electrical activity, several studies have examined these properties in TS and found specific abnormalities. Orth et al. (1992) measured motor thresholds, input-output (I/O) curves, SICI, and cortical silent periods (SP) with TMS in 20 untreated TS participants (12 uncomplicated, four with comorbid ADHD, 4 with comorbid OCD) and 24 healthy subjects. Tics were rated with standard clinical scales and detailed video analysis. There was lower corticospinal excitability at rest in TS compared to controls, speculating that this may be an adaptive response to reduce release of unwanted movements [68]. In another study measuring voluntary motor drive (VMD), the presence of distal motor tics correlated with lower VMD compared to non-distal motor tics and controls [69]. More dramatically, electrical activity measured in cortical brain areas in TS suggests lack of maturation of neurocircuitry, as evidenced by increased resting motor thresholds and variability of motor evoked potential responses in children with TS compared to adults [70]. In addition, lack of neuroplastic neuronal reserves has also been postulated to underlie the lack of natural compensatory ability to suppress and eliminate tics seen in healthy individuals. In order to test the neuroplasticity potential with TMS, Jackson et al. (2013) measured cortical excitability, using TMS-induced MEPs, and found them to be significantly reduced in the TS group in the period immediately preceding a finger movement [71]. These changes in electrical overactivity and lack of neuroplasticity in TS suggest a combination of a glutamatergic imbalance [44], a GABA interneuron deficiency [72], and given recent genetic data, an intrinsic synaptic function abnormality [73]. The latter hypothesis may lead to further studies in neuroplasticity effects of TMS in TS, based on the the long-term potentiation (LTP)-like and long-term depression (LTD)-like effects of brain stimulation [74], as shown additionally by in vitro organotypic preparations, in which repetitive magnetic stimulation induces long-lasting strengthening of glutamatergic synapses and remodeling of dendritic spines [25].

6. Transcranial Magnetic Stimulation as a Therapeutic Tool in Tourette Syndrome

Clinical trials using both transcranial direct current stimulation (tDCS; the delivery of constant, low direct current delivered via electrodes on the cranium) and rTMS have explored the safety and efficacy of noninvasive brain stimulation as a treatment for children and adults with TS. While tDCS has not been studied extensively as a therapeutic option in TS, it has shown promise for its possible efficacy and confirmed safety [75]. Sparse data include a clinical case study using tDCS targeting the
left motor cortex in two adult patients with TS, in which tDCS resulted in a significant reduction in tics compared to sham treatment [76].

The use of rTMS as a treatment for TS has been studied far more extensively. Several early studies examined the efficacy of 1Hz rTMS targeting the left motor cortex and left pre-motor cortex and found no significant reduction in TS tic symptoms overall [17,77]. Subsequently, Chae et al. tested rTMS efficacy targeting the left motor cortex and left prefrontal cortex, and found that while there was no significant reduction in tic severity overall, there was a significant reduction of tic severity in the TS/OCD comorbid subgroup [78], suggesting that the presence of comorbidities might influence results. The positive effect of TMS in comorbid TS was also identified by Bloch et al., who targeted the SMA using deep rTMS. In that study, a significant reduction in tic severity was only seen in the TS/OCD comorbid subgroup [79].

In children, an open-label pilot study tested the efficacy of low frequency rTMS targeting the SMA in ten males with TS ages 9–14 years. Over a 12-week period, the average resting motor threshold rose significantly, while tic severity decreased, although no significant effects on ADHD, anxiety, or depression were seen [80]. A similar study on the effects of daily rTMS targeting the SMA in children with TS also produced lower scores on the Yale Global Tic Severity Scale. A higher resting motor threshold post-treatment was found, in conjunction with lower scores on the Children’s Depression Inventory for depression and (Swanson, Nolan and Pelham) SNAP-IV inventory for ADHD [81]. The safety of TMS is documented by a review of 48 studies in children with various neurological conditions, which showed only minor side effects [82]. (Table 1).
Table 1. Clinical Trials of Transcranial Magnetic in Children and Adults with Tourette Syndrome.

