

Bone-anchored annular closure following lumbar discectomy reduces risk of complications and reoperations within 90 days of discharge

Peter Douglas Klassen¹
Derek Thomas Bernstein²
Hans-Peter Köhler³
Mark P Arts⁴
Bradley Weiner²
Larry E Miller⁵
Claudius Thomé⁶

On behalf of the Annular Closure RCT Study Group

¹Department of Neurosurgery, St. Bonifatius Hospital, Lingen, Germany; ²Department of Orthopaedic Surgery, Methodist Hospital, Houston, TX, USA; ³Department of Neurosurgery, Asklepios Westklinikum Hamburg, Hamburg, Germany; ⁴Department of Neurosurgery, Haaglanden Medical Center, The Hague, The Netherlands; ⁵Miller Scientific Consulting, Inc., Asheville, NC, USA; ⁶Department of Neurosurgery, Medical University Innsbruck, Innsbruck, Austria

Correspondence: Larry E Miller
Miller Scientific Consulting, Inc., 1854
Hendersonville Road, #231, Asheville,
NC 28803, USA
Tel +1 828 450 1895
Email larry@millerscientific.com

Purpose: The purpose of this study was to evaluate perioperative complications of lumbar discectomy with or without bone-anchored annular closure device (ACD) implant in patients at high risk of recurrent disc herniation.

Methods: This was a post hoc analysis of a randomized controlled trial that compared outcomes of lumbar discectomy with or without additional placement of an ACD. Patients presented with imaging evidence of lumbar disc herniation and radicular pain that was unresponsive to conservative care. Randomization occurred intraoperatively following discectomy completion and confirmation of annular defect width ≥ 6 mm. Main outcomes included serious adverse events (SAEs) from any cause, device- or procedure-related SAEs, and reoperations at the index level. The perioperative period included all outcomes occurring between the day of surgery and 90 days following hospital discharge.

Results: Analyses were performed on a modified intention-to-treat population consisting of 272 patients treated with ACD and 278 patients treated with lumbar discectomy only (controls). Mean patient age was 44 years, 59% were men, and mean body mass index was 26 kg/m². Baseline patient characteristics and operative outcomes were comparable between groups. The risks of all-cause SAE (9.7% vs 16.3%, $p=0.056$), device- or procedure-related SAE (4.5% vs 10.2%, $p=0.02$), and index-level reoperation (1.9% vs 5.4%, $p=0.03$) were lower with ACD vs controls. In multivariable logistic regression, control group assignment and female gender were independently associated with higher risk of device- or procedure-related SAE and index-level reoperation, respectively.

Conclusion: In patients undergoing lumbar discectomy to treat symptomatic intervertebral disc herniation, adjunctive placement of an ACD reduces the risk for perioperative complications occurring through 90 days following hospital discharge.

Keywords: annular closure, complications, herniation, lumbar discectomy, randomized controlled trial, reoperation

Introduction

Lumbar discectomy for intervertebral disc herniation is one of the most commonly performed spine procedures.¹ While lumbar discectomy is generally considered safe and provides initial symptomatic relief in most patients, the risk of perioperative complications ranges from 13% to 15%.² As patients with large annular defects following lumbar discectomy have higher rates of recurrent disc herniation over long-term follow-up,³ it is plausible that these patients may experience greater risk of complications during the perioperative period. Perioperative complications place a large financial burden on patients, providers, hospitals, commercial payers, and federal health care agencies.⁴

Hospitals are highly incentivized to reduce complication rates, both in-hospital and following discharge.⁵ Treatment paths that reduce perioperative complication risks following lumbar discectomy have potential to improve patient outcomes and lower health care costs. We report here a post hoc analysis from a randomized controlled trial of a bone-anchored annular closure device (ACD) in patients at high risk of recurrent disc herniation after lumbar discectomy. We hypothesized that ACD implant following lumbar discectomy would reduce the risk for perioperative complications and associated reoperations occurring through 90 days following hospital discharge.

