Locomotor adaptation on a split-belt treadmill can improve walking symmetry post-stroke

Darcy S. Reisman,1,2 Robert Wityk,3 Kenneth Silver4 and Amy J. Bastian2,3

1Department of Physical Therapy, University of Delaware, Newark, DE, 2Kennedy Krieger Institute, Departments of 3Neurology and 4Physical Medicine and Rehabilitation, Johns Hopkins University School of Medicine, Baltimore, MD, USA

Correspondence to: Darcy S. Reisman, PhD, PT, Department of Physical Therapy, University of Delaware, 322 McKinley Laboratory, Newark, DE 19716, USA
E-mail: dreisman@udel.edu

Human locomotion must be flexible in order to meet varied environmental demands. Alterations to the gait pattern occur on different time scales, ranging from fast, reactive adjustments to slower, more persistent adaptations. A recent study in humans demonstrated that the cerebellum plays a key role in slower walking adaptations in interlimb coordination during split-belt treadmill walking, but not fast reactive changes. It is not known whether cerebral structures are also important in these processes, though some studies of cats have suggested that they are not. We used a split-belt treadmill walking task to test whether cerebral damage from stroke impairs either type of flexibility. Thirteen individuals who had sustained a single stroke more than 6 months prior to the study (four females) and 13 age- and gender-matched healthy control subjects were recruited to participate in the study. Results showed that stroke involving cerebral structures did not impair either reactive or adaptive abilities and did not disrupt storage of new interlimb relationships (i.e. after-effects). This suggests that cerebellar interactions with brainstem, rather than cerebral structures, comprise the critical circuit for this type of interlimb control. Furthermore, the after-effects from a 15-min adaptation session could temporarily induce symmetry in subjects who demonstrated baseline asymmetry of spatiotemporal gait parameters. In order to re-establish symmetric walking, the choice of which leg is on the fast belt during split-belt walking must be based on the subject’s initial asymmetry. These findings demonstrate that cerebral stroke survivors are indeed able to adapt interlimb coordination. This raises the possibility that asymmetric walking patterns post-stroke could be remediated utilizing the split-belt treadmill as a long-term rehabilitation strategy.

Keywords: stroke; motor; locomotion; movement

Abbreviations: IRED = infrared emitting diode

tied at the same speed). In contrast, step lengths also are initially asymmetric, but an adaptive response occurs during split-belt walking that acts to re-establish symmetry via feedforward changes in phasing between legs. This adaptation induces an after-effect causing walking asymmetry when returned to normal treadmill conditions. A similar adaptive phenomenon occurs during circular treadmill locomotion, resulting in curved walking trajectories as after-effects (Gordon et al., 1995; Weber et al., 1998). When wearing a robotic gait orthosis that provides viscous resistance to the leg, subjects modulate the swing phase through feedback, but adapt feedforward control of the leg just prior to swing (Lam et al., 2006).

Which brain structures or pathways underlie these processes? Recent work in humans suggests that the cerebellum is required for slower adaptive changes in locomotion, but may not be as critical for reactive changes (Morton and Bastian, 2006). In that study, subjects with cerebellar disorders were able to immediately adjust stance times to split-belt treadmill speeds, but could not adapt interlimb phasing and showed no after-effects, suggesting that reactive changes rely on different descending commands or the interaction of spinal neural networks with the mechanical oscillation of the legs (Lacquaniti et al., 2002; Courtine and Schieppati, 2004). Similar cerebellar deficits in adaptation have been shown in circular treadmill locomotion (Earhart et al., 2002). In cats, nitric oxide deprivation, which is thought to play a role in long-term depression in the cerebellum, abolishes locomotor adaptive capacity but not reactive ability (Yanagihara and Kondo, 1996).

Studies of spinal and decerebrate cats suggest that both forms of flexibility occur in quadrupedal locomotion without cerebral inputs (Kulagin and Shik, 1970; Forssberg et al., 1980; Yanagihara and Kondo, 1996). Yet, the flexibility and stability requirements of human, bipedal gait are generally presumed to require some level of cerebral control (Armstrong, 1986). Evidence from adult humans with stroke or spinal cord injury suggest that locomotor interlimb coordination is strongly influenced by cerebral inputs (Dietz et al., 2002; Kautz and Patten, 2005; Kautz et al., 2006). Here we attempt to understand the role of cerebral structures in reactive modifications and/or adaptation of the locomotor pattern. Based on our previous results in subjects with cerebellar damage, we hypothesize that if cerebellar–cerebral motor connections play a predominant role in the adaptation then cerebral damage should impair adaptation. Alternatively, if plasticity at multiple neural levels or connections between the cerebellum and other motor areas are more important, then adaptation should be relatively normal.

