The importance of different learning stages for motor sequence learning after stroke

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Abstract
The task of learning predefined sequences of interrelated motor actions is of everyday importance and has also strong clinical importance for regaining motor function after brain lesions. A solid understanding of sequence learning in stroke patients can help clinicians to optimize and individualize rehabilitation strategies. Moreover, to investigate the impact of a focal lesion on the ability to successfully perform motor sequence learning can enhance our comprehension of the underlying physiological principles of motor sequence learning. In this article, we will first provide an overview of current concepts related to motor sequence learning in healthy subjects with focus on the involved brain areas and their assumed functions according to the temporal stage model. Subsequently, we will consider the question of what we can learn from studies investigating motor sequence learning in stroke patients. We will first focus on the implications of lesion location. Then, we will analyze whether distinct lesion locations affect specific learning stages. Finally, we will discuss the implications for clinical rehabilitation and suggest directions for further research.

KEYWORDS
motor cortex, motor rehabilitation, plasticity, sequence learning, stroke

1 | INTRODUCTION

Most aspects of human behavior are composed of series of interrelated actions (sequences). This applies not only to motor actions but also to sequences of words, thoughts and memories. Motor skill acquisition is a particularly intuitive example of the ability of the human brain to learn sequences by training. Typical examples of motor sequence learning include learning to play piano and learning to dance. Even motor sequences that appear very difficult at first may be carried out effortlessly given enough time and training. Throughout training, movement parameters are optimized, resulting in efficient and accurate performance. Training also leads to a decreasing need for active attention to movement performance. The changes in resource utilization during the course of training have led to the proposition of different, distinguishable stages of sequence learning. In particular, multiple authors have suggested a division into an early learning phase, a consolidation phase and a slow learning or retention phase. In addition to this temporal model, there is also broad consensus that motor sequence learning takes place in two different modes, mainly distinguished by the presence or absence of conscious awareness of the learned sequence (explicit vs. implicit learning).

Due to the simple accessibility of motor performance and the general importance of the motor system for our quality of life, the motor system is often investigated as a representative of principles underlying sequence learning in general. For this purpose, some standardized tests have been established and will be introduced below.

It is important to note that most complex motor skills consist of sequences of movements (Diedrichsen & Kornysheva, 2015; Hikosaka, Nakamura, Sakai, & Nakahara, 2002; Willingham, 1998). To regain a lost motor skill (e.g., lost through a stroke) not only every
single movement of this skill, but also its implementation as a sequence has to be relearned. The inseparable connection between learning motor skills and sequences makes the understanding of sequence learning important for clinical rehabilitation. Disabilities resulting from stroke are particularly significant as they constitute the leading cause of long-term disability in industrial nations. Hence, there is an urgent need for a solid understanding of sequence learning in stroke patients, which could help clinicians optimize and individualize rehabilitation strategies. Moreover, the impact of a focal lesion on the ability to successfully perform motor sequence learning can help us gain insights into—and an improved comprehension of—the basic physiological principles of motor sequence learning.

In the following sections, we will first provide an overview of important concepts related to motor sequence learning, such as explicit and implicit motor learning, and the dissociation between sequential and motor components. We will then briefly introduce the current methods used to investigate sequential motor learning. Next, we will review the current state of knowledge regarding important brain areas and their assumed functions in motor sequence learning as derived from studies in healthy subjects. Subsequently, we will consider the question of what we can learn from studies investigating motor sequence learning in stroke patients. We will first investigate the evidence of whether patients’ capacity for motor sequence learning is preserved after stroke in general. Next, we will review all studies concerning the implications of stroke for different learning stages according to the temporal stage model. Finally, we will discuss the implications for clinical rehabilitation and suggest directions for further research.

2 | BASIC PRINCIPLES OF MOTOR SEQUENCE LEARNING

Motor sequence learning refers to the process in which a pre-determined ordered list (sequence) of motor actions is performed with increasing spatial and temporal accuracy. This learning process can occur with or without conscious awareness of the learning process itself and the sequential order of elements. The divergences between conscious and unconscious learning have led to the well-accepted concept of distinguishing two different modes—the explicit and implicit modes of learning.

2.1 | Explicit and implicit motor learning

Explicit and implicit motor learning are fundamental concepts for our understanding of sequence learning. The differentiation is mainly based on awareness of the learned skill. While explicit learning requires active attention and some form of rules that can be stated, implicit motor learning takes place without awareness and in the absence of verbal knowledge of the performed motor task (Kleynen, Braun, et al., 2014; Kleynen, Wilson, et al., 2014). Hence, implicit or procedural learning is defined as a learning process that is both incidental and unconscious (Reber, 1993). The classic example of implicit motor learning is to learn to walk as a child or to ride a bicycle. These learning tasks can be accomplished without instructions. Learned skills are retrieved from implicit memory. However, some authors object that implicit learning also comes along with conscious inputs, which then lead to unconscious implications that are learned (Baars, 2010). This conscious input is called explicit or declarative knowledge. Explicit motor learning was defined in a previous consensus paper as follows: “learning which generates verbal knowledge of movement performance (e.g., facts and rules), involves cognitive stages within the learning process and is dependent on working memory involvement” (Kleynen, Braun, et al., 2014; Kleynen, Wilson, et al., 2014). A typical example of explicit learning would be to learn a prescribed sequence of dance steps.

2.2 | Dissociation between sequential and motor components

Another important concept in understanding motor sequence learning is the dissociation between the spatial-sequential order of a movement sequence and its motor control components (Doya, 2000; Ghilardi, Moisello, Silvestri, Ghez, & Krakauer, 2009; Hikosaka et al., 1999; Penhune & Steele, 2012). The spatial-sequential task component concerns the order of movements in space and time, while the movement dynamics comprise the sensorimotor integration of movements, such as the speed of single movements and the timing of movements.

It has been proposed that these components are learned in parallel, with different contributions by specific brain areas. Accordingly, learning the sequential/spatial order of movement follows a different time course than learning the motor components. It is thought that in the early phase of motor learning, improvements are dominated by learning the sequential/spatial characteristics of the movements, while the implementation and optimization of the required motor components dominate improvements in later stages of learning (Ghilardi et al., 2009; Savion-Lemieux & Penhune, 2005).

2.3 | Methods to investigate motor sequence learning

The most commonly used method to examine motor sequence learning is the serial reaction time task (SRTT). Participants receive visual stimuli (cues) appearing at different locations on a screen. The participants are asked to respond to each cue by pressing the correct button as fast as possible. The time between cue presentation and motor response defines the reaction time. A decrease in reaction time is considered learning. However, this decrease in reaction time can be influenced by multiple contaminating factors, such as motivation and visuomotor associations. Therefore, the reaction time for pre-determined sequences is contrasted with the reaction time for random sequences. The difference between sequential and random sequences in the decrease in reaction time is often used as a measure of sequence-specific motor learning. The SRTT paradigm investigates implicit sequence motor learning if the participants do not become aware of the predefined sequence. If, however, participants gain knowledge about this sequence, this indicates an explicit component of motor sequence learning. The SRTT is used in several variants.
Other paradigms for assessment of motor sequence learning are the continuous tracking task (CTT, tracking a target moving in a specific pattern across a screen), finger tapping tasks (e.g., the manual sequence task, or MST, which consists of typing a predefined numeral sequence on a finger pad; Karni et al., 1995), and sequences of reaching (Ghilardi et al., 2009) or eye movements (Albouy et al., 2008).