Methods

This prospective multicenter randomized controlled trial compared outcomes after lumbar discectomy with or without additional placement of an ACD. The primary objective of the study was to determine safety and effectiveness of adjunctive ACD placement through 2-year follow-up in patients at high risk of herniation recurrence.⁶ We report here a post hoc analysis of perioperative complication data from this study. The study was approved by the local ethics committee at each site and all patients provided written informed consent. The trial was prospectively registered at ClinicalTrials.gov (NCT01283438).

Baseline assessments included physical examination, neurologic examination with straight leg lift test, and anteroposterior, lateral, and flexion–extension x-rays. Disc herniation was confirmed with magnetic resonance imaging with T1- and T2-weighted axial and sagittal images and low-dose multiplanar computed tomography. Eligible patients were adults aged 21–75 years with single-level posterior or posterolateral disc herniation between L1 and S1, confirmation of neural compression on magnetic resonance imaging, positive straight leg raise or femoral stretch test, and radicular pain (visual analog scale ≥ 40) and back dysfunction (Oswestry Disability Index ≥ 40) that was unresponsive to at least 6 weeks of conservative treatment. Main exclusion criteria included disc height < 5 mm at the index level, grade II or higher spondylolisthesis, or prior surgery at the index level. Complete inclusion and exclusion criteria in this study have been described elsewhere.⁶

All patients underwent single-level limited lumbar discectomy.⁷ Nuclear material removed from within or outside the disc was placed dry in a syringe and the volume was measured and recorded. Following the discectomy, intraoperative verification of annular defect between 4 and 6 mm in height and 6 and 10 mm in width was required; the latter

requirement has been reported to increase recurrence risk.^{3,8} Eligible patients were then randomized (1:1) to additional ACD implant or discectomy only (controls). Randomization assignments were obtained intraoperatively through a web-based platform. Patients randomized to the control group underwent a standard wound closure procedure with no treatment of the annulus. Patients randomized to ACD underwent device implantation under fluoroscopic guidance as per the manufacturer's surgical technique and instructions for use. Neither patients nor surgeons were blinded to treatment assignment, except for patients in the Netherlands (comprising 15% of the sample) who were blinded because of regional regulatory requirements.

The ACD (Barricaid, Intrinsic Therapeutics, Inc., Woburn, MA, USA) is a permanent implant with two components – an occlusion component and an anchor component (Figure 1). The occlusion component consists of a flexible polymer mesh that is designed to prevent reherniation by physically blocking the annulus at the postsurgery defect and a platinum–iridium radiopaque marker to enable radiographic visualization. The anchor component consists of titanium alloy that is placed into either the caudal-adjacent or cranial-adjacent vertebral body to resist migration. The device is preassembled onto a single-use delivery tool.

Anteroposterior and neutral lateral radiographs were obtained prior to discharge. Immediate postoperative care, including discharge instructions, ambulation restrictions, and physical therapy recommendations, was provided at investigator discretion. Patients returned for planned in-clinic visits at 6 weeks and 3 months post-treatment. Complications and reoperations that occurred in hospital and through 90 days following hospital discharge were recorded and independently monitored for accuracy. An independent data safety monitoring board reviewed and adjudicated all adverse events, monitored study stopping rules, and had the authority to stop the study at any point.

A Bayesian adaptive sample size procedure to test for superiority of two co-primary endpoints at 2-year follow-up has been described elsewhere.⁶ Analyses were performed using a modified intention-to-treat population, which included all randomized control patients and all randomized ACD patients where the delivery tool was inserted through the skin. Continuous data were reported as mean and standard deviation unless otherwise noted. Categorical data were presented as counts and percentages. Group comparisons of procedural data were performed using independent-samples *t*-test or Wilcoxon–Mann–Whitney test for continuous data and Fisher's exact test for categorical data. Time-to-event data

