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The Association between Body Mass Index (BMI) and Back or Leg Pain in Patients with Spinal Conditions: Results from the Genodisc Study

--Manuscript Draft--

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The Association between Body Mass Index (BMI) and Back or Leg Pain in Patients with Spinal Conditions: **Results from the Genodisc Study**

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The Genodisc Study was conducted in accordance with the ethical standards of United Kingdom National Research Ethics Service (ethical approval number in the UK 09/H0501/95) and with the Helsinki Declaration of 1975, as revised in 2000. The study was also approved by local Research Ethics Committees in each recruiting country.

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ABSTRACT

Study design: Prospective observation study

Objective: To identify the relationship between obesity, quantified by body mass index (BMI), and both back and leg pain in spinal patients.

Summary of Background Data: Obesity and back pain are massive public health problems. Given the poor correlation between pain and an pathological change in the spine in the spine, further investigation is required into other, non-pathological, predictors such as obesity.

Methods: The Genodisc Study was one of the largest cross-sectional studies of patients presenting to tertiary spinal units and recruited from six centres in four European countries. In total, 2636 patients were recruited over a 5-year period between 2008 and 2013. Both back and leg pain were scored by patients in the range of 0 to 10. Linear regression was used to model the relationship between BMI and pain. Potential confounders included in the model were: age, Zung Depression score, episodes of sport, gender, disability benefit, family history, previous surgery, smoking status, work type, clinical diagnosis and relevant co-morbidities. Back and leg pain outcomes were modelled separately.

Results: The study included 1160 men and 1349 women with a mean age of 50.9 years and mean BMI of 27.2kg/m². In our fully adjusted model, a 5-point increase in BMI was associated with greater leg (0.19 units [95% CI 0.08,0.31]) but not back (0.10 units [95% CI -0.02,0.22]) pain scores. Although this relationship was statically significant, given the small magnitude of the relationship, the clinical significance is limited. Similarly, female gender, heavy workload, rheumatoid arthritis, previous spine surgery and depression were associated with higher back and leg pain.

Conclusion: In this large observational study of spine patients presenting to tertiary European centres, obesity, as measured by increased body mass index, was associated with greater leg pain.

Key Words: low back pain; sciatica; leg pain; obesity; body mass index; epidemiology

KEY POINTS

1. An increase in BMI was associated with higher back (non-significant) and leg pain (significant) scores.
2. Other factors that were associated with greater BP or LP were female gender, previous spine surgery, heavy workload, rheumatoid arthritis and depression.
3. Back pain was associated with a greater number of significant predictors than LP possibly because it is an umbrella term for poorly defined conditions.

MINI ABSTRACT

The analysis of 2636 patients presenting to tertiary European spine centres showed an increase in BMI was associated with higher back (non-significant) and leg pain (significant) scores. Other factors that were associated with greater BP or LP were female gender, previous spine surgery, heavy workload, rheumatoid arthritis and depression.

INTRODUCTION

Obesity and back pain are massive public health problems. The 2010 Global Burden of Diseases Study found low back pain to be the leading cause of disability worldwide with a global point prevalence of 9.4-11.9%.[1, 2] As an outcome of the Global Burden of Diseases Study, there has been an urgent call for further research to understand the predictors of low back pain.[1, 3]

Given the poor correlation between pain and an pathological change in the spine in the spine,[4] further investigation is required into other, non-pathological, predictors such as obesity. From population-based studies, it has been established that BMI increases the odds of low back and leg pain.[5, 6] However, these studies consider pain as a binary outcome and provide little information for the effect of obesity upon the severity of pain. Furthermore, there is limited information as to the relationship between BMI and pain in the patient population seen in a tertiary care setting. It is important to understand the contributors to back and leg pain in this population, as these are the patients who present to surgeons, rheumatologists and physiotherapists daily.