3 | CURRENT KNOWLEDGE REGARDING THE CEREBRAL NETWORK ENGAGED IN MOTOR SEQUENCE LEARNING

Motor sequence learning is performed by a complex and distributed network of brain regions. Essential parts of this network include the primary motor cortex (M1), the premotor cortex (PMC), the supplementary motor area (SMA), the basal ganglia (BG), the prefrontal cortex (PFC), the posterior parietal cortex (PPC), and the cerebellum (Figure 1).

In the following section, we will briefly describe the primary functions of these brain areas for motor sequence learning and will then discuss their roles in different phases of the learning process.

The motor cortex, comprising M1, PMC, and SMA, is strongly interconnected with frontoparietal regions as well as the striatum and other parts of the BG and generates significant output to the descending motor system. While M1 stores motor sequence information in a somatotopically organized system, PMC is recognized to play a critical role in visual and sensorimotor integration and movement selection, preparing motor output for M1 (Hardwick, Rottschy, Miall, & Eickhoff, 2013). A major function of the SMA seems to be the planning and self-initiation of voluntary movements. Input arrives from pre-SMA, which serves the function of integrating multimodal information by reducing the output of the motor loop when its signals are inconsistent with those of the visual loop (Hikosaka et al., 1999; Nakahara, Doya, & Hikosaka, 2001).

The cerebellum is known to play a crucial role in the fine-tuning of motor performance via feedback loops with cortical structures which are called cortico-cerebellar loops. Cerebellar functioning is firmly connected to the parietal, premotor and frontal cortex and accomplishes a central role in movement optimization. By integrating efferent and afferent information, the cerebellum creates an internal predictive model of sensory states that guides learning by the trial-and-error principle (Miall, 2010; Penhune & Steele, 2012; Shadmehr, Brandt, & Corkin, 1998). This allows predictive control by a feedforward mechanism in which the current sensory state and cortical motor commands are combined to forecast the future state of the body (Bastian, 2006).

The BG are mainly engaged in the chunking of movements and contribute to the storage of information. It is generally agreed that the BG play a prominent role in implicit learning processes. However, some parts of the BG are also involved in explicit learning. The striatum is considered the most critical BG structure engaged in motor sequence learning. The dorsolateral striatum primarily takes part in sensorimotor circuits by communicating with the parietal and sensorimotor cortices, which supports the idea that it plays a leading role in implicit sequence learning by chunking information (Hikosaka et al., 1999; Penhune & Steele, 2012). Chunking is a process in which a motor sequence is subdivided into shorter segments, known as chunks. By being segmented into these chunks, the information can be retained and restored more easily than unsegmented information. The anterior medial striatum is known to be strongly connected to frontal and premotor areas, which are suggested to play a critical role in explicit learning processes. This connection is also called the “associative” circuit (Janacsek & Nemeth, 2013; Penhune & Steele, 2012). Furthermore, the BG are believed to be involved in reward-based learning, integrating the motivational component of learning (Hikosaka et al., 1999). The complex bilateral interactions between cortical areas and the BG constitute a circuit known as the corticostratial loop, which is believed to work in parallel with the corticocerebellar loop.

The PFC and the PPC are mainly engaged in explicit learning processes that are attentionally demanding and require working memory (Eliassen, Souza, & Sanes, 2001; Lewis & Miall, 2003).

Additionally, the parietal cortex is considered to play a crucial role in the integration of somatosensory and visual information and in
preparing multimodal sensorimotor input for the PMC (Hardwick et al., 2013; Lewis & Miall, 2003). Furthermore, the parietal cortex is known to be engaged in the delayed recall of learned motor sequences (Doyon, Penhune, & Ungerleider, 2003).

Although this section is intended to be a brief general overview of the functions of different brain areas, the engagement of these areas changes with ongoing practice. This change over time has a strong implication for our understanding of the mechanism underlying motor sequence learning and, furthermore, has implications for practical rehabilitation. Therefore, we will review the current knowledge about learning stages in motor sequence learning in the next section and will then endeavor to discuss the contribution of lesion studies to this topic.

4 TEMPORAL STAGE MODEL OF MOTOR SEQUENCE LEARNING

Current theoretical models suggest that the process of motor sequence learning can be divided into different temporal stages (Doyon, 2008; Doyon et al., 2009; Hikosaka et al., 1999; Karni et al., 1998; Penhune & Steele, 2012). These stages differ in behavioral parameters, such as the amount of learning and the contribution of different components to the learning process (spatial-sequential or motor component; Doyon et al., 2009; Karni et al., 1995; Karni et al., 1998; Willingham, 1998; Figure 1). Additionally, the contributions of specific brain areas and brain networks to the learning process differ between learning stages (Figure 2).

Several authors propose a differentiation into an early phase, a consolidation phase and a slow learning or retention phase (Censor, Sagi, & Cohen, 2012; Doyon et al., 2003; Karni et al., 1998; Penhune & Steele, 2012), as we will introduce below.

4.1 Stage I—The early learning phase of motor sequence learning

In the early learning phase, a sequence is acquired quickly and flexibly, which allows rapid performance improvement. This performance improvement is mainly caused by improved encoding of the movement in spatial coordinates and therefore affects the spatial-sequential component of the task (Doya, 2000; Hikosaka et al., 2002).

Essential brain regions involved in this early learning process comprise the DLPFC and the PPC. Both areas maintain strong connections to the anterior part of the BG, especially the head of the caudate nucleus (Hikosaka et al., 1999; Hikosaka et al., 2002; Janacsek & Nemeth, 2013; Penhune & Steele, 2012). This corticostratial loop is called the “associative” circuit (Penhune & Steele, 2012) and is particularly engaged in early learning and explicit learning tasks, when the spatial-sequential component is consciously learned and active attention and working memory are needed (Eliassen et al., 2001; Hikosaka et al., 1999; Janacsek & Nemeth, 2013; Jenkins, Brooks, Nixon, Frackowiak, & Passingham, 1994). Accordingly, deficits in learning new sequences were demonstrated in monkeys after excitotoxic lesions of the PFC (Collins, Roberts, Dias, Everitt, & Robbins, 1998) and after reversible blockage of the anterior striatum (Miyachi, Hikosaka, Miyaishita, Kárádi, & Rand, 1997).

The corticostratial loop (or associative circuit) works in parallel with the cortico-cerebellar loop, which is also of critical importance in early learning. There is evidence that cerebellar connections to the parietal, premotor and frontal cortex are important for successful learning in this stage (Clower, Dum, & Strick, 2005; Diedrichsen, Balsters, Flavell, Cussans, & Ramnani, 2009).

As described above, the cortico-cerebellar loop allows predictive control by a feedforward mechanism in which the current sensory state and cortical motor commands are combined to forecast the future state of the body (Bastian, 2006). A correct forecast is essential for performing a context-dependent sequence of movements. The involvement of the cortico-cerebellar loop and the number of necessary corrections scale with subjective task difficulty and the rate and magnitude of errors. Accordingly, the cerebellum is mainly activated in the early stages of learning, when the internal model to forecast the next body state/movement within the sequence is error-prone (Doyon et al., 2002; Lehéricy et al., 2005; Wu, Kansaku, & Hallett, 2004).

