Mechanisms of motor deficits (e.g. hemiparesis and hemiplegia) secondary to stroke and traumatic brain injury remain poorly understood. In early animal studies, a unilateral lesion to the cerebellum produced postural asymmetry with ipsilateral hindlimb flexion that was retained after complete spinal cord transection. Here we demonstrate that hindlimb postural asymmetry in rats is induced by a unilateral injury of the hindlimb sensorimotor cortex, and characterize this phenomenon as a model of spinal neuroplasticity underlying asymmetric motor deficits. After cortical lesion, the asymmetry was developed due to the contralesional hindlimb flexion and persisted after decerebration and complete spinal cord transection. The asymmetry induced by the left-side brain injury was eliminated by bilateral lumbar dorsal rhizotomy, but surprisingly, the asymmetry after the right-side brain lesion was resistant to deafferentation. Pancuronium, a curare-mimetic muscle relaxant, abolished the asymmetry after the right-side lesion suggesting its dependence on the efferent drive. The contra- and ipsilesional hindlimbs displayed different musculo-articular resistance to stretch after the left but not right-side injury. The nociceptive withdrawal reflexes evoked by electrical stimulation and recorded with EMG technique were different between the left and right hindlimbs in the spinalized decerebrate rats. On this asymmetric background, a brain injury resulted in greater reflex activation on the contra- versus ipsilesional side; the difference between the limbs was higher after the right-side brain lesion. The unilateral brain injury modified expression of neuroplasticity genes analysed as readout of plastic changes, as well as robustly impaired coordination of their expression within and between the ipsi- and contralesional halves of lumbar spinal cord; the effects were more pronounced after the left side compared to the right-side injury. Our data suggest that changes in the hindlimb posture, resistance to stretch and nociceptive withdrawal reflexes are encoded by neuroplastic processes in lumbar spinal circuits induced by a unilateral brain injury. Two mechanisms, one dependent on and one independent of affrent input may mediate asymmetric hindlimb motor responses. The latter, deafferentation resistant mechanism may be based on sustained muscle contractions which often occur in patients with central lesions and which are not evoked by afferent stimulation. The unusual feature of these mechanisms is their lateralization in the spinal cord.
Keywords: brain injury; hindlimb motor deficits; lateralization; postural asymmetry; nociceptive withdrawal reflexes

Abbreviations: AIC/I = the contra/ipsilesional asymmetry index; AIL/R = the Left/Right asymmetry index; AISN = the asymmetry index in the spike number; AIThr = the asymmetry index in stimulation threshold; EDL = extensor digitorum longus; HL-PA = hindlimb postural asymmetry; HPDCI = highest posterior density continuous intervals; Int = interrossei; MPA = magnitude of postural asymmetry; NWR = nociceptive withdrawal reflex; PA = probability to develop asymmetry; PC = probability to flex the contralesional hindlimb; PL = peroneus longus; SN = the number of spikes; TBI = traumatic brain injury; Thr = stimulation threshold; UBI = a unilateral ablation brain injury; W = work

Introduction

Traumatic brain injury (TBI) and stroke cause sensorimotor impairments including muscle weakness, spasticity, and stretch and effort-unrelated sustained muscle contractions (Gracies, 2005; Murphy and Corbett, 2009; Grau, 2014; Rousseaux et al., 2014; Kerzoncuf et al., 2015; Pin-Barre and Laurin, 2015; Feldman and Levin, 2016; Jones, 2017; Marinelli et al., 2017; Pin-Barre and Laurin, 2015). In patients, stroke and TBI are mostly unilateral and often result in formation of postural asymmetry with contralateral hemiplegia and hemiparesis. Clinically well-recovered stroke survivors still have a substantial postural asymmetry in the relative contribution of each leg to body sway control (Roelofs et al., 2018). Plastic rearrangements in supraspinal and spinal neurocircuitries induced by aberrant asymmetric activity of descending neural tracts may underlie motor impairments or contribute to motor recovery. In contrast to adaptive changes in the brain, knowledge of brain injury-induced spinal plasticity is limited (Patterson, 2001; Tan et al., 2012; Wolpaw, 2012; Sist et al., 2014; Tennant, 2014; Pin-Barre and Laurin, 2015). Evidence for asymmetric ‘spinal memory’ was presented in early studies by DiGiorgio (1929, 1942) and other researchers (Chamberlain et al., 1963; Patterson, 2001). They demonstrated that a hemicerebellar lesion causes lasting changes in spinal cord functions manifested as the ipsilesional hindlimb flexion and that the hindlimb postural asymmetry (HL-PA) is retained after complete spinal transection. Consistently, monosynaptic and polysynaptic reflexes are enhanced on the ipsilateral side after lateral hemisection of the spinal cord, and this change is sustained after complete spinal transection performed caudally to the hemisection level (Hultborn and Malmsten, 1983a, b; Malmsten, 1983; Frigon et al., 2009; Rossignol and Frigon, 2011; Gossard et al., 2015). The HL-PA is an attractive animal model to unravel mechanisms of pathological ‘spinal memory’, which may underlie asymmetric motor deficits including hemiplegia and hemiparesis.