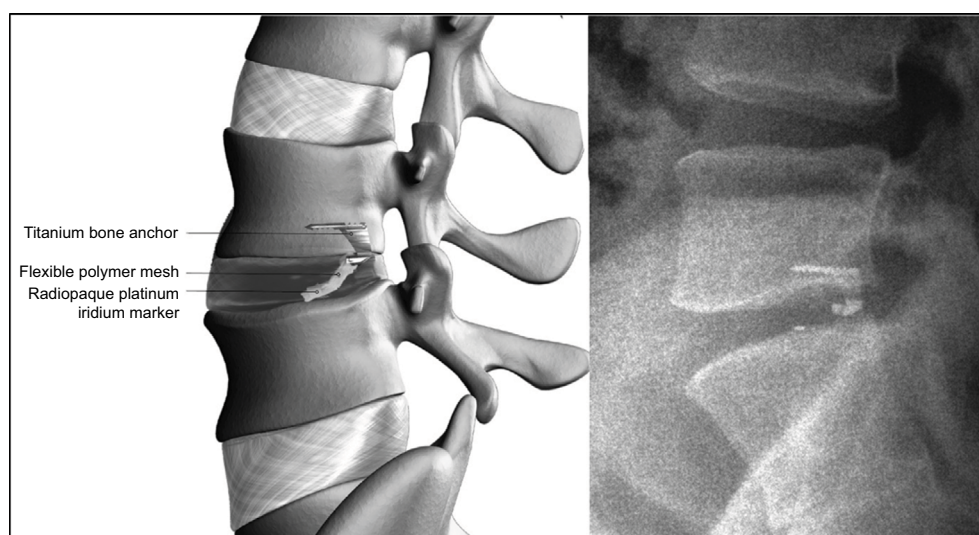


Figure 1 The Barricaid annular closure device.

Notes: Graphic representation of the annular closure prosthesis, with a titanium bone anchor holding the polyester mesh in place (left panel); lateral radiograph showing implanted device at L4–L5 (right panel).

through 90 days following hospital discharge were analyzed using Kaplan–Meier methods; group comparisons were performed with a log-rank test. Univariate logistic regression was performed to determine the association of baseline patient characteristics with risk of perioperative complications. Independent variables included treatment group, age, gender, body mass index, smoking status, and index level. Variables that entered the univariate model at $p < 0.2$ were considered for inclusion in a multivariate model. The multivariate model utilized a forward–backward stepwise elimination variable selection process that optimized the Akaike information criterion by assessing model fit penalized for the number of estimated parameters.⁹ Statistical analyses were performed using SAS v9.4 (SAS Institute, Cary, NC, USA) and R v3.3.2 (R Foundation for Statistical Computing, Vienna, Austria).

Results

A total of 647 patients met the initial study entry criteria and underwent lumbar discectomy. After exclusion of 93 patients because of ineligible annular defect dimensions, 554 patients were randomly allocated to ACD ($n=276$) or control ($n=278$) at 21 centers in six European countries between December 2010 and October 2014. A list of participating centers and investigators is provided in Table S1. Four patients randomized to ACD did not undergo device implantation because of anatomical considerations; therefore, the modified intention-to-treat population consisted of 272 patients treated with ACD and 278 patients treated with limited discectomy only. Follow-up compliance through the 3-month visit was 98% with ACD and 95% with control (Figure 2).

Baseline patient characteristics were comparable between groups (Table 1). Mean patient age was 44 years, 59% were men, and mean body mass index was 26 kg/m². Surgery was most frequently performed at L5–S1 (56%) or L4–L5 (41%). Intraoperative findings were similar between groups including mean volume of nucleus removed (1.3 mL) and mean annular defect area (39 mm²). Surgery time (skin-to-skin) was longer in the ACD group owing to the additional time required for implant placement (70 vs 52 minutes, $p < 0.001$). Median blood loss (50 mL) and hospital stay (3 days) were identical between groups.

Through 90 days following hospital discharge, the cumulative probability of all-cause serious adverse event (SAE) was lower with ACD vs control (9.7% vs 16.3%, $p=0.056$). The cumulative probability of a device- or procedure-related SAE through 90 days following hospital discharge was 4.5% with ACD and 10.2% with control ($p=0.02$) (Figure 3). The most common of these SAEs was herniation at the index level, reported in six (2.2%) ACD patients and 19 (6.8%) control patients ($p=0.01$) (Table 2).

