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The role of inflammation in sciatica: the contradictory effect of macrophages

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Chapter 3

Lumbar disc extrusions reduce faster than bulging disc due to an active role of macrophages in sciatica

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“Have no fear of perfection, you’ll never reach it”
(Salvador Dali)

Abstract

Objective

This retrospective observational histological study aims to associate the size and type of disc herniation with the degree of macrophage infiltration in disc material retrieved during disc surgery in patients with sciatica.

Methods

Disc tissue of 119 sciatica patients was embedded in paraffin and stained with haematoxylin and CD68. Tissue samples were categorized as mild ($0-10/\text{cm}^2$), moderate ($10-100/\text{cm}^2$), and considerable ($>100/\text{cm}^2$) macrophage infiltration. All 119 patients received an MRI at baseline, and 108 received a follow-up MRI at one-year. MRIs were reviewed for the size and type of the disc herniations, and for Modic Changes in the vertebral end plates.

Results

Baseline characteristics and duration of symptoms before surgery were comparable in all macrophage infiltration groups. The degree of macrophage infiltration was not associated with herniation size at baseline, but significantly associated with reduction of size of the herniated disc at one-year post surgery. Moreover, the degree of macrophage infiltration was higher in extrusion in comparison to bulging (protrusion) of the disc. Results were comparable in patients with and without Modic changes.

Conclusion

macrophage infiltration was positively associated with an extruded type of disc herniation as well as the extent of reduction of the herniated disc during one-year follow-up in patients with sciatica. This is an indication that the macrophages play an active role in reducing herniated discs. An extruded disc herniation has a larger surface for the macrophages to adhere to, which leads to more size reduction.

Introduction

Herniation of the intervertebral disc is a highly prevalent disease [11], characterized by radiating pain symptoms due to a compression of the nerve root by the herniated disc. Since 1934 it has been widely accepted that a disc may herniate due to a disruption of the annulus fibrosus or due to its attachments to the adjacent endplate [16]. This disruption of the annulus fibrosus can result in a foreign body reaction aimed at the nucleus pulposus that induces neovascularization [13], which is accompanied by macrophage infiltration [15, 23, 26].

This macrophage infiltration is believed to exacerbate pain symptoms through secretion of pro-inflammatory cytokines such as IL-6, IL8 and TNF-alfa [1, 20, 29]. In contrast, macrophage infiltration may also have a positive effect on symptoms through inducing a phagocytic resorption process, mediated by anti-inflammatory cytokines such as IL-4 and IL-10 [27, 29]. This discrepancy in effect of macrophage infiltration is reflected in the inconsistent findings regarding the correlation between macrophage infiltration and clinical symptoms [14, 23-25, 28]

A possible factor that may reflect the type of macrophage is the presence of Modic Changes (MC), also known as vertebral end-plate signal changes (VESC) on MRI [17, 18]. Recent studies have shown that patients with MC recover more slowly from their herniated discs than those without MC [2, 26]. In addition, Dudli et al. showed that both Modic type 1 and type 2 changes are associated with inflammatory dysmyelopoiesis and fibrogenic changes [4], which infers that MC reflect a factor that plays a role in the inflammatory process in the disc. Hence, it is to be expected that this factor in the inflammatory process, reflected by presence of MC, interferes with the resorption process, possibly by changing macrophage differentiation away from the anti-inflammatory type and towards the pro-inflammatory type. Until now, the interaction between MC and macrophage infiltration on the resorption of the herniated disc (size reduction) remains unknown.

Furthermore, if a herniated disc extrudes instead of bulges (protrudes), it is likely to have a higher exposure to the systemic circulation through more neovascularization, which in turn may result in more macrophage infiltration. However, the evidence supporting this theory is limited [15]. In addition, recent studies have found that MC are associated with less neovascularization [12, 26]. It is thus hypothesized that the presumed association between macrophage infiltration and disc extrusion will be less strong in patients with MC.

In order to test the previously mentioned theories, this study aims to associate macrophage infiltration with the type and size of disc herniation as well as the reduction in disc herniation at one-year follow-up in patients with sciatica. We will also explore this association in patient with and without MC separately. Exploring these associations will provide a more profound understanding of the roles of macrophage infiltration and MC in patients with sciatica due to a lumbar disc herniation.

Materials and Methods

Study population

This retrospective study was performed using participants from the Sciatica Trial [22], a multicentre RCT with 283 patients who suffered from sciatica for 6-12 weeks and had a disc herniation as assessed by means of MRI. 141 patients were randomized to surgery and 125 patients actually underwent surgery (16 recovered before surgery could be performed). The other 142 patients were randomized to prolonged conservative care, of which 55 patients underwent surgery within 1 year, with a mean time to surgery of 15 weeks after randomization. Thus, in the first year after randomization a total of 180 patients underwent surgery for sciatica. Out of the 180 patients, 120 disc samples were available for analysis. Missing samples were due to multiple reasons: not collected during surgery, got lost after surgery, not preserved properly, or got lost after preservation. All surgeries were performed between November 2002 and Feb 2005. The protocol, which included analysis of the disc material, was approved by the medical ethics committees at all participating hospitals.

