

Proceeding Paper

Quantitative Data to Evaluate Clinical Pilates Efficacy in Chronic Low Back Pain Using Inertial Measurement Units [†]

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Abstract: Chronic non-specific low back pain (CNLBP) affects a significant percentage of the global population, with therapeutic exercise being a key management strategy. This study evaluated the effectiveness of the clinical Pilates method in 22 patients with CNLBP. Lumbar spine range of motion (ROM) and completion times for three functional tests (prone plank, side bridge, and supine bridge) were measured before and after a six-week rehabilitation program. Motion data were collected using two inertial measurement units. Results showed statistically significant improvements in kinematic patterns, execution times, and ROM. These findings support the effectiveness of clinical Pilates in improving lumbar spine functionality and patient outcomes.

Keywords: CNLBP; therapeutic exercise; clinical pilates; bridge tests; IMUs



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1. Introduction

Chronic non-specific low back pain (CNLBP) is a widely prevalent condition [1] that is strongly associated with disability and work absences globally [2,3]. Although the causes of CNLBP are complex and multifactorial, they are directly linked to etiological factors including social and demographic characteristics, lifestyle habits, and both physical and psychosocial influences [4].

Lumbar muscle atrophy, reduced strength and endurance, and increased fatigue are commonly observed in association with CNLBP. The stability of the core muscles, which depends on both performance and coordination, is a crucial element in assessing these patients. The supine bridge, prone bridge, and side bridge tests (left and right) are among the most frequently used methods for evaluating core muscle strength and stability [5]. While these tests provide valuable information for patient evaluation, quantitative measurements of the lumbar spine kinematics during these assessments remain limited. CNLBP patients establish a distinctive and unique lumbar motion signature by completing standardized multiplanar, lumbar-specific functional motion tasks. The motion signature is composed of the three-dimensional (3D) lumbar range of motion. In combination, these lumbar-specific motion signature datasets offer an insight into the patients' musculoskeletal system and can be used as measures of functional health [6].

Inertial measurement units (IMUs) are reliable for capturing the kinematic data associated with movement patterns in these clinical tests [5]. These devices are increasingly integrated into routine clinical practice, with the literature supporting their reliability and validity in assessing lumbar spine kinematics [5].

According to current guidelines for managing chronic non-specific low back pain, interventions for this patient population should include a structured therapeutic exercise program [7]. Therapeutic exercise aids in restoring the functionality of these patients [8,9], as it is a targeted form of exercise that has been shown to alleviate pain and reduce disability [10]. Quantitative data are essential for evaluating the effectiveness of therapeutic exercise programs. The validated accuracy of IMU devices makes them well-suited for assessing kinematics as they capture three-dimensional movement data [5,11].

Several studies have shown that clinical Pilates reduces pain and improves functionality [12–17]. The objective of this study was to evaluate the efficacy of clinical Pilates in enhancing the functional capacity of patients with CNLBP. To achieve this, we incorporated the quantitative metrics obtained from IMUs with three clinical assessment tests: the supine bridge test (SUBT), the side bridge test (SBT), and the prone plank bridge test (PBT). This is the first time, to the extent of our knowledge, that the lumbar motion signature, a critical indicator of functional health, has been combined with functional test data to assess the efficacy of rehabilitation programs from a detailed biomechanical perspective.

2. Materials and Methods

A total of 41 individuals (14 males and 27 females) identified as having chronic low back pain were originally screened. The inclusion criteria comprised the existence of non-specific lumbar spine pain persisting for over 6 months, the absence of pathological radiological findings on MRI of the lumbar spine, negative neurodynamic tests of the lower limbs, and positive response to the written consent form. Exclusion criteria included chronic low back pain resulting from structural or abnormal spinal disorders (spondylolysis, spondylolisthesis, scoliosis, bone fractures, neurological symptoms) and pathologies such as cancer or kidney disease associated with reported low back pain originating from the spine.

Seven patients were excluded from the study due to hip pain, one due to spondylolysis, four due to radiological findings of scoliosis deformity, one due to spinal fracture in the past, one due to osteoporosis, and four due to the presence of spondyloarthropathy in the lumbar spine (Modic type II).

A total of 22 patients with non-specific low back pain participated in this study, comprising 8 men and 14 women.