A number of clinical studies focused on changes in monosynaptic reflexes induced by stroke or TBI whereas there is a paucity of studies that analyse polysynaptic reflexes (Delwaide and Oliver, 1988; Thilmann et al., 1991; Wilson et al., 1999; Aymard et al., 2000; Condliffe et al., 2005; Dietz et al., 2009; Hubli et al., 2012). The polysynaptic nociceptive withdrawal reflexes...
(NWRs) may be useful tools for the investigation of pathological changes in the spinal cord, given that these changes are due to converging inputs from the peripheral afferents and descending motor commands to spinal neurons (Crenna and Frigo, 1984; Duysens et al., 1990; Duysens et al., 1992; Rossi and Decchi, 1994; Andersen et al., 1995; Spaich et al., 2004; Sandrini et al., 2005; Spaich et al., 2006; Emborg et al., 2009). Only a few studies have investigated changes in NWR excitability in post-stroke hemiparetic patients with limb motor deficits (Spaich et al., 2006; Serrao et al., 2012; Rueterbories et al., 2013; Spaich et al., 2014). The NWR-related EMG kinematic responses were increased and NWR modulation impaired in stroke patients with hemiparesis (Serrao et al., 2012; Alvisi et al., 2018). The NWR-based functional electrical therapy was found to be useful in the rehabilitation of gait in severely impaired hemiparetic patients by improving post-treatment walking velocity and gait symmetry (Spaich et al., 2014; Gervasio et al., 2018). There is a deficit of mechanistic animal studies that analysed pathological changes in the hindlimb NWRs and their role in sensorimotor impairments induced by brain injury.

Several neurological phenomena are differently developed after injury to the left and right hemisphere. The right side compared to the left-side stroke results in poorer postural responses in quiet and perturbed balance suggesting a prominent role of the right hemisphere in efferent control of balance (Fernandes et al., 2018). Hemineglect and bias of the subjective vertical are general causes of postural asymmetry and instability (Tasseel-Ponche et al., 2015). The serious postural impairment—the contraversive pushing called ‘Pusher syndrome’, which is linked to biased verticality and hemispatial neglect, is more common among patients with the right rather than left-hemisphere lesions (Perennou et al., 2008). Components in trunk control may be impaired depending on the side of the lesion. ‘Postural instability’ is significantly more frequent among patients with right-hemisphere lesions, while ‘apraxic responses’ predominate among those with left-hemisphere injury (Spinazzola et al., 2003). Consistently, animal studies showed that changes in locomotor behaviour and correlated region-specific differences in turnover of neurotransmitters after somatosensory TBI were dependent on the side of brain injury (Robinson, 1979; Pearlson and Robinson, 1981; Spinazzola et al., 2003; Perennou et al., 2008; Sainburg, 2014; Sainburg et al., 2016; Ocklenburg et al., 2017; Fernandes et al., 2018; Stancher et al., 2018). Right but not left somatosensory lesions produced behavioural hyperactivity and bilaterally decreased cerebral and locus ceruleus norepinephrine concentrations.

In this study, the HL-PA phenomenon was characterized as a model of asymmetric hindlimb motor deficits induced by a unilateral injury of the hindlimb representation area of the sensorimotor cortex. We examined whether a unilateral ablation brain injury (UBI) induces HL-PA; whether the fore limbs are also affected; whether the contra- or ipsilesional leg is flexed; and whether HL-PA is encoded at the spinal level and persists after complete spinal cord transection. The suction lesion was performed to restrict a damaged area to the hindlimb sensorimotor cortex with the aim to examine specific changes in hindlimb motor functions and lumbar spinal circuits. Effects of the left- and right-side lesions were studied because of lateralization of the sensorimotor system and its differential responses to the left- and right-side injuries (Robinson, 1979; Pearlson and Robinson, 1981; Spinazzola et al., 2003; Perennou et al., 2008; Sainburg, 2014; Sainburg et al., 2016; Ocklenburg et al., 2017; Fernandes et al., 2018; Stancher et al., 2018). In parallel with the HL-PA, we evaluated whether the hindlimb NWRs are affected by the UBI. We also examined whether HL-PA formation depends on sensory feedback and spinal motor output. Expression of plastic genes was analysed as a readout of neuroplastic changes in the lumbar spinal cord induced by injury of the hindlimb sensorimotor cortex. Genes selected for analysis were transcriptional regulators of synaptic plasticity (cFos, Egr1 and Nfkbia), regulators of axonal sprouting, synapse formation, neuronal survival and neuroinflammation (Arc, Bdnf, Dlg4, Homer-1, Gap43, Syt4 and Tgfβ1), essential components of the glutamate system critical for neuroplastic responses (GluR1, Grin2a and Grin2b), and all are expressed in the spinal cord.

**Materials and methods**

All methods and materials are described in detail in Supplementary material section.

Adult male Sprague Dawley rats (Taconic, Denmark) weighing 170–430 g (see Supplementary Table 1) were used in the study. The animals received food and water ad libitum, and were kept in a 12-h day–night cycle at a constant environmental temperature of 21°C (humidity: 65%) and randomly assigned to their respective experimental groups. Approval for animal experiments was obtained from the Malmo/Lund ethics committee on animal experiments (No.: M7-16), and the ethics committee of the Faculty of Medicine of Porto University and Portuguese Direção-Geral de Alimentação e Veterinária (No. 0421/000/000/2018). All animals were analysed in a single test excluding 11 right-side sham and 14 right-side UBI rats analysed in both HL-PA and EMG experiments (Supplementary Table 1).

**Brain surgery and histology**

The rats were anaesthetized with 3% isoflurane (Abbott, Norway) in a mixture with 65% nitrous oxide and 35% oxygen. The hindlimb sensorimotor cortex was removed by gentle suction through a pipette (tip diameter 0.5 mm) (Fig. 1) (for details, see ‘Supplementary materials’). Control (sham) animals underwent the same procedure but cortex was not aspirated.

The localization and size of the lesion were analysed after perfusion with 4% paraformaldehyde. Following an
overnight post-fixation in the same fixative, the brain tissue was soaked in phosphate-buffered saline with 30% sucrose for 48 h, dissected into blocks which were then sliced into 50-μm sections with a freezing microtome. Every fourth Nissl-stained section across the lesion site was photographed and the rostrocaudal respective mediolateral extension as well as lesion volume were calculated.