Additionally, it is unknown if individuals with cerebral damage have the potential to alter their impaired walking pattern through short-term motor adaptations that result in after-effects. Damage to cerebral structures caused by stroke leaves many individuals with an asymmetric walking pattern (Olney et al., 1994; Hsu et al., 2003; Lamontagne and Fung, 2004). If stroke survivors can show after-effects that reduce their walking asymmetry, even temporarily, following adaptive training on the split-belt treadmill, it would demonstrate that their compromised nervous system is still capable of producing a more normalized (i.e. symmetric) pattern. This then raises the possibility that certain components of gait asymmetry might be remediated with specific rehabilitation strategies using the split-belt treadmill. Thus, a secondary purpose of this study is to investigate whether after-effects following split-belt treadmill walking lead to improvements in gait symmetry in subjects following stroke. Preliminary findings were published in abstract form (Reisman et al., 2005b).

Material and Methods

Subjects

Thirteen individuals who had sustained a single stroke more than 6 months prior to the study (four females and nine males) and 13 age- and gender-matched healthy control subjects were recruited to participate in the study (Table 1). All subjects gave their informed consent prior to participating and a human studies committee approved the study. Subjects were excluded if they had other neurological conditions, orthopaedic conditions affecting the legs or back, uncontrolled hypertension, pacemaker or automatic defibrillator, active cancer, radiological and/or physical examination evidence of damage to the cerebellum or were unable to complete the task. Subjects who customarily wear an ankle-foot orthosis (AFO) were allowed to wear it during testing.

Clinical examination

Subjects underwent a clinical examination including measurement of fast walking speed, the lower extremity portion of the Fugl-Meyer, a test of coordination, reflexes and the ability to move in and out of synergy (Fugl-Meyer et al., 1975), and tests of pressure sensitivity and proprioception. Fast walking speed was measured as the average of three trials along a 25-foot walkway. Sensation of the great toe was tested using graded monofilaments. The lowest filament that could correctly be detected on four out of five trials was recorded. Proprioception was tested at the great toe, ankle, knee and hip by moving the joint ~10° and asking the subject to determine the direction of movement. The number of correct responses out of five trials was recorded and the percentage of trials correct at the great toe was used for subsequent analysis.

Testing paradigm

Subjects were asked to walk on a custom-built treadmill (Woodway USA, Waukesha, WI) comprised of two separate belts, each with its own motor, that permitted the speed of each belt (i.e. each leg) to be controlled independently. During different testing periods, subjects walked on the treadmill with the two belts either moving at the same speed (‘tied’ configuration) or different speeds (‘split-belt’ configuration). During the tied configuration, treadmill belt speeds were either ‘slow’ (0.5 m/s) or ‘fast’ (1.0 m/s). In the split-belt configuration, one treadmill belt was set at the slow speed while the other was set at the fast speed.
Stroke subjects participated in two testing sessions. In the first session, the leg assigned to the fast belt during the split-belt period was randomly determined as either the paretic or non-paretic leg. In the second session, the contralateral leg was tested on the fast belt during split-belt walking. Matched controls were randomly assigned to either their right or left leg on the fast belt and participated in only one testing session. Each session consisted of three testing periods. In the Baseline period, the treadmill belts were split (one belt fast, the other belt slow). In the Post-adaptation period, the belts were returned to the tied slow configuration. Figure 1A illustrates the experimental paradigm. The durations of each testing period were: Baseline 2 min (tied slow), 2 min (tied fast), 2 min (tied slow), Adaptation 15 min, and Post-adaptation 6 min (Fig. 1A). Subjects were given standing or seated rest breaks every 5 min during Adaptation, or more frequently as requested. Three of the stroke survivors were only able to complete 6 min of Adaptation due to fatigue.

Prior to data collection, subjects walked on the treadmill (in the tied condition) briefly at both the fast and slow speeds. They were not given any practice in the split-belt configuration, though they were told that the two belts would move at two different speeds at some point during the testing. For safety purposes while walking on the treadmill, all subjects held onto a front handrail and wore a ceiling mounted safety harness around the upper chest. The harness did not support body weight or interfere with subjects’ walking.

