Correlation between lumbar sagittal alignment, Modic changes, and endplate erosion in patients with degenerative disease

CURRENT STATUS: POSTED

Bin Lv
The affiliated people's with Jiangsu University
dr_lvbin@sina.com
ORCiD: https://orcid.org/0000-0002-4368-0232

Haosheng Wang
Jilin University

10.21203/rs.3.rs-18890/v1

SUBJECT AREAS
Orthopedics

KEYWORDS
lumbar sagittal alignment, Modic change, endplate defect, clinical outcomes
Abstract
Purpose To explore the role of lumbar sagittal alignment in the occurrence of Modic changes and endplate defects (MC&ED) development in patients with a spinal degenerative disease, and the relationship between lumbar sagittal alignment and patient-report outcomes.

Background Increasing attention has been focused on MC&ED as playing a potential role in the etiopathogenesis of lumbar degeneration. The precise understanding of the mechanisms leading to progression of MC&ED is lacking. Hence, we investigated how lumbar sagittal alignment influences the MC&ED.

Patients and methods Ninety-six consecutive asymptomatic or symptomatic patients with Modic changes or endplate defect were retrospectively recruited in this study from August 2016 to December 2018. MC&ED were observed in 76 patients and not observed in 20 patients, representing two groups for comparison. The lumbar sagittal parameters were measured, including lumbar lordosis (LL), pelvic incidence (PI), sacral slope (SS), and pelvic tilt (PT). The lumbar lordosis index (LLI) and idea LL were then calculated. Clinical outcomes were assessed using a visual analog scale (VAS) and a Oswestry Disability Index (ODI) before and after operation.

Results There were no significant differences in the distribution of demographics and baseline clinical variables between both groups. Mean age and BMI showed a significant difference between both groups (P<0.05). There were significant correlations between LL, LLI, Lossof LL, and Level 1 (r=0.281, 0.230, and 0.284, P<0.05) Also, PI, PT were significantly correlated with Level 4 (r=0.249, 0.202, P<0.05). Compared with presurgery scores, an improvement was seen in postoperative VAS and ODI scores (P<0.05). Further, the postoperative scores at 24 months in the without Modic or end plate defect group showed greater improvements compared with the with Modic or endplate defect group (P<0.05).

Conclusion This analysis indicated that maintaining lumbar sagittal alignment was related to a lower risk of Modic changes in patients with the spinal degenerative disease. The lumbar sagittal alignment might be a factor that influenced the posterior inclination of the pelvis in symptomatic lumbar disease.
Introduction
Degenerative pathologies of the lumbar spine, including Modic changes and endplate defect, are common in the elderly population.[1] Modic changes (MCs) are shown as MRI signal variation in vertebral and subchondral bone marrow adjacent to the endplate.[2] MCs are assigned into three subtypes: Type I, II, and III according to T1- and T2-weighted MRI.[3, 4] Radiological and histological examination analysis have revealed that different subtypes of MCs respectively displayed inflammatory responses, fatty marrow, and subchondral bone sclerosis. MCs are frequently observed in pathological structure changes like disc degeneration, disc herniation, and endplate defect.[5–7] A recent study demonstrated that there is an association between endplate defect and disc degeneration in a large-scale, population-based study. Several theories have been proposed with respect to the etiology of MCs and endplate defect. Recently, clinical studies have paid attention to endplate defects as acting as a concomitant role in the aetiopathogenesis of MCs.[8] Korres et al. demonstrated that the lumbar spine acts a role in maintaining the upright posture and forming the body trunk shape via maintaining proper sagittal alignment.[9] In fact, sagittal alignment has been related to various lumbar conditions. There are several reports demonstrated that sagittal malalignment contributes to lumbar disc degeneration. Buckland et al. reported that worsening sagittal malalignment associated with lumbar stenosis on stereoradiographs.[10] Ogura et al. showed that sagittal spinopelvic alignment as a significant impact on health-related quality of life and clinical outcomes after surgery.[11] Recent studies have demonstrated that patients with lumbar malalignment are frequently present sympathetic symptoms such as low back pain and. The motion segment is composed of adjoining vertebrae, the lumbar disc, and adjoining ligaments between them, of which the endplate, located between the vertebrae and the intervertebral disc, is considered to be the most vulnerable structure.[12] It is obvious that both MCs and endplate defect is related to the lumbar degeneration. Therefore, it is considered and the lumbar sagittal malalignment as a transitory stage of a lumbar degenerative process. However, the role of lumbar sagittal alignment in the progression of Modic changes and endplate defects has not been illuminated. In the current study, we supposed that lumbar sagittal alignment
patterns may be associated with the development of either Modic changes or endplate defects. Thus, we explored whether differences in sagittal parameters and especially spinopelvic parameters, could predict the development of either Modic changes or endplate defects.