Analysis of HL-PA

The magnitude of postural asymmetry (MPA) and the side of the flexed limb were assessed as described previously (Bakalkin and Kobylyansky, 1989). Briefly, the measurements were performed under isoflurane anaesthesia, or in unanaesthetized decerebrate spinalized rats. The rat was placed in ventral decubitus (prone) position on a millimetre-grid paper sheet. To minimize effects of tactile stimulation during the analysis three threads were glued to the nails of the middle three toes of each limb. The fore or hindlimbs were gently pulled by the threads for 5–10 mm to reach the same level, then set free and the MPA was measured in millimetres as the length of the projection of the line connecting symmetric limb points (digits 2–4) on the longitudinal axis of the rat (Fig. 2A). The procedure was repeated six times in immediate succession, and the mean HL-PA value for a given rat was calculated and used in statistical analyses. For analysis in the supine position, the rat was placed in a V-shaped trough, a 90°—angled frame located on a levelled table surface with the 1-mm grid sheet; otherwise, the procedure was the same as for the prone position. The threshold of 2 mm that was the 94th MPA percentile in the sham rats was applied to define the asymmetric rats. A limb displaying shorter projection was regarded as flexed.

Analysis of hindlimb resistance to stretch

Stretching force was analysed after spinal cord transection using the micromanipulator-controlled force meter device constructed in the laboratory. The resistance was measured as the amount of mechanical work $W_{\text{Left}}$ and $W_{\text{Right}}$ to stretch the left- and right hindlimbs, where $W$ was stretching force integrated over stretching distance interval from 0 to 10 mm.

EMG experiments

The EMG activity of the extensor digitorum longus (EDL), interrossei (Int) and peroneus longus (PL) muscles from both hindlimbs were recorded as described previously (Schouenborg et al., 1992; Weng and Schouenborg, 1996).

Analysis of gene expression

The rats were exposed to the left- or right-side UBI or the left- or right-side sham surgery (four groups; $n = 10/
group). The rats were sacrificed by decapitation on Day 3 after the injury, the lumbar spinal cords were dissected into the left and right halves. RNA purification, quality evaluation, cDNA synthesis and quantitative RT-PCR were described elsewhere (Kononenko et al., 2018).

mRNA levels of 13 neuroplastic genes (Supplementary Table 2) were normalized to geometric mean of expression levels of two control genes Actb and Gapdh.

Statistical analysis

Postural asymmetry, stretching force and EMG data were analysed using Bayesian methods. Predictors and outcomes were centred and scaled before we fitted Bayesian regression models via full Bayesian framework by calling Stan 2.19.2 (Carpenter et al., 2017) from R 3.6.1 R Core Team (2018) using the brms 2.10.0 (Bürkner, 2017) interface. Models used Student’s t response distribution with identity link function unless and had no intercepts with indexing approach to predictors (McElreath, 2020). Default priors were provided by the brms according to Stan recommendations (Gelman, 2019). Intercepts, residual SD and group-level SD were determined from the weakly informative prior student_t(3, 0, 10). P-values, adjusted using the multivariate t distribution with the same covariance structure as the estimates, were produced by frequentist summary in emmeans 1.4 (Searle et al., 1980) together with the medians of the posterior.
distribution and 95% highest posterior density continuous intervals (HPDCI). The asymmetry and contrast between groups were defined as significant if both 95% HPDCI did not include zero and adjusted P-value was ≤0.05.

Differences in gene expression were assessed by two-way ANOVA (normally distributed data) or Kruskal–Wallis test of ranks (data with non-normal distribution). The Bonferroni procedure was used for multiple testing adjustments. Post hoc analysis was performed by Tukey’s HSD test (normally distributed data).

For analysis of intra- and inter-regional gene co-expression patterns, Spearman’s rank correlation coefficient was calculated for all gene pairs in each area (N = 78) and between the areas (N = 169). Sample size for each correlation coefficient was n = 10 animals. Statistical comparison of gene–gene coordination strength between the animal groups was performed using Wilcoxon signed-rank test to the set of absolute values of all correlation coefficients. Differences in the proportion of positive and negative correlations between animal groups were assessed using the Fisher’s exact test with 2×2 contingency table.

Details are shown in Supplementary materials.

Data availability

Data supporting the findings of this study are available within the article, its Supplementary material or upon request.

Results

Formation of postural asymmetry in UBI rats

In the first set of experiments, we examined whether the UBI induces formation of forelimb or HL-PA, whether the ipsi- or contralesional limb is flexed, and whether the asymmetry is retained after complete spinal cord transection and decerebration (Figs 1 and 2; Supplementary Fig. 1; Supplementary Tables 1, 3 and 4). The lesion produced by unilateral ablation by suction was confined to the hindlimb motor area (Fig. 1; Supplementary methods). The MPA, the probability to develop asymmetry (P_A), and the probability to flex the contralesional hindlimb (P_C) in the prone and supine positions were analysed 1–14 days after UBI or sham surgery. The UBI-induced asymmetry in hindlimb posture (HL-PA) was evident in anaesthetized animals before spinalization, and in unanaesthetized spinalized at high thoracic level and decerebrate rats (Fig. 2C). The hindlimb MPA was substantially (i.e. ~3-fold) higher in the UBI compared to sham-operated rats analysed before and after spinal cord transection and decerebration in the prone position; and before the transection and decerebration in the supine position. In the UBI group, correlations were significant between MPA analysed (i) before and after spinalization and decerebration in prone position; and (ii) in the prone and supine positions before these operations (Supplementary Table 4). The MPA in the UBI group was stable during the observation period from the 1st to 19th day after the injury; a day of measurement included as confounder in statistical models showed no time effects. No apparent forelimb asymmetry was developed.