Histological analysis

Disc material of all operated patients was collected and fixed in 4% formaldehyde solution after surgery and were subsequently stored for future analysis. For the purpose of this retrospective study, samples were embedded in blocks of paraffin and stained with CD68 to evaluate macrophage infiltration. A detailed description of the protocol was published in our previous work [3].

The evaluation was done by two independent investigators, who were blinded to clinical information and MRI data. The training of these independent researchers was carried out by a senior pathologist. The number of macrophages on each sample was counted and estimated. Using this method, the tissue samples were categorized based on the extent of macrophage infiltration. The categories consisted of mild (0-10 /cm²), moderate (10-100 /cm²), and considerable (>100 /cm²) macrophage infiltration. Subsequently, a consensus score between the independent researchers was determined. An acceptable consensus score was predefined as 60%. It was pre-defined that all samples would be re-assessed with a senior pathologist if consensus was less than 60%.

MRI protocol and Image evaluation of the sciatica trial

MRI scans were performed at baseline by a 1.5 Tesla scanner, and both sagittal T1- and T2-weighted images of the lumbar spine were obtained. Image evaluation of MC was according to the criteria of Modic et al [17, 18]. Image evaluation was done according to a predefined protocol (Supplementary Table S1)[9]. Definitions of imaging characteristics were based on the recommendations from the combined task forces of the North American Spine Society, the American Society of Spine Radiology, and the American Society of Neuroradiology for classification of lumbar disc pathology. MRIs were evaluated by 2 neuroradiologists and 1 neurosurgeon. All three were blinded to histological data and clinical information. The readers were not involved in the selection or treatment of the patients included. Inter-observer agreement analyses was published earlier by el Barzouhi et al.[6, 7]. For the statistical analyses, the majority opinion of the three independent researchers (answer by at least two of the three MRI assessors) was used.

Type of disc herniation as quantified by MRI was categorized as follows: bulging (or protrusion), extrusion, or not applicable, Based on characteristics described by Fardon and Milette [8]. The definition of protrusion as defined by the protocol is more commonly known as 'bulging' in daily clinical practice. Therefore, this term is used in this article. The size of the herniated disc was measured as the surface in mm². The decrease in size of the disc herniation between baseline and 1 year after randomization was measured as a percentage reduction in surface. MC were scored as Type 1, Type 2, or Type 3 according to Modic et al. [17, 18]. The inter-observer agreement was substantial for the MC (69-97%), as described by el Barzouhi (2014) [7], no kappa calculated because the prevalence of MC type 1 and 3 were too low. At last, inter-observer agreement was substantial regarding disc type (kappa = 0.62) [6].

Statistical analysis

The categorized histological findings were associated with the MRI variables (type and size of the herniated disc, and disc herniation reduction at one year follow-up). In addition, histological findings were split into a group with, and a group without MC at the herniated disc level and comparisons with all MRI variables were repeated for these subgroups. The association between the histological findings and disc type and was tested using X² tests for categorical data. The association between the histological findings and disc size was done using a one-way ANOVA, At last, the association between the histological findings and disc size reduction was assessed using a Kruskal Wallis test. P-values of < 0.05 were regarded as significant.

Results

Demographics

Of the 180 patients that underwent surgery, 119 patients disc samples were preserved and analyzed. Missing samples were due to multiple reasons: samples were either not collected during surgery, got lost after surgery, were not preserved properly, or got lost after preservation. 103 of the 119 patients received an MRI at one-year follow-up. The baseline characteristics age, gender, BMI and duration of sciatica prior to surgery of the three macrophage infiltration groups were comparable (Table 1).

The histological data analysis

CD68 staining to identify macrophages resulted in the following distribution: 47 (39.5%) patients had mild-, 45(37.8%) patients moderate, and 27 (22.7%) considerable macrophage infiltration (Table 2).The consensus score was excellent (0.96). (Table 2). Examples of the CD68 samples and their categories are shown in Figure 1.

The MRI data analysis

Forty-one patients demonstrated bulging (protrusion) of the disc, 74 an extrusion, and in 2 patients discs were scored as not evidently protruding; in 2 cases information was missing. The disc size was measured in 116 patients with a mean of $77.9 \text{ mm}^2 \pm 3.6 \text{ SE}$ (2 samples were missing). The reduction in disc size was measured over 108 patients, which had a mean of $66.6\% \pm 4.1 \text{ SE}$, of which 46 disc herniations completely disappeared. Furthermore, a total of 2 patients with MC type 1, 32 patients with MC type 2, no patients with MC type 3, and 85 patients had no MC at the vertebrae adjacent to the herniated disc. Because only two patients showed MC type 1, these samples were excluded from the additional statistical analysis in which only patients with MC were assessed. Thus, in this additional analyses, only MC type 2 were taken into account.