The risk of reoperation at the index level over this period was also lower with ACD (1.9% vs 5.4%, $p=0.03$) (Figure 4). In the control group, 18 index-level reoperations were performed in 15 patients. These included discectomy ($n=8$), discectomy with ACD implant ($n=2$), hematoma evacuation ($n=3$), wound revision ($n=1$), and multiple (four) wound revisions ($n=1$). In the ACD group, six index-level reoperations were performed in five patients including discectomy ($n=3$), decompression with subsequent fusion ($n=1$), and implant removal ($n=1$). The risk of reoperation for recurrent

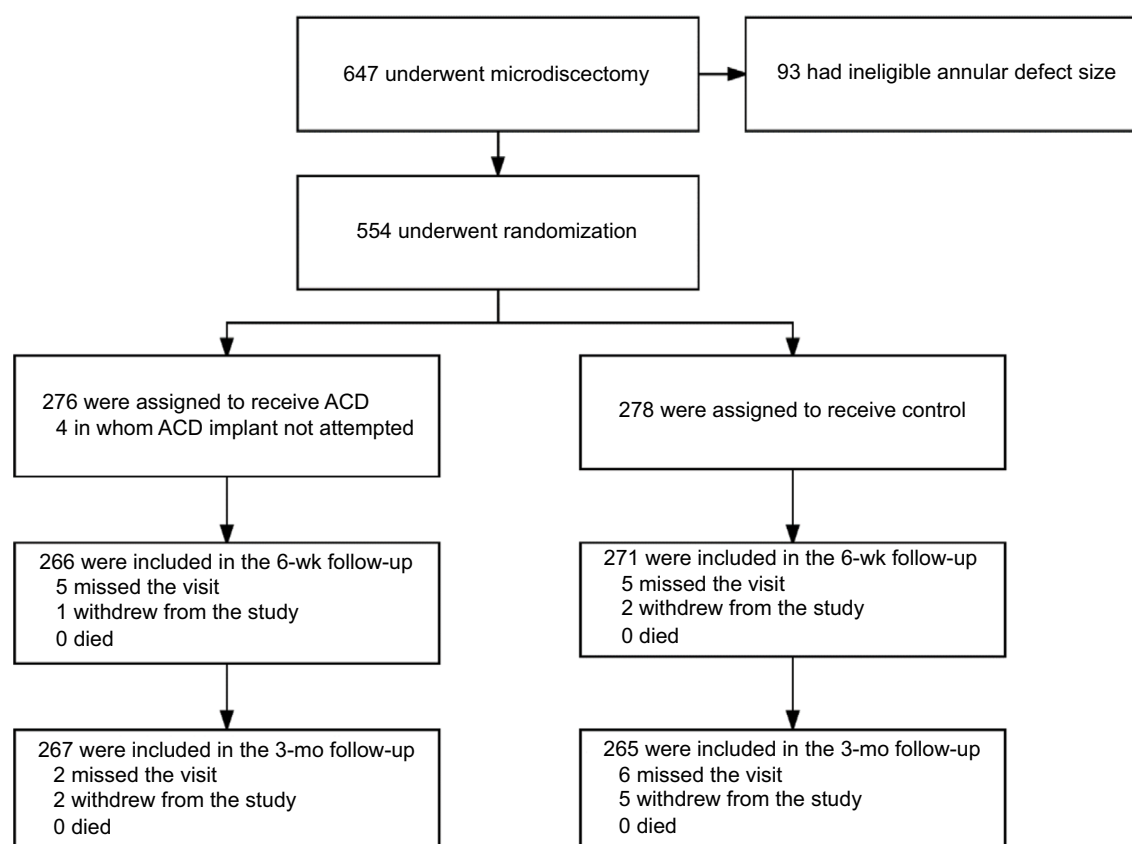


Figure 2 CONSORT patient flow diagram.