The rats showing a MPA >2 mm were regarded as asymmetric; the 2 mm threshold corresponded to the 94th percentile of hindlimb MPA in the sham surgery group (Fig. 2; Supplementary Fig. 1). The P_A was significantly higher in the UBI rats compared to the sham-operated animals when they were analysed before and after spinal cord transection and decerebration in the prone position; and before these operations in the supine position (Fig. 2D). The P_C for asymmetric UBI rats was significantly greater than 50% contra-/50% ipsilesional flexion ratio for the prone position before and after transection and decerebration; and for the supine position before these operations (Fig. 2E). Figure 2 shows data for the right-side UBI; the left-side UBI similarly produced HL-PA with contralesional hindlimb flexion (MPA, P_A and P_C did not differ between the left- and right-side UBI groups: n = 10 and 16, respectively; P > 0.4 for rat groups analysed either in the prone or supine positions, and either before or after spinalization). Thus the unilateral focal injury to the hindlimb representation area of the sensorimotor cortex resulted in formation of the HL-PA with contralesional flexion whereas forelimbs were not apparently affected. The asymmetry persisted after spinalization and decerebration suggesting that the UBI-induced spinal neuroplastic response underlies HL-PA maintenance.

Effects of deafferentation and neuromuscular signalling blockade on HL-PA

We next sought to determine whether afferent somatosensory input is required for the HL-PA formation in the UBI rats. The HL-PA was analysed before and after bilateral rhizotomy from the L1 to S2 spinal levels in the spinalized rats received earlier either the left- or right-side UBI. After the left-side UBI, the MPA and P_A were strongly, 9.0- and 3.6-fold, respectively, reduced after rhizotomy (Fig. 3A–C; Supplementary Figs 2 and 3). In contrast, in the right-side UBI rats the MPA and P_A remained stable after rhizotomy and were substantially, 8.0- and 2.6-fold, respectively, higher than those in the left-side UBI group. These data suggest that the HL-PA maintenance in spinalized rats may depend on (i) afferent somatosensory input after the left-side UBI, and on (ii) activity of motoneurons signalling to hindlimb muscles or changes in neuromuscular system after the right-side injury.
To examine whether the HL-PA induced by the right-side UBI is maintained by central or peripheral (neuromuscular) mechanism, pancuronium (pavulon), a curare-mimetic muscle relaxant was administered in rats displaying HL-PA to block the neuromuscular junction (Fig. 3D–F; Supplementary Fig. 4). Pancuronium infusion in the anaesthetized rats that received the right-side UBI 3 days before the experiment, resulted in rapid decrease of the MPA and then completely abolished HL-PA. The MPA was stable in UBI rats given saline for at least 120 min ($n = 9$).

**Stretching resistance analysis**

We next studied whether the UBI may differently affect biomechanical properties of the contra- and ipsilesional hindlimbs by analysing the passive hindlimb musculo-articular resistance to stretching in the anaesthetized spinalized rats using the micromanipulator-controlled force meter device (Fig. 4). The asymmetry in the resistance was assessed using (i) the left/right asymmetry index for the work ($W$), $AI_W = \log_2 (W_{\text{Left}}/W_{\text{Right}})$, where $W_{\text{Left}}$ and $W_{\text{Right}}$ were the $W$ applied to stretch the left- and right hindlimbs (Fig. 4D and E); and (ii) the index for difference in the work between the left- and right hindlimbs $\Delta W = (W_{\text{Left}} - W_{\text{Right}})$ (Fig. 4F and G). The indexes were integrated for a 10-mm stretching distance. Both the $AI_W$ and $\Delta W$ were statistically analysed because they may depend differently on the resistance force and the distance over which they are stretched.

The right (contralesional) versus left (ipsilesional) hindlimb demonstrated greater resistance after the left-side
UBI (Fig. 4D and F). The produced changes were significant in both the AIW [median of the posterior distribution as estimate (median) = −0.611, 95% HPD CI = (−1.023, −0.216), adjusted P-value (P) = 0.008, the Left to Right ratio in resistance or L/R ratio was 1.85-fold larger for right-side UBI] (Fig. 4E) and the ΔW (ΔAIW) (E) and ΔW (ΔΔW) (G) were assessed in Bayesian framework and are reported as significant if their 95% HPD CI did not include zero and adjusted P-values were ≤0.05 (shown by *).

UBI effects on hindlimb NWRs

 Along with changes in posture, the UBI may differentially affect the NWRs of the contra- and ipsilesional hindlimbs producing an asymmetric pattern. We addressed this
question by analysis of the hindlimb NWRs evoked by electrical stimulation of symmetric digit spots on left and right hind paws, and recorded using EMG techniques in unanaesthetized UBI rats after their decerebration and complete spinal cord transection (Figs 5 and 6; Supplementary Tables 1 and 5). EMG responses were recorded from the EDL, Int and PL muscles of the contra- and ipsilesional hindlimbs in the rats exposed to the left- or right-side UBI, and the left- or right-side sham surgery (Figs 5 and 6). EMG activity was recorded in response to 18 stimulations (Fig. 5A); most responses were stable across the stimulation train, however, wind-up was sporadically observed. Peristimulus histograms denote that a main fraction of the responses was recorded in the 0.2–1.0 s interval. This fraction was included in statistical analysis (Fig. 5B) and was likely elicited through activation of the C-fibres.

