Evidence of Neuroplasticity With Robotic Hand Exoskeleton Study for Post-Stroke Rehabilitation: A Randomized Controlled Trial

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Research

Keywords: Stroke, Neurological rehabilitation, Wrist, Metacarpophalangeal joint, Robotic exoskeleton, Transcranial Magnetic Stimulation, cortical-excitability

Posted Date: November 11th, 2020

DOI: https://doi.org/10.21203/rs.3.rs-67841/v2

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Version of Record: A version of this preprint was published at Journal of NeuroEngineering and Rehabilitation on May 6th, 2021. See the published version at https://doi.org/10.1186/s12984-021-00867-7.
Abstract

Background: A novel electromechanical robotic-exoskeleton was designed in-house for rehabilitation of wrist joint and Metacarpophalangeal (MCP) joint.

Objective: The objective was to compare the rehabilitation effectiveness (clinical-scales and neurophysiological-measures) of robotic-therapy training-sessions with dose-matched control in patients with stroke.

Methods: A pilot prospective parallel single-blinded randomized controlled study at clinical-settings was designed with patients with stroke within 2 years of chronicity. Patients were randomly assigned to receive an intervention of 20 sessions of 45-minutes each, five days a week for four-weeks, in Robotic-therapy Group (RG) (n=12) and conventional upper-limb rehabilitation in Control-Group (CG) (n=11). We hypothesized to evaluate the exoskeleton based therapy for the effects on functionability of upper-limb and cortical-excitability in patients with stroke as compared to conventional-rehabilitation. Clinical-scales— Modified Ashworth Scale, Active Range of Motion, Barthel-Index, Brunstrom-stage and Fugl-Meyer scale (Shoulder/Elbow and Wrist/Hand component), and neurophysiological-measures of cortical-excitability (using Transcranial Magnetic Stimulation) –Motor Evoked Potential and Resting Motor-threshold, were acquired pre and post-therapy. No side effects were noticed in any of the patients.

Results: RG and CG showed significant (p<0.05) improvement in all clinical motor-outcomes except Modified Ashworth Scale in CG. RG showed significantly (p<0.05) higher improvement over CG in Modified Ashworth Scale, Active Range of Motion and Fugl-Meyer (FM) scale and FM Wrist-/Hand component). Increase in cortical-excitability in ipsilesional-hemisphere was found to be statistically significant (p<0.05) in RG over CG, as indexed by decrease in Resting Motor-Threshold and increase in amplitude of Motor Evoked Potential. No significant changes were shown by the contralesional-hemisphere. Interhemispheric RMT-asymmetry evidenced significant (p<0.05) changes in RG over CG indicating increased cortical-excitability in ipsilesional-hemisphere along with interhemispheric changes.

Conclusion: Neurophysiological-changes in RG could most likely be a consequence of plastic-reorganization and use-dependent plasticity. Robotic-exoskeleton training could significantly improve motor-outcomes and cortical-excitability in patients with stroke.

Introduction

Stroke is one of the leading causes of mortality and morbidity worldwide (1). Flexor hypertonia of wrist is one of the commonest presentation. The ability to actively initiate extension movements at wrist and nger against exor-hypertonia is one of the key indicators of motor recovery (2),(3). Regaining hand-function and Activities of daily-living (ADL) is particularly impervious to therapy or rehabilitation pertaining to the complexity of motor-control needed for distal-joints (4). Conventional rehabilitation-therapy is time taking, labour-intensive and subjective, which with high clinical-load and absence of skilled resources gets difficult for the present medical and healthcare-system to provide appropriate or effective rehabilitation services (5).

Although rehabilitation with neuro-rehabilitation robots has shown encouraging clinical-results (5,6,15,7–14), it is currently limited to a very few hospitals and not widely used because of associated high-cost and an infrastructural-requirement to station these large and complex devices, high set-up time, safety and usability restricting its success (16),(17),(18). Rehabilitation-strategies need to take into account the multifaceted nature of disability, which itself changes with time elapsed post-stroke and address with a multimodal-approach. Hence, the device needs to be flexible enough to accommodate a large patient-population. An effective rehabilitation device for hand should be able to facilitate a specific pattern of movements mirroring complex inter-joint coordination of hand with a patient-specific impairment, currently not integrated in the available devices.

In our previous work, we have designed a robotic-hand exoskeleton for rehabilitation of the wrist and MCP (Metacarpophalangeal) joint, to synchronize wrist-extension with finger-flexion and wrist-flexion with finger-extension, mimicking movement of ADL (19). With simple and easy-to-operate exoskeleton for low-resource settings, the exoskeleton targets spasticity through a synergy-based rehabilitation approach while also maintaining patient-initiated therapy through residual muscle-activity for maximizing voluntary effort. The lightweight and portable device has shown evidence of improvement in quantitative motor clinical-outcomes in patients with chronic stroke (19).
The aim of the present study was twofold. The first objective was to assess the clinical effectiveness of the novel robotic-exoskeleton device (19) and the second is comparison of its clinical-effectiveness with conventional upper-limb rehabilitation. We hypothesized that the exoskeleton based rehabilitation therapy might show better clinical outcome for distal-function of wrist and cortical-excitability in patients with stroke as compared to conventional-rehabilitation.

Materials And Methods

More than 300 patients (n>300) were screened in the out-patient clinic of the Department of Neurology, AIIMS, New-Delhi over 3 years from July-2016 to January-2019. Stroke diagnosis was established clinically in all patients. All clinical-assessments and standard-care were given to the patients with stroke by a trained physiotherapist. Institutional Review Board (IRB) at All India Institute of Medical Sciences (AIIMS), New-Delhi, India, approved the study under protocol-number IEC/NP-99/13.03.2015 and was registered with clinical trial number ISRCTN95291802. All the patients signed the written informed consent before enrolment.