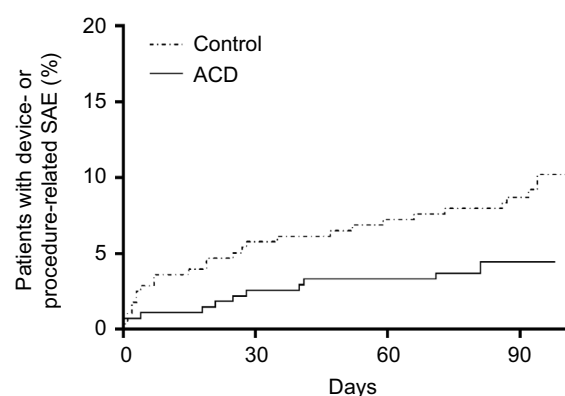
Notes: Modified intention-to-treat population consisted of 272 patients with attempted ACD implant and 278 patients assigned to control. Compliance with clinical follow-up at 3 months follow-up was 98% with annular closure device (ACD) and 95% with control.

Abbreviations: ACD, annular closure device; mo, months; wk, weeks.

Table 1 Baseline patient characteristics

Characteristic	Annular closure (N=272)	Control (N=278)
Age – years – mean ± SD	43±11	44±10
Male gender – n (%)	156 (57)	171 (62)
Body mass index – kg/m ² – mean ± SD	26±4	26±4
Smoking history – n (%)	173 (64)	175 (63)
Medical history – n (%)*		
Musculoskeletal	95 (35)**	91 (33)***
Head and neck	62 (23)**	54 (20)***
Gastrointestinal	53 (20) ^a	59 (21)**
Cardiovascular	49 (18)**	48 (17)**
Genitourinary	39 (14)**	35 (13)**
Skin	29 (11)**	30 (11)**
Respiratory	28 (10)**	44 (16)**
Index level – n (%)		
L2–L3	2 (1)	1 (<1)
L3–L4	8 (3)	5 (2)
L4–L5	123 (45)	101 (36)
L5–S1	139 (51)	171 (62)

Notes: *Medical history variables reported with frequency of 10% or more in either group. **Data from two patients not reported. ***Data from one patient not reported. ^aData from three patients not reported.



No. at risk

ACD	272	263	261	259
Control	278	261	254	249

Figure 3 Cumulative probability of device- or procedure-related SAE through 90 days following hospital discharge.

Notes: Day 0 is surgery date. Kaplan–Meier estimate is 4.5% with ACD and 10.2% with control (log-rank p -value = 0.02).

Abbreviations: ACD, annular closure device; SAE, serious adverse event.

herniation was lower in the ACD group vs controls (0.7% vs 4.0%, $p=0.02$).

In univariate logistic regression, treatment group was the only variable that was associated with occurrence of a device- or procedure-related SAE (control vs ACD: odds

ratio [OR]=2.2, $p=0.03$). Control group assignment (OR=3.1, $p=0.03$) and female gender (OR=2.8, $p=0.03$) were associated with greater risk of index-level reoperation in the univariate model (Table 3). In each multivariable model, control group assignment and female gender were independently associated with greater complication risks (Table 4). Comparing women to men in the control group, the probabilities were 13.1% vs 7.0% for a device- or procedure-related SAE and 8.4% vs 3.5% for index-level reoperation. In the ACD group, the probabilities were 6.0% vs 3.2% for a device- or procedure-related SAE and 3.4% vs 0.6% for index-level reoperation.

Discussion

The results of this post hoc analysis from a randomized controlled trial demonstrated that reherniation was the most common perioperative complication following lumbar discectomy for disc herniation. Further, adjunctive use of an ACD following lumbar discectomy reduced the risk for perioperative complications and associated reoperations compared to lumbar discectomy alone. To the authors' knowledge, this is the first study to characterize perioperative complications in patients with large annular defects following lumbar discectomy.