| Authors            | Design                  | Age (Range) | N  | MF | Comorbidities                  | Intervention | Comparison Groups | Targeted Brain Area | Results                                                                 |
|--------------------|-------------------------|-------------|----|----|--------------------------------|--------------|------------------|---------------------|-------------------------------------------------------------------------|
| **Pediatric TS TMS Clinical Trials**                                                                                                                                  |
| Kwon, 2011 [80]    | Open label pilot study  | 9.57 ± 2.75 (9–14) | 10 | 10:0 | OCD (n = 1), ADHD (n = 3) | rTMS         | None             | SMA                 | Tic symptoms improved significantly, no improvement in ADHD, anxiety or depression. |
| Le, 2013 [81]      | Open label              | 10.61 ± 2.18 (7–16) | 25 | 22:3 | Not specified                  | rTMS         | None             | SMA                 | Tic severity significantly lowered, also hyperactivity, attentional deficits, depression and anxiety lower with treatment. |
| **Adult TS TMS Clinical Trials**                                                                                                                                        |
| Bloch, 2016 [79]   | Open label              | 32.6 ± 12.7 (20–61) | 12 | 1:1 | OCD (n = 6), ADHD (n = 4) | Deep rTMS    | None             | SMA                 | Tics did not improve overall but comorbid TS + OCD significant improvement tic severity. |
| Chae, 2004 [78]    | Randomized, blinded crossover | 34.9 ± 16.4 (13–60) | 8  | 5:3 | OCD (n = 4), ADHD (n = 3) | rTMS         | None             | L MC, L PFC         | Tic and OCD symptoms improved significantly. |
| Mantovani, 2006 [18]| Open label              | 38.9 ± 11.9 (18–70) | 10 | 8:2 | OCD only (n = 5), OCD + TS (n = 2) | Low frequency rTMS | None             | SMA                 | Tics and OCD symptoms significantly reduced up to 3 months. |
| Munchau, 2002 [77]| Single-blinded, placebo control | 38.0 ± 13.2 (unknown) | 16 | 3:1 | OCD (n = 7)                  | rTMS         | Sham/Placebo     | L Premotor and L Motor Cortex | No significant improvement. |
| Orth, 2005 [17]    | Single-blinded, placebo control | 29.0 ± SD (19–52) | 5  | 4:1 | ADHD (n = 2)                  | rTMS         | Sham             | Left (and Bilateral) Premotor Cortex | rTMS no significant effect on global tic severity. |
| Landeros-Weisenberger, A., 2015 [83] | Randomized double-blind sham-control | 33.7 ± 12.2 (unknown) | 20 | 4:1 | OCD (n = 5), ADHD (n = 8) | rTMS         | Sham             | SMA                 | No significant reduction of tic severity. |

rTMS = repetitive transcranial magnetic stimulation; SMA = supplementary motor area; MC = Motor Cortex; PFC = Prefrontal Cortex; OCD; ADHD; PFC; SD.
7. Predictors of Transcranial Magnetic Stimulation Efficacy in Tourette Syndrome

An examination of predictors of efficacy of TMS for TS has resulted in preliminary considerations. In line with the mechanism of TMS therapeutics, the degree of brain electrical activity in individuals with TS might constitute a predictor variable. For example, ADHD has been associated with abnormal electrical activity (excess theta activity) [84], so that in those individuals with comorbid ADHD and TS, brain electrical abnormalities might be cumulative. In turn, TMS may be more efficacious in those subgroups of TS with greater electrical abnormalities (i.e., TS + ADHD). A study of deep TMS as an add-on treatment for intractable TS in adults showed no significant effect on tic severity overall in TS alone, but a significant decrease in comorbid TS [79]. Similarly, as described above, Chae et al. found that rTMS was more effective in comorbid TS compared to TS alone [78]. In both studies, TS was comorbid with OCD and ADHD (TS+OCD+ADHD), a complex phenotype that has shown to “breed true” as a TS subtype [6]. Overall, the few extant trials of TMS in TS suggest that the most improvement with TMS is shown in TS patients who are: younger, have comorbid ADHD/OCD and use SMA as a target (Table 1).