2.1 Study-Design

A pilot prospective parallel single-blinded randomized controlled study at clinical-settings was designed which included pre and post clinical-outcome measures of therapy-intervention. Once enrolled, patients were then randomized under two groups- Robotic-therapy group (RG) and Control-Group (CG). Patients in RG underwent total 20 therapy-sessions, 5 days a week for 4-weeks, along with conventional physiotherapy-training. Patients in CG underwent conventional physiotherapy-sessions determined based on individual clinical-requirements. The same therapist provided therapy-sessions to all patients in both groups.

2.2 Patient Enrolment

Patients were enrolled based on inclusion-criteria, age 18–70 years, having ischemic / hemorrhagic stroke within 3-24 months, Mini-Mental Scale = 24-30; Brunnstrom-stage (BS) = 3-5; Modified Ashworth Scale (MAS) = 1, 1+, 2 (Figure-1). All enrolled patients continued to have their routine medication and standard medical care during the therapy-sessions, as advised by the neurologist. Patients with contra-indication to Transcranial Magnetic Stimulation (TMS) and any other progressive neurological or cognitive disorders were excluded from the study. The enrolled patients were allocated a pre-defined allocation sequence. Simple randomization was performed using opaque envelopes within which color cards signified the groups. Patients were instructed to choose the opaque envelopes in the pre-defined sequence. The cards they choose signified the group they were enrolled in for the study. Randomization, outcome measurements, and data analysis were performed by different individual not involved in the intervention.

2.3 Data Collection

2.3.1 Clinical scale measures (primary outcomes)

All the participants underwent clinical-assessment (pre and post-therapy) for measuring the level of spasticity by Modified Ashworth Scale (MAS 0-4), range of voluntary wrist-movement defined in terms of Active Range of motion of wrist (AROM 0°-70°) as measured by a goniometer, stage of stroke recovery Brunnstrom-Stages (1-7), ADL measured by Barthel-Index (0-100) and functional and sensorimotor-control of upper-limb as measured by Fugl-Meyer Scale (0-66).

2.3.2 Cortical-excitability measures using TMS (secondary outcomes)

Patients were allowed to sit comfortability in the chair, kept forearm pronated, elbow-joint at 90–120° flexion, wrist-joint at a neutral position and fingers at rest. Single-pulse TMS was given to evoke the Motor Evoked Potential (MEP) signal, using a flat 70mm figure-of-eight coil (type D70 (AC), serial no. 0326, Magstim Rapid2, Magstim, UK), at the cortical-representation of the Extensor Digitorum Communis (EDC) muscle (on contralateral motor-cortex with reference to the EEG cap) of the ipsilesional and contralesional-hemisphere. Cortical-excitability was measured in terms of Resting motor-threshold (RMT) and Motor Evoked Potential (MEP)-amplitude using TMS over ipsilesional and contralesional-hemisphere according to the standard-protocol (20). Resting Motor-threshold was defined as the minimum intensity of TMS required to elicit a MEP in target contralateral-muscle in
5/10 trials, recorded in EMG, over the muscle cortical-representation in the primary motor-cortex. MEP encapsulates information relevant to the cortical-excitability of the brain, conduction, and functional-integrity of the corticospinal-tract (21). MEP should be ≥ 50µv peak-to-peak amplitude at the hotspot in 5/10 consecutive trials.

2.4 Robotic therapy-sessions

An electromechanical robotic-exoskeleton was developed for rehabilitation of wrist-joint and fingers-joint (19) (Figure-2). It synchronizes wrist-extension with finger-flexion and wrist-flexion with finger-extension, a common pattern encountered in ADL. Device is actively-initiated by Electromyogram (EMG) and provides interactive adaptive performance-biofeedback in real-time. Device was safe, user-friendly and patient-centric with customizable motion-parameters, as per the clinical-presentation: (i) initial-position for range of motion (ROM), (ii) final-position for ROM, (iii) speed, (iv) residual muscle-activity and (v) height of finger-support. All sessions were given at the hospital set-up under the supervision of an expert clinician. Therapy-protocol consisted of total 20 sessions, each of 45-min over 4-weeks. Each 45-minute robotic-therapy session consisted of approximately 250-trials of 10-seconds each. Patients were advised to take a 5-min break for rest in between the therapy if there is a feeling of pain or fatigue. For details on exoskeleton design and protocol, please refer (19).

2.5 Data analysis

Data analysis was performed in MATLAB R2018a (MATHWORKS®). The data were tested for normality using Shapiro-Wilk test and was found that clinical-measures were not normally distributed in CG. Hence, non-parametric Wilcoxon signed-rank were used for intragroup-comparison of differences in post–pre-therapy within the group and non-parametric Man-Whitney tests were used for intergroup-comparison of RG and CG. Interhemispheric-asymmetry for pre and post-therapy measures were calculated and was tested using Wilcoxon signed-rank test. Two-way repeated measure ANOVA was applied to assess the effect of time (two levels-pre and post) and side (two levels-ipsilesional and contralesional) on RMT. Regression and correlation-analyses were performed to investigate the relationship of recovery parameters TMS neurophysiological-parameter with clinical-outcome. A p-value < 0.05 was considered as statistically significant. MAS score of 1, 1+, 2, 3, 4 was mapped as 1, 1.5, 2, 3, 4 for all statistical calculation purpose, respectively as suggested by Rong et. al (13).