The early spatial-sequential part of learning proceeds in an effector-unspecific manner. Sequential movements are transferred into a spatial coordinate frame, and the sequence is learned mainly independently of the body part that executes it (Censor, 2013; Hikosaka et al., 2002). Storage of motor information in the later learning stages, however, seems to be effector-specific (Japikse, Negash, Explicit/motor component

FIGURE 2 Schematic representation of the involvement and the interaction of brain areas in the three stages of motor sequence learning. Abbreviations: M1, primary motor cortex; PMA, premotor area; PPC, posterior parietal cortex; pre-SMA, pre-supplementary motor area; SMA, supplementary motor area)
Howard, & Howard, 2003). Therefore, the effectiveness of the transfer of learning between the two hands is most significant in this early learning phase and decreases with ongoing learning (Japikse et al., 2003).

4.2 | Stage II—The consolidation phase of motor sequence learning

The second phase is characterized by stabilization of the learned motor sequence and is called the "consolidation" phase. Performance improves incrementally by training and gains increased resistance to interference. This process is highly dependent on the ability to chunk sequences into smaller subsequences, which is accomplished by the BG.

The transition from the early learning phase to the consolidation phase is characterized by a slowing down of the learning speed (Doyon & Benali, 2005; Hikosaka et al., 1999).

The pre-SMA is an important node associated with the transition from Stage I to Stage II. The major function of the pre-SMA seems to be mediation between spatial sequence processing and sensorimotor movement integration (Nakahara et al., 2001). Hence, the pre-SMA is active when one must rely on spatial processes and when motor output must be determined and updated by incoming sensory input (Hikosaka et al., 1999; Nakahara et al., 2001), linking conditional rules to actions (Hardwick et al., 2013). This assumption is consistent with anatomical connections projecting from the DLPFC via the pre-SMA to the SMA (Luppino, Matelli, Camarda, & Rizzolatti, 1993; Tanji, 1994).

The consolidation process primarily includes the encoding of motor associations and the forming of motor chunks and is based mainly on a corticostriatal loop circuit comprising parts of the BG (especially the putamen) and the premotor–motor cortex (especially, the SMA; Hikosaka et al., 1999). In particular, the posterior parts of the putamen and caudate seem to play a central role in the chunking of information. Chunking simplifies the storage and recall of learned sequenced tasks, which is necessary due to the limited capacity of working memory (Fonollosa, Neftci, & Rabinovich, 2015). In addition to the practice-related consolidation of motor sequences, the BG seem to have a special role in offline consolidation. This concept is supported by imaging studies demonstrating that activation of the BG (especially the putamen) increases after sleep following a motor sequence learning task (Debas et al., 2010). Likewise, it was shown that offline motor consolidation during sleep was associated with increased integration within the corticostriatal network (Albouy et al., 2008; Debas et al., 2014).

During the transition to Phase II, cerebellar activation decreases, while striatal activation increases (Doyon et al., 2002; Lehéricy et al., 2005; Wu et al., 2004). The shift in activity from the cerebellar–cortical network toward a striatal–cortical network is thought to correspond to an improved cerebellar model to forecast movements and an associated decrease in the necessity for error correction and model updates.

4.3 | Stage III—The retention phase of motor sequence learning

The transition between Stage II and Stage III is very fluent, which makes a clear differentiation difficult. At some point, however, learning enters a late, slow phase, which is called the "automatization" phase. At this point, performance will be optimized with a slow learning rate. The movement execution becomes "automatic" and can be performed with decreasing attention or entirely implicitly (Karni et al., 1998).

In this process, the motor cortex is of particular importance. The M1, the PMC, and the PPC are considered important hubs in the late stage of motor sequence learning. It has been shown that all three areas are engaged in the delayed recall of learned motor sequences (Doyon et al., 2003). While the PMC is associated with the integration of visual and sensorimotor information and the formation of motor routines, the primary motor cortex is thought to regulate and store use-dependent muscle synergies, creating "motor maps" to optimize and precise sequential movements (Censor et al., 2012; Hardwick et al., 2013; Penhune & Steele, 2012). Hence, the motor cortex is activated when a sequence is learned and a movement becomes "automatic." After the motor map is optimized, the information becomes robust and steady.

Furthermore, it was shown in monkeys that neurons of the SMA have come to represent a learned sequence after extensive training (Tanji, 1994). The involvement of the SMA in the representation of well-learned motor sequences is also supported by studies in humans comparing well-learned sequences to those that are not well learned (Gordon, Lee, Flament, Ugurbil, & Ebner, 1998; Hund-Georgiadis & von Cramon, 1999).

Beyond the motor cortex, there is some evidence that even in this late learning stage, cerebellar loop structures might contribute by correcting errors. Despite the well-established finding that cerebellar activation is mainly present during the fast learning phase and decreases over time (Doyon et al., 2002; Lehéricy et al., 2005; Wu et al., 2004), some studies also demonstrated that the dorsal parts of the dentate nucleus are especially active in the late stage of learning. Hikosaka and coworkers hypothesized a feedback mechanism for fine-tuning of velocity, force and timing in the late stage of well-learned movements; this mechanism was hypothesized to be implemented by a loop circuit between the anterior cerebellum and the motor cortices (Hikosaka et al., 1999). However, there seems to be no direct storage of information or motor memory in cerebellar structures (Doyon et al., 2003; Penhune & Steele, 2012).

4.4 | Summary

Early stages of sequence learning require a mainly frontoparietal "associative" network (especially, prefrontal and parietal cortex), which is interconnected with the anterior BG on the one hand (corticocerebellar loops) and with the cerebellum on the other hand (corticocerebellar loops). While corticostriatal loops are essential in active attention and working-memory demands, the cerebellar loop is
believed to play an important role due to its function in predictive control and adaptive optimization of movement performance.

Over the course of training, the activation shifts to a sensorimotor circuit linking the posterior part of the BG with various motor cortical areas (especially, the SMA). These connections are thought to be mainly responsible for encoding motor associations and forming motor chunks. Hence, BG function is crucial for the consolidation phase of learning. When a sequence is well learned, storage seems to take place in primary motor cortex areas (M1) and the SMA, which are therefore responsible for delayed recall.

The transitions between the different learning stages are fluent, and to a certain extent, the learning processes can occur in parallel. The proportions of activation during the different stages are also dependent on task demands. Even if the learning modes blend together, it can be observed that early and fast learning requires predominantly explicit processes, while the slow or late learning stage is mainly dependent on implicit learning.

5 | CONTRIBUTIONS OF STROKE STUDIES TO THE PRESENT UNDERSTANDING OF MOTOR SEQUENCE LEARNING

Based on our current knowledge regarding the functioning and interaction of individual brain areas within the cerebral network, one might expect that localized lesions of specific brain regions might interfere with different aspects of motor sequence learning. However, the question of how a lesion might impair motor sequence learning is highly nontrivial because a lesion affects not only the function of the lesioned area but also the function of all directly or indirectly connected areas. It is therefore impossible to predict the behavioral consequences of brain lesions based purely on our understanding of network functioning in healthy subjects. This question can be answered only by investigating motor sequence learning in patients after structural brain damage. Studying the impact of lesions on network functioning is not only crucial for our pathophysiological understanding but also of immense clinical relevance, especially concerning poststroke rehabilitation strategies.