8. Future Directions

As discussed above, the comorbidity of TS with ADHD, and possibly OCD, plausibly confers additional abnormal cortical excitability compared with TS alone [67], which might account for the greater response to TMS treatment in “complex TS”, compared to those instances of TS without ADHD [78,79]. The hyperconnectivity between insula (an interoceptive center) and SMA (a motor planning center) observed in functional neuroimaging studies of TS further support the use of the SMA as a preferred target for TMS trials in TS. Future research in TMS trials in TS might additionally benefit from translational approaches measuring brain connectivity pre- and post-neuroimaging as well as neuroplasticity probes. Ultimately, the use of TMS in TS holds promise for an improved understanding of the pathophysiology and enhanced therapeutics in this debilitating condition.

Author Contributions: For research articles with several authors, a short paragraph specifying their individual contributions must be provided. The following statements should be used “Conceptualization, M.G.; Methodology, L.D.-S. and R.H.; Writing-Original Draft Preparation, L.D.-S., R.H. and M.G.; Writing-Review & Editing, L.D.-S., R.H. and M.G.

Funding: This research received no external funding.

Conflicts of Interest: The authors declare no conflict of interest.

References
1. De la Tourette, G.G. Étude sur une affection nerveuse caractérisée par de l’incoordination motrice accompagnée d’écholalie et de coprolalie (jumping, latah, myriachiti). Archives de Neurologie 1885, 9, 158–200.
2. Pauls, D.L.; Cohen, D.J.; Heimbuch, R.; Detlor, J.; Kidd, K.K. Familial pattern and transmission of gilles de la tourette syndrome and multiple tics. Arch. Gen. Psychiatry 1981, 38, 1091–1093. [CrossRef] [PubMed]
3. Groth, C.; Mol Debes, N.; Rask, C.U.; Lange, T.; Skov, L. Course of tourette syndrome and comorbidities in a large prospective clinical study. J. Am. Acad. Child Adolesc. Psychiatry 2017, 56, 304–312. [CrossRef] [PubMed]
4. Robertson, M.M. The gilles de la tourette syndrome: The current status. Br. J. Psychiatry J. Ment. Sci. 1989, 154, 147–169. [CrossRef]
5. Comings, D.E.; Comings, B.G. Ts, learning, and speech problems. J. Am. Acad. Child Adolesc. Psychiatry 1994, 33, 429–430. [CrossRef] [PubMed]
6. Grados, M.A.; Mathews, C.A. Latent class analysis of gilles de la tourette syndrome using comorbidities: Clinical and genetic implications. Biol. Psychiatry 2008, 64, 219–225. [CrossRef] [PubMed]
7. Mathews, C.A.; Grados, M.A. Familiality of tourette syndrome, obsessive-compulsive disorder, and attention-deficit/hyperactivity disorder: Heritability analysis in a large sib-pair sample. J. Am. Acad. Child Adolesc. Psychiatry 2011, 50, 46–54. [CrossRef] [PubMed]
8. Comings, D.E. A controlled study of tourette syndrome. Vii. Summary: A common genetic disorder causing disinhibition of the limbic system. Am. J. Hum. Genet. 1987, 41, 839–866. [PubMed]
9. Nag, A.; Bochukova, E.G.; Kremeyer, B.; Campbell, D.D.; Muller, H.; Valencia-Duarte, A.V.; Cardona, J.; Rivas, I.C.; Mesa, S.C.; Cuartas, M.; et al. Cnv analysis in tourette syndrome implicates large genomic rearrangements in col8a1 and nrnx1. PLoS ONE 2013, 8, e59061. [CrossRef] [PubMed]

10. Wong, D.F.; Brasic, J.R.; Singer, H.S.; Schretlen, D.J.; Kuwabara, H.; Zhou, Y.; Nandi, A.; Maris, M.A.; Alexander, M.; Ye, W.; et al. Mechanisms of dopaminergic and serotonergic neurotransmission in tourette syndrome: Clues from an in vivo neurochemistry study with pet. Neuropsychopharmacology 2008, 33, 1239–1251. [CrossRef] [PubMed]