In the current economic climate with heightened scrutiny on health care resource utilization, hospitals continually face a delicate balance between limiting unnecessarily long hospitalizations and reducing the risk of complications and readmissions following discharge. With Diagnosis-Related Group-based flat payment schedules¹⁰ and financial penalties for higher-than-average readmission rates set forth by the Hospital Readmission Reduction Program,¹¹ adoption of therapies that reduce in-hospital and postdischarge complications may improve patient outcomes and potentially increase hospital margins. Given that 480,000 discectomy procedures are performed annually in the USA¹² and that adjunctive ACD implant reduced perioperative complication risk by over 50%,

Table 2 Listing of device- or procedure-related serious adverse events through 90 days following hospital discharge

Characteristic	Annular closure (N=272)	Control (N=278)
Patients reporting at least one SAE	12 (4.4%)*	26 (9.4%)**
Index-level herniation	6	19
Wound complication	2	6
Epidural hematoma	0	2
Anchor/mesh dislocation	2	0
Back/leg pain	2	0
Coronary heart disease	0	1
Dural tear	0	1
Hypesthesia	1	0

Notes: *13 events in 12 patients, **29 events in 26 patients.

Abbreviation: SAE, serious adverse event.

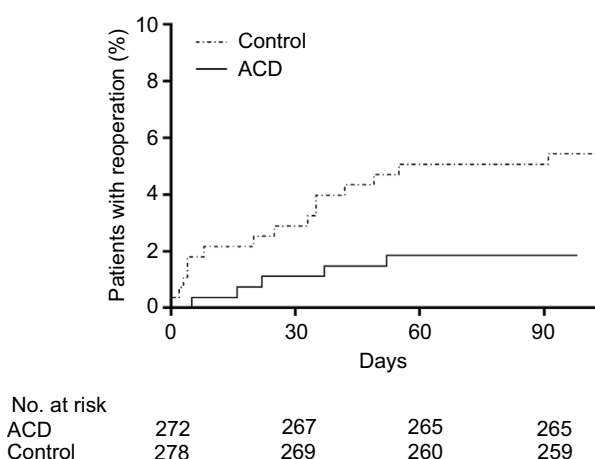


Figure 4 Cumulative probability of index-level reoperations through 90 days following hospital discharge.

Notes: Day 0 is surgery date. Kaplan-Meier estimate is 1.9% with ACD and 5.4% with control (log-rank p -value =0.03).

Abbreviation: ACD, annular closure device.

Table 3 Univariate logistic regression of predictors of complications through 90 days following hospital discharge

Variable	Unit of measure	Device- or procedure-related SAE			Index-level reoperation		
		OR	95% CI	p-value	OR	95% CI	p-value
Treatment group	Control vs ACD	2.24	1.10, 4.53	0.03	3.05	1.09, 8.50	0.03
Gender	Female vs male	1.90	0.98, 3.68	0.06	2.83	1.11, 7.21	0.03
Current smoker	Yes vs no	1.80	0.92, 3.50	0.09	2.40	0.94, 6.12	0.07
Age	Per 10 years	1.30	0.96, 1.77	0.09	1.32	0.87, 2.00	0.19
Index level	L5	1	(Reference)		1	(Reference)	
	L4	1.58	0.80, 3.12	0.18	2.43	0.94, 6.28	0.07
	L2 or L3	2.45	0.52, 11.68	0.26	2.88	0.33, 24.90	0.34
Body mass index	Per 5 kg/m ²	1.11	0.75, 1.64	0.61	1.51	0.91, 2.51	0.11

Abbreviations: ACD, annular closure device; OR, odds ratio; SAE, serious adverse event.

Table 4 Multivariate logistic regression of predictors of complications through 90 days following hospital discharge

Variable	Unit of measure	Device- or procedure-related SAE			Index-level reoperation		
		OR	95% CI	p-value	OR	95% CI	p-value
Treatment group	Control vs ACD	2.32	1.14, 4.71	0.02	3.22	1.15, 9.05	0.03
Gender	Female vs male	1.98	1.01, 3.86	<0.05	3.00	1.17, 7.68	0.02

Abbreviations: ACD, annular closure device; OR, odds ratio; SAE, serious adverse event.

it is plausible that significant cost savings may be realized with routine ACD placement in patients undergoing lumbar discectomy who are at high risk for reherniation based on annular defect size.