During testing, subjects were alerted when the treadmill was going to start, but were not instructed about belt speeds or coupling. Belt speeds were never changed while the belts were moving. Subjects were instructed to look straight ahead and refrain from looking down at the belts while walking so that they could not use visual information to determine belt speeds. An examiner stood by to monitor compliance with this instruction. At the beginning of each period, subjects were asked whether they felt the two belts were moving at the same speed or two different speeds. If they thought the belts were moving at different speeds, they were asked to indicate which belt (right or left) was moving faster.

Table I  Subject characteristics.

<table>
<thead>
<tr>
<th>Subject</th>
<th>Age</th>
<th>Lesion location</th>
<th>LE Fugl-Meyer score</th>
<th>Asymmetric Step length = S, Double support = DS</th>
<th>Fast over ground walking speed (m/s)</th>
<th>AFO a</th>
<th>Mono-filament threshold b (g)</th>
<th>Time since stroke (months)</th>
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<tr>
<td>S1</td>
<td>48</td>
<td>R hemisphere</td>
<td>33/34</td>
<td>S, DS</td>
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<td>No</td>
<td>4.31</td>
<td>192</td>
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<td>21/34</td>
<td>S</td>
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<td>No</td>
<td>4.56</td>
<td>98</td>
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<td>47</td>
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<td>15/34</td>
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<td>108</td>
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<tr>
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<td>&gt;6.65</td>
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<td></td>
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<td>4.31</td>
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<tr>
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<td>62</td>
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<td>Yes</td>
<td>4.56</td>
<td>38</td>
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aAnkle-foot orthoses.
bNormal = 3.61 g.

Fig. 1  (A) Time course for the experimental paradigm showing Baseline, Adaptation, and Post-adaptation periods. (B) Illustration of marker locations and the method used to calculate limb angle. (C) Illustration of parameter calculations. Stride and step length depicted in over ground walking with forward progression. IC = initial contact.

Data collection
Computerized gait analysis was performed using OPTOTRAK (Northern Digital, Waterloo, ON) sensors that were used to...
record 3D position data from both sides of the body (Fig. 1B). Infrared emitting diodes (IREds) were placed bilaterally (Fig. 1B) on the foot (5th metatarsal head), ankle (lateral malleolus), knee (lateral joint space), hip (greater trochanter), pelvis (iliac crest) and shoulder (acromion process). Foot contacts were determined using four contact switches per foot; two placed on the forefoot and two on the heel. Voltages reflecting treadmill belt speeds were recorded directly from treadmill motor output. Marker position and analogue data (foot switches and treadmill speed) were synchronized and sampled simultaneously using OPTOTRAK software at 100 and 1000 Hz, respectively.

Data analysis
Three-dimensional marker position data were low-pass filtered at 6 Hz. Custom software written in MATLAB (The MathWorks Inc., Natick, MA) was used for all subsequent analyses. Based on our previous work, we measured spatial and temporal walking parameters that were expected to change rapidly using online feedback control and parameters that were expected to change more gradually using adaptive mechanisms (Reisman et al., 2005a). The rapidly changing parameters were stride length and the percent time in stance phase; the adaptive parameters were step length and the percent time in double support (Fig. 1C). Stride and step lengths are spatial gait parameters while percent time in stance and double limb support are temporal measures. We calculated a third adaptive parameter, limb angle phasing, to quantify the phase relationship between the two limbs. All of these measures were calculated for both limbs. Stride length was the distance travelled by the ankle marker in the anterior–posterior direction from contact to lift-off of one limb (Fig. 1C). This method of calculation could alter stride length if subject’s anterior–posterior position on the treadmill translated during walking. However, because subjects were required to hold on to the handrail, this was not a factor. The percent time in stance phase (the time from contact to lift-off) was expressed as a percentage of the stride time (the time from contact to next contact) for that limb. Step length was the anterior–posterior distance between the ankle markers at the time when each foot contacted the ground (Fig. 1C). The percent time in double limb support was the time that both feet were in contact with the floor expressed as a percentage of the stride time for each leg. To calculate limb angle phasing, we first measured limb angle for each leg using the vector connecting the hip and metatarsal markers in the sagittal plane (Fig. 1B). Phasing between the legs was calculated as the time difference (expressed as a percent of stride time) at the peak overlap of the cross-correlation between the two limb angles over three consecutive stride cycles.