Stimulation thresholds (Thr) and the number of spikes (SN) of EMG responses to current stimulation were compared between the left and right limbs using the Left/Right asymmetry index \[ AI = \log_2(L/R) \], were L and R were values for the left- and right-side muscles] for the threshold (\( A_{\text{Thr}} \)) or the spike number (\( A_{\text{SN}} \)) (Fig. 6D–G; the number of analysed animals is shown in Supplementary Table 5). No asymmetry in the sham surgery and UBI groups, and no significant differences between the groups were revealed for the threshold values (Fig. 6D and F). Analysis of the sham surgery groups revealed significant asymmetries of EMG responses in the number of spikes for the EDL in the right-side sham surgery group [median = −1.088, 95% HPDCI = (−1.898, −0.265), \( P = 0.034 \), L/R ratio = 0.47] and for the Int in the left-side sham group [median = −1.914, HPDCI = (−3.070, −0.725), \( P = 0.005 \), L/R ratio = 0.27] (Fig. 6E). The median \( A_{\text{SN}} \) of each of these muscles in each the left and right sham group was less than zero. Analysis of the combined sham surgery group demonstrated that the number of spikes was higher on
the right compared to the left side [EDL: median = −1.049, 95% HPD CI = (−1.693, −0.418), P = 0.007; Int: median = −0.923, 95% HPD CI = (−1.581, −0.236), P = 0.040)].

After the brain injury, significant asymmetry in the number of evoked spikes was identified for the EDL muscle in the right-side UBI rats [median = 1.147, HPD CI = (0.356, 1.910), P = 0.015, L/R ratio = 2.2].
Responses were revealed after the left-side UBI. Effects of professional versus ipsilesional side. No significant asymmetric expression of 13 neuroplasticity genes that were assessed whether the left- and right-side UBI affects expression in the lumbar spinal cord. Differences between the right-side UBI and sham surgery groups were analysed. Two-way ANOVA with operation type (UBI versus sham) and measurement side (left versus right) as factors revealed significant differences between UBI and sham surgery groups for Grin2a (P adjusted by Bonferroni correction, adj. \( P = 0.0004 \)), Tgfb1 (adj. \( P = 0.0025 \)) and Dlg4 (adj. \( P = 0.019 \)), and between measurement sides for Grin2b (adj. \( P = 0.037 \)). The left- and/or right-side UBI resulted in a downregulation of Grin2a (post hoc test: left-side UBI, \( P = 0.0007 \); right-side UBI, \( P = 0.028 \)) and Dlg4 (left-side UBI, \( P = 0.0007 \)), and upregulation of Tgfb1 (left-side UBI, \( P = 0.002 \); right-side UBI, \( P = 0.025 \)) (Fig. 7A–C). Grin2b demonstrated higher expression in the left half compared to the right half of the spinal cord (post hoc test: \( P = 0.0029 \)) (Fig. 7D) that was replicated by analysis of intact animals (\( n = 10 \); paired t-test: \( P = 0.0001 \)) (Fig. 7E).

Gene co-expression patterns characterize regulatory interactions within and across tissues (or CNS areas) (Dobrin et al., 2009; Long et al., 2016). We examined whether UBI affects gene–gene (i) inter-area correlations between the left and right halves of the lumbar spinal cord, and (ii) intra-area correlations in these halves. The inter-area positive correlations dominated in both the left and right sham surgery groups, while their proportion was reduced after the UBI; the effect was statistically significant after the left (\( P = 5.8 \times 10^{-7} \)) (Fig. 7F and H), but not right-side brain injury (\( P = 0.081 \)) (Fig. 7G and I). Proportion of the intra-area positive correlations was decreased after the UBI in the left (left-side injury: \( P = 7.7 \times 10^{-4} \); right-side injury: \( 2.0 \times 10^{-3} \)) and right (left-side injury: \( P = 1.4 \times 10^{-4} \); right-side injury: \( 4.1 \times 10^{-4} \)) halves of lumbar spinal cord (Supplementary Fig. 5). Furthermore, the coordination strength of gene co-expression patterns was significantly decreased after the left UBI in the contralesional (right side) lumbar domain (\( P = 1.8 \times 10^{-11} \)) but not in the ipsilesional (left side) domain (Supplementary Fig. 5); effects of the right-side UBI were not significant.

Thus the UBI induced changes in expression of neuroplasticity genes and impaired coordination of their expression between and within the ipsi- and contralesional halves of lumbar spinal segments; the effects were more pronounced after the left-side compared to the right-side brain injury.

**Discussion**

The principal findings of this study are the formation of the HL-PA with flexion on the contralesional side, and the asymmetric changes in hindlimb NWRs and in resistance to stretching, all induced by the unilateral focal injury to the hindlimb representation area of sensorimotor cortex. These effects were evident in the rats spinalized at the high thoracic level suggesting the brain injury-induced rearrangement of lumbar spinal circuits prior to the transection. This notion is supported by the UBI-induced changes in expression and co-expression patterns of neuroplasticity genes in the lumbar spinal cord. Peripheral biomechanical mechanisms may likely be excluded because HL-PA was blocked by pancuronium. Formation of the contralesional flexion along with higher resistance to stretching and activation of NWRs on the contralesional

(Fig. 6E). Representative examples of the UBI-induced asymmetry for the EDL muscle is shown on Fig. 6B and C. The UBI and sham surgery groups were significantly different in the AI5N the EDL after the right-side injury [median = 2.238, HPDCl = (1.111, 3.354), \( P = 0.001 \), 4.7-fold difference in the L/R ratio] (Fig. 6G). No significant differences were revealed after the left-side UBI.