The lower spine's range of motion (ROM) was measured during the course of four functional tests: the bridge test in both supine and prone postures, and the side bridge test from the right and left sides. Two MetaMotionR+ inertial measurement units (IMUs) were employed to evaluate the movement characteristics of the lumbar spine. The MetaMotionR+ is engineered for the acquisition and transmission of sensory data and carries CE certification. The raw data are recorded and transmitted at 400 Hz and 100 Hz, respectively, via Bluetooth. The raw data were captured and transferred at frequencies of 400 Hz and 100 Hz, respectively, using Bluetooth. Two clinicians acquired and examined the CSV file with the data on a local computer. The data synthesis combined information from a 3D gyroscope, a 3D magnetometer, and a 3D accelerometer to generate a stable directional position system known as quaternion or Euler angles. Furthermore, the sophisticated software seamlessly integrated the raw data from the three devices to optimize the performance of each. This approach encompassed displacement correction algorithms for each sensor, status monitoring for standardization, and the use of a Kalman filter to yield reliable and

accurate direction values. The MMR+ is a scientifically validated apparatus utilized for capturing human movement data and the monitoring of movement patterns. A list of its technological features includes:

- 6-axis accelerometer BMI160 + gyroscope;
- LTR-329ALS ambient light sensor;
- 3-axis magnetometer BMM150;
- 9-axis sensor fusion by BOSCH;
- 8 MB memory;
- Rechargeable lithium-ion battery;
- Vibration motor;
- Low-power Bluetooth, CPU, power button, LED, and GPIO.

A Bluetooth unit integrated into each IMU wirelessly transmitted the data to the patient's smartphone. The IMUs and Bluetooth units were powered by a 3.6V battery. The data collection process included six distinct phases.

2.1. Phase 1—Calibration

Accurate movement capture using an IMU needs the correct alignment between the device's reference frame and the body's orientation. Prior to installation and activation, the units were calibrated on a level horizontal plane to ensure that each unit maintained an identical starting position in accordance with the coordinates of the Cartesian system. Adhering to a strict calibration procedure is essential for acquiring trustworthy data. Calibration was conducted using the Metawear application installed on a smartphone (MbleintLab Inc., 2758 Lantz ave San Jose, CA 95124, USA. software version 2.0.1).

2.2. Phase 2—Unit Installation

The sensors were placed to precisely identify the orientation and location of the device frame according to the natural movement axis of the lumbar spine joints. The first sensor was positioned on the L1 spinous process, and the second on the L5 spinous process. Both units were attached to the skin using non-allergenic two-sided adhesive tape.

2.3. Phase 3—Bluetooth Protocol and Data Transmission

Subsequent to each recording of the specified functional tests, the acquired data were transferred over Bluetooth to a computer for further analysis. The analysis included pre- and post-filtering of the accelerometer, gyroscope, and magnetometer's three-dimensional data in the sensors' local coordinate system.

2.4. Phase 4—Initial Recording Observation

Despite the accuracy of the portable devices and the use of the root mean square error (RMSE), a computer-based body model allows the clinician to verify the correctness of motion reconstruction and data acquisition. Should inaccuracies or erroneous motion records be identified, the process of calibration and measurements would need to be reiterated to guarantee the capture of representative data.

2.5. Phase 5—Data Collection for Functional Tests

Following a successful initial testing, the patients were instructed to execute the four functional bridge assessments. Each participant received detailed directions concerning the process and conducted two practice trials prior to the actual data collection.

The smartphone was kept adjacent to them during the measurement process to avert signal connection issues while capturing and transferring data from the units to the smartphone. Upon completion of the data collection, the clinical physical therapist

detached the sensors from the patients and shared the data obtained from the smartphone to the computer.

2.6. Clinical Pilates Intervention Program

The intervention program consisted of exercises aimed at strengthening the transverse abdominis, rectus abdominis, upper body muscles, erector spinae, hip muscles, and glutes. Specifically, the exercises included the following:

Hundreds lvl 1, Clam lvl 1, Single leg stretch lvl 1, lvl 2, Side kick lvl 1, lvl 2, Double leg stretch lvl 1, Side lift adduction lvl 1, Scissors lvl 1, lvl 2, Dart lvl 1, Single leg circle lvl 1, Swan dive lvl 1, Shoulder bridge lvl 1, Swimming lvl 1, lvl 2, Hip twist lvl 1, lvl 2.

The intensity of the exercises was determined as follows:

Exercises at stage 1 (lvl 1) were primarily easy and performed in a single-axis direction. Following these were exercises at stage 2 (lvl 2), which were more challenging than those at stage 1 due to additional movement steps (cognitively demanding), but still performed in a single-axis direction [8].