Results

Twenty-seven patients who met the eligibility criteria were randomized and allocated into two groups- RG (n=13) and CG (n=14). One patient (n=1) in RG and three patients (n=3) in CG could not complete the therapy, thus, the data were excluded from further analysis. All patients in RG (n=12) and CG (n=11) (all right-handed patients with stroke, age=41.9±11.1 years, Male:Female=19:4) (Table-1) completed successfully the therapy-sessions in 30-35 days. There were no significant (p > 0.05) differences in the pre-therapy measures in terms of clinical-scales and MEP signal among both the intervention-groups. At pre-therapy measurements, MEP was evoked only for 9 patients (RG=4, CG=5) out of total 23 in ipsilesional-hemisphere, and for all patients in contralesional-hemisphere. No side effects or adverse effects were noticed in any of the patients.

3.1 Comparison of Clinical-scales

Post-therapy, all clinical-scales in both groups did show significant changes in improvement, except for MAS in CG. However, all clinical-scales (MAS, AROM, FM, and FMW/H) in RG showed statistically significant changes over CG. MAS in RG changed from 1.75±0.26 to 1.29±0.33 and in CG from 1.86±0.5 to 1.59±0.66 showing a significant decrease in spasticity at wrist-joint in RG and not in CG (RG-p=0.0009, CG-p=0.12) with significant (p=0.03) intergroup changes (Table-2). ADL, as indexed by AROM and BI, in both groups showed statistically significant differences. ADL significantly improved in both the groups, from 150±9.70 to 34.580±14.50 in RG (p=0.0004) and from 13.60±7.70 to 200±8.060 in CG (p=0.002). However, RG manifested statistically significant sensorimotor-scores as compared to CG as intergroup-comparison did evidence significant differences (p=0.02) (Table-2). BI changed from 74.17±12.4 to 89.17±7.93 in RG (p=0.0009) and from 69.54±12.9 to 82.72±14.3 in CG (p=0.0009); the intergroup-comparison did not show any significant differences (p=0.82). BS showed statistically significant differences in both
groups, RG changed from 3.67±0.78 to 4.83±0.94 (p=0.0004) and CG changed from 3.72±1 to 4.45±1.2 (p=0.015). The intergroup-comparison did not show significant differences between both groups (p=0.311) (Table-2).

FMU/L scores measure sensorimotor-control gain in both groups after the intervention. FMU/L for RG changed from 36±7.78 to 50.25±6.59 (p=0.0004) and from 37.45±9.1 to 45.45±9.7 for CG (p=0.0009). RG manifested statistically significant improvement in sensorimotor-scores as compared to CG with significant (p=0.039) differences in intergroup-comparison (Table-2). For the proximal-part-Shoulder/Elbow component of FMS/E, both groups showed statistically significant increase, RG changing from 26.2±5.6 to 33.5±3.8 (p=0.0009) and from 26±7.07 to 29.8±7.08 in CG (p=0.002). However, the intergroup-comparison did not show any significant (p=0.13) differences. For the distal-part Wrist/Hand component of FM (FMW/H), both groups showed statistically significant increase, in RG changing from 9.7±2.7 to 16.6±4.3 (p=0.0004) and in CG changing from 11.45±2.9 to 15.18±3.6 (p=0.0009). RG manifested statistically significant (p=0.012) sensorimotor-improvement in intergroup-comparison over CG (Table-2).

3.2 Comparison of Cortical-excitability

3.2.1 Ipsilesional-hemisphere

MEP, in some patients with stroke, is not recordable even after delivering TMS-stimuli at the highest possible intensity due to lower cortical-excitability. In those patients with no MEP recorded, RMT is taken as value of 100, as has been suggested in literature. Change in RMT showed statistically significant differences in post-therapy in RG as compared to CG. Post-therapy, in RG showed a significant decrease in RMT from 95.3±7.87.23 to 79.58±14.38 (p=0.0039) (indicating increase in cortical-excitability), whereas CG showed a decrease from 89 to 85.18±17.9 (p=0.12) (Table-2). RG also evidenced a significant increase in MEP-amplitude (indicating increase in cortical-excitability) from 39.4±60.4 µv to 94.3±63.2 µv (p=0.048). In CG, MEP almost remained same (~38 µv) pre-to-post-therapy (p=0.312). RG, thus, demonstrated a significant increase in cortical-excitability in intergroup-comparison over CG, as evidenced by decrease in RMT (p=0.037) and increase in MEP-amplitude (p=0.0142) (Table-2).

In this study, ~54% of patients in CG did not evoke MEP. Approximately ~67% of patients in RG too did not evoke MEP at the pre-therapy measurements, most of which later evoked post-therapy showing the therapeutic-effectiveness of the exoskeleton. It is worth-noting that in RG, measurable MEP was evoked only in 4/12 patients at the pre-therapy measurements, however, post-therapy MEP was observed in 9/12 patients, indicating a considerable increase in cortical-excitability in 5 patients. In CG, MEP was evoked only in 5/11 patients at the pre-therapy measurements and was observed in 6/11 patients post-therapy.