Multiple studies have investigated the effects of structural lesions on motor sequence learning using the affected and unaffected hands (see Table 1). The benefit of investigating the affected hand is the clinical importance. It allows us to investigate the difficulties that patients encounter in learning a motor sequence task in its full complexity, and it also allows us to test the effectiveness of possible interventions. Most of the available studies have investigated sequence-specific learning as defined by the difference in reaction time or error rate between random and repeated sequences. A major problem in interpreting the results of such studies is the differentiation of those deficits that are attributable to pure motor impairment and the inability to optimize motor parameters and those deficits that result from an impaired ability to learn the spatial/sequential components. Therefore, multiple studies have investigated motor sequence learning after stroke not in the affected hand but in the unaffected hand. Although primary motor abilities and learning of simple motor responses are highly lateralized functions, learning of spatial/sequential parameters is considered a more global function with a high level of transfer of sequence knowledge from one hand to the other (Japikse et al., 2003). Therefore, utilizing the unaffected hand allows researchers to investigate impairments of motor sequence learning per se (largely independent of pure motor impairments and impairments in the optimization of motor components).

5.1 | Is motor sequence learning preserved after stroke?—A general overview

As introduced above, several studies have investigated motor sequence learning in the affected or the unaffected hand.

Concerning the affected (contralesional) hand, the results are heterogeneous. Some studies conclude that motor sequence learning is preserved (Meehan et al., 2011; Shin et al., 2005; Wadden et al., 2015), while other studies find an impairment of motor sequence learning for the affected hand after stroke (Dirnberger et al., 2013; Exner et al., 2001; Fleming et al., 2018; Gomez Beldarrain, 1999; Gomez Beldarrain et al., 2002; Wadden et al., 2017). All available examples of these studies investigated patients in the chronic phase of stroke.

While two studies mainly addressed explicit learning (Fleming et al., 2018; Zimerman et al., 2012), one study investigated both learning modes (Gomez Beldarrain et al., 2002), and the majority of studies tested implicit learning capacity using the SRTT or the CTT.

Concerning explicit learning, two of the abovementioned studies revealed an impairment in explicit motor learning after stroke (Fleming et al., 2018; Gomez Beldarrain et al., 2002). One study reported a beneficial effect of transcranial direct current stimulation on the impaired capacity for explicit motor learning (Zimerman et al., 2012).

Regarding implicit sequence learning, the results are more divergent than those for explicit learning. Six studies demonstrated an impairment in implicit sequence learning for the affected hand (Dirnberger et al., 2013; Exner et al., 2001; Gomez Beldarrain, 1999; Gomez Beldarrain et al., 2002; Wadden et al., 2017), while three others reported that implicit motor learning was preserved after stroke (Meehan et al., 2011; Shin et al., 2005; Wadden et al., 2015). Additionally, a recent meta-analysis from 2017 investigated the question of whether implicit motor learning is preserved after stroke. The authors found insufficient evidence that stroke patients can implicitly learn with the affected side. No significant difference was found between implicit and explicit sequential motor learning after stroke (Kal, Houdijk, et al., 2016; Kal, Winters, et al., 2016).

Regarding the effect of stroke on the unaffected, ipsilesional hand, there is some evidence that implicit motor sequence learning is possible after stroke (Boyd & Winstein, 2003; Dovern et al., 2015; Dovern et al., 2017; Orrell et al., 2007; Pohl et al., 2001; Shin et al., 2005). This result was also found in the meta-analysis by Kal et al., who stated that stroke patients are able to learn implicitly on the
| Author, year                          | Lesion location          | n | Lesion side       | Stroke stage          | Method               | Tested hand | Endpoint | Learning type | Outcome/conclusion                                                                 |
|--------------------------------------|--------------------------|---|-------------------|-----------------------|----------------------|-------------|----------|---------------|------------------------------------------------------------------------------------|
| Boyd & Winstein, 2001                | SMC, pons                | 12| Both, unilateral  | Chronic (>6 months)   | SRTT, with/without EI| Unaffected  | RT change| Implicit and explicit          | Impaired implicit motor-sequence learning in stroke, EI prior to physical practice benefits implicit learning |
| Boyd & Winstein, 2003                | SMC                      | 10| Both, unilateral  | Chronic (>6 months)   | SRTT, with/without EI| Unaffected  | RT change| Implicit and explicit          | Preserved implicit sequence learning, but EI degraded both performance and learning |
| Boyd & Winstein, 2004                | BG                       | 10| Both, unilateral  | Chronic (>6 months)   | CTT, with/without EI | Unaffected  | RMSE changes| Implicit and explicit          | No improvement of implicit learning by providing EI |
| Boyd, Quaney, Pohl, & Winstein, 2007 | Cortical/subcortical/both| 28| Both, unilateral  | Chronic (>6 months)   | SRTT/SHMT            | Unaffected  | RT change| Implicit          | Impaired implicit sequence learning in moderate-stroke group, performance of mild-stroke and HC dependent on type of task |
| Boyd et al., 2009                    | BG                       | 13| Both, unilateral  | Chronic (>6 months)   | SRTT                 | Unaffected  | RT change| Implicit          | Impaired implicit sequence learning and chunking in basal ganglia stroke |
| Dovern et al., 2013                  | Cerebellum               | 10| Both, uni- or bilateral | Chronic (>6 months)   | SRTT                 | Both, simultaneously | RT change| Implicit          | Impaired implicit sequence learning in cerebellar stroke patients |
| Dovern et al., 2015                  | MCA (no further information) | 24| Left              | Subacute/chronic (>4 days) | SRTT                 | Unaffected  | RT change| Implicit          | Preserved implicit sequence learning only when both spatial and temporal sequence information are provided |
| Dovern et al., 2017                  | MCA (no further information) | 12| Left              | Subacute/chronic (129 ± 187d) | SRTT variant | Unaffected  | RT change| Implicit          | Preserved implicit sequence learning in stroke, even if no predictable temporal information is available, generally slower RT in stroke |
| Exner, Weniger, & Irle, 2001         | Thalamus                 | 15| Both, uni- or bilateral | Chronic (>6 months)   | SRTT variant         | "Right or left hand" | RT change| Implicit          | Impaired implicit learning in thalamus as well as in BG lesions (esp. lesions located in ventral portions of the thalamus) |
| Fleming, Newham, & Rothwell, 2018    | Cortical/subcortical/infratentoriell | 12| Both, unilateral  | Chronic (>6 months)   | Reaching for targets on a monitor | Affected  | OT, PL, speed| Explicit          | Impaired sequence specific learning in stroke |
| Author, year | Lesion location | n  | Lesion side | Stroke stage | Method | Tested hand | Endpoint | Learning type | Outcome/conclusion |
|-------------|----------------|----|-------------|--------------|--------|-------------|----------|---------------|-------------------|
| Gomez Beldarrain, 1999 | Prefrontal cortical | 22 | Both, unilateral | Immediately after diagnosis/chronic | SRTT variant | Both, simultaneously | RT change, errors | Implicit | Impaired implicit sequence learning in PFC lesion, larger lesions (>2 cm) significantly more impaired |
| Gomez Beldarrain, Gafman, Ruiz de Velasco, Pascual-Leone, & Garcia-Monco, 2002 | Prefrontal cortical | 25 | Both, uni- or bilateral | Chronic (>24 months), 1x < 12 months | SRTT / PTT | Both, simultaneously | Change of RT (SRTT) / RMSE (PTT) | Implicit and explicit | Impaired implicit and explicit sequence learning in PFC lesioned patients |
| Gomez Beldarrain et al., 2008 | Prefrontal | 14 | Both, uni- or bilateral | Chronic (>12 months) | SRTT variant before and after sleep | “Preferred” | RT change | Implicit | Impaired sequence learning in PFC and parietal lesion, benefit from a night of sleep only in patients with prefrontal lesions |
| Meehan, Randhawa, Wessel, & Boyd, 2011 | Subcortical | 9 | Right | Chronic (>12 months) | CTT | Affected | RMSE change | Implicit | Preserved implicit sequence motor learning, but more errors in stroke group |
| Orrell, Eves, Masters, & MacMahon, 2007 | “Anterior circulation system” | 7 | Right | Chronic (>12 months) | SRTT variant | Unaffected | RT change | Implicit | Preserved implicit sequence learning but generally slower RT in stroke |
| Pohl, McDowd, Filion, Richards, & Stiers, 2001 | No information given | 47 | Both, unilateral | Chronic (>6 months) | Hand movement sequence | Unaffected | RT change | Implicit | Preserved implicit sequence learning, but generally slower RT in stroke |
| Pohl, McDowd, Filion, Richards, & Stiers, 2006 | No information given | 37 | No information given | Subacute (18–45 d) | Hand movement sequence | Unaffected | RT change | Implicit | Preserved implicit sequence learning in mild and moderate stroke, generally slower RT in moderate stroke group |
| Rösser et al., 2008 | Cortical / subcortical | 18 | No information given | Chronic (>12 months) | SRTT variant | Affected | RT change | Implicit | L-Dopa treatment improves procedural learning capacity in chronic stroke patients |
| Shin, Aparicio, & Ivy, 2005 | BG | 4 | Both, unilateral | No information given | SRTT | Both, separately | RT learning score (latency and accuracy) | Implicit | Preserved implicit sequence learning (spatial and temporal) in BG stroke for ipsi- and contralateral hand |
| Siensukon & Boyd, 2009 | Cortical / subcortical | 40 | Both, unilateral | Chronic (>12 months) | CTT with / without night of sleep | Unaffected | RMSE change | Implicit and explicit | Sleep improved offline consolidation in stroke patients in explicit and implicit tracking task |
| Vakil, Kahan, Huberman, & Osimani, 2000 | BG | 16 | Both, unilateral | No information given | SRTT (motor + non-motor) | RH (five affected/11 unaffected) | RT change | Implicit | Impaired implicit sequence learning (motor and non-motor) |