The primary aim of our study was to define a relationship between obesity and both low back (BP) and leg pain (LP) scores separately in a large population of patients presenting to tertiary spine centres. We also present data on other predictors of low back and leg pain.

PATIENTS AND METHODS

Genodisc Study Design

Patients were recruited as part of the pan-European Genodisc study[7] All patients presenting to six tertiary spine care centres in four countries; UK, Hungary, Slovenia and Italy, were invited to participate. In total, 2636 patients were recruited over a 5-year period (2008-2013).

Pain Scores

The Genodisc Participant Survey collected patient reported information for both back and leg pain. It specifically asked participants to rate their pain experienced in the preceding week and score it on an scale ranging from 0, meaning no pain, to 10 being the worst pain imaginable. Back pain was defined as pain in the lower back. Participants were asked to score LP that went below their knee, in an attempt to differentiate true radicular LP from other causes, such as hip osteoarthritis.

Other Participant Data

Demographic and patient reported information including age, gender, height and weight (from which BMI was calculated), co-morbidities, smoking status, occupation, family history, previous surgery and disability benefit were also collected. As an assessment of mood, participants were also asked to complete the Zung Self Reported Depression Score (Zung).

Statistical Methods

Stata 13.1 (Stata, College Station, Tx, USA) was used for all statistical analysis. Univariate analysis was initially performed. For the multivariate models, the predictor variable was BMI. The outcome variables, BP and LP scores were modelled separately as continuous variables. Other confounders included in the model were age, Zung Depression score, number of sporting activities per week, gender, disability benefit, family history, previous surgery, smoking status, work type, clinical diagnoses (disc herniation, spinal stenosis, spondylolisthesis and non-specific back pain) and co-morbidities (rheumatoid arthritis, osteoarthritis, osteoporosis, fibromyalgia, migraine, irritable bowel syndrome, anxiety, hypertension, diabetes and cancer). Multiple imputation has been shown to be a valid method in overcoming missing data [8] and was used to overcome incomplete data.

RESULTS

Table 1 describes the Genodisc population which contained information on 2636 patients. BMI was normally distributed with a mean of 27.11kg/m². The median (interquartile range [IQR]) duration of symptoms was 10 (4-24) months for BP and 8 (3-20) months of LP. Underweight (BMI<18 kg/m²) and morbidly obese (BMI>35 kg/m²) patients presented after a longer duration of both back and leg pain when compared to other patients (Figure 1a).

Increasing BMI was associated with higher unadjusted BP and LP scores (Table 2). It is important to note the marked increase in the prevalence of hypertension and diabetes with increasing BMI.

Table 3 shows the main result of the univariate and multivariate linear regression models. Multivariate adjusted model 1 for BP shows that a 5-point increase in BMI was associated with a non-significant increase in BP score of 0.20 units (95% CI -0.02, 0.22). However in the univariate model, this association was significant. To investigate where this relationship was lost in our multivariate model, we sequentially removed individual confounders. When hypertension and diabetes were removed (Back pain multivariate adjusted model 2), the effect of BMI upon BP remained significant at 0.15 units (95% CI 0.04,0.27) with minimal change to the coefficients of the other confounders.

In the multivariate model for LP, a 5-point increase in BMI was associated with a 0.19 unit (95% CI 0.15,0.31) increase in pain (Table 3). There was very little change in this effect with the removal of hypertension and diabetes from our regression model (Leg pain multivariate adjusted model 2; Table 3). Figure 2 graphically represents the adjusted linear effect of increasing BMI upon back and leg pain.

The regression coefficients with associated confidence intervals from multivariate model for both back and leg pain are illustrated in Figure 3. Female gender, depression (Zung Depression Score),

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rheumatoid arthritis, heavy workload and previous surgery were all significant positive predictors of both increased BP and LP. Sport was a significant negative predictor for both BP (-0.07 [95% CI -0.14,-0.01]) and LP (-0.10 [95% CI -0.14,-0.04]). Specifically, for BP, the strongest positive predictors were receiving benefit for disability (0.67 units [95% CI 0.30, 1.04]), a heavy workload (0.67 units [95% CI 0.33,1.01]) and rheumatoid arthritis (0.72 units [95% CI 0.33,1.11]). By contrast, the strongest association for LP was a diagnosis of lumbar disc herniation (1.08 units [95% CI 0.84,1.33]).