11. Church, J.A.; Fair, D.A.; Dosenbach, N.U.; Cohen, A.L.; Miezin, F.M.; Petersen, S.E.; Schlaggar, B.L. Control networks in paediatric tourette syndrome show immature and anomalous patterns of functional connectivity. Brain 2009, 132, 225–238. [CrossRef] [PubMed]

12. Scahill, L.D.; Leckman, J.F.; Marek, K.L. Sensory phenomena in tourette’s syndrome. Adv. Neurol. 1995, 65, 273–280. [PubMed]

13. Craig, A.D. Interception: The sense of the physiological condition of the body. Curr. Opin. Neurobiol. 2003, 13, 500–505. [CrossRef]

14. Neuner, I.; Werner, C.J.; Arrubla, J.; Stocker, T.; Ehlen, C.; Wegener, H.P.; Schneider, F.; Shah, N.J. Imaging the where and when of tic generation and resting state networks in adult tourette patients. Front. Hum. Neurosci. 2014, 8, 362. [CrossRef] [PubMed]

15. Tinaz, S.; Malone, P.; Hallett, M.; Horovitz, S.G. Role of the right dorsal anterior insula in the urge to tic in tourette syndrome. Mov. Disord. 2015, 30, 1190–1197. [CrossRef] [PubMed]

16. Strigo, I.A.; Craig, A.D. Interception, homeostatic emotions and sympathovagal balance. Philos. Trans. R. Soc. Lond. Ser. B Biol. Sci. 2016, 371. [CrossRef] [PubMed]

17. Orth, M.; Kirby, R.; Richardson, M.P.; Snijders, A.H.; Rothwell, J.C.; Trimble, M.R.; Robertson, M.M.; Munchau, A. Subthreshold rtms over pre-motor cortex has no effect on tics in patients with gilles de la tourette syndrome. Clin. Neurophysiol. 2005, 116, 764–768. [CrossRef] [PubMed]

18. Mantovani, A.; Lisanby, S.H.; Pieraccini, F.; Ulivelli, M.; Castrogiovanni, P.; Rossi, S. Repetitive transcranial magnetic stimulation (rtms) in the treatment of obsessive-compulsive disorder (ocd) and tourette’s syndrome (ts). Int. J. Neuropsychopharmacol. 2006, 9, 95–100. [CrossRef] [PubMed]

19. Biot, J.-B.; Savart, F. Note sur le magnetisme de la pile de volta. Ann. Chim. Phys. 1820, 15, 222–223.

20. Faraday, M. Experimental researches in electricity. Philos. Trans. R. Soc. Lond. 1832, 122, 125–162. [CrossRef]

21. George, M.S.; Wassermann, E.M.; Post, R.M. Transcranial magnetic stimulation: A neuropsychiatric tool for the 21st century. J. Neuropsychiatry Clin. Neurosci. 1996, 8, 373–382. [PubMed]

22. Pascual-Leone, A.; Gates, J.R.; Dhuna, A. Induction of speech arrest and counting errors with rapid-rate transcranial magnetic stimulation. Neurology 1999, 41, 697–702. [CrossRef] [PubMed]

23. Pascual-Leone, A.; Grafman, J.; Hallett, M. Modulation of cortical motor output maps during development of implicit and explicit knowledge. Science 1994, 263, 1287–1289. [CrossRef] [PubMed]

24. Viachos, A.; Muller-Dahlhaus, F.; Rosskopf, J.; Lenz, M.; Ziemann, U.; Deller, T. Repetitive magnetic stimulation induces functional and structural plasticity of excitatory postsynapses in mouse organotypic hippocampal slice cultures. J. Neurosci. 2012, 32, 17514–17523. [CrossRef] [PubMed]

25. Lenz, M.; Viachos, A. Releasing the cortical brake by non-invasive electromagnetic stimulation? Rtms induces ltd of gabaergic neurotransmission. Front. Neural Circuits 2016, 10, 96. [CrossRef] [PubMed]

26. Barker, A.T.; Jalinous, R.; Freeston, I.L. Non-invasive magnetic stimulation of human motor cortex. Lancet 1985, 1, 1106–1107. [CrossRef]