Materials And Methods

Study Population

This prospective study enrolled 96 symptomatic volunteers (age: mean 56, range 27–83; 44 males and 52 females) from August 2016 to December 2018. Written informed consent was got from all volunteers. This study was approved by the ethics board of the Affiliated people's Hospital of Jiangsu University. All volunteers were diagnosed with low back pain and underwent an MRI examination according to standardized protocols. Exclusion criteria were: (1) history of lumbar surgery; (2) old vertebral body fracture; (3) abnormal bone metabolism including cancer, infection, and trauma; and (4) high-grade spondylolisthesis. Inclusion criteria were: (1) age ≥ 25 years; (2) suffered from chronic low back pain more than 3 months after conservative medication; and (3) confirmed by lumbar MRI. 76 patients diagnosed with MC&ED were enrolled in the MC&ED groupand the remaining 20 without MC&ED were enrolled in a MC&ED group. Demographic data including age, gender, body mass index (BMI), smoking status, and Charlson comorbidity index (CCI) were collected.

Lumbar sagittal parameters

Sagittal and axial images were obtained on MRI scans from T12 upper endplate margin to S1 lower endplate margin. The measurements of the lumbar and pelvis sagittal parameters were: (1) lumbar lordosis (LL), LL was measured from the L1 superior endplate to S1 superior; (2) pelvic incidence (PI): the angle between the line perpendicular to the sacral plate and the line connecting the midpoint of the sacral plate to the bicoxofemoral axis; (3) sacral slope (SS): the angle between the S1 superior endplate and a horizontal line; (4) pelvic tilt (PT): the angle between a vertical line originating at the center of the bicoxofemoral axis and a line drawn between the same point and the middle of the superior endplate of S1; (5) lumbar lordosis index (LLI) is equal to LL divided by PI; (6) idea LL = (PI × 0.5481 + 12.7) × 1.087 + 21.61, Loss of LL = ideaLL - LL. (Fig. 1)

Assessment of endplate defects
The loss or disruption of the complete appearance was defined as endplate defects. Change of endplate defects visible on sagittal MR images was categorized into three grades according to Feng et al.[13] Type 1 endplate defect, focal endplate defect, was observed as a discontinuity or hollow on the endplate. Type 2 endplate defect, corner endplate defect, was observed as a legible corner on the endplate. Type 3 endplate defect, erosive endplate defect, observed as the extensively irregular, serrated, or worm-eaten appearance.[14] Inter-rater agreement for the coding of endplate defect was calculated using Cohen's weighted kappa and Pearson's correlation.

Assessment of MCs

MCs were categorized into type 0, I, II and III using sagittal multi-positional MRI image. Type 0 MCs showed normal endplate, type I MCs showed low T1 and high T2 signal, type II MCs showed high T1 and T2 signals, and type III MCs showed low T1 and T2 signals.

Statistical analysis

All statistical analyses were performed using SPSS (version, 21.0 SPSS Inc, Chicago, IL). Demographic characteristics were expressed as means ± standard deviations (x ± s) for continuous variables. Categorical variables were expressed as proportions and percentages. Mann-Whitney U test or chi-square test was used to compare parameters between groups. Moreover, Pearson correlation and linear regression analysis were also performed to analyze the relationship between lumbar sagittal alignment and the clinical outcomes. P less than 0.05 was considered statistically significant.