Associations between number of macrophages and type and size of disc herniation

The type of disc herniation at baseline showed a significant association with the degree of macrophage infiltration, suggesting that patients with more macrophage infiltration are more likely to have an extruded disc as compared to a bulging (protruded) disc ($p = 0.05$). The presence of MC type 2 was not of significant influence on this association (no MC: $p = 0.07$, MC type 2: $p = 0.68$) (Figure 2). In addition, MC2 was not associated with either macrophage infiltration ($p=0.53$) or type of disc herniation ($p=0.32$) (Supplementary Figure S1). This verifies that the association between extrusion and macrophage infiltration was not confounded by MC2.

Disc herniation size at baseline was not associated with the degree of macrophages infiltration ($p = 0.43$). Again, the presence or absence of MC type 2 changes did not significantly influence this finding (no MC: $p = 0.91$, MC type 2: $p = 0.11$) (Figure 3).

Associations between number of macrophages and reduction at one-year follow-up

The timing of surgery was not equal for all 119 patients. In the sciatica RCT about 40% of patients that were randomized to conservative treatment, crossed over to surgical treatment because of unbearable symptoms. These patients had a longer timespan between start of the complaints and surgical intervention, varying from approximately the same waiting time as the surgical group (mean = 15 days after randomization) to 18 months after randomization. In our sample group, 34 of 119 patients were originally randomized to conservative care and crossed over. Because the follow-up MRI was made at one-year after randomization and not after surgery, not all of the crossed over patients were comparable with the patients randomized for surgery. Therefore, crossed over patients with waiting time for surgery longer than 6 months were excluded from this analyses, which resulted in exclusion of 7 of the 34 patients.

Reduction of disc herniation, one year after surgery, was positively associated with the degree of macrophage infiltration at baseline ($p=0.01$) (Figure 4). When testing for the presence and absence of MC Type 2 separately, no significant associations were found (no MC: $p = 0.06$, MC type 2: $p = 0.23$, Figure 4).

In addition, we found that disc extrusion were larger than bulging discs and we found that extruded discs were significantly associated with a larger reduction in size at one year ($p = 0.001$). Since macrophage infiltration, disc extrusion and relative reduction in size at one-year follow-up showed multicollinearity, the association between macrophage infiltration and relative reduction at one year could be confounded by extrusion. For example, because extruded discs could be easier to resect during surgery. In order to falsify this possibility, an additional analysis was performed with the data from the conservative group of the Sciatica trial [5]. If this was due to surgery, extrusion would not associate with reduction in the conservative group: The additional analysis showed that extrusion in the conservative group also significantly associated with size reduction at follow-up in the quantitative analyses (T-test: p -value = 0.03). Both additional analyses are shown in Supplementary Figure S2.

Discussion

The most important finding of this study is the association between the number of macrophages present at baseline and the decrease in size of the disc herniation at one-year follow-up in patients with sciatica due to a lumbar disc herniation. This is accompanied by an association between a higher number of macrophages, in extruded as compared to bulging disc herniations. Since extrusion of disc material was demonstrated to resorb faster, patients with a disc extrusion rather than a bulging disc are more likely to benefit from prolonged conservative treatment.

The association between macrophage infiltration and disc extrusion was in line with previous findings of Lohr et al. [14]. This could be explained by the likelihood that disc extrusion as compared to protrusion leads to a larger exposure to systemic circulation, which makes it easier for macrophages to infiltrate the disc during the foreign-body reaction [13]. When performing a separate analysis for patients with and without MC, an MRI feature that has been shown to interact with disc inflammation in recent literature [18], no significant differences were found. However, only in patients without MC an almost significant association between macrophage infiltration and disc extrusion ($p=0.07$) was demonstrated, while in patients with MC no such trend was seen ($p = 0.68$). This lack of significance could be explained by the small sample sizes. Hence, this finding could indicate that MC type 2 interfere with the normal course of macrophage infiltration, possibly through a disruption in neovascularization [12, 26]. Nevertheless, due to the small sample sizes, these findings should be interpreted with caution.

Furthermore, even though the size of the disc herniation was not associated with the degree of macrophage infiltration at baseline, we did find that the number of infiltrated macrophages was associated with a larger regression of herniated disc material at one-year follow-up. This indicates the importance of a resorption process induced by macrophages in cleaning up the herniated disc material [13, 21]. Since this study was the first that analyzed this association, no comparison with previous literature could be made.

The additional analyses performed with the data from the conservative group of the Sciatica trial showed that the association between macrophage infiltration and size reduction of the herniated disc was not confounded by surgery. Hence we conclude that the reduction was likely to be induced by the infiltrated macrophages.