27. Bunse, T.; Wobrock, T.; Strube, W.; Padberg, F.; Palm, U.; Falkai, P.; Hasan, A. Motor cortical excitability assessed by transcranial magnetic stimulation in psychiatric disorders: A systematic review. Brain Stimul. 2014, 7, 158–169. [CrossRef] [PubMed]

28. Peters, S.K.; Dunlop, K.; Downar, J. Cortico-striatal-thalamic loop circuits of the salience network: A central pathway in psychiatric disease and treatment. Front. Syst. Neurosci. 2016, 10, 104. [CrossRef] [PubMed]

29. Haber, S.N. The primate basal ganglia: Parallel and integrative networks. J. Chem. Neuroanat. 2003, 26, 317–330. [CrossRef] [PubMed]

30. Smith, Y.; Bevan, M.D.; Shink, E.; Bolam, J.P. Microcircuitry of the direct and indirect pathways of the basal ganglia. Neuroscience 1998, 86, 353–387. [PubMed]

31. Mink, J.W. The basal ganglia: Focused selection and inhibition of competing motor programs. Prog. Neurobiol. 1996, 50, 381–425. [CrossRef]
32. Cheon, K.A.; Ryu, Y.H.; Namkoong, K.; Kim, C.H.; Kim, J.J.; Lee, J.D. Dopamine transporter density of the basal ganglia assessed with [123i]ipt spect in drug-naive children with tourette’s disorder. Psychiatry Res. 2004, 130, 85–95. [CrossRef] [PubMed]

33. Serra-Mestres, J.; Ring, H.A.; Costa, D.C.; Gacinovic, S.; Walker, Z.; Lees, A.J.; Robertson, M.M.; Trimble, M.R. Dopamine transporter binding in gilles de la tourette syndrome: A [123i]fp-cit/spect study. Acta Psychiatr. Scand. 2004, 109, 140–146. [CrossRef] [PubMed]

34. Ernst, M.; Zametkin, A.J.; Jons, P.H.; Matochik, J.A.; Pascualvaca, D.; Cohen, R.M. High presynaptic dopaminergic activity in children with tourette’s disorder. J. Am. Acad. Child Adolesc. Psychiatry 1999, 38, 86–94. [CrossRef] [PubMed]

35. Ernst, M.; Zametkin, A.J.; Jons, P.H.; Matochik, J.A.; Pascualvaca, D.; Cohen, R.M. High presynaptic dopaminergic activity in children with tourette’s disorder. J. Am. Acad. Child Adolesc. Psychiatry 1999, 38, 86–94. [CrossRef] [PubMed]

36. Albin, R.L.; Koeppe, R.A.; Bohnen, N.I.; Nichols, T.E.; Meyer, P.; Wernette, K.; Minoshima, S.; Kilbourn, M.R.; Frey, K.A. Increased ventral striatal monoaminergic innervation in tourette syndrome. Neurology 2003, 61, 310–315. [CrossRef] [PubMed]

37. Singer, H.S.; Morris, C.; Grados, M. Glutamatergic modulatory therapy for tourette syndrome. Med. Hypotheses 2010, 74, 862–867. [CrossRef] [PubMed]

38. Floresco, S.B.; Todd, C.L.; Grace, A.A. Glutamatergic afferents from the hippocampus to the nucleus accumbens regulate activity of ventral tegmental area dopamine neurons. J. Neurosci. 2001, 21, 4915–4922. [CrossRef] [PubMed]

39. Nordstrom, E.J.; Bittner, K.C.; McGrath, M.J., 3rd; Burton, F.H. “Hyperglutamatergic cortico-striato-thalamo-cortical circuit” breaker drugs alleviate tics in a transgenic circuit model of tourettes syndrome. Brain Res. 2003, 1629, 38–53. [CrossRef] [PubMed]

40. Anderson, G.M.; Pollak, E.; Sattler, R.; Vidensky, S.; Rothstein, J.D.; Singer, H.; Wang, T. Genetic and functional studies of a missense variant in a glutamate transporter, slc1a3, in tourette syndrome. Psychitr. Genet. 2011, 21, 90–97. [CrossRef] [PubMed]