3.2.2 Contralesional-hemisphere

There were no significant changes shown by the contralesional-hemisphere. Both RG and CG evidenced minimal differences in RMT (mean increase of ~2% in both groups) (Table-2). RG showed decrease from 67.33±10.07 to 65.08±11.12 (p=0.051) and CG from 68.09±11.7 to 66.18±12.57 (p=0.052). Intergroup-comparison too did not show statistically significant differences (p=0.87) (Table-2). MEP-amplitude in RG decreased from 506.33±247 to 355.3±191.5 µv (p=0.33) and from 200.2±77 µv to 185.4±268.3 µv in CG (p=0.41). MEP-amplitude observed a considerably higher decrease (mean~151 µv) in RG indicating changes in cortical-excitability, as compared to CG (mean~15 µv). The inter-group comparison however, was not statistically significant (p=0.51) between both the groups (Table-2).

3.2.3 Inter-hemispheric differences and asymmetries

The effect of robotic-exoskeleton training on cortical-excitability was assessed within both hemispheres. RG showed statistically significant differences between ipsilesional and contralesional-sides as one factor and time points- pre and post-therapy as another factor on RMT (p=0.049, F=4.08), evidencing the dependence of time and hemisphere sides on each other. However, CG did not show any statistical differences (p=0.06, F=3.68).

RG also evidenced a statistically significant reduction in interhemispheric-RMT asymmetry as measured by the ratio of RMT for two hemispheres (RMT_{asymm} = RMT Ipsilesional / RMT contralesional) from pre-to-post-therapy (Table-2). RG showed a decrease
in $RMT_{asymm}$ from 1.43±0.21 to 1.25±0.31 (mean decrease of 0.18, p=0.012), however, CG showed a decrease from 1.33±0.30 to 1.30±0.28 (mean decrease of 0.03, p=0.59), indicating a trend of normalization of RMT-asymmetry ($RMT_{asymm}$ should decrease as ipsilesional RMT should be decreased from pre-to-post) over the course of intervention in RG. RG also manifested statistically significant (p=0.028) changes in intergroup-comparison over CG (Table-2). The relative change in interhemispheric-RMT asymmetry-ratio ($\Delta RMT_{asymm-ratio}$) changed with RG having a mean increase of 0.12±0.14 and CG a mean increase of 0.01±0.1 (p=0.028), indicating the extent of normalization of RMT-asymmetry over the duration of intervention in RG as compared to CG (Table-2).

3.3 Relationship between TMS neurophysiological-measures and clinical-outcome

The recovery parameters from TMS-measures denoting the change from pre-to-post-therapy were observed to be strongly correlated with the relative change/improvement in distal motor-outcome ($\Delta FMW/H$). The first parameter, relative change in RMT in the ipsilesional-hemisphere ($\Delta RMT_{ipsi}$) was significantly (p=0.0235) different for both the groups with a mean increase of 0.16±0.12 in RG and 0.04±0.09 in CG. The linear-regression analysis indicated that $\Delta RMT_{ipsi}$ (as predictive/independent-variable) could correlate with $\Delta FMW/H$ (as dependent-variable) and could predict $\Delta FMW/H$ in RG ($r=0.64, F=7.24, p=0.022$) (Figure-3a), indicating that increased cortical-excitability in the ipsilesional-hemisphere was correlated with the functional clinical-outcome. This correlation was not found in CG ($r=0.47, F=2.62, p=0.13$) (Table-2, Figure-3a). The relationship between $\Delta RMT_{ipsi}$ and $\Delta FMW/H$ for both groups is shown in scatter-plot in figure-3a. The distal functional-outcome $\Delta FMW/H$ also showed significantly (p=0.012) different results for both groups with a mean increase of 0.73±0.45 in RG and 0.33±0.14 in CG.

The second parameter, relative change in RMT-ratio ($\Delta RMT_{asymm-ratio}$) was significantly (p=0.028) different for both the groups. Similar to above, $\Delta RMT_{asymm-ratio}$ (as predictive/independent-variable) was observed to be strongly correlated $\Delta FMW/H$ (as dependent-variable) ($r=0.6, F=5.77, p=0.03$) (Figure-3b), indicating that tendency towards extent of normalization RG could correlate and predict the clinical-outcome in RG. This correlation was not found in CG ($r=0.29, F=0.83, p=0.38$) (Figure-3b). The relationship between $\Delta RMT_{asymm-ratio}$ and $\Delta FMW/H$ in both groups is shown in scatter-plot in figure-3b.

Discussion

The study demonstrated clinical and neurophysiological-changes in response to the robotic-exoskeleton (19) training compared to the conventional-rehabilitation. The clinical-scales showed improvement in both RG and CG, however, increased cortical-excitability in the ipsilesional-hemisphere was shown only in RG. Five patients in RG, with the absence of MEPs at the pre-therapy measurements, showed the appearance of MEPs in the ipsilesional-hemisphere post-therapy. The improvement in RMT in ipsilesional-hemisphere showed a trend of normalization over the intervention and were also correlated with sensorimotor-functional improvement.