(Continues)
However, multiple studies revealed impairment in implicit motor sequence learning (Boyd et al., 2007; Boyd & Winters, 2003; Dirnberger et al., 2013; Exner et al., 2001; Gomez Beldarrain, 1999; Gomez Beldarrain et al., 2002; Vakil et al., 2000).

With respect to explicit motor learning, the meta-analysis found no significant difference between implicit and explicit sequential motor learning for the unaffected hand (Kal, Houdijk, et al., 2016; Kal, Winters, et al., 2016).

There are some available data suggesting a correlation between stroke severity and the degree to which motor sequence learning is impaired (Boyd et al., 2007; Gomez Beldarrain, 1999; Pohl et al., 2001). This correlation might also affect the ipsilesional hand. In multiple studies, patients with different degrees of stroke severity (“mild” vs. “moderate”, assessed by scores on the Orpington Prognostic Scale) showed different amounts of absolute improvement in the reaction time of trained, patterned sequences (Boyd et al., 2007; Pohl et al., 2001). In contrast, another study found no difference in sequence-specific learning between mildly and moderately affected patients, while there was an overall slowing of reaction time in the more severely affected patients (Pohl et al., 2006). Limitations of these studies include the lack of information on the exact location and extent of each lesion. The effects of stroke severity on implicit learning might also be influenced by task context. Mildly affected patients showed greater improvement in the serial hand movement task than in the SRTT, while patients with moderate stroke did not show any difference between the two test modalities (Boyd et al., 2007). The various results of different test modalities emphasize the impact of the type of motor task that is tested, with the tasks possibly assessing different qualities of motor learning that make different demands on working memory.

Despite the heterogeneity of the results, there is a tendency for motor sequence learning to be more impaired in the affected hand than in the unaffected hand, while the unaffected hand also demonstrates at least partly impaired function in some studies. However, a general finding of nearly all studies is a decreased reaction time and a permanently increased error rate in stroke patients compared to healthy controls, irrespective of the lesion location, the investigated side or the learning stage (Boyd et al., 2007; Boyd & Weinstein, 2003; Dovern et al., 2017; Fleming et al., 2018; Gomez Beldarrain et al., 2002; Meehan et al., 2011; Orrell et al., 2007; Pohl et al., 2006; Siengsukon & Boyd, 2009; Vakil et al., 2000; Wadden et al., 2015).

### 5.2 Contributions of stroke studies to the temporal stage model of motor sequence learning

In this section, we will analyze the effects of cerebral lesions on motor sequence learning in further detail. We are particularly interested in the relationship between the location of the lesion and its impact on motor sequence learning. However, as described above, the function and importance of a particular brain area for motor sequence learning change across different learning stages. Lesions might, therefore, compromise motor sequence learning differently in different learning stages.
stages. It is crucial to differentiate the behavioral consequences of specific lesions in relation to the stages of learning. This differentiation is especially challenging because the available studies do not address the effects of lesion location on learning stages as their primary research questions.

5.2.1 Stage I—Contributions of stroke studies to knowledge of the early learning phase of sequential motor learning

The PFC is known to be one of the core areas involved in early learning. Accordingly, early learning was found to be significantly reduced in patients with prefrontal lesions (Gomez Beldarrain, 1999; Gomez Beldarrain et al., 2002; Gomez Beldarrain, Astorgano, Gonzalez, & Garcia-Monco, 2008). Prefrontal lesions caused disturbances in explicit learning tasks as measured by a pursuit tracking task (Gomez Beldarrain et al., 2002) as well as in tasks that were mainly implicit (Gomez Beldarrain, 1999). Overall, impairments in motor sequence learning were more severe in the contralateral hand than in the ipsilateral hand and were correlated with lesion size (Gomez Beldarrain, 1999).

Deficits in implicit learning were found to extend beyond the very early learning stage (Gomez Beldarrain, 1999; Gomez Beldarrain et al., 2002). The authors suggested not only that the involvement of the PFC extends beyond the early learning stage but also that different parts of the PFC contribute diversely to the different learning modes and learning stages (Gomez Beldarrain et al., 2002). While the dorsolateral PFC is heavily involved in the network subserving working memory (Manoach et al., 1999), the frontopolar PFC is heavily involved in planning (Koechlin, Basso, Pietrini, Panzer, & Grafman, 1999), suggesting that deficits in early and explicit learning are more pronounced after frontopolar than dorsolateral PFC lesions, while the reverse is true of deficits in implicit learning and during later stages.

Current knowledge regarding lesions in the PFC suggests that every learning task, even if it mainly involves implicit learning, requires prefrontal activation with active attention in the very beginning. If this early learning stage is disturbed, the transition into the consolidation phase may also be impaired and delayed. Accordingly, patients with prefrontal lesions showed greater benefit than healthy controls from a night of sleep (Gomez Beldarrain et al., 2008).

A reduction in early sequence-specific motor learning was also demonstrated in patients with lesions in the ventral portion of the thalamus, an area that is strongly connected to the PFC (Exner et al., 2001). The authors speculated about the importance of individual thalamic nuclei, suggesting that they might be specific to sequence learning. However, given the importance of thalamic nuclei for cortico-cortical communication, we suggest instead that these results simply reflect an impairment of the thalamic contribution to driving and modulating cortico-thalamo-cortical information transfer.