Baseline asymmetry in the group of stroke survivors was determined individually for each subject. A stroke survivor was deemed asymmetric if the average of five strides in the second slow Baseline period exceeded two standard deviations of the mean asymmetry (0.02 m for step length and 1.0% double support) in the healthy control subjects during the same period. These calculations were completed separately for double support and step length.

Statistical analyses were completed using the averages of the first five strides (after the treadmill reaches steady state speed) in each of the Baseline periods, the first (after the treadmill reaches steady state speed) and last five strides of the Adaptation period (early and late Adaptation, respectively) and the first (after the treadmill reaches steady-state speed) and last five strides of the Post-adaptation period (early and late Post-adaptation, respectively). Statistical comparisons were completed using MATLAB and Statistica (StatSoft, Tulsa, OK) software. To compare results between groups we used a repeated measures analysis of variance (ANOVA) with a between-subjects factor of group (control and stroke) and a within-subjects (repeated measures) factor of testing period (late Baseline, early and late Adaptation and early and late Post-adaptation). To compare differences between testing sessions in the paretic group we completed a repeated measures ANOVA with a within-subjects factor of leg (leg tested on the fast belt during split-belt walking). When the ANOVA yielded significant results, post hoc analyses were completed using Tukey’s honestly significant different test. Paired t-tests were used to determine changes in symmetry of double support and step length between Baseline and Post-adaptation for the sub-group of asymmetric stroke survivors. Pearson product–moment correlations were performed to test for relationships between impairment measures (LE Fugl-Meyer score, sensation and proprioception) and the magnitude of step length difference after-effect. For the limb angle phasing values, the Watson U² statistic was used to test for differences across testing periods (Batschelet, 1981). The level of statistical significance for all measures was set at $P<0.05$.

Results
For all results, we refer to the legs on the slower and faster moving belts during the split-belt portion of the paradigm as the ‘slow’ and ‘fast’ legs, respectively. Overall, results of the control group were similar to that previously described (Reisman et al., 2005a). While there were individual differences in the responses of the stroke survivors, the group as a whole performed similarly to healthy controls.

Reactive changes in walking
Figure 2 illustrates the walking parameters that normally change rapidly during the split-belt period, and show no after-effects. Stride lengths are plotted stride-by-stride for a typical control subject and for a stroke subject in both testing conditions (Fig. 2A). All are symmetric during the Baseline period, and rapidly change during early Adaptation so the fast stride is longer and the slow stride is shorter. These alterations were maintained throughout Adaptation, and switched back to baseline levels immediately during the Post-adaptation period. Group data for stride length showed a main effect of testing period ($P<0.0001$), but no group differences or group by testing period interactions (Fig. 2C). All groups made comparable, rapid changes in stride symmetry when going from Baseline to Adaptation (i.e. split belt) conditions ($P<0.001$), but then immediately returned to Baseline levels of symmetry during Post-adaptation, with no difference between those two periods ($P=0.87$, stroke; $P=0.78$, control).

Figure 2B shows percent stance time for a control and for both sessions from a stroke subject. Some stroke subjects showed a stance time asymmetry during the Baseline period
with a shorter stance time on the paretic leg. Just as was observed for stride length, subjects rapidly changed the time spent in the stance phase in response to changes in the treadmill speeds. During Adaptation, stance time on the fast leg quickly decreased and stance time on the slow leg quickly increased. In Post-adaptation, both legs' stance time rapidly returned to Baseline levels. Group data also show the stance time asymmetry of stroke survivors (Fig. 2D); they spent less time on the paretic leg. As such, there was a main effect of group ($P < 0.01$) in addition to the main effect of testing period ($P < 0.0001$). There was also a group $\times$ testing period interaction ($P < 0.01$), due to a reduced ability to increase the paretic stance time when it was the slow leg during Adaptation (Fig. 2D, open triangles). However, the basic reactive pattern still held in the Adaptation and Post-adaptation periods for all groups. There was a significant difference between Baseline and early Adaptation periods ($P < 0.001$), but only slight, non-significant differences between the Baseline and early Post-adaptation periods ($P = 0.98$, stroke; 0.30, control). Furthermore, there were no differences between early and late Adaptation periods ($P = 1.0$, stroke; 0.21, control). Thus, both groups rapidly changed percent stance time on each leg in response to split belts and then almost immediately returned to baseline levels when the belts were returned to the same speeds (Fig. 2D).