41. Lemmon, M.E.; Grados, M.; Kline, T.; Thompson, C.B.; Ali, S.F.; Singer, H.S. Efficacy of glutamate modulators in tic suppression: A double-blind, randomized control trial of d-serine and riluzole in tourette syndrome. Pediatr. Neurol. 2015, 52, 629–634. [CrossRef] [PubMed]

42. Johnson, J.; Tian, N.; Caywood, M.S.; Reimer, R.J.; Edwards, R.H.; Copenhagen, D.R. Vesicular neurotransmitter transporter expression in developing postnatal rodent retina: Gaba and glycine precede glutamate. J. Neurosci. 2015, 23, 1519–1522. [CrossRef] [PubMed]

43. Han, B.; Bellemer, A.; Koelle, M.R. An evolutionarily conserved switch in response to gaba affects development and behavior of the locomotor circuit of caenorhabditis elegans. Genetics 2015, 199, 1159–1172. [CrossRef] [PubMed]

44. Selten, M.; van Bokhoven, H.; Nadif Kasri, N. Inhibitory control of the excitatory/inhibitory balance in psychiatric disorders. F1000Reserch 2018, 7, 23. [CrossRef] [PubMed]

45. Furukawa, M.; Tsukahara, T.; Tomita, K.; Iwai, H.; Sonomura, T.; Miyawaki, S.; Sato, T. Neonatal maternal separation delays the gaba excitatory-to-inhibitory functional switch by inhibiting kcc2 expression. Biochem. Biophys. Res. Commun. 2017, 493, 1243–1249. [CrossRef] [PubMed]
51. Tomiyasu, M.; Aida, N.; Shibasaki, J.; Umeda, M.; Murata, K.; Heberlein, K.; Brown, M.A.; Shimizu, E.; Tsuji, H.; Obata, T. In vivo estimation of gamma-aminobutyric acid levels in the neonatal brain. *NMR Biomed.* **2017**, *3*, e3666. [CrossRef] [PubMed]

52. Kalanithi, P.S.; Zheng, W.; Kataoka, Y.; DiFiglia, M.; Grantz, H.; Saper, C.B.; Schwartz, M.L.; Leckman, J.F.; Vaccarino, F.M. Altered parvalbumin-positive neuron distribution in basal ganglia of individuals with tourette syndrome. *Proc. Natl. Acad. Sci. USA* **2005**, *102*, 13307–13312. [CrossRef] [PubMed]

53. Bode, C.; Richter, F.; Sprote, C.; Brigadski, T.; Bauer, A.; Fietz, S.; Fritschy, J.M.; Richter, A. Altered postnatal maturation of striatal gabaergic interneurons in a phenotypic animal model of dystonia. *Exp. Neurol.* **2017**, *287*, 44–53. [CrossRef] [PubMed]

54. Mahone, E.M.; Puts, N.A.; Edden, R.A.E.; Ryan, M.; Singer, H.S. Gaba and glutamate in children with tourette syndrome: A (1)h mr spectroscopy study at 7t. *Psychiatry Res.* **2018**, *273*, 46–53. [CrossRef] [PubMed]

55. Puts, N.A.; Harris, A.D.; Crocetti, D.; Nettles, C.; Singer, H.S.; Tommerdahl, M.; Edden, R.A.; Mostofsky, S.H. Reduced gabaergic inhibition and abnormal sensory symptoms in children with tourette syndrome. *J. Neurophysiol.* **2015**, *114*, 808–817. [CrossRef] [PubMed]

56. Draper, A.; Stephenson, M.C.; Jackson, G.M.; Pepes, S.E.; Jackson, S.R. Inhibition, disinhibition, and the control of motor excitability in tourette syndrome. *Curr. Biol.* **2014**, *24*, 2343–2347. [CrossRef] [PubMed]

57. Koos, T.; Tepper, J.M. Inhibitory control of neostriatal projection neurons by gabaergic interneurons. *Nat. Neurosci.* **1999**, *2*, 467–472. [CrossRef] [PubMed]

58. Jackson, G.M.; Draper, A.; Dyke, K.; Pepes, S.E.; Jackson, S.R. Inhibition, disinhibition, and the control of action in tourette syndrome. *Trends Cognit. Sci.* **2015**, *19*, 655–665. [CrossRef] [PubMed]