Results

Demographics details

Distribution of demographic variables such as age, weight, smoking status, and duration of symptoms in the two groups was significantly different (P > 0.05); there was no significant difference in gender and BMI (P < 0.05). Moreover, Modic changes and/or endplate defects observed in 79.16% of volunteers in group A (76/96), and in 20.83% of volunteers in group B (20/96), indicating no statistical difference in occurrence among groups. (Table 1)
Table 1
Demographics details of the with/without MCs or endplate defect groups

| Parameter          | with MC&ED | without MC&ED | P-value |
|-------------------|------------|---------------|---------|
| Number            | 76         | 20            |         |
| Mean age (years)  | 57.25 ± 14.02 | 49.7 ± 10.25  | < 0.05  |
| Gender            |            |               | > 0.05  |
| Male              | 34         | 10            |         |
| Female            | 42         | 10            |         |
| Body weight (kg)  | 67.15 ± 10.81 | 68.45 ± 7.91  | > 0.05  |
| Height (cm)       | 166.07 ± 7.56 | 164.80 ± 8.35 | > 0.05  |
| BMI (kg/m²)       |            |               | < 0.05  |
| < 30              | 13         | 6             |         |
| > 30              | 63         | 14            |         |
| smoking consumption | 17     | 5             | > 0.05  |
| Alcohol consumption | 23    | 6             | > 0.05  |
| CCI               | 2.11 ± 0.89 | 2.87 ± 1.04   | > 0.05  |

Lumbar sagittal parameters

We further compared several lumbar sagittal parameters between both groups. PI in group A was 51.46°±10.29° and in group B was 49.7°±10.26° (P > 0.05). PT in group A was 17°±6.97° and in the group B was 16.9°±6.9° (P > 0.05). SS in group A was 34.48°±8.07° and in group B was 32.81°±8.35° (P < 0.05). LL in group A was 40.50°±11.20° and in group B was 37.90°±10.43° (P < 0.05). LLI in group A was 0.79 ± 0.17 and in group B was 0.77 ± 0.20 (P < 0.05). Loss of LL in group A was 25.59°±8.77° and in the group, B was 27.12°±8.61° (P < 0.05). (Table 2)

Table 2
Comparison of the lumbar sagittal alignment between the with MCs or endplate defect and without MCs or endplate defect groups

| Parameter | with MC&ED         | without MC&ED        | P-value |
|-----------|--------------------|----------------------|---------|
| PI        | 51.46 ± 10.29      | 49.7 ± 10.26         | > 0.05  |
| PT        | 17 ± 6.97          | 16.9 ± 6.9           | > 0.05  |
| SS        | 34.48 ± 8.07       | 32.81 ± 8.35         | < 0.05  |
| LL        | 40.50 ± 11.20      | 37.90 ± 10.43        | < 0.05  |
| LLI       | 0.79 ± 0.17        | 0.77 ± 0.20          | > 0.05  |
| Loss of LL| 25.59 ± 8.77       | 27.12 ± 8.61         | > 0.05  |

Clinical outcomes

The postoperative VAS and ODI scores at 3, 6, 12 and 24 months differed significantly from preoperative scores in two groups (P < 0.05). However, 24-month postoperative VAS and ODI scores demonstrated improvement in the with Modic or endplate defect group compared with the without Modic or endplate defect group. VAS and ODI scores decreased at 3 and 6 months postoperative and then fell back slightly at 12 and 24 months postoperative (P < 0.05; Table 3).
### Table 3
clinical outcomes before and after surgery

| Time                  | VAS score          | ODI score          |
|-----------------------|--------------------|--------------------|
| **with MCs or endplate defect** |                    |                    |
| Presurgery            | 5.74 ± 4.55*       | 24.12 ± 2.62*      |
| 3 months postsurgery  | 2.88 ± 2.55*       | 21.47 ± 3.31*      |
| 6 months postsurgery  | 2.25 ± 2.36*       | 17.03 ± 3.46*      |
| 12 months postsurgery | 2.71 ± 2.45*       | 11.25 ± 1.61*      |
| 24 months postsurgery | 3.43 ± 1.31*       | 14.93 ± 1.86*      |
| **without MCs or endplate defect** |                    |                    |
| Presurgery            | 4.84 ± 4.77**      | 21.43 ± 2.25**     |
| 3 months postsurgery  | 2.53 ± 2.43**      | 19.47 ± 2.62**     |
| 6 months postsurgery  | 1.78 ± 1.89**      | 13.03 ± 2.76**     |
| 12 months postsurgery | 2.54 ± 2.45**      | 10.75 ± 1.32**     |
| 24 months postsurgery | 3.02 ± 2.26**      | 12.93 ± 2.02**     |