A different possible confounder on the rate reduction of disc herniation, could be the duration of symptoms before surgery. For example, a longer chronic inflammation process could possibly lead to a different result as compared to a short lived inflammation process [19]. However, since the treatment duration was equal for all macrophage groups, this possibility was excluded.

This study has several strong points. Both the histological and MRI analyses were performed by multiple evaluators and demonstrated substantial inter agreement or consensus scores. Moreover, this study was the first to compare macrophage infiltration to disc parameters on MRI between patients with and without MC type 2. This study also has some limitations. The samples that were evaluated for the histological analysis were old, which might have reduced their quality. Nevertheless, the CD68 staining showed clearly identifiable macrophages and the quality was validated by a senior pathologist. Hence, we assume that the age of the samples did not alter our result. Also, macrophage infiltration does not cover the entire inflammatory picture including the different effects of other inflammatory- cells, cytokines and proteins that also infiltrate the disc as a consequence of neovascularization. However, previous findings have identified macrophages as the main type of inflammatory cells found in discus samples [10, 14]. Thus, it is likely that they contribute the most to the overall inflammation response. Lastly, because all patients already received surgery, it remains unclear whether the reduction of the herniated disc at one year is also associated with macrophage infiltration in a population without surgical intervention. Due to ethical considerations, this could not be studied.

In conclusion, macrophage infiltration was positively associated with an extruded type of disc herniation as well as the extent of reduction of the herniated disc during one-year follow-up in patients with sciatica. In order to evaluate our findings, our results should be repeated by a study with a larger sample size.

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Conflict of interest

None of the authors has any conflict of interest. No funding was received for the conductance of this study.

Bibliography

1. Jordan J, Konstantinou K, and O'Dowd J, *Herniated lumbar disc*. BMJ Clin Evid, 2011. 2011: 1118
2. Mixer WJ BJ, *Rupture of Intervertebral Disc with Involvement of Spinal Canal*. New England Journal of Medicine, 1934. 211(210): p. 5.
3. Kobayashi S, Meir A, Kokubo Y et al, *Ultrastructural analysis on lumbar disc herniation using surgical specimens: role of neovascularization and macrophages in hernias*. Spine (Phila Pa 1976), 2009. 34(7): p. 655-62.
4. Lotz JC, Fields AJ, and Liebenberg EC, *The role of the vertebral end plate in low back pain*. Global Spine J, 2013. 3(3): p. 153-64.
5. Rotherl R, Woertgen C, Holzschuh M et al, *Macrophage tissue infiltration, clinical symptoms, and signs in patients with lumbar disc herniation. A clinicopathological study on 179 patients*. Acta Neurochir (Wien), 1998. 140(12): p. 1245-8.
6. Shan Z, Fan S, Xie Q et al, *Spontaneous resorption of lumbar disc herniation is less likely when modic changes are present*. Spine (Phila Pa 1976), 2014. 39(9): p. 736-44.
7. Arango Duque G and Descoteaux A, *Macrophage cytokines: involvement in immunity and infectious diseases*. Front Immunol, 2014. 5: p. 491.
8. Pedersen LM, Schistad E, Jacobsen LM et al, *Serum levels of the pro-inflammatory interleukins 6 (IL-6) and -8 (IL-8) in patients with lumbar radicular pain due to disc herniation: A 12-month prospective study*. Brain Behav Immun, 2015. 46: p. 132-6.
9. Zu B, Pan H, Zhang XJ et al, *Serum Levels of the Inflammatory Cytokines in Patients with Lumbar Radicular Pain Due to Disc Herniation*. Asian Spine J, 2016. 10(5): p. 843-849.
10. Wang K, Bao JP, Yang S et al, *A cohort study comparing the serum levels of pro- or anti-inflammatory cytokines in patients with lumbar radicular pain and healthy subjects*. Eur Spine J, 2016. 25(5): p. 1428-1434.
11. Rotherl RD, Woertgen C, and Brawanski A, *Pain resolution after lumbar disc surgery is influenced by macrophage tissue infiltration. A prospective consecutive study on 177 patients*. J Clin Neurosci, 2002. 9(6): p. 633-6.
12. Rotherl RD, Woertgen C, Holzschuh M et al, *Is there a clinical correlate to the histologic evidence of inflammation in herniated lumbar disc tissue?* Spine (Phila Pa 1976), 1998. 23(11): p. 1197-200; discussion 1200-1.
13. Woertgen C, Rotherl RD, and Brawanski A, *Influence of macrophage infiltration of herniated lumbar disc tissue on outcome after lumbar disc surgery*. Spine (Phila Pa 1976), 2000. 25(7): p. 871-5.
14. Lohr M, Lebenheim L, Berg F et al, *Gadolinium enhancement in newly diagnosed patients with lumbar disc herniations are associated with inflammatory peridiscal tissue reactions - Evidence of fragment degradation?* Clinical Neurology and Neurosurgery, 2014. 119: p. 28-34.
15. Modic MT, Masaryk TJ, Ross JS et al, *Imaging of degenerative disk disease*. Radiology, 1988. 168(1): p. 177-86.
16. Modic MT, Steinberg PM, Ross JS et al, *Degenerative disk disease: assessment of changes in vertebral body marrow with MR imaging*. Radiology, 1988. 166(1 Pt 1): p. 193-9.
17. Ding L, Teng X, Fan S et al, *The Association Between Modic Changes of Lumbar Endplates and Spontaneous Absorption of Herniated Intervertebral Discs*. Cell Biochem Biophys, 2015. 71(3): p. 1357-63.
18. Dudli S, Sing DC, Hu SS et al, *ISSLS PRIZE IN BASIC SCIENCE 2017: Intervertebral disc/bone marrow cross-talk with Modic changes*. Eur Spine J, 2017. 26(5): p. 1362-1373.