DISCUSSION

In this large group of spine patients presenting to European tertiary spinal units, an increase in BMI was associated with higher back (non-significant) and leg pain (significant) scores (Figure 2 and Figure 3). Other factors that were associated with greater BP or LP were female gender, previous spine surgery, heavy workload, rheumatoid arthritis and depression (Zung Depression Score) (Figure 3). It is important to note that due to the cross-sectional nature of this study, we can only infer associations not causation.

Back pain was associated with a greater number of significant predictors than LP possibly because it is an umbrella term for poorly defined conditions (Figure 3). Leg pain, below the knee, however, usually has an underlying disc herniation. This is supported in the LP model, where the diagnosis of disc herniation had a strong positive association increasing LP score by 1.08 units (95% CI 0.84,1.33) (Figure 3).

In our analysis, when we considered hypertension and diabetes as confounders, we found that the effect of BMI upon BP score, but not LP score, was markedly weakened with minimal change in any of the other confounders.

1 The relationship between the cardiovascular risk factors and spine related pain is unclear as it is
2
3 difficult to identify a direct effect of hypertension and diabetes upon back and leg pain. Certain
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5 authors have described an atherosclerotic hypothesis to explain a causal relationship between
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7 hypertension and diabetes and BP. This relationship has been established primarily in cadaveric
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9 studies[9, 10] and also in an occupation-based epidemiological study.[11] Within this occupational
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11 study, the authors showed that over a 28-year period patients with higher blood pressure at baseline
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13 had increased risk of BP. This could explain why hypertension and diabetes only confound the
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15 relationship between BMI and BP, and not LP.
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23 It is well recognised that both hypertension and diabetes are associated with obesity as part of the
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25 metabolic syndrome. Our data supports a positive association between increasing prevalence of both
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27 diabetes and hypertension with increasing BMI (Table 2). Given this, we believe, these confounders
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29 most likely lie on the causal pathway between obesity and BP and hence adjusting for these may not
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31 be appropriate. As such we interpret the effect of hypertension and diabetes to be primarily related
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33 to BMI.
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40 A recent meta-analysis, by Shiri et al found that people with increased BMI had greater odds of
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42 developing BP, increased prevalence of BP and were more likely to have chronic BP.[6] Importantly,
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44 this relationship was dose dependant, with obese people having higher levels of pain than those who
45
46 were overweight. Another recently published meta-analysis, by the same group, showed a similar
47
48 dose dependant relationship between increasing BMI and self reported LP symptoms.[5] A limitation
49
50 of these meta-analyses and of the original studies is that both BP and LP were considered as binary
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52 “yes/no” variables. Our study adds to this by showing a similar relationship exists in BP and LP for
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54 patients seeking tertiary care and by providing a more detailed understanding on the effect of BMI
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56 upon the severity of both BP and LP.
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Shiri also noted that overweight or obese patients were more likely to “seek care” for their BP.[6] Based on this information we would have expected our group of patients to have a higher BMI than the general population. However, when we compare the mean BMI within our population (27.2 kg/m²) to those reported in population based studies such as the English Longitudinal Study of Aging (27.9 kg/m²),[12] the United Kingdom Biobank (27.4 kg/m²)[13] or a large Hungarian cohort (25.9 kg/m²),[14] there are minimal differences. This is important, as although overweight and obese people from the study of Shiri et al[6] appear more likely to seek primary care, based on the mean BMI in this study, it appears that these patients do not filter through to tertiary care. It seems unlikely that this is because their symptoms are less severe as we have found increased BMI is associated with more severe pain (Figure 2) and also with longer duration of pain symptoms, suggesting a longer period before tertiary care consultation (Figure1a). A hypothetical explanation could be negative institutional attitudes for the health care of the obese[15] could possibly lead to restriction in their access to tertiary level care for back and leg pain, similar to that seen in osteoarthritis.[16]