Note: *P < 0.05 vs. pre-surgery; †P < 0.05 vs. with MCs or endplate defect group.

### Table 4
Spearman’s rank correlation coefficients between the lumbar sagittal parameters and clinical outcomes

| PI  | PT  | SS  | LL  | LLI | Loss of LL | Level 1 | Level 2 | Level 3 | Level 4 | Level 5 | Level S1 | ED  |
|-----|-----|-----|-----|-----|-----------|---------|---------|---------|---------|---------|---------|-----|
| PI  | R   | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.174 | 0.375 | 0.113     | 0.352   | 0.855   | 0.014   | 0.805   | 0.080   | 0.473   |     |
| PT  | R   | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| SS  | R   | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| LL  | R   | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| LLI | R   | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Loss of LL | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Level 1 | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Level 2 | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Level 3 | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Level 4 | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   |     |
| Level S1 | R | 1   |     |     |           |         |         |         |         |         |         |     |
| P   | 0.000 | 0.000 | 0.000 | 0.000 | 0.000     | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000   | 0.000 |

Note: **P < 0.01, *P < 0.05.

Abbreviations: PI, pelvic incidence; PT, pelvic tilt; SS, sacral slope; LL, lumbar lordosis; LLI, lumbar lordosis index; ED, Endplate defect.

**Discussion**
In this study, we showed two novel findings. First, patients with improper lumbar sagittal alignment had a higher occurrence of Modic and endplate defect.

Previous studies regarding sagittal balance have demonstrated that lumbar malalignment after spine surgery may cause poor clinical outcomes. Patients should be provided proper correction of the spinal alignment in order to prevent complications to occur. However, there is no studies focus attention on the effect of lumbar sagittal alignment on the progression of Modic changes and endplate defects before surgery until now. Thus, we conducted a study that mainly focused on the effect of lumbar sagittal alignment on the occurrence of Modic changes and endplate defects.

We think that the mechanism regarding the association between lumbar sagittal alignment and Modic changes or endplate defects is multifaceted. The lumbar segment with MCs or endplate defects tends to lumbar disk degeneration, the latter of which destabilizes the lumbar spine in turns and ultimately form a vicious cycle. Moreover, the vertebral endplate fulfills the critical function of disc nutrient transport. MCs decrease nutrition transmit ability of endplate, infection factors serving as initial pathogenesis. In addition, morphological changes such as fatty infiltration or change in cross-sectional area of lumbar muscles were observed in patients with low back pain. Patients with low back pain demonstrate a significantly higher prevalence of MCs and endplate defect with an unclear mechanism. We hypothesize that the change of lumbar sagittal alignment leads to lumbar muscles of these MCs patients fail to support spinal stability and finally led to fatty infiltration or change in cross-sectional area of lumbar muscles.

The unstable lumbar spine is able to enlarge the range of movement, which accelerates the degeneration of endplate and lumbar disc. Finally, the interaction between the two makes the disease to develop into a serious direction. In this study lumbar sagittal parameter was measured in patients with low back pain and we found that lumbar sagittal parameter in volunteers with MCs or endplate defect was larger than those in subjects without MCs or endplate defect. A significant correlation between lumbar sagittal parameter, MCs, and endplate defect was found in logistic regression analysis. We speculated that increased lumbar sagittal alignment could increase lateral shear force, leading to the destroyed structure of endplate and formation of MCs.
In terms of the Modic and endplate defect, the prevalence of Modic and endplate defect was 79.16% (76/96) in with Modic and endplate defect group and 21.5% (20/96) in without Modic and endplate defect group. Meanwhile, differences in the presurgical lumbar sagittal alignment were observed in two groups. Thus, we hypothesize that presurgical lumbar sagittal alignment, measured by PI, PT, SS, LL, and PI-LL, was related to the incidence of Modic and endplate defect in symptomatic subjects.