19. Karamouzian S, Eskandary H, Faramarzee M et al, *Frequency of lumbar intervertebral disc calcification and angiogenesis, and their correlation with clinical, surgical, and magnetic resonance imaging findings*. Spine (Phila Pa 1976), 2010. 35(8): p. 881-6.
20. Peul WC, van Houwelingen HC, van den Hout WB et al, *Surgery versus prolonged conservative treatment for sciatica*. N Engl J Med, 2007. 356(22): p. 2245-56.
21. Djuric N, Yang X, Barzouhi AE et al, *Gadolinium Enhancement is not Associated with Disc Inflammation in Patients with Sciatica*. Spine (Phila Pa 1976), 2019.
22. Fardon DF, Williams AL, Dohring EJ et al, *Lumbar disc nomenclature: version 2.0: Recommendations of the combined task forces of the North American Spine Society, the American Society of Spine Radiology and the American Society of Neuroradiology*. Spine J, 2014. 14(11): p. 2525-45.
23. El Barzouhi A, Vleggeert-Lankamp CL, Lycklama ANGJ et al, *Magnetic resonance imaging interpretation in patients with sciatica who are potential candidates for lumbar disc surgery*. PLoS One, 2013. 8(7): p. e68411.
24. el Barzouhi A, Vleggeert-Lankamp CL, van der Kallen BF et al, *Back pain's association with vertebral end-plate signal changes in sciatica*. Spine J, 2014. 14(2): p. 225-33.
25. Fardon DF, Milette PC, Combined Task Forces of the North American Spine Society ASoSR et al, *Nomenclature and classification of lumbar disc pathology. Recommendations of the Combined task Forces of the North American Spine Society, American Society of Spine Radiology, and American Society of Neuroradiology*. Spine (Phila Pa 1976), 2001. 26(5): p. E93-E113.
26. el Barzouhi A, Vleggeert-Lankamp CL, Lycklama a Nijeholt GJ et al, *Magnetic resonance imaging in follow-up assessment of sciatica*. N Engl J Med, 2013. 368(11): p. 999-1007.
27. Peluffo H, Solari-Saquieres P, Negro-Demontel ML et al, *CD300f immunoreceptor contributes to peripheral nerve regeneration by the modulation of macrophage inflammatory phenotype*. J Neuroinflammation, 2015. 12: p. 145.
28. Oort J and Scheper RJ, *Histopathology of acute and chronic inflammation*. Agents Actions Suppl, 1977(3): p. 25-30.
29. Habtemariam A, Gronblad M, Virri J et al, *A comparative immunohistochemical study of inflammatory cells in acute-stage and chronic-stage disc herniations*. Spine (Phila Pa 1976), 1998. 23(20): p. 2159-65; discussion 2166.

Table 1. Baseline characteristics of the three histologically defined inflammation groups

	Mild (N=48)	moderate (N=45)	considerable (N=27)
Age	40.4±9.6	42.1±10.3	43.7±6.6
Male gender	36 (75.0%)	28 (62.2%)	22 (81%)
Body-mass index	26.0±4.1	25.8±3.5	25.9±3.2
Duration of sciatica in weeks	9.6±1.9	9.7±2.3	8.7±2.1

Values for gender are n and (%) or means ± SD.