Our findings for BP are similar to those of Fanuele et al, who, though not looking at pain directly, used the American National Spine Network data to model the effect of BMI upon disability arising from low back pain.[17] Here, the authors found that obesity was associated with decreased functional status and increased disability. Unfortunately, given the nature of their dataset the authors were not able to correct for depressive symptoms, a significant confounder in our study, nor did they consider LP as an outcome in their adjusted model. Similarly, Heuch et al recently showed, in a longitudinal study, that a BMI of greater than 30kg/m² increases the odds of developing BP, further supporting the deleterious relationship between obesity and BP.[18]

As there appears a direct relationship between BMI and the severity of pain (Figure 2); the next question we must ask is if weight loss help reduce the pain. Recently, three separate small case-series of back pain patients have found that post bariatric surgery with resulting rapid weight loss, obese patients have less BP[19-21] and reduced spine related disability.[19, 20] Surgical weight loss represents an extreme form of weight loss and is a treatment that may not be widely available but the results support the idea that weight loss could lead to decreased pain and this requires further investigation.

As well as weight loss, our results suggest that exercise is associated with less back pain (Figure 3). Within a population setting, Smuck et al, in addition to finding a dose dependant increase of BP with BMI, showed that moderate physical was protective from back pain in an overweight and ultraobese population, but not in the obese group.[22] However, when BMI was considered continuous, this study found physical activity conferred only a small protective effect. The Nord-Trøndelag Health study also found a small protective effect of physical activity upon back pain.[18] To an extent, these results are in keeping with our study where lower BP and LP scores were associated with greater physical activity (Figure 3). More specifically, the greatest decrease in pain score was noted between patients involved in only one episode of sport a week as compared to those who did none (Figure 1b). Similarly, with increasing BMI the associated number of episodes of sporting activity decreases suggesting these patients are less likely to partake in potentially beneficial exercise (Table 2). Alternatively, patients with greater pain may be less willing to exercise so whether exercise is protective cannot be fully assessed from this cross-sectional data.

The link between obesity and pain is usually thought to be mechanical but biochemical pathways may also operate. Obesity is known to cause a systemic low grade inflammatory milieu and there is growing evidence that there is a biochemical link between obesity, degeneration of musculoskeletal tissues and pain.[23] Leptin, the prototypical adipokine, is reported to increase the synthesis of pro-