4.1 Comparison of Clinical-scales of Robotic-therapy group with control-group

The robotic-therapy was effective in releasing spasticity at the wrist-joint with ~26% (p=0.03) improvement over ~14% in CG. The regain in normal muscle-tone is considered as a predictor of recovery or the first-step in recovery (22) followed by an increase in muscle-strength and improvement in functional movements or ADL, finally leading to muscle-strength. In this study, AROM and Barthel-Index have been measured as the indicators of ADL (Table-2). Both groups showed significant improvement of AROM, however, RG showed significantly (p=0.02) higher improvement of 130% over 47% in CG. AROM is one of an important parameters in evaluating ADL and increase in AROM of wrist could lead to greater participation in ADL (23). For Barthel-Index, both groups showed similar (~20%) improvement (p=0.82). BI is a non-specific crude discrete measure of ADL, measuring discretely for independence, partial-dependence and fully-dependence hence, even a minimal improvement will lead to an increase in the score by 5 units. A significant improvement in BI was observed for both the groups (RG & CG). BI is not much reliable as scores might get affected depending on dominant/non-dominant side, thus, is not alone predictor for therapeutic-outcomes (24). Both groups showed significant (intergroup p=0.31) improvement for BS, however, RG showed~32% improvements over ~20% in CG (Table-2). As BS measure the synergy pattern of joints, it depends mainly on various factors e.g. proximal and/or distal-joint condition of recovery, MAS and chronicity etc.
FMU/L, stroke-specific scale, is most reliable measure of sensorimotor-functionalities of the whole-arm (25). RG established significantly higher improvements of ~40% over ~21% in CG (p=0.039). In FMS/E, RG showed ~28% improvement over ~15% in CG. RG did not show significantly different results (p=0.13) as expected as the intervention was not focused on proximal-component. However, for FMW/H distal-component, both groups showed significant improvement, where but RG showed significantly higher improvement of ~72% over 32% in CG (p=0.012) because of intensive and repetitive-training of wrist and MCP. Similar improvement has also been reflected in proximal-joints in along with distal-joints robotic-training possibly because of compensatory muscle-activities from proximal-joints (12).

Thus, RG shows an overall increase in sensorimotor-ability and functionality evidenced by increase in FM, AROM and decrease in spasticity (MAS). Patients in RG showed more participation in ADL, reflected by an increase in BI and AROM of the wrist (23). Among the available wrist rehabilitative-devices and dose-matched conventional-intervention for rehabilitation of wrist in stroke, the reported gain in FMW/H is < 4 on a scale of 24 (12),(8),(26–28). In our study, the improvement in the FM score was ~14.25 (W/H~7). Any direct comparison with literature is, however, inappropriate considering different factors like chronicity of patients, lesion site, age of the patients, etc. in this study. HWARD (29) also showed improvement in FMW/H (~3.8), indicating the usefulness of synchronizing both wrist and MCP-joint movement in grasp and release.

4.2 Comparison of Cortical-excitability of Robotic-therapy with the control-group

Cortical-excitability in pre-therapy measurements was found to be lower in patients with stroke as observed by higher RMT and lower MEP (30),(31),(32),(33),(34). In some patients due to low cortical-excitability, MEP is not recordable even after delivering TMS-stimuli at the highest possible intensity (Maximum Stimulator Output at 100%) (30),(31),(32),(33),(34). In those patients with no MEP recorded, RMT is taken as value of 100, as suggested in literature (35),(36). This study investigated specific impact of therapeutic-interventions on TMS neurophysiological-parameters for cortical-pathways in the context of functional-gains of the hand motor-function.

4.2.1 Ipsilesional and Contralesional-hemisphere changes

At pre-therapy measures in the ipsilesional-hemisphere, only ~39% of the total patients’ cohort in the study evoked MEP (Table-2). In RG, 67% of patients (8/12) did not evoke MEP. Out of these 8 (67%) patients, 5 (62%) patients later evoked MEP showing the therapeutic-effectiveness of the exoskeleton in RG. RG showed a significant (p=0.0039) decrease in RMT from pre-to-post-therapy-sessions which was not observed in CG (p=0.12). RG showed ~16% improvement as compared to ~4% improvement in CG (p=0.037). Interestingly, RG showed significantly (p=0.048) higher increase in MEP-amplitude post-therapy with increase of ~140% (mean=54.9 µv), whereas CG showed no such improvement. In the contralesional-hemisphere, MEP-amplitude showed a considerable decrease in both groups, though not significant, RG evidencing a considerable decrease of ~30% (mean=151.03 µv) and CG a decrease of only ~7% (mean=14.8 µv) with no significant differences (p=0.51) (Table-2).

Cortical-excitability measures are used as an objective investigative tool to measure the treatment responsiveness and prognostication as it provides insights into membrane-excitability of neurons, conduction, and functional-integrity of corticospinal-tract and neuromuscular-junctions (37). The significant decrease in RMT and increase in MEP-amplitude in the ipsilesional-hemisphere demonstrates significant amount of increase in cortical-excitability (38), as was demonstrated in the RG versus CG. It can be interpreted that recovery of motor-function could most likely be a consequence of plastic-reorganization and use-dependent plasticity (38). Cortical-excitability and corticospinal-tract integrity have also been shown to be correlated with functional recovery potential in patients with chronic stroke (31) and exoskeleton-training appears to be beneficial in activating the ipsilesional-hemisphere for chronic patients (13.8±9.1 months). Activation of ipsilesional-hemisphere could indicate either vicariation of the loss of neural circuits or unmasking of pre-existing synapses or recruitment of perilesional areas in ipsilesional-hemisphere or exploitation of the preserved functional recovery reservoir in ipsilesional-hemisphere (35),(39),(40),(41). Further, a ~30% decrease in MEP-amplitude in contralesional-hemisphere over the duration of intervention might indicate a decrease in cortical-excitability, evidencing a trend towards restoring the Inter-Hemisphere Inhibition (IHI) balance in the motor-network between the two hemispheres(39),(40), however, it needs to be further evaluated in a larger cohort.
The cortical-excitability measures, usually, are acquired in pre and post-intervention mostly involving brain-stimulation studies. Examples are repetitive TMS, Transcranial Direct-Current Stimulation (tDCS)(42),(43), etc. or in a combination of brain-stimulation with other neuro-rehabilitation strategies like Constrain Induced Movement-Therapy (CIMT) (44) or mirror-therapy (45) or training(46),(47). However, studies evaluating the therapeutic-changes in the cortical-excitability using robotic-training intervention are very rare. To best of our knowledge, only two studies attempt to evaluate the effect of active robotic-training on changes in cortical-excitability, using commercially available devices, such as Lokomat-robot (lower-limb) (48) and ARMEO (upper-limb) (49).