When comparing results from the two sessions in stroke subjects (for both stride length and stance time), the statistical analyses revealed a main effect for testing period ($P < 0.0001$ for both stride length and stance time), and for
leg tested ($P<0.01$, stance time only) but no interaction between testing period and leg tested, indicating that the stroke survivors showed similar patterns of performance for the rapidly changing parameters, regardless of which leg walked on the fast belt during the Adaptation period ($P>0.1$ for both).

In summary, during split-belt treadmill locomotion, subjects with cerebral damage were able to rapidly change their walking pattern to accommodate different belt speed relationships. This occurred regardless of whether the paretic or non-paretic leg walked on the fast belt during the Adaptation period.

**Slowly adapting changes in walking**

Figure 3 shows the walking parameters that we expected to change more slowly over the course of Adaptation: step length and double support time. Step length changes show the same general pattern in controls and people with hemiparesis, regardless of which leg is made to go faster during adaptation (Fig. 3A). Here we report the difference in step length (fast minus slow leg). In the Baseline period, the control subject showed step length differences at or near zero indicating symmetry. The stroke subject shown in this example (Fig. 3A) tended to take shorter steps with the paretic leg, which is why Baseline values are not zero. However, some stroke subjects show the opposite step length asymmetry (see ‘Changes in asymmetry’ section later). During split-belt Adaptation, the example subjects initially had asymmetric step lengths that slowly adapted towards symmetry (i.e. zero). With the return to tied belts in Post-adaptation, both subjects initially showed the reverse asymmetry (negative after-effect) which slowly returned to baseline levels. Figure 3C shows group data indicating only
a significant effect of testing period \((P<0.0001)\). Thus, control and stroke groups performed similarly over the different testing periods. Post hoc tests showed significant changes from baseline to early Adaptation, improved symmetry over the course of adaptation and significant after-effects \((Baseline \text{ versus early Post-adaptation, } P<0.001)\).

Figure 3B shows double support times, also expressed as differences (fast minus slow). Both subjects had double support time differences near zero during the Baseline period, indicating symmetry. Some stroke subjects showed double support differences that were not zero, indicating an asymmetry of the two periods of double support. When this occurred, the double support period at the end of paretic asymmetry of the two periods of double support. When this occurred, the double support period at the end of paretic leg stance was always longer than double support at the end of non-paretic stance (see ‘Changes in asymmetry’ section later). In Adaptation, the subjects initially showed substantial asymmetry that gradually was adjusted towards symmetry. In Post-adaptation, the subjects demonstrated the expected reverse asymmetry (negative after-effect), which was also gradually corrected to baseline levels. Group data showed main effects for group \((P<0.01)\) and testing period \((P<0.0001)\), but no interaction. The group effect was due to the fact that stroke survivors had asymmetries in double support time as described earlier. However, this bias did not affect the general pattern of adaptation and after-effects. Post hoc analyses showed that both groups changed double support difference from Baseline to early Adaptation \((P<0.05)\), modified it over the course of Adaptation, and showed significant after-effects \((Baseline \text{ versus early Post-adaptation, } P<0.05 \text{ stroke, } P<0.001, \text{ control})\).

### Interlimb phasing

Our previous work in healthy subjects showed that a small phase shift in limb motions can abolish asymmetries of step length and double support observed in early Adaptation \((Reisman et al., 2005a)\). Figure 4A shows the limb angles of the fast and slow legs during slow Baseline, early Adaptation and early Post-adaptation periods for a control and a stroke subject. When the interlimb phase is 0.5, the legs are moving reciprocally and the widths of the light and dark grey shaded areas are equal (top trace for both subjects). In early Adaptation, both subjects showed a phase shift whereby the fast leg was phase advanced relative to the slow leg (light < dark grey region). This phase advance is slowly corrected stride by stride (Fig. 4B). In early Post-adaptation there is a negative after-effect with the opposite phase shift (dark < light grey bars).

Interlimb phase relationship was determined using a cross-correlation function. Figure 4B depicts the lag time at the peak in the cross-correlation function calculated between the two limb angles for the same control and stroke subjects shown in Fig. 5A. Throughout the Baseline period, the limb angle interlimb phasing values for both subjects were near 0.5. In early Adaptation, both subjects initially showed a phase shift reflecting the phase advancement of the fast leg, which was gradually corrected over the course of Adaptation and showed a negative after-effect in Post-adaptation. Group results depicted in Fig. 4C show that limb phasing changed from Baseline to early Adaptation \((P<0.05)\) and the new phase relationship was stored, resulting in an after-effect in the Post-adaptation period \((compare \text{ Baseline to early Post-adaptation Fig. 5C, } P<0.01)\). Thus, individuals with stroke appear to use the same means for adapting interlimb coordination as control subjects.