Concerning lesions of the BG, some available data suggest that early implicit sequence learning is not seriously impaired for the unaffected hand (Dovern et al., 2015; Shin et al., 2005; Vakil et al., 2000) or the affected hand (Shin et al., 2005; Vakil et al., 2000). Accordingly, the administration of levodopa to stroke patients did not improve implicit learning in the very early stages, while improvements were found in later stages (Rösser et al., 2008). These findings are in line with a growing body of literature suggesting that the BG are mainly active in the consolidation of learned sequences. In contrast, one study reported a diminished learning rate in patients with chronic BG lesions, even in the early acquisition phase, as measured by diminished RT changes for repeated sequences (Boyd et al., 2009).

The effect of cerebellar lesions on early learning has not been thoroughly studied. One study found that implicit learning was preserved in the early learning stage, whereas impairments were found in later stages (Dimberger et al., 2013). This finding seems counterintuitive given what is known about the function of the cerebellum in healthy subjects. One possible explanation might be the diversity of cerebellar lesion locations, possibly mainly affecting regions that are known to be involved in late learning stages, such as the dentate nucleus. The authors further noted that there is a disturbance especially in perception-based mechanisms (Dimberger et al., 2013), underlining the role of the cerebellum in the “predictive” mechanism of movement control (Bastian, 2006).

Several studies were unable to find significant impairments of sequential motor learning in the early learning phase after stroke, either for the affected hand (Fleming et al., 2018; Meehan et al., 2011; Siensksukon & Boyd, 2009) or for the unaffected hand (Dovern et al., 2017; Orrell et al., 2007; Pohl et al., 2001; Pohl et al., 2006). One important reason for this lack of a significant effect might be the selection of patients. Multiple studies have selected patients on the basis of anamnestic information on stroke and clinical impairments. These studies have often used clinical scores such as the Orpington Prognostic Scale (Pohl et al., 2001, Pohl et al., 2006), the Fugl-Meyer Assessment (Fleming et al., 2018), or the MRC Scale (Rösser et al., 2008). These scores, however, do not test for difficulties in learning. They are mainly sensitive to lesions in brain areas responsible for gross motor movements and are particularly altered by lesions of the primary motor cortex itself or its downstream connections. Thus, the lesion locations in the cited studies are strongly biased toward lesions of these areas. The negative results therefore suggest that brain lesions of the motor cortex or its downstream connections exert limited influence on early motor learning. This was observed mainly in the chronic stages of stroke but also seemed to apply to patients in the subacute stroke stage (<45 days after stroke), who also showed no difference in sequence-specific implicit learning compared to healthy controls (Pohl et al., 2006).

Overall, many studies reported fully or at least partly preserved motor sequence learning. Interestingly, there are no reports of patients with a complete inability to achieve motor learning. Hence, it can be assumed that there is no “all-or-nothing principle,” even when one important part of the responsible network is destroyed. This further indicates that sequential motor learning in the early stages is performed by a widespread network of brain regions without a central region mandatory for learning success and that lesions to brain areas engaged in this process can be (at least partly) compensated.
5.2.2 | Stage II—Contributions of stroke studies to knowledge of the consolidation phase of sequential motor learning

Multiple studies have investigated motor sequence learning after stroke during the consolidation phase using an extended version of a sequence task or by investigating sequence practice over several days.

Impaired implicit and explicit motor sequence learning was consistently found in patients with lesions of the BG (Siengsukon & Boyd, 2009; Valkil et al., 2000). To further elucidate the underlying mechanisms, analyses have focused on the individual organization of learned responses and the transition from one sequence element to the next. The analysis of this transition allows the identification of subsequences (chunks) of movements (Miller, 1994; Povel & Collard, 1982; Rosenbaum, Kenny, & Derr, 1983). Studies have found substantial differences between patient and control groups in organizing sequential responses (Boyd et al., 2009; Dovern et al., 2015). It was demonstrated that the ability to chunk elements of a repeated, implicitly presented sequence into functional subsequences of movements is impaired after BG lesions (Boyd et al., 2009). Patients organized sequences to a lesser extent than healthy controls, and the chunks were also shorter (Boyd et al., 2009). This impairment was demonstrated for the affected as well as the unaffected hand and even if the patients did not differ from healthy controls during the early learning stage (Boyd et al., 2009; Valkil et al., 2000).

A further study augmented these results by investigating the ability to learn a motor sequence in patients after left MCA lesions by giving them implicit multimodal (spatial and temporal) information (Dovern et al., 2015). The learning progress did not differ between patients and controls. However, patients demonstrated an impaired ability to form independent spatial and temporal sequence representations. The underlying subchunks were found to be more rigid and inflexible than those of healthy controls (Dovern et al., 2015). Imaging analyses revealed that diminished learning scores were strongly associated with lesions of the striatum. These studies demonstrate that the chunking of subsequences might depend on the complexity of the given multidimensional information, which stroke patients learn in a more rigid way than healthy controls. The associations between multiple information dimensions were further investigated by a study that demonstrated preserved implicit learning effects in the event of consistently missing temporal information (Dovern et al., 2017). This further suggests that patients with BG lesions do not show decreased learning effects due to missing additional information but that they are particularly susceptible to context disruptions, which then decrease learning effects (Dovern et al., 2017).

The critical role of the BG in chunking is supported by the finding of faster responses on learned sequences (but not random sequences) in the consolidation phase after administration of levodopa than without levodopa in stroke patients (Rösser et al., 2008).

Results similar to those of BG lesions were also found in lesions of the ventral thalamic nuclei (Exner et al., 2001), with decreased implicit learning not only in the early learning stage but also in the consolidation phase (Exner et al., 2001). However, the effects of thalamic lesions are less investigated than those of BG lesions. It is tempting to speculate that this effect is caused by the high level of interconnection between the thalamus and the BG (Alexander, DeLong, & Strick, 1986; Middleton & Strick, 2000).

Interestingly, when additional explicit information is given to patients with lesions affecting the BG or sensorimotor cortical areas, this information seems to have an adverse effect on implicit learning during the consolidation phase (Boyd & Weinstein, 2003; Boyd & Weinstein, 2004). Contrary to these findings, a further investigation found an improvement in response to additional explicit information in chronic unilateral stroke patients (11 sensorimotor areas, 1 pontine; Boyd & Weinstein, 2001). To explain these findings, the authors suggested differences according to the exact lesion location, particularly with respect to M1, the PMC, and the SMA. In particular, the PMC might play a central role in the integration of explicit information into implicit motor performance due to its strong connections to prefrontal areas (e.g., the DLPFC) and to the caudate nucleus. Hence, a lesion that strongly affects the PMC might interfere with the benefit of additional explicit information for an implicitly formed motor plan.

It was further suggested that the additional explicit information given to the subjects imposed an unmanageable working memory load that might interfere with the implicitly made movement plan. However, the small sample sizes in the patient groups that received explicit knowledge (n = 4 [Boyd & Weinstein, 2001]; n = 5 [Boyd & Weinstein, 2003]; n = 5 [Boyd & Weinstein, 2004]) might have also contributed to these divergences.