4.2.2 Specific five-patients in RG

A very critical outcome of the therapy was that in RG, MEP was evoked in ipsilesional-hemisphere only for 4/12 patients at the pre-therapy measurements, however, MEP was later evoked for 9/12 patients after robotic-therapy. However, in CG, MEP was evoked only for 5/11 patients and was later evoked for 6/11 patients at post-therapy. Five specific patients in RG who did not evoke MEP at pre-therapy (0 µv) and later evoked MEP (mean=136.6±38.48 µv), showed a decrease of RMT in ipsilesional-hemisphere (mean=27±9.64), relative change RMT-ratio (mean=0.22±0.13) and mean increase in clinical-scales (FMW/H: 7.8±2.38, BI: 22±11.72, AROM: 22±2.73). These changes were relatively much higher than the changes in patients who had MEP evoked at pre-therapy measures. The appearance of MEP in five patients indicates that the robotic-therapy has an immense potential of training and reorganization of brain based on use-dependent plasticity, for patients with chronic stroke. The increase in cortical-excitability and normalization of TMS neurophysiological-makers on the ipsilesional-side are also accompanied by greater recovery of hand-function, indexed by sensorimotor and functional recovery (by clinical-scales FMW/H, BI & AROM). Indeed, the appearance of MEPs in ipsilesional-side could be a critical recovery marker in stroke recovery.

4.2.3 Inter-hemispheric differences and asymmetries

The diachisis between ipsilesional-areas and intact neuronal-networks of contralesional-areas may disturb the cortical-excitability and connectivity-patterns of connected, remote, or primary-motor areas of contralesional-hemisphere (via transcallosal-fibers). The effect of robotic-exoskeleton training on cortical-excitability of both hemisphere shows remodelling of the bilateral primary-motor areas in RG (time*sides p=0.049, F =4.08) which is not shown in CG (time*sides p=0.06, F=3.68). The effect of exoskeleton-training shows the potential of the exoskeleton to accelerate the cortical-plasticity phenomena in favour of functional-restoration with changes in both ipsilesional and contralesional-hemispheres.

For cortical excitability to be increased in ipsilesional-hemisphere for patients with stroke, the ipsilesional-RMT should be decreased from pre-to-post-therapy and hence, RMT$_{\text{asymm}}$ (RMT Ipsilesional/RMT contralesional) should decrease/approach normalization (35). Significant differences were observed between the groups when TMS-neurophysiological changes over the intervention was expressed in terms of the interhemispheric-asymmetry ratio RMT$_{\text{asymm}}$ indicating a significantly (p=0.028) greater trend towards the normalization of asymmetry of TMS-measures in RG in response to exoskeleton-training than CG. Also, the extent of normalization i.e. $\Delta$RMT$_{\text{asymm-ratio}}$ showed a mean increase of 0.12±0.14 in RG and CG an increase of mean 0.011±0.1 (Table-2). Normalization might indicate the recruitment of peri-lesional areas in the ipsilesional-hemisphere or exploitation of the preserved functional-recovery reservoir in the ipsilesional-hemisphere (35),(39),(40),(41). Normalization in response to therapy, in terms of TMS measures on the ipsilesional-side, has been shown to have a greater recovery of arm and hand function in acute, sub-acute and chronic stages (35).

4.3 TMS neurophysiological improvement correlating the motor-outcome of both groups

The amount of change in TMS neurophysiological-measures of corticomotor-pathways ($\Delta$RMT$_{\text{ipsi}}$ and $\Delta$RMT$_{\text{asymm-ratio}}$) were found to be associated with the amount of improvement in functional motor-outcome during rehabilitation of the distal-part of upper-limb ($\Delta$FMW/H) (Figure-3). These parameters were significantly different for RG and CG ($\Delta$RMT$_{\text{ipsi}}$ p=0.0235, $\Delta$RMT$_{\text{asymm-ratio}}$ p=0.028 and $\Delta$FMW/H p= 0.012). Greater improvement (decrease) in motor-threshold tend to show greater increases with clinical-outcome and was found to have strong positive statistical correlation with $\Delta$FMW/H in RG ($\Delta$RMT$_{\text{ipsi}}$ r=0.64, p=0.022 and not in CG r=0.47, p=0.13 and $\Delta$RMT$_{\text{asymm-ratio}}$ r=0.6, p=0.03 and not in CG (r=0.29, p=0.38) (Figure-3). The improvement (decrease) in RMT, found to be associated with recovery of motor function (31), were most likely due to increased cortical-
excitability of preserved motor-pathways with earlier studies in sub-acute and chronic stroke demonstrating correlation of improvement in TMS neurophysiological-measures (improvement in RMT and normalization) with functional improvement (35), (50), (51). This also suggests the usefulness of TMS-measures as an index of recovery in the ipsilesional-hemisphere (35). These neurophysiological-measures were obtained specifically from cortical-representation of EDC muscle, a clinically affected muscle, with a specific function which was involved in training with a robotic-exoskeleton, whereas most clinical-measures do not necessarily require a particular muscle-group and measures motor-function in a broader sense.