1 inflammatory cytokines, pain generators and destructive mediators in a knee osteoarthritis model.[24,
2 25] Adipokines can elicit a similar response in the intervertebral disc, the primary joint of the spine[26]
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4 as well as altered pain behaviour in a lumbar nerve root compression model.[27]
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9 Obesity has also been implicated or associated in other pain and psychiatric disorders such as
10 fibromyalgia,[28] migraines[29, 30] and depression.[31, 32] For patients with depression and migraine
11 headaches, a similar dose dependant relationship of increasing BMI with greater symptoms is
12 seen.[29, 30, 32] This linear relationship is less clear in fibromyalgia with authors suggesting an
13 important relationship between obesity, physical activity and symptoms.[28] Furthermore, altered
14 adipokines have also been associated with migraine headaches[30] and fibromyalgia.[33] Taken
15 together, this would suggest increased BMI is an important mediator in main pain related disorders
16 and the effect could be mediated by systemic rather than local mechanisms.
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31 The Genodisc Study is one of the largest populations of patients suffering from spinal conditions with
32 prospectively collected data in a standardised format. As a result, the study carries considerable power
33 and enables us to adjust for many potential participant and clinician reported confounders without
34 limiting the validity of our results. We believe our results are generalisable within the tertiary care
35 setting as patients were recruited from six sites in four countries with a resulting heterogeneous
36 population.
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47 The cross sectional nature of this study raises three important caveats when interpreting the results.
48 Firstly, we cannot establish causation, however within the general population, there is evidence that
49 obesity may be a factor directly leading to back and leg pain.[6, 18]
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57 Secondly, the clinical relevance of these findings requires discussion. For the numerical pain rating
58 scale, as used in this study, a reduction of two points or 30% is generally accepted as a clinically
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1 meaningful difference.[34, 35] From our study, even a very large 15 kg/m² increase in BMI would not
2 show a clinically meaningful difference in pain score. The small coefficients seen in our study may be
3 statistically enhanced by the large sample size.
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9 Although, the definition of a clinically important difference was derived from chronic pain populations
10 including patients with low back and neuralgic pain, it is important to note that such difference
11 represents a change in pain score, which is usually an intra-individual change within a longitudinal
12 cohort or interventional study. Even though our population is similar, the interpretation of clinical
13 relevance is most likely informative rather than prescriptive in our observational study. Similarly, our
14 findings are consistent to what is seen in the literature suggesting a true result rather than a statically
15 anomaly.
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26 Finally, pain is a symptom, which is not constant and can change because of factors other than those
27 relating to a biological or pathoanatomical process. Although we attempted to acquire a more general
28 picture of pain symptoms by asking participants to rate their pain over the previous week, longitudinal
29 studies are required to answer the question of causation.
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41 CONCLUSION

42 This study provides evidence supporting the hypothesis that obesity is independently associated with
43 leg pain and that the effect of increasing BMI upon pain is linear. It also provides information on other
44 clinically important predictors of pain in spine patients, in particular female gender, heavy workload,
45 rheumatoid arthritis, previous spine surgery and depression. Given the growing evidence for a
46 relationship between obesity and both back and leg pain, we need to move forward to understand
47 the underlying biological pathway and also to define evidence-based management modalities for
48 obese patients with spinal conditions.
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REFERENCES

1. Hoy D, March L, Brooks P, et al. (2014) The global burden of low back pain: estimates from the Global Burden of Disease 2010 study. *Ann Rheum Dis* 73:968–974. doi: 10.1136/annrheumdis-2013-204428
2. Hoy D, Bain C, Williams G, et al. (2012) A systematic review of the global prevalence of low back pain. *Arthritis & Rheumatism* 64:2028–2037. doi: 10.1002/art.34347
3. Hoy DG, Smith E, Cross M, et al. (2014) Reflecting on the global burden of musculoskeletal conditions: lessons learnt from the Global Burden of Disease 2010 Study and the next steps forward. *Ann Rheum Dis*. doi: 10.1136/annrheumdis-2014-205393
4. Ishimoto Y, Yoshimura N, Muraki S, et al. (2013) Associations between radiographic lumbar spinal stenosis and clinical symptoms in the general population: the Wakayama Spine Study. *Osteoarthritis Cartilage* 21:783–788. doi: 10.1016/j.joca.2013.02.656
5. Shiri R, Lallukka T, Karppinen J, Viikari-Juntura E (2014) Obesity as a risk factor for sciatica: a meta-analysis. *Am J Epidemiol* 179:929–937. doi: 10.1093/aje/kwu007
6. Shiri R, Karppinen J, Leino-Arjas P, et al. (2010) The association between obesity and low back pain: a meta-analysis. *Am J Epidemiol* 171:135–154. doi: 10.1093/aje/kwp356
7. Battié MC, Lazáry A, Fairbank J, et al. (2014) Disc degeneration-related clinical phenotypes. *Eur Spine J* 23 Suppl 3:S305–14. doi: 10.1007/s00586-013-2903-5
8. Shrive FM, Stuart H, Quan H, Ghali WA (2006) Dealing with missing data in a multi-question depression scale: a comparison of imputation methods. *BMC Medical Research Methodology* 6:57. doi: 10.1186/1471-2288-6-57
9. Kauppila LI (2009) Atherosclerosis and disc degeneration/low-back pain--a systematic review. *Eur J Vasc Endovasc Surg* 37:661–670. doi: 10.1016/j.ejvs.2009.02.006
10. Kauppila LI (1997) Prevalence of stenotic changes in arteries supplying the lumbar spine. A postmortem angiographic study on 140 subjects. *Ann Rheum Dis* 56:591–595. doi:

10.1136/ard.56.10.591

11. Leino-Arjas P, Solovieva S, Kirjonen J, et al. (2006) Cardiovascular risk factors and low-back pain in a long-term follow-up of industrial employees. *Scand J Work Environ Health* 32:12–19. doi: 10.5271/sjweh.971
12. Zaninotto P, Pierce M, Breeze E, et al. (2010) BMI and waist circumference as predictors of well-being in older adults: findings from the English Longitudinal Study of Ageing. *Obesity* 18:1981–1987. doi: 10.1038/oby.2009.497
13. (2014) UK Biobank Data Showcase. In: UK Biobank Data Showcase. <http://biobank.ctsu.ox.ac.uk/crystal/field.cgi?id=21001>. Accessed 12 Apr 2014
14. Vamos EP, Kopp MS, Keszei A, et al. (2008) Prevalence of diabetes in a large, nationally representative population sample in Hungary. *Diabetes Res Clin Pract* 81:e5–8. doi: 10.1016/j.diabres.2008.04.022
15. Puhl R, Brownell KD (2001) Bias, discrimination, and obesity. *Obes Res* 9:788–805. doi: 10.1038/oby.2001.108
16. Judge A, Batra RN, Thomas GE, et al. (2014) Body mass index is not a clinically meaningful predictor of patient reported outcomes of primary hip replacement surgery: prospective cohort study. *Osteoarthritis and Cartilage* 22:431–439. doi: 10.1016/j.joca.2013.12.018
17. Fanuele JC, Abdu WA, Hanscom B, Weinstein JN (2002) Association between obesity and functional status in patients with spine disease. *Spine* 27:306–312. doi: 10.1097/00007632-200202010-00021
18. Heuch I, Heuch I, Hagen K, Zwart J-A (2013) Body mass index as a risk factor for developing chronic low back pain: a follow-up in the Nord-Trøndelag Health Study. *Spine* 38:133–139. doi: 10.1097/BRS.0b013e3182647af2
19. Khoeir P, Black MH, Crookes PF, et al. (2009) Prospective assessment of axial back pain symptoms before and after bariatric weight reduction surgery. *Spine J* 9:454–463. doi: 10.1016/j.spinee.2009.02.003