We next assessed whether the magnitude of the injury-induced changes in the spike number asymmetry significantly differs between the left- and right-side UBI using the contra/ipsilesional asymmetry index \( AI_{C/I} = [AI_{C/I} = \log_{2}(C/I)] \), where C and I were the number of spikes of the contra and ipsilesional muscles. Differences between effects of the right-side UBI \( [\Delta AI_{C/I,R-UBI} = AI_{C/I,R-UBI} - AI_{C/I,R-sham}] \) and left-side UBI \( [\Delta AI_{C/I,L-UBI} = AI_{C/I,L-UBI} - AI_{C/I,L-sham}] \) were computed as \( \Delta AI_{C/I} = (\Delta AI_{C/I,R} - \Delta AI_{C/I,L}) \) (Fig. 6H). The Bayesian analysis demonstrated that the ratio of the \( AI_{C/I} \) for the right-side UBI to the \( AI_{C/I} \) for the left-side UBI is significantly greater compared to the left \( AI_{C/I} \) for the right-side UBI to the \( AI_{C/I} \) for the left-side UBI for the contralateral hindlimb muscles only. Other muscles were not significant asymmetry differences were revealed after the left-side UBI.

Effects of the left-side and right-side UBI on the EDL muscle were significantly different in the \( AI_{C/I} \) magnitude.

**UBI effects on expression of neuroplasticity genes, and their intra- and inter-regional co-expression patterns in lumbar spinal cord**

The HL-PA and asymmetry in the NWRs may persist after spinalization and decerebration due to the UBI-induced neuroplastic response in the lumbar spinal domains underlying by changes in gene expression. We assessed whether the left- and right-side UBI affects expression of 13 neuroplasticity genes that were Arc, Bdnf, cFos, Dlg4, Egr1, Homer1, Gap43, GluR1, Grin2a, Grin2b, Nfkbia, Syt4 and Tgfb1 in the left and right halves of the lumbar spinal cord analysed as a readout of neuroplastic response. Four groups including the left- and right-side UBI, and the left- and right-side sham surgery groups were analysed. Two-way ANOVA with operation type (UBI versus sham) and measurement side (left versus right) as factors revealed significant differences between UBI and sham surgery groups for Grin2a (P adjusted by Bonferroni correction, adj. \( P = 0.0004 \)), Tgfb1 (adj. \( P = 0.0023 \)) and Dlg4 (adj. \( P = 0.019 \)), and between...
The ipsilateral hindlimb flexion was induced by hemispheric lesion (DiGiorgio, 1929, 1942; Alella, 1948; Chamberlain et al., 1963; Patterson, 2001), or hemilabyrinthectomy (Straka and Dieringer, 1995). In contrast, contralateral hindlimb flexion was developed after a focal injury of the hindlimb representation area of the sensorimotor cortex (although a small part of the hindlimb somatosensory cortex might have been aspirated in some rats as also was described elsewhere (Tandon et al., 2007)) or large injury of one hemisphere ((Varlinskaia et al., 1984; Vartanyan and Klementiev, 1991) and the present work). Search for neurotransmitter system involved in HL-PA formation demonstrated that opioid peptides and synthetic opioids (Bakalkin et al., 1980; Chazov et al., 1981; Bakalkin et al., 1986; Bakalkin and Kobylyansky, 1989) along with Arg-vasopressin (Klement’ev et al., 1986; Lukoyanov et al., 2018) may induce HL-PA in spinalized animals. The striking finding was that the response to administration of these substances was developed either on the left or right side of the body. The κ-opioid agonists dynorphin and bremazocine induced flexion of the left hindlimb, while δ-agonist Leu-enkephalin and Arg-vasopressin caused the right hindlimb to flex. Release of the endogenous opioid peptides in rats exposed to stress or pain stimuli also produced HL-PA, which was blocked by administration of the opioid antagonist naloxone (Bakalkin et al., 1982). These side-specific neurohormonal mechanisms may potentially mediate formation of the left or right hindlimb flexion after UBI.

The right-side UBI-induced HL-PA was resistant to bilateral deafferentation suggesting that spinal reflexes do not contribute to asymmetry formation. In contrast, bilateral lumbar rhizotomy after the left-side UBI abolished HL-PA that thereby may be developed due to differences between reflexes on the contra- and ipsilesional sides. In the course of the asymmetry analysis, no nociceptive stimulation was applied and tactile stimulation was negligible; the legs were stretched by pulling the threads glued to nails of the toes. The HL-PA magnitude was the same and correlated between the supine and prone positions for which effects of tactile stimulation, if any, should be different. Application of lidocaine on the toes, which were pulled at the stretching, did not affect the asymmetry formation (unpublished data). It was firmly established that the stretch and postural limb reflexes are abolished immediately and for days after complete spinal cord transection (Miller et al., 1996; Musienko et al., 2010; Frigon et al., 2011) and substantially decreased under anaesthesia (Zhou et al., 1998; Fuchigami et al., 2011). Therefore, the NWRs and stretch reflex likely do not contribute to HL-PA formation and/or its...