Also, these neurophysiological-parameters individually establishes as a significant predictor ($\Delta RMT_{\text{ipsi}} r = 0.64, F=7.24, p=0.022$ and $\Delta RMT\text{asymm-ratio } r=0.6, F=5.77, p=0.03$) of functional rehabilitation-outcome of hand ($\Delta FMW/H$) in RG, indicating that changes in cortical-excitability of ipsilesional-hemisphere could be used to predict the clinical-outcome, hence, emerging as critical recovery parameters to be considered and evaluated in larger data-samples. This might possibly be the plasticity markers predicting the responsiveness of chronic post-stroke patients (49). Hence, the correlation and prediction of improvement in $\Delta FMW/H$ component by these muscle-specific neurophysiological-measures comes as an evidence of task-specific rehabilitation of specific-muscle.

4.4 Changes due to the device

The exoskeleton training in RG induced an evident modulation in ipsilesional and contralesional-hemispheres. However, (significant) changes in CG were found to be limited only to the clinical-scales, and the changes in brain was specifically found only in the RG. An increase in cortical-excitability in the ipsilesional-hemisphere along with interhemispheric-normalization of $RMT\text{asymm}$ could point towards the recruitment of peri-lesional areas in ipsilesional-hemisphere or exploitation of the preserved functional recovery reservoir in the ipsilesional-hemisphere (35),(39),(40),(41). The decrease of RMT and change in RMT asymmetry from distal-muscle was also accompanied by functional markers-FMW/H evidencing sensorimotor-plasticity, functional recovery along with task-dependent rehabilitation. Greater magnitude of the neurophysiological-changes observed in RG, as compared to CG, may be attributed to the unique features of the device e.g. easy donning and doffing of the device for repositioning of hand, maximum finger-extension at baseline position to provide maximum stretch to reduce spasticity; and diverse attributes of device e.g bio-triggered, real-time adaptive performance-feedback, counteracting flexor-hypertonia against gravity, mimicking functional motion, user-friendly, simple design and patient-specific customizable features with different amount of sensory inputs (proprioceptive, visual, tactile) (19). In combination with the above features, the other unique feature of the device is that it allows the facilitation of a specific-pattern of movements mirroring complex inter-joint coordination of hand with a patient-specific impairment, currently not integrated in the available other devices with isolated-joint movements (52). The devices were able to simulate the movement pattern maintaining joint-coordination, especially at the distal-joints which could aid in translating the motor-improvements into ADL.

Limitations: Even though data are promising, the study had few limitations such as small sample-size and no long term follow-up of patients. As most of the patients at our quaternary hospital came from far places across India and it was not possible to follow-up with them once they have left New-Delhi. The outcomes provided critical data to plan a multicentric trial in future to systematically investigate the potential of the exoskeleton.

Conclusion

The robotic-exoskeleton illustrated reduction in spasticity, improvement in clinical scales and increased cortical-excitability in patients with stroke as compared to the control-group. This technology-based intervention has the potential to enhance the recovery of stroke neuro-rehabilitation.

Declarations

Ethical Statement: Institutional Review Board (IRB) at All India Institute of Medical Sciences (AIIMS), New-Delhi, India, approved the study under protocol-number IEC/NP-99/13.03.2015. All the patients signed the written informed consent before enrolment.

Consent for Publication: All the patients signed the written informed consent for publication before enrolment.
Availability of Data and Material: The datasets used and/or analyzed in the current study are available from the corresponding author on reasonable request.

Competing Interest: The Author(s) declare(s) that there is no conflict of interest.

Funding: This work was financially supported by SERB, DST India (YSS/2015/000697).

Authors Contributors: NS and AM conceptualized and designed the study. AM led the study and provided the scientific inputs. MS performed patient recruitment, physiotherapy and data collection. NK and PS provided the scientific inputs, clinical support and clinical resources for experiments. NS performed literature survey, developed device, data analysis, data interpretation, wrote the manuscript. AM reviewed the manuscript at multiple iteration with NS. All authors reviewed and approved the manuscript.

Acknowledgment: The authors thank all patients who participated in the study.

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Tables

Table 1: Details of patients with stroke enrolled in Robotic Group (n=12) and Control Group (n=11)

| Group          | Gender (M/F) | Stroke type | Ipsilesional site, Left/Right | Age (years) | Chronicity (months) |
|----------------|--------------|-------------|-------------------------------|-------------|---------------------|
| Robotic-therapy | 10/2         | 9/3         | 5/7                           | 41.1±12.8   | 13.8±9.1            |
| Control        | 9/2          | 8/3         | 5/6                           | 42.7±9.3    | 10.3±5.0            |