59. Rothwell, J.C.; Day, B.L.; Thompson, P.D.; Kujirai, T. Short latency intracortical inhibition: One of the most popular tools in human motor neurophysiology. *J. Physiol.* **2009**, *587*, 11–12. [CrossRef] [PubMed]

60. Gilbert, D.L.; Bansal, A.S.; Sethuraman, G.; Sallee, F.R.; Zhang, J.; Lipps, T.; Wassermann, E.M. Association of cortical disinhibition with tic, adhd, and ocd severity in tourette syndrome. *Mov. Disord.* **2004**, *19*, 416–425. [CrossRef] [PubMed]

61. Gilbert, D.L.; Salle, F.R.; Zhang, J.; Lipps, T.D.; Wassermann, E.M. Transcranial magnetic stimulation-evoked cortical inhibition: A consistent marker of attention-deficit/hyperactivity disorder scores in tourette syndrome. *Biol. Psychiatry* **2005**, *57*, 1597–1600. [CrossRef] [PubMed]

62. Ziemann, U.; Paulus, W.; Rothenberger, A. Decreased motor inhibition in tourette’s disorder: Evidence from transcranial magnetic stimulation. *Am. J. Psychiatry* **1997**, *154*, 1277–1284. [PubMed]

63. Draper, A.; Jude, L.; Jackson, G.M.; Jackson, S.R. Motor excitability during movement preparation in tourette syndrome. *J. Neuropsychol.* **2015**, *9*, 33–44. [CrossRef] [PubMed]

64. Brandt, V.C.; Niessen, E.; Ganos, C.; Kahl, U.; Baumer, T.; Munchau, A. Altered synaptic plasticity in tourette’s syndrome and its relationship to motor skill learning. *PLoS ONE* **2014**, *9*, e98417. [CrossRef] [PubMed]

65. Martín-Rodriguez, J.F.; Ruiz-Rodriguez, M.A.; Palomar, F.J.; Cáceres-Redondo, M.T.; Vargas, L.; Forcachia, P.; Gómez-Crespo, M.; Huertas-Fernández, I.; Carrillo, F.; Madruga-Garrido, M.; et al. Aberrant cortical associative plasticity associated with severe adult tourette syndrome. *Mov. Disord.* **2015**, *30*, 431–435. [CrossRef] [PubMed]

66. Suppa, A.; Marsili, L.; Di Stasio, F.; Berardelli, I.; Roselli, V.; Pasquini, M.; Cardona, F.; Berardelli, A. Cortical and brainstem plasticity in tourette syndrome and obsessive-compulsive disorder. *Mov. Disord.* **2014**, *29*, 1523–1531. [CrossRef] [PubMed]

67. Orth, M.; Rothwell, J.C. Motor cortex excitability and comorbidity in gilles de la tourette syndrome. *J. Neurol. Neurosurg. Psychiatry* **2009**, *80*, 29–34. [CrossRef] [PubMed]

68. Orth, M.; Munchau, A.; Rothwell, J.C. Corticospinal system excitability at rest is associated with tic severity in tourette syndrome. *Biol. Psychiatry* **2008**, *64*, 248–251. [CrossRef] [PubMed]

69. Heise, C.A.; Wanschura, V.; Albrecht, B.; Uebel, H.; Roessner, V.; Himpel, S.; Paulus, W.; Rothenberger, A.; Tergau, F. Voluntary motor drive: Possible reduction in tourette syndrome. *J. Neural Transm.* **2008**, *115*, 857–861. [CrossRef] [PubMed]

70. Pépés, S.E.; Draper, A.; Jackson, G.M.; Jackson, S.R. Effects of age on motor excitability measures from children and adolescents with tourette syndrome. *Dev. Cognit. Neurosci.* **2016**, *19*, 78–86. [CrossRef] [PubMed]
71. Jackson, S.R.; Parkinson, A.; Manfredi, V.; Millon, G.; Hollis, C.; Jackson, G.M. Motor excitability is reduced prior to voluntary movements in children and adolescents with tourette syndrome. *J. Neuropsychol.* 2013, 7, 29–44. [CrossRef] [PubMed]