We found that in patients with large annular defects following lumbar discectomy for disc herniation, perioperative complication risk was higher in women than men. This is in agreement with other studies that have reported female gender as an independent risk factor for inferior outcomes following lumbar disc surgery.¹³ However, we are unaware of other studies that have examined the influence of gender on perioperative complications following lumbar discectomy. It is plausible that the greater complication risk in women was confounded by preoperative status. In the Swedish Spine Register of over 15,000 patients, women scheduled for lumbar disc surgery reported inferior clinical status compared to men including greater pain severity, inferior walking ability, higher consumption of analgesics, and greater disability.¹⁴ In the current study, no other preoperative variable was independently associated with perioperative complication risk. Still, the possible influence of unmeasured preoperative status variables on the elevated complication risk in women remains a possibility. Despite the higher risk of perioperative complications in women following lumbar discectomy, the risk of device- or procedure-related SAE and index-level reoperation in women remained lower with ACD vs controls.

Strengths of this study included a randomized design, rigorous study entry criteria, and adjudicated complication reporting. There were also several limitations of the study that warrant consideration. First, as investigators and most patients were aware of treatment allocation, the potential for expectation bias must be acknowledged. Second, while SAEs were reported and reviewed in a standardized fashion, the decision to perform surgery was based on patient willingness and surgeon recommendations. Third, the results of this study are applicable to patients with large annular defects following lumbar discectomy. Perioperative complication risk in patients with small vs large annular defects has not been reported to date.

Conclusion

In patients undergoing lumbar discectomy to treat symptomatic intervertebral disc herniation, adjunctive placement of an ACD reduces the risk for perioperative complications occurring through 90 days following hospital discharge.

Disclosure

Intrinsic Therapeutics provided support for the conduct of the study, analysis of the data, and preparation of this manuscript. PDK, CT, and LEM are consultants to Intrinsic Therapeutics. The authors report no other conflicts of interest in this work.