Several studies have failed to show a significant poststroke impairment of motor sequence learning during the consolidation phase. Preserved learning was mainly found in studies that were not restricted to specific lesion locations. For instance, preserved implicit motor learning with the unaffected hand during the consolidation phase was found for lesions in the "anterior circulation" or "MCA" territory or for "cortical/subcortical" infarcts (Orrell et al., 2007; Pohl et al., 2001; Siengsukon & Boyd, 2009), but implicit learning was also preserved for the affected hand after subcortical lesions (Meehan et al., 2011; Wadden et al., 2015).

However, there are also studies available demonstrating an impairment of motor sequence learning during the consolidation phase without specifying the exact lesion location (Dovern et al., 2015; Wadden et al., 2017). In contrast to studies demonstrating preserved sequence learning, these studies used different methods of analysis or design. Instead of comparing the group difference in RT changes between random and predefined sequences, Wadden and colleagues analyzed group differences by fitting the learning progress with an exponential curve and included the behavioral parameter of individual motor impairment in the model (Wadden et al., 2017). They demonstrated impaired implicit motor sequence learning with the affected hand after subcortical lesions in the right hemisphere. However, a careful inspection of the available data suggests that the main difference between patients and controls is attributable more to the early learning phase than to the consolidation phase.
However, consolidation does not occur exclusively during extended practice. There is general agreement that the consolidation process of learned sequence information can also be improved by sleep. This process is called sleep-dependent offline consolidation.

Expectably, BG lesions seem to impair offline consolidation (Dovern et al., 2015). This is consistent with the results of neuroimaging studies in healthy people, which showed increased integration in the corticostriatal network after sleep following a motor sequence-learning task (Debas et al., 2010; Debos et al., 2014). A similar effect of impaired offline consolidation was also found after parietal lesions. In contrast, patients with prefrontal lesions seem to benefit from offline consolidation during sleep more strongly than healthy subjects (Gomez Beldarrain et al., 2008). These findings support the assumption that the parietal cortex is necessary in both the early and late stages of learning, while the PFC is mainly involved in early learning processes.

Without providing exact lesion information (cortical/subcortical), a further study demonstrated improved sleep-dependent offline learning in stroke patients for explicit and implicit tasks compared to healthy subjects (Siengsukon & Boyd, 2009). No significant difference was found without sleep. The authors hypothesized a change in sleep structure after stroke with an impact on stroke recovery. However, no sleep measurements were performed to test this hypothesis.

5.2.3 Stage III—Contributions of stroke studies to knowledge of the late learning phase of sequential motor learning

Very limited data are available regarding potential impairments of stroke patients in the late learning phase of motor sequence learning. Studies have investigated motor sequence learning for a maximum of 5 days of training (Meehan et al., 2011; Wadden et al., 2015; Wadden et al., 2017). However, the question of whether a late learning stage is reached depends not only on the training time but also on the complexity of the task. All three cited studies investigated a CTT. It remains unclear whether, after 5 days of training on a CTT, the late learning stage is reached. The available data give some indications that the process remains in the consolidation phase, although the boundary between these learning stages is uncertain. However, even if those studies reached the late learning stage, the available data do not allow us to draw conclusions regarding specific effects of stroke on that stage.

One study tested delayed retention of the SRTT on day 15 after 2 days of practice in patients with right hemispheric lesions. No difference in retention was found between the stroke group and the control group (Orrell et al., 2007). Despite the prolonged time window of retention, the late learning phase is presumable not reached in only 2 days of practice.

5.2.4 Conclusions

Despite some heterogeneities, the results of studies that investigated patients with motor impairments due to brain lesions indicate impairments of motor sequence learning in the early learning stage by impaired integration of multimodal information for motor sequence learning, slowed general reaction time and impaired explicit learning. Stroke patients have greater difficulty in optimizing the motor control components of sequential movements than the sequential/spatial order of movements. There is consistent evidence that the consolidation phase is particularly impaired after BG lesions. This effect is probably due to impaired chunking of information.Chunks in patients with BG lesions seem to be more rigid, more sensitive to disruptions and less adaptable than those in healthy controls. There are some indications that MCA infarcts in general cause difficulties in integrating information necessary for building chunks of sequences during the consolidation phase. Decreased learning rates after the offline consolidation phase were found in striatal and parietal lesions but not in prefrontal lesions.

However, the heterogeneity and small sample sizes of the available studies render it difficult to draw clear conclusions.

5.3 Methodological considerations

From a theoretical perspective, it seems obvious that motor sequence learning after stroke must be impaired to some degree if lesions affect brain structures that are necessary for motor sequence learning. However, as outlined above, multiple studies failed to find clear evidence of impaired motor sequence learning after stroke (Meehan et al., 2011; Shin et al., 2005; Wadden et al., 2015). A recent meta-analysis found insufficient evidence that stroke patients could implicitly learn with the affected side (Kal, Houdijk, et al., 2016; Kal, Winters, et al., 2016). Although the meta-analysis was thoroughly performed, the underlying studies varied in so many aspects that results are difficult to combine. To better understand the reasons for the divergences in study results, it is necessary to discuss some methodological issues that, in our opinion, affect the interpretation of results and possible conclusions.

First, an important factor contributing to divergent results is the inhomogeneity of lesions across studies. There are multiple studies available that present no information on the method through which lesion location was determined (Dimberger et al., 2013; Dovern et al., 2017; Fleming et al., 2018; Gomez Beldarrain et al., 2008; Meehan et al., 2011; Pohl et al., 2001; Rösser et al., 2008; Shin et al., 2005; Wadden et al., 2017; Zimerman et al., 2012). Some studies stated that infarctions were located in the “MCA territory” or “anterior circulation system” without further information (Orrell et al., 2007; Pohl et al., 2001; Pohl et al., 2006). Other studies provided some information on lesion location while mixing different lesion locations together (cortical/subcortical/both (Boyd et al., 2007), cortical/subcortical/infratentorial (Fleming et al., 2018), and cortical/subcortical (Rösser et al., 2008; Siengsukon & Boyd, 2009). Some studies restricted the lesions to subcortical locations (Meehan et al., 2011; Wadden et al., 2015; Zimerman et al., 2012). Only a few studies investigated lesion locations in a more specific way by analyzing the relationship between lesion location and performance in motor sequence learning (Wadden et al., 2017).

Aside from lesion locations, the available studies also varied in other characteristics, such as severity of stroke, time since stroke, sample size, task types, and duration of training. Moreover, it remains
insufficiently understood how motor impairment of the affected arm interferes with and affects the results of sequence learning.

The chosen method for testing learning outcomes is another problem. Most studies used an SRT task and measured reaction time as the primary outcome parameter. SRTT is generally very susceptible to experimental changes that may explain some of the variability in the results (Willingham, Greenberg, & Thomas, 1997). In addition, there are methodological concerns regarding the interpretation of SRTT results. Stroke patients have been found to have a generally slower reaction time than healthy subjects (Boyd et al., 2007; Orrell et al., 2007; Pohl et al., 2001; Pohl et al., 2006). Patients were able to improve their reaction time by practicing patterned sequences, while the introduction of random sequences increased their reaction time (Pohl et al., 2001, Pohl et al., 2006). Most studies measured motor sequence learning as changes in reaction time for a learned sequence compared to an equal-length random sequence. However, it remains unclear how an improvement in patients’ reaction time can be compared to the improvement of healthy controls when their baseline reaction time is different. As measured by the absolute decrease in reaction time, some patients improved more than the healthy subjects (Pohl et al., 2001, Pohl et al., 2006). As measured by percentage improvement, patients with longer baseline reaction time must improve by a greater absolute quantity to achieve the same percentage decrease, which further complicates comparisons. With respect to these methodological problems, the question of whether and to what degree sequence learning after stroke is impaired cannot be answered definitively in most studies. An important methodological advance can be found in a recent study that compared performance data between groups by fitting them to an exponential function. This method allows the rate of skill acquisition to be estimated and additionally allows an improved comparison of learning outcome parameters (Wadden et al., 2017). The authors found significantly slower rates of improvement in implicit sequence-specific motor performance with the affected hand in stroke patients than in healthy controls.