72. Rapanelli, M.; Frick, L.R.; Pittenger, C. The role of interneurons in autism and tourette syndrome. *Trends Neurosci.* 2017, 40, 397–407. [CrossRef] [PubMed]

73. Huang, A.Y.; Yu, D.; Davis, L.K.; Sul, J.H.; Tsetsos, F.; Ramensky, V.; Zelaya, I.; Ramos, E.M.; Osiecki, L.; Chen, J.A.; et al. Rare copy number variants in nrnx1 and cntn6 increase risk for tourette syndrome. *Neuron* 2017, 94, 1101–1111.e7. [CrossRef] [PubMed]

74. Dayan, E.; Censor, N.; Buch, E.R.; Sandrini, M.; Cohen, L.G. Noninvasive brain stimulation: From physiology to network dynamics and back. *Nat. Neurosci.* 2013, 16, 838–844. [CrossRef] [PubMed]

75. Hameed, M.Q.; Dhamne, S.C.; Gersner, R.; Kaye, H.L.; Oberman, L.M.; Pascual-Leone, A.; Rotenberg, A. Transcranial magnetic and direct current stimulation in children. *Curr. Neurol. Neurosci. Rep.* 2017, 17, 11. [CrossRef] [PubMed]

76. Mrakic-Sposta, S.; Marceglia, S.; Mameli, F.; Dilena, R.; Tadini, L.; Priori, A. Transcranial direct current stimulation in two patients with tourette syndrome. *Mov. Disord.* 2008, 23, 2259–2261. [CrossRef] [PubMed]

77. Munchau, A.; Bloem, B.R.; Thilo, K.V.; Trimble, M.R.; Rothwell, J.C.; Robertson, M.M. Repetitive transcranial magnetic stimulation for tourette syndrome. *Neurology* 2002, 59, 1789–1791. [CrossRef] [PubMed]

78. Chae, J.H.; Nahas, Z.; Wassermann, E.; Li, X.; Sethuraman, G.; Gilbert, D.; Sallee, F.R.; George, M.S. A pilot safety study of repetitive transcranial magnetic stimulation (rtms) in tourette’s syndrome. *Cognit. Behav. Neurol.* 2004, 17, 109–117. [CrossRef]

79. Bloch, Y.; Arad, S.; Levkovitz, Y. Deep tms add-on treatment for intractable tourette syndrome: A feasibility study. *World J. Biol. Psychiatry* 2016, 17, 557–561. [PubMed]

80. Kwon, H.J.; Lim, W.S.; Lim, M.H.; Lee, S.J.; Hyun, J.K.; Chae, J.H.; Paik, K.C. 1-hz low frequency repetitive transcranial magnetic stimulation in children with tourette’s syndrome. *Neurosci. Lett.* 2011, 492, 1–4. [CrossRef] [PubMed]

81. Le, K.; Liu, L.; Sun, M.; Hu, L.; Xiao, N. Transcranial magnetic stimulation at 1 hertz improves clinical symptoms in children with tourette syndrome for at least 6 months. *J. Clin. Neurosci.* 2013, 20, 257–262. [CrossRef] [PubMed]

82. Krishnan, C.; Santos, L.; Peterson, M.D.; Ehinger, M. Safety of noninvasive brain stimulation in children and adolescents. *Brain Stimul.* 2015, 8, 76–87. [CrossRef] [PubMed]

83. Landeros-Weisenberger, A.; Mantovani, A.; Motlagh, M.G.; de Alvarenga, P.G.; Katsovich, L.; Leckman, J.F.; Lisanby, S.H. Randomized sham controlled double-blind trial of repetitive transcranial magnetic stimulation for adults with severe Tourette syndrome. *Brain Stimul.* 2015, 8, 574–581. [CrossRef] [PubMed]

84. Loo, S.K.; McGough, J.J.; McCracken, J.T.; Smalley, S.L. Parsing heterogeneity in attention-deficit hyperactivity disorder using eeg-based subgroups. *J. Child Psychol. Psychiatry Allied Discipl.* 2018, 59, 223–231. [CrossRef] [PubMed]

© 2018 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).