It is important to emphasize that all exercises began at stage 1 difficulty and progressively increased to stage 2 as each individual gained good control and stabilization.

The therapeutic exercise program was conducted three times a week for a duration of 6 weeks.

3. Results

For the statistical analysis, an independent samples t-test was used with a one-tailed approach and allowance for unequal variances (heteroscedasticity) between the two samples. The study's statistical power was calculated using G*Power 3.1.9.7 software for Windows (Heinrich-Heine-Universität Düsseldorf, Düsseldorf, Germany). A power analysis for two independent means was performed with a significance level of $\alpha = 0.05$. The statistical power related to sample size and the observed effect size exceeded 80% for each variable (92.7% for the prone bridge test, 89.3% for the left side bridge test, and nearly maximal at 98.3% for both the supine and right-side bridge tests).

The intraclass correlation coefficient (ICC, two-way random, absolute agreement) with the corresponding 95% confidence interval was used to assess the relative reliability, providing insight into the consistency of within-session measurements across all bridge tests and clinical passive range of motion assessments (ICC > 0.90).

3.1. Supine Bridge Test (SBT)

The position achieved during the performance of the supine bridge test involved lumbar spine extension (compared with the anatomical position), with a mean extension of 11.86 and a standard deviation of 1.80 before the start, and a mean extension of 14.27 and a standard deviation of 1.33 after the completion of the therapeutic exercise program (p -value = 0.0003) (Figure 1).

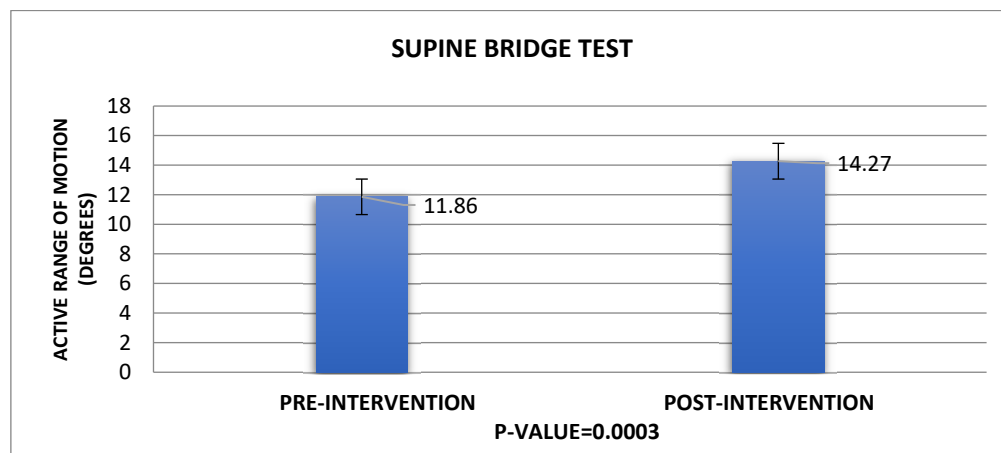


Figure 1. Kinematic results of the supine bridge test.

3.2. Prone Bridge Test (PBT)

The position achieved during the performance of the prone bridge test involved lumbar spine extension (compared with the anatomical position), with a mean extension of 6.23 and a standard deviation of 2.36 before the start, and a mean extension of 3.95 and a standard deviation of 1.17 after the completion of the therapeutic exercise program (p -value = 0.001) (Figure 2).

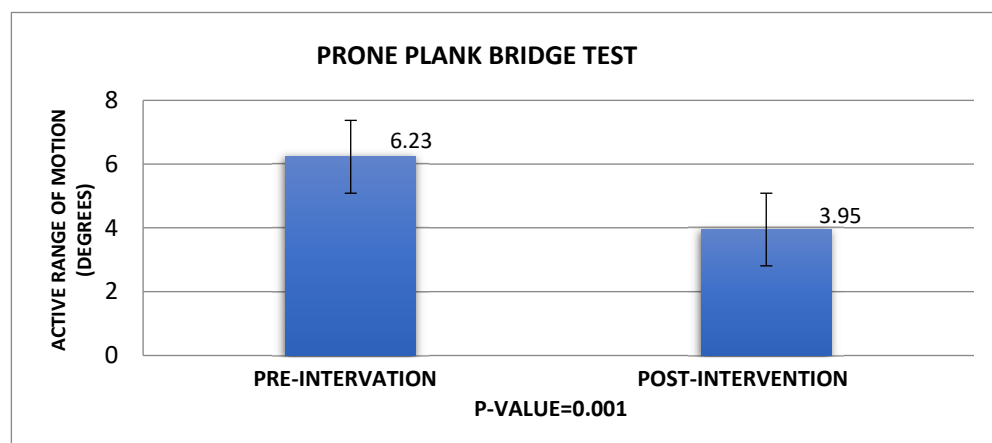


Figure 2. Kinematic results of the prone plank bridge test.