### Changes in asymmetry

Some of our stroke subjects showed asymmetries in step length and/or double support times during baseline walking. Here we tested whether we could induce after-effects that altered those asymmetries. Since after-effects are assessed Post-adaptation (i.e. belts tied at the same speed), alterations in the walking pattern would be due to changes in motor commands; they would not simply be a mechanical phenomenon, as might be seen during split-belt portions of the paradigm. Figure 5A illustrates an improvement in step length asymmetry for a stroke subject who, at baseline, takes a shorter step with his paretic leg (compare open and closed circles in the three Baseline periods). By effectively increasing this asymmetry during early Adaptation period, we present a situation that drives adaptation of motor commands to ultimately reduce the asymmetry. Thus, in the Post-adaptation period we see after-effects that result in symmetrical step lengths on the two legs (compare open and closed circles in the Post-adaptation period).

Figure 5B illustrates a similar case for double support time. In the Baseline periods, this stroke subject spends longer in the double support period at the end of paretic leg stance (compare open and closed circles in the three Baseline periods). Again, this asymmetry is increased during the early Adaptation period and the subject adapts his motor commands. In the Post-adaptation period we again see that the after-effect has improved symmetry (compare open and closed circles in the Post-adaptation period). Similar effects were found in general for the subjects with hemiparesis that demonstrated a baseline asymmetry (Fig. 5D).

People with hemiparesis can show step length asymmetries in either direction \((Olney et al., 1994; Roth et al., 1997; Chen et al., 2005)\). In order to re-establish symmetric walking, the training pattern (i.e. which leg is on the fast belt during Adaptation) must therefore be based on the subject’s initial asymmetry. This is illustrated for all subjects in Fig. 5C. Subjects who take a longer paretic step during Baseline are trained with the paretic leg on the slow belt to induce an after-effect that leads to greater symmetry (grey lines, left figure). Subjects who take a shorter paretic step in Baseline, are trained with the paretic leg on the fast belt to
induce an after-effect that leads to greater symmetry (black lines, right figure). If we train in the wrong direction, we get after-effects that worsen their step length asymmetry. And, if there was no asymmetry to start with, we induce after-effects that cause asymmetry.

We tested whether the after-effects following split-belt treadmill walking could lead to improved symmetry of step length and double support for the group of subjects who demonstrated a baseline asymmetry. Table 1 shows that six subjects met our criteria for asymmetry of step length at baseline, three with a longer paretic step and three with a longer non-paretic step. Seven subjects were found to have baseline double support asymmetries; all subjects spent longer in the double support period at the end of paretic leg stance. We analysed the conditions that would theoretically induce symmetric after-effects. Results of the paired t-test reveal that there was a significant difference between the step length asymmetry in the slow Baseline period compared to the early Post-adaptation period (P = 0.01), with subjects becoming more symmetric in Post-adaptation (left side of Fig. 5D). A similar trend was found for double support, with subjects becoming more
symmetric in Post-adaptation ($P = 0.07$, right side of Fig. 5D).

**Impairment level and after-effect magnitude**

An important question is whether any of the stroke survivors’ impairments relate to their ability to adapt interlimb coordination. Our subjects had widely varying sensory and motor impairments (Table 1). We were surprised that we found no significant correlations between lower extremity impairment scores (i.e. Fugl-Meyer score, sensation, proprioception) and the magnitude of the step length after-effect, which was our measure of adaptive ability (all $P > 0.18$). After-effect magnitudes were also unrelated to subjects’ over ground fastest walking speeds ($P = 0.737$).

**Perceptual after-effect**

During Post-adaptation, all control subjects felt that the opposite leg was moving faster, compared to during Adaptation. This perceptual after-effect is the same as was observed in the previous study (Reisman et al., 2005a). In contrast, only six out of the thirteen stroke subjects demonstrated the same perceptual after-effect. The perceptual after-effect (or lack thereof) in the stroke subjects was not correlated with the magnitude of the motor after-effect ($P = 0.73$).