In terms of the lumbar parameters, the results indicated that PT, SS, and PI-LL in the without Modic and endplate defect group were significantly higher than those in the Modic and endplate defect group, whereas xx and xx in both groups demonstrated no significant difference. To the author's knowledge, no previous literature focuses attention on the relationship between lumbar sagittal alignment and without Modic and endplate defect. It is generally accepted that physiological cervical sagittal balance is important for normal spine function, and the maintenance of and improvements to cervical spine function are bound to have an effect on clinical efficacy and outcomes.25,26 As a result, Modic changes and endplate defect may be caused by malalignment of the lumbar spine. However, previous studies emphasized that there exist a relationship between sagittal alignment and degenerative changes only in the cervical spine.[17] In terms of the lumbar spine, Baron et al. showed that spinopelvic sagittal alignment plays a predisposing role in the pathogenesis of lumbar degeneration.[18]

Even though we couldn't confirm whether these characteristics were the effect or the predictor of lumbar degeneration in the current retrospective study, they have potential clinical implications. Lumbar sagittal alignment plays a crucial role in predicting the development of Modic changes and endplate defects in symptomatic patients. Meanwhile, in the corrective surgery for patients with lumbar deformity, emphasis should be placed on the reconstruction of the lumbar sagittal alignment. Furthermore, lower lumbar muscles dystrophy because of a surgical exposure can contribute to postoperative deformities. Li et al. reported that there is a correlation between spinopelvic alignment and degeneration of lower lumbar paraspinal muscles in elderly patients in a recent study.[19] Hence, the paraspinal muscle-strengthening exercises of lower lumbar levels could slow the progression of Modic changes and endplate defects. Longitudinal studies with subject outcomes need to be
performed to validate our hypotheses.

Several limitations existed in this study. First, bias could not be avoided due to the nature of retrospective studies. Second, the study mainly focused on preoperative lumbar sagittal alignment and prevalence of Modic changes and endplate defects and not postoperative lumbar alignment or changes in lumbar alignment. Third, a high standard deviation of lumbar sagittal parameters reflects the flexibility of the lumbar spine. Thus, we should determine the reproducibility of lumbar sagittal parameters in further research. In addition, the lack of patient medical records made it difficult to verify the differences in clinical scores between patients with the imbalance and normal balance.

Conclusion
Lumbar sagittal alignment is a good predictor of the development of Modic changes and endplate defects for the association between them. Hence, maintaining a proper lumbar sagittal alignment was important in lumbar surgery, which seems to act as a vital role in the improved clinical outcomes.

Declarations
Acknowledgment
This study was sponsored by the Clinical Medicine Science and Technology Development Foundation of Jiangsu University (educational reform series Grant No: 14).

Conflict of interest
The authors declare that they have no conflict of interest.

References
1. Wang Y, Videman T, Battie MC. Modic changes: prevalence, distribution patterns, and association with age in white men. Spine J. 2012;12(5):411-416.
2. Paholpak P, Dedeogullari E, Lee C, et al. Do modic changes, disc degeneration, translation and angular motion affect facet osteoarthritis of the lumbar spine. Eur J Radiol. 2018;98:193-199.
3. Modic MT, Steinberg PM, Ross JS, Masaryk TJ, Carter JR. Degenerative disk disease: assessment of changes in vertebral body marrow with MR imaging. Radiology. 1988;166(1 Pt 1):193-199.
4. Modic MT, Masaryk TJ, Ross JS, Carter JR. Imaging of degenerative disk disease. *Radiology.* 1988;168(1):177-186.

5. Maatta JH, Karppinen JI, Luk KD, Cheung KM, Samartzis D. Phenotype profiling of Modic changes of the lumbar spine and its association with other MRI phenotypes: a large-scale population-based study. *Spine J.* 2015;15(9):1933-1942.