3.3. Side Bridge Test Left (SBTL)

The position achieved during the performance of the left side bridge test involved right lateral flexion of the lumbar spine (compared with the anatomical position), with a mean value of 3.96 and a standard deviation of 1.19 before the start, and a mean value of 2.47 and a standard deviation of 1.21 after the completion of the therapeutic exercise program (p -value = 0.0004) (Figure 3).

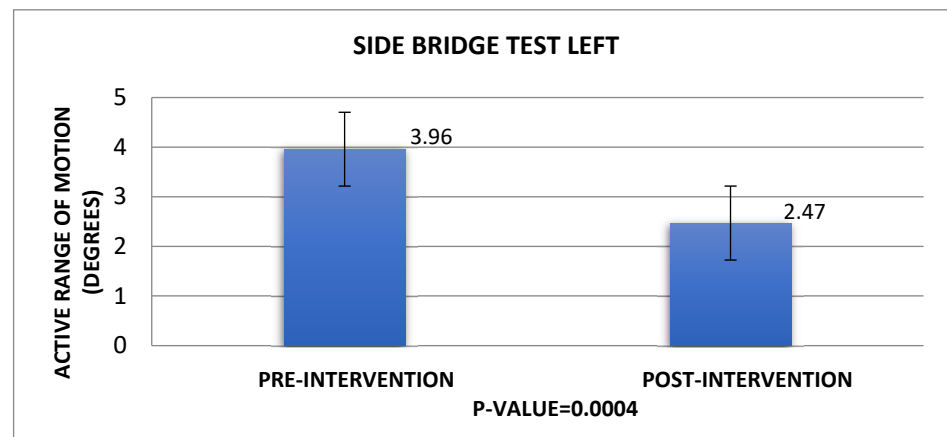


Figure 3. Kinematic results of the side bridge test (left).

3.4. Side Bridge Test Right (SBTR)

The position achieved during the performance of the right-side bridge test involved left lateral flexion of the lumbar spine (compared with the anatomical position), with a mean value of 3.52 and a standard deviation of 0.81 before the start and a mean value of 2.27 and a standard deviation of 1.49 after the completion of the therapeutic exercise program (p -value = 0.00001) (Figure 4).

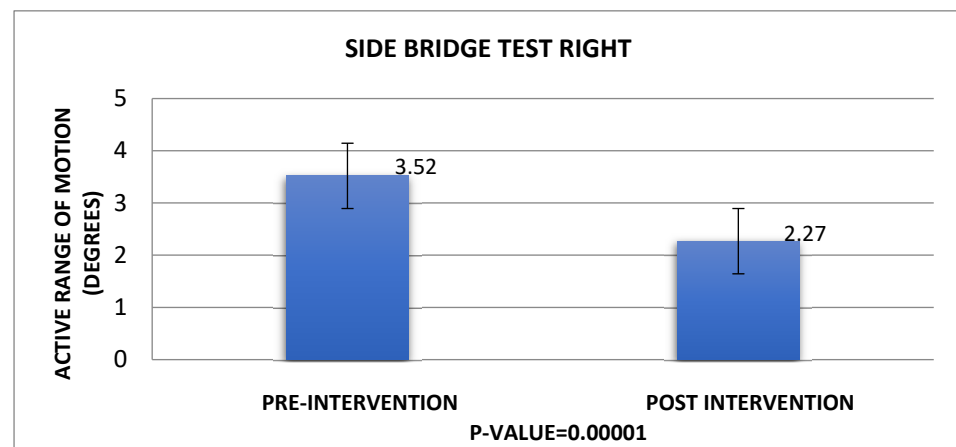


Figure 4. Kinematic results of the side bridge test (right).