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**Figure 7** Expression of neuroplasticity genes (A–C), lateralized expression (D, E), and inter-area co-expression patterns (F–I) in the lumbar spinal cord of the rats exposed to the left- and right-side UBI or sham surgery. Four groups of rats (n = 10/group) were sacrificed on Day 3 after the injury for tissue dissection. Shown are log2-transformed levels of Grin2a (A), Tgfb1 (B), Dlg4 (C) and Grin2b (D, E) mRNAs in the lumbar spinal cord. In (A–C), L- and R- stand for the left- and right-side UBI or sham surgery. In (D and E), Left and Right denote the left and right spinal half. In (D), the data combined for two UBI and two sham surgery groups (n = 40) and post hoc P are shown; these four groups did not differ between each other in the Grin2b expression in either the left or right domain; therefore, the data were pooled for statistical analysis. (E) Data for intact rats (n = 10) and P-value for left–right difference (paired t-test). The horizontal line in the box represents the median; the box hinges represent the first (Q1) and third quartiles (Q3). Upper and lower whiskers extend from the hinge to the highest/lowest value that lies within the 1.53 interquartile range (IQR) of the hinge. Adjusted P-values are shown. (F–I) Heatmaps with Spearman coefficients for correlations in expression levels between the left and right lumbar domains for all gene pairs in the left- (F) and right- (G) side sham operation, and the left- (H) and right- (I) side UBI groups.
maintenance in the preparations of the unanaesthetized decerebrate spinalized UBI rats, and spinalized UBI rats under anaesthesia analysed in the present study. On the other hand, responses mediated by the Group II muscle afferents may remain active after spinalization in acute experiments (Jankowska, 1992; Valero-Cabre et al., 2004; Lavrov et al., 2015). Together with these observations, our findings suggest that the HL-PA is a complex phenomenon that is developed either due to (i) a persistent asymmetric activity of lumbar motoneurons not stimulated by afferent input; or due to (ii) discharge of proprioceptive neurons activated perhaps by Group II muscle afferents, which are tonically active and maintain muscle tone. These two mechanisms may differentially contribute to formation of the left and right hindlimb flexion. The second mechanism is supported by human studies demonstrated that the Group II muscle afferents are involved in the exaggerated reflexes in spastic hemiplegic and paraplegic patients (Eriksson et al., 1996; Marque et al., 2001).

Our data indicate that NWRs of EDL and Int muscles are lateralized in control animals. On this left-right asymmetric background, the left- and right-side brain lesions were not mirror symmetric in their effects on the hindlimb NWRs. After the right-side UBI, sign of the AI SN for the EDL was changed from minus to plus, but the NWR was still significantly asymmetric—different from zero. Thus, the right-side UBI resulted in greater activation of withdrawal EDL responses on the left (contralateral) side compared to the right (ipsilateral) side. The left-side UBI did not produce significant changes in the EDL, Int and PL NWRs. Muscles which may be affected by the left-side injury remain to be identified. The NWRs have a modular organization in which each module controls a single or a few synergistic muscles and each module performs a comprehensive and functionally adapted sensorimotor transformation resulting in appropriate activation of the target muscle(s) (Schouenborg, 2002). Effects of the left- and right-side UBI may be interpreted in the context of this mechanism; different sets of NWR modules may be targeted by the left- and right-side brain injury.

Dependence of the right hindlimb flexion formation on afferent stimulation corroborates asymmetry of mono- and polysynaptic segmental reflexes in intact rats and cats (Hultborn and Malmsten, 1983a, b; Malmsten, 1983) and asymmetry of NWRs in control rats, which all display greater activity on the right body side. The uncovered lateralization in the spinal cord may be relevant as the physiological basis for several neurological phenomena differently developed after injury to the left and right hemisphere (see Introduction section; Spinazzola et al., 2003; Fernandes et al., 2018). The hemisphere specific motor effects laid the basis for the general concept of motor lateralization in which the left hemisphere assures mechanical efficiency under predictable conditions, while the right hemisphere imparts robustness under unpredictable situations (Sainburg, 2014). Our results are in line with the motor lateralization concept. They suggest that the left hindlimb motor cortex might enable tonic, afferent stimulation independent control of the right hindlimb postural responses while its lesion eliminates this function. Complementarily, the right cortex might control the afferent-dependent postural reactions of the left hindlimb, and its injury could diminish segmental reflex-mediated motor responses.

Besides left-right differences in segmental reflexes (Hultborn and Malmsten, 1983a, b; Malmsten, 1983; and this study), physiological, anatomical and molecular asymmetries in the spinal cord (Bakalkin et al., 1980; Chazov et al., 1981; Bakalkin et al., 1986; Bakalkin and Kobylansky, 1989; Nathan et al., 1990; de Kovel et al., 2017; Kononenko et al., 2017; Ocklenburg et al., 2017; Kononenko et al., 2018; Woytowicz et al., 2018) along with general lateralization of motor functions (Sainburg, 2014; Sainburg et al., 2016; Knebel et al., 2018; Stancher et al., 2018) have been reported. Three-quarters of cervical spinal cords are asymmetric with larger right side (Nathan et al., 1990). Spinal neural circuits controlling activity of the left and right hindlimb muscles may be differentially affected by opioid agonists (Bakalkin et al., 1980; Chazov et al., 1981; Bakalkin et al., 1986; Bakalkin and Kobylansky, 1989). Expression of opioid receptors is lateralized in the spinal cord, while their inter-regional co-expression patterns are side-specific and intra-regional (between the dorsal and ventral domains) co-expression profiles are differently affected by the left- and right-side body injury (Kononenko et al., 2017). In human embryos transcriptional maturation rates are different between left and right spinal cord suggesting that spinal cord lateralization is not directed from the forebrain (de Kovel et al., 2017). The present study demonstrates that the Grin2b gene coding for the NMDA receptor subunit, which is involved in circuit formation and synaptic plasticity, is expressed at higher levels in the left compared to the right lumbar spinal cord. Interestingly, the lateralized expression of this gene underlies structural and functional left–right asymmetries in the hippocampus (Kawakami et al., 2003; Wu et al., 2005; Kohl et al., 2011).

Expression of the Grin2a, Dlg4 and Tgfb1 neuroplasticity genes in the lumbar spinal segments was altered after the unilateral brain lesion. The Grin2a subunit of the glutamate receptors regulates formation of neural circuits and their plasticity. The Dlg4 gene codes for PSD95 involved in AMPA receptor-mediated synaptic plasticity and post-NMDA receptor activation events (Won et al., 2016). Tgfb1 gives rise to transforming growth factor β1 regulating inflammation, expression of neuropeptides and glutamate neurotoxicity (Buisson et al., 2003; Santibanez et al., 2011). The glutamate receptors may modulate spinal NWRs (You et al., 2004) and flexion response in a model of spinal learning (Joynes et al., 2004), and mediate HL-PA formation induced by electrical limb
stimulation (Anderson and Winterson, 1995). These findings suggest that glutamate receptors may have a role in the UBI-induced formation HL-PA and modifications of hindlimb NWRs.