6. Mok FP, Samartzis D, Karppinen J, Fong DY, Luk KD, Cheung KM. Modic changes of the lumbar spine: prevalence, risk factors, and association with disc degeneration and low back pain in a large-scale population-based cohort. *Spine J.* 2016;16(1):32-41.

7. Zehra U, Cheung JPY, Bow C, Lu W, Samartzis D. Multidimensional vertebral endplate defects are associated with disc degeneration, modic changes, facet joint abnormalities, and pain. *J Orthop Res.* 2018.

8. Rade M, Maatta JH, Freidin MB, Airaksinen O, Karppinen J, Williams FMK. Vertebral Endplate Defect as Initiating Factor in Intervertebral Disc Degeneration: Strong Association Between Endplate Defect and Disc Degeneration in the General Population. *Spine.* 2018;43(6):412-419.

9. Vazifehdan F, Karantzoulis VG, Igoumenou VG. Sagittal alignment assessment after short-segment lumbar fusion for degenerative disc disease. *International Orthopaedics.* 2018.

10. Buckland AJ, Ramchandran S, Day L, et al. Radiological lumbar stenosis severity predicts worsening sagittal malalignment on full-body standing stereoradiographs. *Spine J.* 2017;17(11):1601-1610.

11. Ogura Y, Shinozaki Y, Kobayashi Y, et al. Impact of sagittal spinopelvic alignment on clinical outcomes and health-related quality of life after decompression surgery without fusion for lumbar spinal stenosis. *J Neurosurg Spine.* 2019:1-6.

12. Zhao FD, Pollintine P, Hole BD, Adams MA, Dolan P. Vertebral fractures usually affect
the cranial endplate because it is thinner and supported by less-dense trabecular bone. *Bone.* 2009;44(2):372-379.

13. Feng Z, Liu Y, Yang G, Battie MC, Wang Y. Lumbar Vertebral Endplate Defects on Magnetic Resonance Images: Classification, Distribution Patterns, and Associations with Modic Changes and Disc Degeneration. *Spine.* 2018;43(13):919-927.

14. Zehra U, Cheung JPY, Bow C, Lu W, Samartzis D. Multidimensional vertebral endplate defects are associated with disc degeneration, Modic changes, facet joint abnormalities and pain. *J Orthop Res.* 2018.

15. Cho JH, Joo YS, Lim C, Hwang CJ, Lee DH, Lee CS. Effect of one- or two-level posterior lumbar interbody fusion on global sagittal balance. *Spine J.* 2017;17(12):1794-1802.

16. Farrokhi MR, Yadollahikhales G, Gholami M, Mousavi SR, Mesbahi AR, Asadi-Pooya AA. Clinical Outcomes of Posterolateral Fusion vs. Posterior Lumbar Interbody Fusion in Patients with Lumbar Spinal Stenosis and Degenerative Instability. *Pain physician.* 2018;21(4):383-406.

17. Tamai K, Romanu J, Grisdela P, Jr., et al. Small C7-T1 lordotic angle and muscle degeneration at C7 level were independent radiological characteristics of patients with cervical imbalance: a propensity score-matched analysis. *Spine J.* 2018;18(9):1505-1512.

18. Zarate-Kalfopulos B, Reyes-Tarrago F, Navarro-Aceves LA, et al. Characteristics of Spinopelvic Sagittal Alignment in Lumbar Degenerative Disease. *World Neurosurg.* 2019.

19. Li Q, Sun J, Cui X, Jiang Z, Li T. Analysis of correlation between degeneration of lower lumbar paraspinal muscles and spinopelvic alignment in patients with osteoporotic vertebral compression fracture. *Journal of back and musculoskeletal rehabilitation.* 2017;30(6):1209-1214.
Figures
Figure 1

The lumbar sagittal alignment of the lumbar spine. Abbreviations: PI, pelvic incidence; PT,
pelvic tilt; SS, sacral slope; LL, lumbar lordosis.

Figure 2

VAS and ODI scores before and after surgery. Note: *P<0.05 vs. pre-surgery; † P<0.05 vs. with MC&ED group.