20. Melissas J, Kontakis G, Volakakis E, et al. (2005) The effect of surgical weight reduction on functional status in morbidly obese patients with low back pain. *Obes Surg* 15:378–381. doi: 10.1381/0960892053576703
21. Lidar Z, Behrbalk E, Regev GJ, et al. (2012) Intervertebral disc height changes after weight reduction in morbidly obese patients and its effect on quality of life and radicular and low back pain. *Spine* 37:1947–1952. doi: 10.1097/BRS.0b013e31825fab16
22. Smuck M, Kao M-CJ, Brar N, et al. (2014) Does physical activity influence the relationship between low back pain and obesity? *Spine J* 14:209–216. doi: 10.1016/j.spinee.2013.11.010
23. Lago F, Dieguez C, Gomez-Reino J, Gualillo O (2007) Adipokines as emerging mediators of immune response and inflammation. *Nat Clin Pract Rheumatol* 3:716–724. doi: 10.1038/ncprheum0674
24. Vuolteenaho K, Koskinen A, Kukkonen M, et al. (2009) Leptin enhances synthesis of proinflammatory mediators in human osteoarthritic cartilage--mediator role of NO in leptin-induced PGE2, IL-6, and IL-8 production. *Mediators of Inflammation* 2009:345838. doi: 10.1155/2009/345838
25. Koskinen A, Vuolteenaho K, Nieminen R, et al. (2011) Leptin enhances MMP-1, MMP-3 and MMP-13 production in human osteoarthritic cartilage and correlates with MMP-1 and MMP-3 in synovial fluid from OA patients. *Clinical and experimental rheumatology* 29:57–64.
26. Segar A, Edwards K, Fairbank J (2014) Adipokines and the intervertebral disc: does a biochemical link exist between obesity and back pain? *Osteoarthritis and Cartilage*. doi: 10.1016/j.joca.2014.02.553
27. Lim G, Wang S, Zhang Y, et al. (2009) Spinal leptin contributes to the pathogenesis of neuropathic pain in rodents. *J Clin Invest*. doi: 10.1172/JCI36785
28. Okifuji A, Donaldson GW, Barck L, Fine PG (2010) Relationship between fibromyalgia and obesity in pain, function, mood, and sleep. *J Pain* 11:1329–1337. doi: 10.1016/j.jpain.2010.03.006

29. Bigal ME, Liberman JN, Lipton RB (2006) Obesity and migraine: a population study. *Neurology* 66:545–550. doi: 10.1212/01.wnl.0000197218.05284.82
30. Bigal ME, Lipton RB (2006) Obesity is a risk factor for transformed migraine but not chronic tension-type headache. *Neurology* 67:252–257. doi: 10.1212/01.wnl.0000225052.35019.f9
31. Peterlin BL, Rapoport AM, Kurth T (2010) Migraine and obesity: epidemiology, mechanisms, and implications. *Headache* 50:631–648. doi: 10.1111/j.1526-4610.2009.01554.x
32. Luppino FS, de Wit LM, Bouvy PF, et al. (2010) Overweight, obesity, and depression: a systematic review and meta-analysis of longitudinal studies. *Arch Gen Psychiatry* 67:220–229. doi: 10.1001/archgenpsychiatry.2010.2
33. Ursini F, Naty S, Grembale RD (2011) Fibromyalgia and obesity: the hidden link. *Rheumatol Int* 31:1403–1408. doi: 10.1007/s00296-011-1885-z
34. Farrar JT, Young JP, LaMoreaux L, et al. (2001) Clinical importance of changes in chronic pain intensity measured on an 11-point numerical pain rating scale. *Pain* 94:149–158.
35. Childs JD, Piva SR, Fritz JM (2005) Responsiveness of the numeric pain rating scale in patients with low back pain. *Spine* 30:1331–1334.

TABLES

Table 1: Characteristics of the Genodisc participants. Data are n (%), mean (standard deviation [SD]) or median (interquartile range [IQR])

Characteristic:		Missing (n)
Age (years), mean (SD)	50.85 (14.59)	111
Women, n (%)	1349 (53.8%)	127
BMI (kg/m ²), mean (SD)	27.22 (4.80)	169
Pain Score (units), mean (SD)		
Back	6.18 (2.88)	211
Leg	6.66 (2.91)	217
Duration of Symptoms (months), median (IQR)		
Back	10 (4-24)	432
Leg	8 (3-20)	436
Zung Depression Score, mean (SD)	39.83 (9.00)	942
Sport per Week (episodes), median (IQR)	0 (0-2)	133
Disability Benefit, n (%)	296 (12.1)	183
Family History, n (%)	747 (30.3)	172
Previous Surgery, n (%)	715 (27.9)	69
Smoking Status, n (%)		264
Never	1079 (45.5)	
Previous	622 (26.2)	
Current	671 (28.3)	
Work Type, n (%)		234
Sedentary	684 (28.5)	
Light	605 (25.2)	
Medium	563 (23.4)	
Heavy	550 (22.9)	
Clinical Diagnosis, n (%)		
Lumbar disc herniation	1413 (55.5)	90
Spinal stenosis	968 (39.1)	161