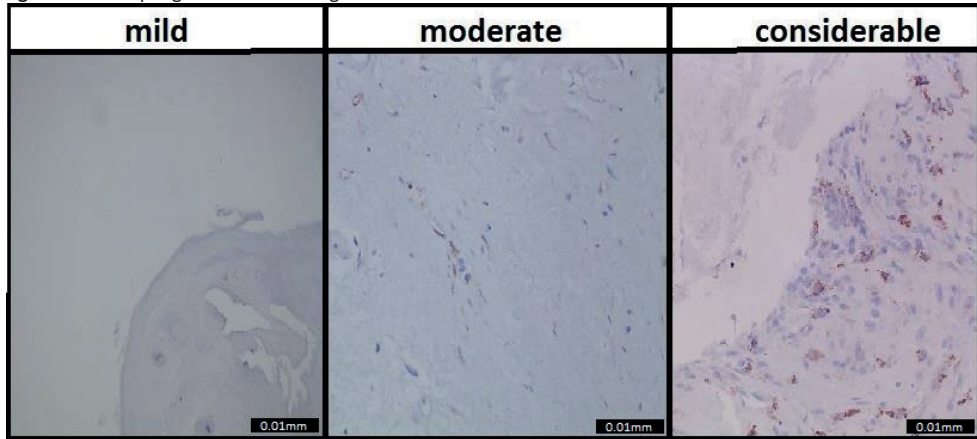
No significant baseline differences were observed between the three inflammation groups.

Table 2. Consensus score of the pathological findings

	A vs. B % agreement
Inflammation at baseline (3 categories)	95.8
Mild	96.8
Moderate	94.6
considerable	96.2

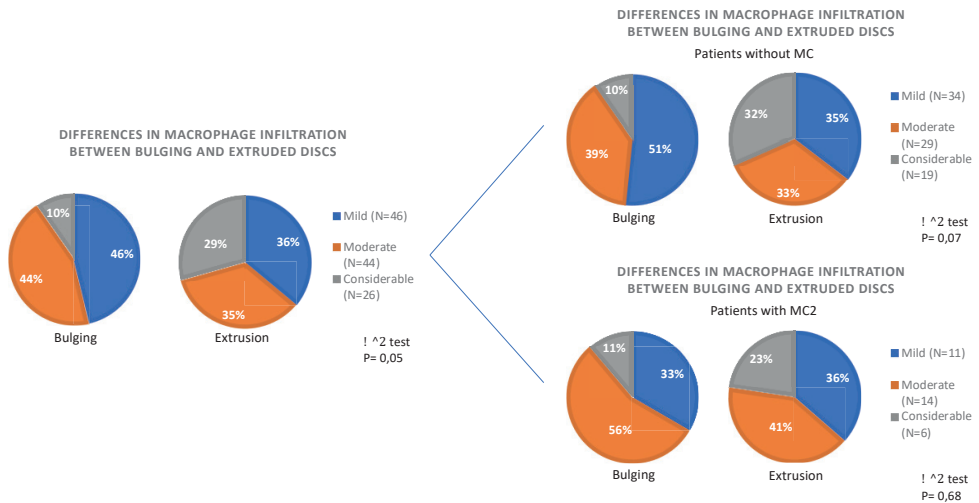
A and B both represent independent observers. The agreement was assessed after the consensus reading.

Figure 1. Macrophage infiltration categories.



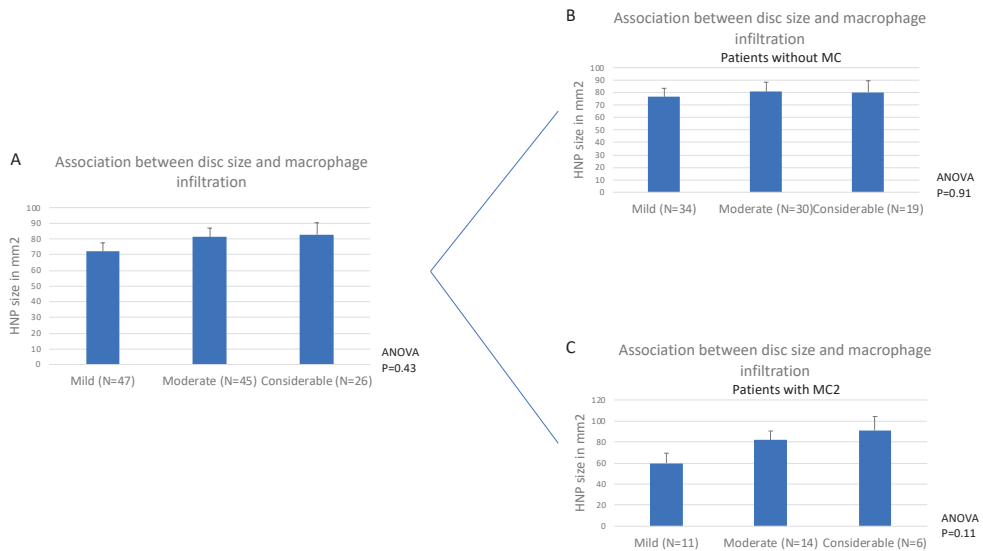
Examples of mild (0-10 macrophages per cm²), moderate (10-100 macrophages per cm²), and considerable (>100 macrophages per cm²) inflammation

Figure 2. Association between the type of disc herniation and macrophage infiltration at baseline.