Theoretical considerations and the limited available data suggest that impairments of sequential motor learning are influenced by stroke severity. However, the available stroke studies are biased concerning the severity of stroke, with a preference for mildly to moderately affected patients, while the ability of severely affected patients to learn motor sequences remains unknown. This is because it is quite difficult to investigate severely impaired patients, as the severity of impairment might interfere with the results in many dimensions (compromised execution due to motor impairment as well as additional neurological disabilities such as neglect, aphasia, apraxia, cognitive impairment, and general slowing of movements). There are also logistical concerns to address.

6 | PRACTICAL CONCLUSIONS FOR PATIENT REHABILITATION

One of the most important concepts for practical rehabilitation is the differentiation between explicit and implicit motor learning, as these two forms of motor learning require different processing pathways and cerebral resources. Accordingly, cerebral lesions can affect one system more strongly than the other (Reber & Squire, 1998). For example, severe deficits in explicit learning have been reported after medial temporal lobe damage and in patients with amnesia, while implicit learning capabilities remained intact (Reber & Squire, 1998; Shadmehr et al., 1998). With respect to explicit learning deficits, there are multiple available studies demonstrating that the main process in explicit learning (the shift from using declarative knowledge and active attention toward using procedural knowledge) can be disturbed after stroke (Kal, Houdijk, et al., 2016; Kal, Winters, et al., 2016; Kleynen et al., 2013; Orrell, Masters, & Eves, 2009).

This is particularly problematic because deficits in the level of conscious control of movements are associated with the extent of functional impairment after stroke (Orrell et al., 2009; Stapleton, Ashburn, & Stack, 2001). Accordingly, the more severe the impairments, the less they are able to profit from explicit motor learning.

This problem affects motor rehabilitation, as most physical therapists provide mainly verbal or physical instructions on how a movement should be performed to improve motor performance, thereby engaging explicit motor learning (Durham, Van Vliet, Badger, & Sackley, 2009; Johnson, Burridge, & Demain, 2013). A disturbance in explicit motor learning might prevent patients from reaching a high level of automated motor performance for motor tasks that are initially highly dependent on working memory and conscious control.

The current studies on implicit motor learning are promising because they suggest that this process is preserved in most patients. However, a major problem here is that implicit motor learning after stroke was mostly investigated with the SRTT, which has little or no relevance to practical rehabilitation. It remains unclear whether the positive findings for implicit learning can be extended to more complex motor tasks, in which not only the sequence of movements but also the optimization of the movement parameters of the motor sequence must be considered. The quality of the available studies is mostly insufficient to derive precise recommendations for practical rehabilitation. Nevertheless, the available data suggest that rehabilitation strategies should be better adapted to individual needs based on lesion location and consecutive functional impairments.

Patients suffering from infarcts affecting frontoparietal regions with impairment in the early learning phase should be trained by using slow steps of explicit additional information with little attentional demand. Sequences should be simple so that the consolidation phase can be reached as fast as possible. In addition, rehabilitation approaches for these patients should also use learning strategies that involve implicit learning, rather than relying solely on explicit instructions. Possible and promising approaches are dual-task learning, analogy learning or errorless learning (Kal, Houdijk, et al., 2016; Kal, Winters, et al., 2016; Kleynen, Braun, et al., 2014; Kleynen, Wilson, et al., 2014; Orrell, Eves, & Masters, 2006). There are few studies exploring these implicit approaches in stroke patients. Orrell et al. investigated the errorless learning strategy by using a dynamic balancing task (Orrell et al., 2006) and found a benefit of implicit motor learning techniques compared to the traditional discovery
learning based on explicit rules. The analogy learning approach was investigated in a little pilot study by Kleynen et al. who concluded that analogy learning might be a feasible and useful intervention in stroke rehabilitation (Kleynen, Braun, et al., 2014; Kleynen, Wilson, et al., 2014). Nevertheless, further research concerning stroke patients is needed.

Patients suffering from BG lesions are impaired mainly in the consolidation phase. It remains unclear whether the use of the available attentional resources is beneficial or whether these patients should focus on simple sequences for consolidation. Additional time should be given to the chunking of information to avoid the usage of inefficiently short subchunks.

It also remains unclear whether the potential for interhemispheric transfer of sequence knowledge in healthy subjects can be used to improve motor learning in patients. If sequence learning for the affected hand is impaired, the patient can attempt to learn the sequence with the other hand, at least in the early phase of motor learning. Afterward, the training might be further practiced with the affected hand. However, there are no studies available that investigate this hypothesis. There are also no data available that investigate the impact of patient motivation and the functioning of the reward system on motor sequence learning.

7 | PERSPECTIVES AND IMPLICATIONS FOR FUTURE RESEARCH

In the last decade, the body of knowledge regarding the effects of structural brain lesions on motor sequence learning has grown substantially. There is increasing understanding of the effects of a non-localized stroke with mild to moderate clinical impairments on the capacity for explicit and implicit motor sequence learning. However, as outlined above, we can further increase our understanding through studies that are less heterogeneous than the existing literature. First, there is an urgent need for studies that investigate motor sequence learning in patients with highly localized lesions. In this respect, the most consistent studies are available for BG lesions. These studies have contributed substantially to our understanding of BG function in motor sequence learning and have laid groundwork for our understanding of deficits in the chunking of information. Similar increases in understanding can be expected from studies investigating lesions in other key regions of the brain network responsible for motor sequence learning.

Most studies used tasks in which the complexity of the sequence was contrasted by the simplicity of its motor implementation (e.g., pressing of buttons). It has been shown in healthy subjects that sequence learning is strongly affected by the complexity of the stimulus–response association. The performance of a well-synchronized but complex task requires the optimization of multiple parameters such as timing, velocity and force that contribute to performance. The effect of increased complexity of the stimulus–response association on motor sequence learning in stroke patients remains unknown but could provide important insights for the practice of rehabilitation.

It is also unknown whether impaired sequence learning with the affected hand can be improved by training with the unaffected hand in the early learning stages.

Another important question that we are only beginning to investigate is the influence of stroke on different learning stages. There are currently no studies available that focus on the role of lesion location on different learning stages. In particular, there are no studies available that use different learning strategies in different learning stages.

The influence of the structural integrity and functioning of the reward system on sequential motor learning in stroke patients is unknown.

Moreover, in the current state of research, studies without clear descriptions of the lesion locations do not contribute substantially to further insights. Instead, there is a desperate need for studies investigating more complex stimulus–response associations with more precise lesion mapping across different learning stages. Such studies have the potential to reveal the mechanisms underlying impaired sequence learning and insufficient rehabilitation results. Such studies might also contribute to the development of new therapeutic strategies that are more individualized than what is currently available.

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CONFLICT OF INTERESTS

The authors declare that there are no conflicts of interest.

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