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Supplementary materials

Table S1 List of Participating Centers, Ethics Committees, and Investigators

Site name	Ethics committee name	Ethics committee address	Principal investigator	Sub-investigator(s)
LKH Graz (Graz, Austria)	Ethikkommission: Medizinische Universität Graz	Auenbruggerplatz 2, A-8036 Graz	Sandro Eustacchio	Karin Pistracher Martin Trummer
ZNA Middleheim (Antwerp, Belgium)	ZNA/OCMW Antwerpen	Lindendreef 1, 2020 Antwerpen	Robert Hes*	Guido Dua
OLVG-location West (formerly: Sint Lucas Andreas Ziekenhuis) (Amsterdam, The Netherlands)	St. Lucas Andreas Ziekenhuis	Postbus 9243, 1006 AE Amsterdam	Gerrit J Bouma	Michiel B Lequin P Richard Schuurman
St. Bonifatius Hospital GmbH (Lingen, Germany)	MHH (Hannover Medical School)	MHH Ethics Committee Org, Unit 9515, 30623 Hannover	Peter Douglas Klassen	Bert Baume
Klinikum Augsburg (Augsburg, Germany)	BLÄK (Ethics Committee of the Bavarian Regional Medical Association) MHH (Hannover Medical School)	Mühlbauerstrasse 16, 81677 Munich MHH Ethics Committee Org, Unit 9515, 30623 Hannover	Volkmar Heidecke	
Charite-Virchow-Klinikum (Berlin, Germany)	Charité Ethikkommission	Charitéplatz 1, 10117 Berlin	Peter Vajkoczy	Johannes Woitzik Marcus Czabanka Charlotte Flüh
University Medical Center Schleswig-Holstein (Kiel, Germany)	Medizinische Fakultät Der Christian-Albrechts- Universität Zu Kiel	Universitäts-Kinderklinik, Schwanenweg 20, 24105 Kiel	Hubertus Mehdorn† Senol Jadik	Isabel Lübbling Geoffrey Lesage
OLV Ziekenhuis (Aalst, Belgium)	O.L.Vrouwziekenhuis Aalst Ethisch Comité ZNA/OCMW Antwerpen	Lindendreef 1, 2020 Antwerpen Lindendreef 1, 2020 Antwerpen	Frederic Martens	Djaya Kools
Universitätsmedizin Mannheim (Mannheim, Germany)	Med. Ethik-Kommission II	Maybachstr. 14, 68169 Mannheim	Aldemar Hegewald	Jason Perrin
University Hospital Düsseldorf (Düsseldorf, Germany)	Heinrich Heine Universität Düsseldorf Ethikkommission der Medizinischen Fakultät	Moorenstrasse 5, 40225 Düsseldorf	Richard Bostelmann	Jan Cornelius
Donauislar Klinikum Deggendorf (Deggendorf, Germany)	BLÄK (BLAK defers to MHH after initial approval) MHH (Hannover Medical School)	Mühlbauerstrasse 16, 81677 Munich MHH Ethics Committee Org, Unit 9515, 30623 Hannover	Stefan Rath	Adisa Kuršumović
Asklepios Westklinikum Hamburg (Hamburg, Germany)	Ethik-Kommission Der Ärztchamber Hamburg	Ärztchamber Hamburg, Postfach 76 01 09, 22051 Hamburg	Hans-Peter Köhler	Dirk Lympius Ekkehard von Saldern
Isala Ziekenhuis (Zwolle, The Netherlands)	Isala Klinieken	Locatie Sophia, Gebouw B, Kamer 1.43, Postbus 10400, 8000 GK Zwolle	Willem Art van den Brink	Dharmin Nanda
Medical University Innsbruck (Innsbruck, Austria)	Ethikkommission der Medizinischen Universität Innsbruck	Geschäftsstelle, Innrain 43, 1. Stock, A-6020 Innsbruck	Claudius Thomé	Pierre-Pascal Girod
A.Z Nikolaas Hospital (St. Nikolaas, Belgium)	Commissie voor Medische Ethiek, A.Z. Nikolaas ZNA/OCMW Antwerpen	Moerlandstraat 1, 9100 Sint- Niklaas Lindendreef 1, 2020 Antwerpen	Erik Van den Kelft	David Van der Planken
Haaglanden Medical Center: Antoniusshove (Den Haag, The Netherlands)	METC Zuidwest Holland	Postbus 5011, 2600 GA Delft	Mark Arts	
Haaglanden Medical Center: Westeinde (Den Haag, The Netherlands)	METC Zuidwest Holland	Postbus 5011, 2600 GA Delft	Jasper Wolfs	

(Continued)

Table S1 (Continued)

Site name	Ethics committee name	Ethics committee address	Principal investigator	Sub-investigator(s)
Kantonsspital Aarau (Aarau, Switzerland)	Kanton Aargau, Kantonale Ethikkommission	Bachstrasse 15, 5001 Aarau	Javier Fandino	Jenny Kienzler
Knappschafts-Krankenhaus Bochum-Langendreer (Bochum, Germany)	Ethics Committee of the Medical Faculty of the Ruhr University	Gesundheitscampus 33, 44801 Bochum	Martin Barth	Christopher Brenke
Centre Hospitalier Régional Universitaire de Lille (CHRU), Hôpital Roger Salengro (Lille, France)	Comité de Protection des Personnes (CPP)	1 place de Verdum, 59045 Lille Cedex	Richard Assaker	Fahed Zairi
Orthopädische Klinik und Poliklinik Universitätsmedizin Rostock (Rostock, Germany)	Ethikkommission an der Medizinischen Fakultät der Universität Rostock	Universitätsmedizin Rostock, PF 10 08 88, 18055 Rostock	Susanne Fröhlich	Dorit Panser-Schulz

Notes: *PI moved to AZ Klina, Brasschaat, Belgium. EC approval at this site to follow patients enrolled at original site. PI at ZNA Middleheim now Guido Dua. †PI Mehdorn retired. Former Co-Investigator Jadik now PI.

Abbreviations: EC, ethics committee; PI, principal investigator.

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