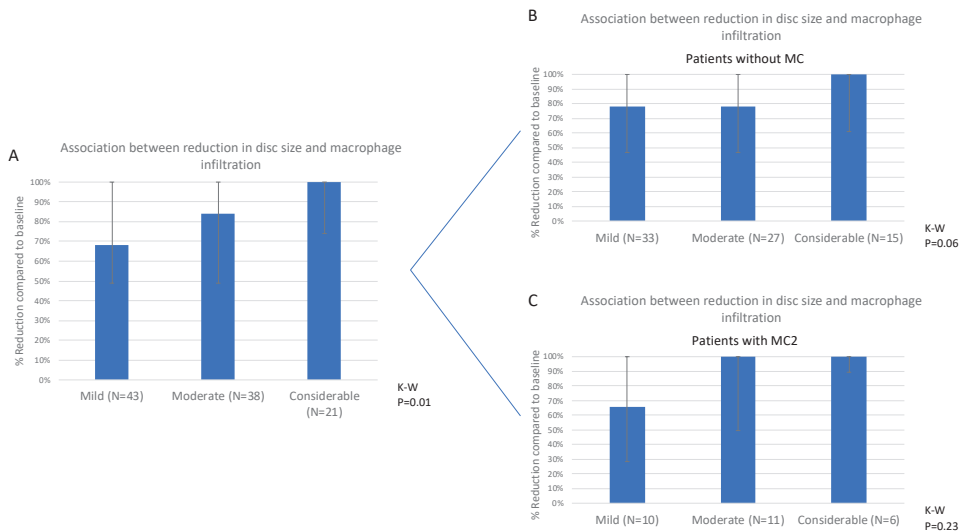
Pie charts display the distribution of the macrophage infiltration groups in percentages, a χ^2 test was performed to assess the significance in distribution between bulging and extruded discs, and p values are given. Comparison a for the whole population, 2B for patients without MC, and 2C for patients with MC

Figure 3. Association between the size of disc herniation and macrophage infiltration at baseline.



Macrophage infiltration groups are shown on the X axis, values on the Y axis are mean HNP surface size in squared. 3B for patients without MC, and 3C for patients with MC

Figure 4. Association between the degree of macrophage infiltration and the percentage surface reduction of the HNP between baseline and 1-year follow-up on MRI.



Macrophage infiltration groups are shown on the X axis, values on the Y axis are median percentages of axial surface millimeter, error bars are SEs and p values for the one-way ANOVA test are provided. Comparison a for the whole population, 4B for patients without MC, and 4C for patients with MC