**Discussion**

In this study, we have demonstrated that cerebral and subcortical strokes causing a range of sensory and motor deficits did not impair a person’s ability to make immediate reactions or slower adaptations during split-belt treadmill locomotion. Importantly, we found that stroke subjects could temporarily store new interlimb relationships, demonstrating that the compromised nervous system is still capable of producing a more normal pattern. This ability was not correlated with any sensory or motor impairment measures that were collected, nor did it appear to depend on perception of the after-effects. The results of
the present study are in contrast to our previous findings in people with focal cerebellar damage, who are impaired or unable to adapt interlimb coordination, despite being able to make immediate reactive changes (Morton and Bastian, 2006).

Equally important is the finding that these adaptations can induce after-effects that temporarily restore symmetry in stroke survivors. This is a critical finding, given that many previous studies have shown that changes in coordination and symmetry during locomotor activities can be difficult to achieve, even with training (Silver et al., 2000; Kautz et al., 2005; Den Otter et al., 2006). The fact that our after-effects are short-lived is not surprising given that training was only 15 min. Yet, this demonstration suggests that the nervous system of cerebral stroke survivors is capable of flexibility which could be capitalized on with the appropriate long-term training paradigm.

**Role of supraspinal inputs in locomotor adaptation**

A previous study of decerebrate cats showed that they could adapt interlimb coordination when walking on a split-belt treadmill (Yaniagihara and Kondo, 1996). When function of the cerebellum was altered through nitric oxide deprivation, adaptation was impaired. This suggests that cerebellar, rather than cerebral, structures are more involved in this process. The circuit in the cat likely involves cerebellar influences on brainstem structures contributing to the vestibulo- and reticulo-spinal pathways. Our human work has also shown that cerebellar damage disrupts interlimb adaptation (Morton and Bastian, 2006).

But, we hypothesized that in humans, the cerebellum could be involved through its projections to cerebral motor areas, via thalamus or through its brainstem projections. In our prior study, we found that not all cerebellar lesions produced the same effect on split-belt adaptation: damage causing balance and gait impairments disrupted adaptation, whereas damage causing voluntary leg control deficits did not (Morton and Bastian, 2006). Balance and gait deficits are more linked to damage of midline cerebellar structures, which project to, and receive input from, the brainstem (Chambers and Sprague, 1955; Morton and Bastian, 2003). This coupled with the results of the current work suggest that cerebellar projections to brainstem motor areas might be more important than projections to cerebral motor areas for this adaptive process.

Although cerebral involvement in the adaptive process cannot be definitively ruled out, our results show that many types of cerebral and subcortical lesions do not impair this adaptive ability. The subjects in this current study had damage to many structures, including: frontal, parietal and temporal regions of cerebral cortex, subcortical white matter (e.g. internal capsule), the basal ganglia and the pons that resulted in a wide range of sensory and motor impairments. For example, one subject (S9) had a large parietal lesion that caused profound loss of proprioceptive sense from his paretic leg, yet he was able to adapt normally. Our original thought was that proprioceptive information from the limbs would be important for this adaptation. We surmise that adaptation still occurred since proprioceptive information reached the cerebellum through the dorsal and ventral spino-cerebellar tracts, and that conscious perception of limb position might not be required for this adaptation. Another subject (S5) had damage to the pons, which might cause an adaptation deficit given the cortico-ponto-cerebellar projection through the middle cerebellar peduncle. Yet, he adapted quite normally. One interpretation of this result is that the damage did not sufficiently affect the inferior cerebellar peduncle, which carries information to and from brainstem motor regions.

While the results of the present study suggest that cerebral structures are not critical for the split-belt locomotor adaptation studied here, there is evidence to suggest that they are important for other visually driven adjustments like obstacle avoidance. Recordings from pyramidal track and rubrospinal neurons in the cat demonstrate an increased discharge of these neurons when stepping over an obstacle (Drew, 1993; Widajewicz et al., 1994; Lavoie and Drew, 2002). These neurons appear to respond when the animal is using visual information to control the trajectory of paw and are not thought to be responding to peripheral feedback (Drew, 1993; Drew et al., 1996). In addition, induced lesions of the motor cortex in cats lead to an inability to step over obstacles attached to a moving belt (Drew et al., 1996). Finally, stroke survivors were unable to adequately modify their stepping pattern to successfully avoid obstacles during treadmill walking, leading to a much higher failure rate than healthy controls (Den Otter et al., 2005). One interpretation of all of these results is that cerebral regions are less important for peripherally driven locomotor adaptations, but critical for visually guided gait modifications, though these issues require further study.