Obesity and back/leg pain

Spondylolisthesis	400 (16.3)	186	Table 2: Categorised
Non-specific Back Pain	359 (14.7)	198	
Comorbidities , n (%)		0	

	Underweight (<18.50)	Normal Weight ($18.50-24.99$)	Overweight ($25.00-29.99$)	Obese ($30.00-34.99$)	Severe Obese (≥ 35.00)
Rheumatoid Arthritis			233 (8.8)		
Osteoarthritis			298 (11.3)		
Number of Patients	37	780	952	451	153
Osteoporosis			180 (6.8)		
Back pain score mean (95% CI)	5.35 (4.39,6.31)	6.00 (5.80,6.21)	6.16 (5.98,6.35)	6.24 (5.98,6.50)	7.24 (6.83,7.65)
Fibromyalgia			21 (0.8)		
Leg pain score mean (95% CI)	6.27 (5.10,7.44)	6.42 (6.21,6.64)	6.58 (6.39,6.76)	6.97 (6.72,7.21)	7.70 (7.36,8.03)
Celiac disease			581 (14.5)		
Irritable Bowel Syndrome			162 (6.1)		
Episodes of sport per week mean (95% CI)	1.08 (0.39,1.78)	1.06 (0.93,1.18)	1.01 (0.90,1.12)	1.01 (0.84,1.18)	0.72 (0.48,0.96)
Anxiety			233 (8.8)		
Hypertension			715 (27.1)		
Type 2 diabetes mellitus (%)	0.0	3.9	7.4 214 (8.1)	15.0	21.3
Hypertension (%)	10.8	14.9	27.6 298 (11.3)	45.6	53.8
Cancer					

unadjusted mean low back pain and leg pain score, participation in sport and prevalence of hypertension and diabetes for each group. BMI with

Table 3: Univariate and multivariate regression coefficients for the effect of BMI upon back and leg pain scores. Multivariate model 1 includes hypertension and diabetes as confounders, multivariate model 2 does not. Coefficient is the regression coefficient. Each coefficient represents a change in pain score for a 5-unit increase in BMI. A positive coefficient represents increased pain. Models were fitted to 50 multiple imputation datasets (n=2636). Multivariate model 1 is adjusted for all confounders as described in the main text. In multivariate model 2, hypertension and diabetes were excluded.

	Back Pain Score			Leg Pain Score		
	Coefficient	(95% CI)	p value	Coefficient	(95% CI)	p value
Univariate	0.26	(0.14,0.37)	0.0	0.35	(0.23,0.47)	0.0
Multivariate model 1	0.10	(-0.02,0.22)	0.10	0.19	(0.08,0.31)	0.0
Multivariate model 2	0.15	(0.04,0.27)	0.01	0.22	(0.10,0.33)	0.0

FIGURE LEGENDS

Figure 1: a) Mean duration of pain (months) experienced by patients categorised by BMI (body mass index). b) Mean pain scores categorised by episodes of sport. Markers represent mean and 95% confidence intervals. ** $p < 0.01$ when compared to a) normal BMI or b) no sport

Figure 2: Adjusted mean back and leg pain score for each BMI (body mass index) category. Bars show mean and 95% confidence Interval

Figure 3: Confidence interval plots showing regression coefficients of multivariate model 1 (adjusted for hypertension and diabetes) for both the back and leg pain. Models fitted to 50 multiple imputation datasets ($n=2636$). Coefficient is the regression coefficient. A positive coefficient represents higher pain and a negative coefficient represents lower pain. The solid diamond represents the effect and the error bars the 95% confidence interval. If the confidence interval does not cross the “No Effect” dotted line the predictor is significant.

Figures 1

Figures 2

Figures 3

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