Table 2: Comparison of clinical-scales, cortical-excitability (in ipsilesional and contralateral-hemisphere) and interhemispheric parameters in Robotic Group with Control Group
|                      | Robotic-therapy Group | Intergroup p-value | Control Group |                  |                  |                  |                  |                  |
|----------------------|-----------------------|--------------------|---------------|-----------------|-----------------|-----------------|-----------------|-----------------|
|                      |                        | Mean+ Standard deviation |                      | Mean+ Standard deviation |                  |                  |                  |                  |
|                      | Pre-Therapy | Post-Therapy | Difference of mean | Robotic group p-value | Pre-Therapy | Post-Therapy | Difference of mean | Control group p-value |
| AS                   | 1.75±0.26             | 1.29±0.33           | 0.46           | 0.0009          | 0.03*           | 1.86±0.5        | 1.59±0.66       | 0.27             | 0.12             |
| OMI                  | 150±9.70              | 34.58±14.50         | 19.580         | 0.0004          | 0.02*           | 13.60±7.70      | 200±8.060       | 6.40             | 0.002            |
| SI                   | 74.17±12.4            | 89.17±7.93          | 15             | 0.0009          | 0.82            | 69.54±12.9      | 82.72±14.3      | 13.18            | 0.0009           |
| S                    | 3.67±0.78             | 4.83±0.94           | 1.16           | 0.0004          | 0.311           | 3.72±1          | 4.45±1.2        | 0.73             | 0.015            |
| UL                   | 36±7.78               | 50.25±6.59          | 14.25          | 0.0004          | 0.039*          | 37.45±9.1       | 45.45±9.7       | 8                | 0.0009           |
| W/H                  | 9.7±2.7               | 16.6±4.3            | 6.9            | 0.00048         | 0.012*          | 11.45±2.9       | 15.18±3.6       | 3.73             | 0.0009           |
| S/E                  | 26.2±5.6              | 33.5±3.8            | 7.3            | 0.0009          | 0.13            | 26±7.07         | 29.8±7.08       | 3.8              | 0.002            |
| IW/H                 | 0.73±0.45             | 0.012*              | 0.33±0.14      |                  |                  |                  |                  |                  |

### Clinical Scales

|                      |                      |                      |                  |                  |                  |                  |                  |                  |
|----------------------|----------------------|----------------------|------------------|------------------|------------------|------------------|------------------|------------------|
| MAS (max 4)          | AROM (max 70)        | Active Range of Motion |                  |                  |                  |                  |                  |                  |
| BI (max 100)         | BS (max 7)           | Brunstrom Stage       |                  |                  |                  |                  |                  |                  |
| FMU/L (max 66)       | FMW/H (max 24)       | Fugl-Meyer Wrist Hand |                  |                  |                  |                  |                  |                  |
| FMS/E (max 42)       | MEP A (µv)           | MEP Amplitude RMT (%) |                  |                  |                  |                  |                  |                  |
| IL: Ipsilateral, CL: Contralateral | RMT<sub>asymm</sub> = (RMT Ipsilesional / RMT contralesional), |                  |                  |                  |                  |                  |                  |                  |
| ∆RMT<sub>asymm</sub> = (Post RMT<sub>asymm</sub> - Pre RMT<sub>asymm</sub>) / Pre RMT<sub>asymm</sub> | Relative improvement in RMT ratio |                  |                  |                  |                  |                  |                  |                  |
| ∆RMT<sub>ipsi</sub> = (Pre RMT<sub>ipsi</sub> - Post RMT<sub>ipsi</sub>) / Pre RMT<sub>ipsi</sub> | Relative decrease/improvement in ipsilesional RMT |                  |                  |                  |                  |                  |                  |                  |
| ΑFMW/H = (Post FMW/H - Pre FMW/H) / Pre FMW/H | Relative improvement in Fugl-Meyer (W/H) |                  |                  |                  |                  |                  |                  |                  |

*shows the statistical significance differences (p < 0.05) between RG and CG

**Figures**
Figure 1
Patient Enrolment Consort

Control Group (CG) (n=14)
45 min/day, 5 times a week, 4 weeks
(Total twenty sessions of Physiotherapy)

Robotic Group (CG) (n=13)
45 min/day, 5 times a week, 4 weeks
(Total twenty sessions of Robotic-Therapy)

Data Acquisition:
CG (n=11) & RG (n=12)
Clinical Scales: MAS, BI, BS, FM, AROM
Cortical Excitability: RMT & MEP

Data Analysis & Interpretation

Screened for Eligibility (n>300)
Clinical Assessment (n=27)
Randomized

Inclusion Criteria:
Ischemic / Haemorrhagic, single lesioned, Cortical / Subcortical stroke, stroke onset = 3-24 months, Age=18-70 Y, MAS=1, 1+, 2, BS=3-5, MMSE Score=24-30, MEP≥10 μV in Affected Hemisphere

Figure 2
Motor
Gearbox
Four bar linkage
LED Dot Matrix for Performance feedback
Green LED to signal for Voluntary feedback
PCB in black box
EMG electrodes
Forearm attachment
Whole set-up of exoskeleton with performance biofeedback, voluntary cue and PCB in the black control-box which also works as user-interface (refer Singh et al (19))

**Figure 3**

Scatter-plot showing relationship between improvements in RMT in the ipsilesional-hemisphere and improvements in functional performance of the distal-component pre-to-post-therapy for individual patient’s data. Greater decreases in motor-threshold tend to show greater increases in FMW/H. Red Line (RG) and the blue line (CG) represents a linear-trend in improvement in distal motor-outcome (ΔFMW/H) score as a function of change in the ipsilesional-hemisphere (ΔRMTipsi) in RG and CG pre-to-post-therapy in which RG shows a significantly strong correlation (r=0.64, F=7.24, p=0.022), Scatter-plot showing relationship between change in RMT asymmetry-ratio (ipsilesional/contralesional) pre-to-post-therapy and functional performance of distal-component for individual patient’s data. Greater decreases in motor-threshold tend to show greater increases in FMW/H. Red line (RG) and the blue line (CG) represents a linear-trend in improvement in distal motor-outcome (ΔFMW/H) score as a function of change in RMT-ratio (ΔRMTasymm-ratio) in RG and CG pre-to-post-therapy in which RG shows a significantly strong correlation (r=0.6, F=5.77, p=0.03)

**Supplementary Files**

This is a list of supplementary files associated with this preprint. Click to download.

- consortchecklist.docx