**Locomotor flexibility post-stroke**

An important finding of this study is that stroke survivors retain sufficient adaptability of the central nervous system to alter spatiotemporal interlimb relationships. Previous research has suggested that a major deficit in motor control following stroke is the inability to flexibly produce different motor patterns in response to changing demands. In a pedaling task, Brown and colleagues found that stroke subjects could increase muscle activity in response to increased workload demand, but produced the same abnormal muscle activation pattern (Brown et al., 1996). In a reaching task, stroke survivors could not decouple shoulder abduction and elbow extension torque when reaching to different target locations (Beer et al., 2004). In contrast, our results demonstrate that stroke survivors...
can adjust interlimb control during walking in response to changing demands. This is consistent with a previous study of stroke locomotion which found that when stroke subjects walked substantially faster than their normal speed, the symmetry of double support was improved (Lamontagne and Fung, 2004). Our results further this finding by demonstrating that not only can stroke survivors flexibly adjust spatiotemporal gait parameters to changing demands, they can also store the new pattern and produce it even when the new task demands are removed.

**Locomotor adaptation and gait asymmetry post-stroke**

After only 15 min of split-belt treadmill walking, stroke subjects demonstrate after-effects in double support and step length that improve the symmetry of these variables. While this effect is short-lived, it is not trivial. Previous studies have shown that traditional locomotor training has little or no influence on interlimb coordination or symmetry post-stroke (Silver et al., 2000; Kautz et al., 2005; Den Otter et al., 2006). These results occur despite improvements in walking speed (Kautz et al., 2005; Den Otter et al., 2006) and have led to the suggestion that more targeted rehabilitation may be needed (Kautz et al., 2005). The results of novel treadmill training interventions on gait symmetry post-stroke have been mixed. Some previous studies using body-weight supported treadmill training or fast treadmill training have demonstrated improvements in symmetry post-stroke (Hesse et al., 1999; Lamontagne and Fung, 2004), while others have shown no improvement (Hesse et al., 2001). Our study demonstrates that stroke subjects retain the ability to produce a symmetric gait pattern following short-term exposure to novel interlimb coordination demands like the split-belt treadmill. The results also illustrate that, because stroke subjects show heterogeneous patterns of asymmetry, in order to re-establish symmetric walking, the training pattern (i.e. which leg is on the fast belt during Adaptation) must be based on the subject’s initial asymmetry. Whether long-term training with a split-belt treadmill followed by over ground walking to practise the symmetric pattern can lead to more permanent changes in symmetry in stroke subjects remains to be tested. However, our results support the idea that the impaired nervous system following a stroke is at least capable of the flexibility required to produce shorter-term changes in gait symmetry.

**Utilizing motor adaptations as a rehabilitation intervention**

A legitimate question is whether after-effects from adaptive training are useful for rehabilitation. Other studies have utilized after-effects of a visuomotor adaptation to improve task performance in persons post-stroke (Rossetti et al., 1998; Patton et al., 2006). Rossetti and colleagues have shown that the after-effects following prism adaptation can improve the right-deviated pointing errors normally made by stroke survivors with left hemispatial neglect (Rossetti et al., 1998). Using a robot and force field to produce disturbing forces to the hand during reaching, Patton and colleagues found that the after-effects produced following training in the force field could improve pointing errors in a group of stroke survivors (Patton et al., 2006). Just as in the present study, this group also found that the magnitude of the after-effects were not correlated with the degree of clinical impairment (Patton et al., 2006). This is potentially important as it suggests that regardless of the severity of sensorimotor deficits, subjects with cerebral damage following stroke retain the ability to make motor adaptations. Especially striking here is the lack of relationship between fast walking speed and the ability to make motor adaptations. Previous work has suggested that walking speed in stroke is correlated with level of motor impairment (Brandstater et al., 1983). However, our study reveals that neither the level of motor impairment or walking speed is correlated to the ability to make a locomotor adaptation. This makes the utility of this type of motor adaptation training more appealing, as it suggests that it may be successfully utilized across a broad spectrum of patients. However, in order to determine the role of split-belt locomotor adaptation in rehabilitation, further research is needed to investigate the effects of long-term training, and how these effects transfer to real-world tasks like walking over ground.

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**References**