Remarkably the co-expression patterns of neuroplasticity genes were strongly dysregulated after the UBI as follows from the decline in the coordination strength and decrease in the proportion of positive inter- and intra-area correlations. Robust changes in the inter- and intra-regional gene co-expression patterns is a novel phenomenon suggesting that the UBI abolishes coordination of gene expression between the left and right halves of anatomically symmetric spinal cord, and within each of these halves.

Ten neuroplastic genes which individual expression levels were not significantly affected by the UBI were nevertheless dysregulated as the components of the inter- and intra-area co-expression networks. These genes were a neuronal activity-dependent transcription factor cFos; Gap-43 coding for growth-associated protein Gap-43 that regulates axonal growth and neural network formation; brain-derived neurotrophic factor Bdnf regulating synaptogenesis; activity-regulated cytoskeletal Arc gene implicated in numerous plasticity paradigms; Egr1 regulating transcription of growth factors, DNA damage and ischaemia genes; Nfkbia (I-Kappa-B-Alpha) that inhibits NF-kappa-B/REL complexes regulating activity-dependent inhibitory and excitatory neuronal function; GluR1 and Grin2b coding for the glutamate ionotropic receptor AMPA type subunit 1 and NMDA receptor subunit, respectively, both involved in glutamate signalling and synaptic plasticity; Homer-1 giving rise to Homer Scaffold Protein 1, a component of glutamate signalling involved in nociceptive plasticity; and Syt4 (Synaptotagmin 4) playing a role in dendrite formation and synaptic growth and plasticity (Hayashi et al., 2000; Adkins et al., 2006; O’Mahony et al., 2006; Tappe et al., 2006; Vavrek et al., 2006; Larsson and Bromman, 2008; Dolan et al., 2011; Grasselli and Strata, 2013; Harris et al., 2016; Epstein and Finkbeiner, 2018).

Changes in the co-expression patterns were likely mediated through the epigenetic and transcriptional mechanisms affected by the UBI, and therefore characterization of the co-regulation phenomenon does not require analysis of protein products of analysed genes. We did not study UBI effects on the levels of neuroplastic proteins. However, the identified increase in the Tgfβ1 mRNA levels corroborated with elevation in the levels of Tgfβ1 protein in the spinal lumbar segments after the TBI, and elevation of neurotrophins and inflammatory cytokine expression in the spinal cord after cortical stroke (Sist et al., 2014; Tennant, 2014; Kononenko et al., 2019). It is interesting to examine a role of protein products of three neuroplastic genes affected by the UBI in the brain lesion-induced changes in hindlimb functions.

Clinical studies demonstrated that a large fraction of subjects with stroke and cerebral palsy do not relax their muscles—they are tonically constricted without any voluntary command. This phenomenon is called spastic dystonia and defined as ‘stretch- and effort-unrelated sustained involuntary muscle activity following central motor lesions’ (Gracies, 2005; Lorentzen et al., 2018). This clinical phenomenon can alter posture at rest thereby contributing to the hemiplegia (Lorentzen et al., 2017) and is regarded as a form of efferent muscle hyperactivity (Gracies, 2005; Baude et al., 2019). Spastic dystonia may have a central mechanism which does not depend on afferent input in contrast to spasticity based on exacerbated reflex excitability (Sheean and McGuire, 2009). It was postulated that rhizotomy which abolished sensory input and diminish stretch reflexes would have no effect on spastic dystonia. The example is the absence of effects of dorsal rhizotomy on development of muscle contractures in children with cerebral palsy (Tedorff et al., 2015). A cause of spastic dystonia was experimentally addressed in early studies by Denny-Brown (1966, 1980) who demonstrated that monkeys with lesions of motor cortex display involuntary muscle activity. It persisted following disruption of sensory input to the spinal cord suggesting that a central mechanism but not exaggerated reflexes are involved. Sustained involuntary muscle activity developed in the absence of stretch and voluntary stimulation contributes to body deformities in stroke patients, and thereby it is critical to develop experimental approach for preclinical analysis of spastic dystonia (Gracies, 2005; Marinelli et al., 2017; Pingel et al., 2017; Lorentzen et al., 2018; Baude et al., 2019). The HL-PA that is induced by lesion to the hindlimb sensorimotor cortex and that is not affected by rhizotomy may represent a small animal model for investigation of this phenomenon.

In conclusion, the HL-PA resistance to deafferentation in the spinalized rats may be developed due to the UBI-induced neuroplastic changes in the efferent hindlimb neuromuscular circuits. This finding is concordant with the pathological non-stretch and non-effort-related sustained involuntary muscle activity observed after central motor lesions in clinical situations (Gracies, 2005; Sheean and McGuire, 2009; Tedroff et al., 2015; Marinelli et al., 2017; Lorentzen et al., 2018; Baude et al., 2019). The unusual feature of this phenomenon is its association with the right-side brain lesion. The lateralization of mono- and polysynaptic reflexes (Hultborn and Malmsten, 1983a, b; Malmsten, 1983) including NWRs and the asymmetry in stretching force support hypothesis on asymmetric organization of the spinal cord. These findings represent a subject for further analyses essential for understanding of motor deficits secondary to stroke and TBI, and basic mechanisms of sensorimotor integration in the spinal cord.
Supplementary material

Supplementary material is available at Brain Communications online.

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Competing interests

The authors report no competing interests.

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