

THE EFFECTIVENESS AND COST-EFFECTIVENESS OF THE COADMINISTRATION OF MINDMOTIONPRO PLUS STANDARD PRACTICE VERSUS STANDARD PRACTICE IN EARLY POST-STROKE UPPER LIMB REHABILITATION:



THE MOVE-REHAB STUDY

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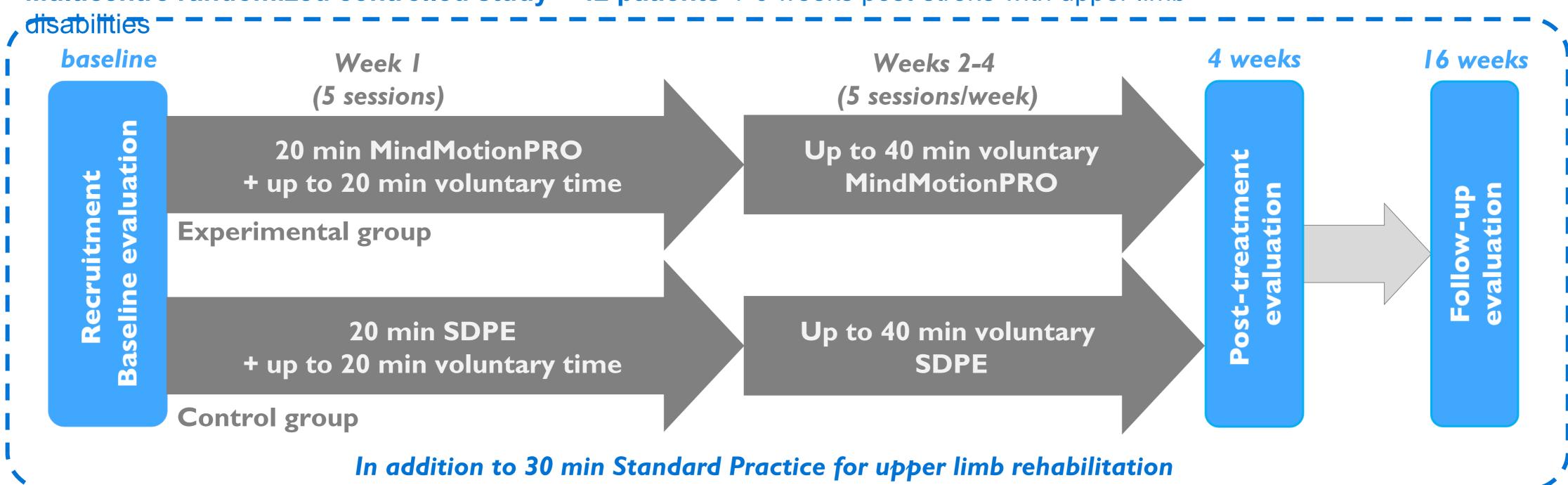
BACKGROUND

- •Stroke survivors often experience motor disabilities, limiting their independence in performing daily activities. Recent studies suggested that effective post-stroke rehabilitation should be challenging, repetitive, task-specific, motivating and intensive to foster greater neuroplasticity.
- •Virtual Reality (VR) based training has been shown to be an effective complement to standard rehabilitation, especially in intensifying the therapy dose and maintaining patient motivation.
- •MindMotionPRO is a Class I medical device providing immersive VR-based training for upper extremities with 3D motion tracking and real-time visual feedback.

OBJECTIVE

•To compare the **rehabilitation dose** between an experimental group using standard rehabilitation plus MindMotionPRO and a control group following Self-Directed Prescribed Exercises (SDPE) in addition to standard rehabilitation.

Multicentre randomized controlled study – 42 patients 1-6 weeks post-stroke with upper limb





INTERVENTIONS

- •MindMotionPRO: Upper extremity rehabilitation exercises in a virtual reality environment, with integrated concepts of constraint-induced therapy, mirror therapy, action-observation therapy, and motor imagery
- •Self-Directed Prescribed Exercise (SDPE): Upper extremity rehabilitation exercises following the Graded Repetitive Arm Supplementary Program (GRASP)

OUTCOMES

- •Primary Outcome: Time spent in active rehabilitation and number of exercises performed by the patient
- Secondary Outcomes:
 - Motor function and clinical assessments (Fugl-Meyer Assessment for Upper Extremities, streamlined Wolf Motor Function Test, Barthel Index, Stroke Impact Scale, NIH Stroke Scale and Motor Activity Log)
 - Motivation (Intrinsic Motivation Inventory)
 - Resources utilization (therapist time)

INCLUSION CRITERIA

- •Male/Female > 18 years
- •First anterior or middle cerebral artery ischemic or hemorrhagic stroke
- •1 to 6 weeks post-stroke
- •Motor difficulties in using paretic arm with FMA-UE between 20 and 40
- •Stroke severity according to NIHSS score between 5 and 14
- •Able to provide informed consent
- •Not currently participating in other intervention studies

EXCLUSION CRITERIA

- •Any medical condition compromising the safety or ability to take part in the study, including upper limb conditions not linked to stroke and co-morbidities
- •Recurrent and moderate-to-high upper limb pain limiting rehabilitation dose
- •More than one or uncontrolled epileptic seizures since stroke onset
- •Mild to severe cognitive impairment (Mini Mental State Exam < 24/30)
- •Depression (Hospital Anxiety and Depression Scale > 8/21)
- •Moderate to severe hemispatial neglect (Bells tests > 6 errors)

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