3.5. Wearable Sensors—Functional Test Duration Data

A statistically significant improvement was observed in the time that the final position was maintained in all tests (and was not abandoned due to pain or fatigue). Specifically, during the performance of the supine bridge test, patients maintained the final position for $72.3/\pm 29.43$ (mean value (s)/ \pm standard deviation) before the intervention, while after the intervention, the recorded values increased to $162.5/\pm 31.24$ (mean value (s)/ \pm standard deviation), p -value = 0.001.

During the performance of the prone bridge test, patients maintained the final position for $34.31/\pm 20.51$ (mean value (s)/ \pm standard deviation) before the intervention, increasing these values after the intervention to $65.3/\pm 25.45$ (mean value (s)/ \pm standard deviation), p -value = 0.00003.

Regarding the performance of the left side bridge test, the recorded values for the final position were $10.2/\pm 8.49$ (mean value (s)/ \pm standard deviation) before the intervention, while after the intervention, the values improved to $27.42/\pm 4.33$ (mean value (s)/ \pm standard deviation), p -value = 0.002.

Finally, during the performance of the right-side bridge test, the recorded values for the final position were $9.22/\pm 7.34$ (mean value (s)/ \pm standard deviation) before the intervention and $29.19/\pm 4.45$ (mean value (s)/ \pm standard deviation) after the intervention, respectively, p -value = 0.0001 (Table 1).

Table 1. Time results pre- and post-intervention for each assessment test.

Assessment Test	Pre-Intervention (Average Value (s)/ \pm Standard Deviation)	Post-Intervention (Average Value (s)/ \pm Standard Deviation)	p -Value
SUBT	72.3/ \pm 29.43	162.5/ \pm 31.24	0.001
PBT	34.31/ \pm 20.51	65.3/ \pm 25.45	0.00003
SBT(L)	10.2/ \pm 8.49	27.42/ \pm 4.33	0.002
SBT(R)	9.22/ \pm 7.34	29.19/ \pm 4.45	0.0001

4. Discussion

This study explored the effectiveness of clinical Pilates intervention for individuals with chronic non-specific low back pain, focusing on improvements in lumbar spine functionality and mobility. CNLBP is among the most prevalent musculoskeletal conditions globally, significantly diminishing quality of life due to the pain and functional limitations it causes. Key factors influencing the condition's progression include psychosocial and physical elements and the reduced endurance of core muscles [17].

In this context, clinical Pilates therapeutic exercises offer a personalized approach to reduce pain and enhance physical function [1]. Clinical Pilates has shown greater effectiveness than other treatments, such as pharmacotherapy and traditional lumbar stabilization exercises, in addressing CNLBP [18–21]. This study used inertial measurement units (IMUs) to accurately record lumbar spine movements during four functional bridge tests in various positions: supine, prone, and side bridges (right and left). IMUs provided precise data on lumbar spine mobility across three axes, contributing to a reliable assessment of movement [5,22–24].

The study's results demonstrated significant enhancements in hold duration and lumbar spine mobility subsequent to the intervention.

The findings indicated increased lumbar extension, decreased flexion, and greater ability to sustain proper trunk position during the tests. These outcomes suggest improved muscular endurance and superior spinal muscle control.

Moreover, women exhibited increased susceptibility to CNLBP [25–28], while individuals with a higher BMI presented greater vulnerability to the condition [29].

The study's limitations included a limited sample size and potential IMU errors, possibly due to skin movement (artifacts) [5,30–32]. Despite these limitations, the findings suggest that clinical Pilates interventions are beneficial in the management of CNLBP, highlighting the necessity for additional research to determine optimal exercise protocols and evaluate the intervention's efficacy across various age groups [33].

5. Conclusions

Therapeutic exercise, particularly clinical Pilates, has been proven to be effective in alleviating pain and enhancing functional capacity in individuals with CNLBP, as supported by the existing literature [19,34]. This study further validates these outcomes through temporal and kinematic data collected via IMUs, which, due to their compact size, ease of use, and mobility features, provide reliable and objective assessment data for various clinical tests [5]. Following a six-week intervention, notable improvements were observed in spinal mobility

and core muscle endurance. Ongoing research is essential to examine the applications of wearable sensors in chronic conditions combined with surface electromyography [35] to assess the long-term effects and specific types of therapeutic exercise.

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Institutional Review Board Statement: The study was conducted in accordance with the Declaration of the University of Peloponnese and approved by the Research Ethics Committee of the School of Health Sciences, University of Peloponnese (protocol code 6367 and date of approval 19 March 2024).

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study. Written informed consent was obtained from the patients to publish this paper.

Data Availability Statement: The data of this research could be made available upon request.

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