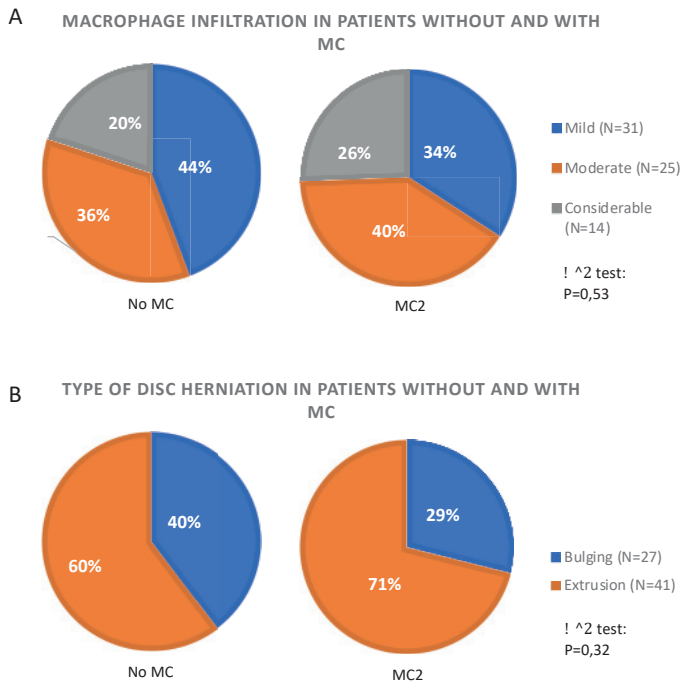
Supplementary Appendix

Supplementary Table S1. MRI study variables

Disc level	Variable	Category
Disc level with the most severe nerve root compression	Disc level	1. Not applicable: no nerve root compression 2. L2L3 3. L3L4 4. L4L5 5. L5S1
	Disc contour at this level	1. Bulging: presence of disc tissue circumferentially (50-100%) beyond the edges of the ring apophyses 2. Herniation: localized displacement of disc material beyond the normal margins of the intervertebral disc space
	Certainty about the presence of disc herniation	1. Definite about the presence: no doubt about the presence 2. Probable about the presence: some doubt but probability > 50% 3. Possible about the presence: reason to consider but probability < 50% 4. Definite about the absence: no doubt about the absence
	Loss of disc height at this level	1. Yes 2. No
	Signal intensity of nucleus pulposus on T2 images at this level	1. Hypointensity 2. Normal 3. Hyperintensity
	Certainty about the presence of nerve root compression	1. Definite about the presence: no doubt about the presence 2. Probable about the presence: some doubt but probability > 50% 3. Possible about the presence: reason to consider but probability < 50% 4. Definite about the absence: no doubt about the absence
	Spinal canal stenosis	1. Yes 2. No
	Disappearance of epidural fat	1. Completely disappeared 2. Partly disappeared 3. No disappearance
	Presence of vertebral end plate changes and its extent	1. No VESC (Vertebral Endplate Signal Changes) 2. VESC type 1: hypointense on T1-weighted sequences and hyperintense on T2-weighted sequences 3. VESC type 2: increased signal on T1 weighted sequences and isointense or slightly hyperintense signal on T2 weighted sequences 4. VESC type 3: hypointense both on T1- and T2-weighted sequences 5. VESC type 1 and 2
		1. Yes 2. No

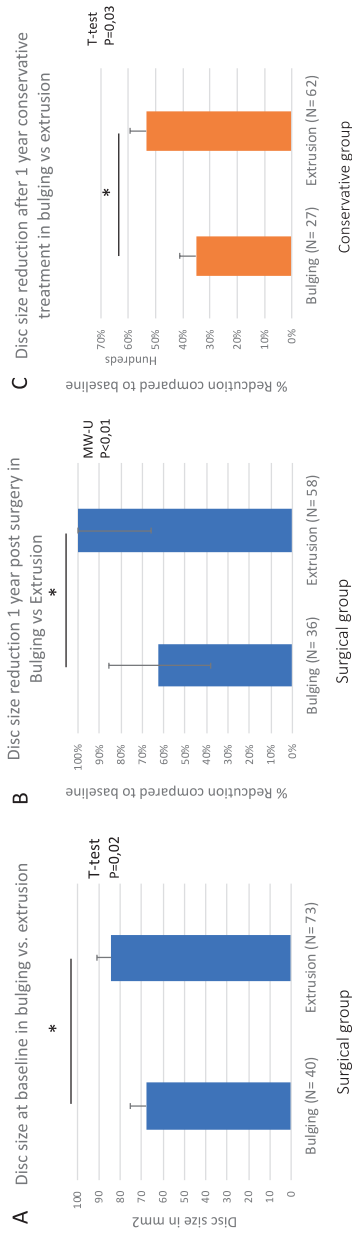
Disc level	Variable	Category
If a disc herniation is considered	Presence of impaired discs at more than one level	<ol style="list-style-type: none"> 1. Central zone: zone within the vertebral canal between sagittal planes through the medial edges of each facet 2. Sub-articular zone: zone, within the vertebral canal, sagittally between the plane of the medial edges of the pedicles and the plane of the medial edges of the facets, and coronally between the planes of the posterior surfaces of the vertebral bodies and the under anterior surfaces of the superior facets. 3. Foraminal zone: zone between planes passing through the medial and lateral edges of the pedicles 4. Extra-foraminal zone: the zone beyond the sagittal plane of the lateral edges of the pedicles, having no well-defined lateral border
	Location	1. Right 2. Left 3. Right and left
	Size disc herniation in relation to spinal canal	<ol style="list-style-type: none"> 1. Large stenosing, size >75% of the spinal canal 2. Large, size 50-75% of the spinal canal 3. Average, size 25-50% of the spinal canal 4. Small, size <25% of the spinal canal
	Form disc herniation	<ol style="list-style-type: none"> 1. Protrusion: localized displacement of disc material beyond the intervertebral disc space, with the base against the disc of origin broader than any other dimension of the protrusion. 2. Extrusion: localized displacement of disc material beyond the intervertebral disc space, with the base against the disc of origin narrower than any one distance between the edges of the disc material beyond the disc space measured in the same plane, or when no continuity exists between the disc material beyond the disc space and that within the disc space.

Figure S1. Associations between MC and macrophage infiltration, and between MC and the type of disc herniation at baseline.



S1A: Pie charts display the distribution of the macrophage infiltration groups in percentages. **S1B:** Pie charts display the distribution of the bulging and extruded discs in percentages, χ^2 tests were performed to assess the significance in distribution between patients without and with MC, p values are given.

Figure S2. Association between disc type and percentage reduction in axial surface at one year.



SZA compares disc size in the surgical group: disc type is shown on the X axis and values on the Y axis are mean size at baseline, error bars are SE. **SZB** compares axial surface reduction in the surgical group: disc type is shown on the X axis and values on the Y axis are median percentages of axial surface reduction compared to baseline, error bars are interquartile ranges. **SZC** compares axial surface reduction in the conservative group: disc type is shown on the X axis and values on the Y axis are percentages of axial surface reduction compared to baseline error bars are SE's. p values for T-tests or Mann Whitney-U tests are provided accordingly