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Automated Percutaneous Discectomy and Percutaneous Endoscopic Discectomy

Table of Contents
Coverage
Policy Guidelines
Description
Rationale
Coding
References
Policy History

Related Policies (if applicable)
None

Disclaimer

Carefully check state regulations and/or the member contract.

Each benefit plan, summary plan description or contract defines which services are covered, which services are excluded, and which services are subject to dollar caps or other limitations, conditions or exclusions. Members and their providers have the responsibility for consulting the member's benefit plan, summary plan description or contract to determine if there are any exclusions or other benefit limitations applicable to this service or supply. **If there is a discrepancy between a Medical Policy and a member's benefit plan, summary plan description or contract, the benefit plan, summary plan description or contract will govern.**

Coverage

Automated percutaneous discectomy is **considered experimental, investigational and/or unproven** as a technique of intervertebral disc decompression in individuals with back pain and/or radiculopathy related to disc herniation in the lumbar, thoracic, or cervical spine.

Percutaneous endoscopic discectomy is **considered experimental, investigational and/or unproven** as a technique of intervertebral disc decompression in individuals with back pain and/or radiculopathy related to disc herniation in the lumbar, thoracic, or cervical spine.

NOTE: This policy does not address endoscopic foraminotomy alone (without discectomy).

Policy Guidelines

None.

Description

Surgical management of herniated intervertebral discs most commonly involves discectomy or microdiscectomy, performed manually through an open incision. Automated percutaneous discectomy involves placement of a probe within the intervertebral disc under image guidance with aspiration of disc material using a suction cutting device. Endoscopic discectomy involves the percutaneous placement of a working channel under image guidance, followed by visualization of the working space and instruments through an endoscope, and aspiration of disc material.

Background

Back pain or radiculopathy related to herniated discs is an extremely common condition and a frequent cause of chronic disability. Although many cases of acute low back pain and radiculopathy will resolve with conservative care, surgical decompression is often considered when the pain is unimproved after several months and is clearly neuropathic in origin, resulting from irritation of the nerve roots. Open surgical treatment typically consists of discectomy in which the extruding disc material is excised. When performed with an operating microscope, the procedure is known as microdiscectomy.

Minimally invasive options have also been researched, in which some portion of the disc material is removed or ablated, although these techniques are not precisely targeted at the offending extruding disc material. Ablative techniques include laser discectomy and radiofrequency (RF) decompression which are addressed in separate policies. Intradiscal electrothermal annuloplasty is another minimally invasive approach to low back pain. In this technique, RF energy is used to treat the surrounding disc annulus which is addressed in a separate policy.

Herein, this policy addresses automated percutaneous and endoscopic discectomy, in which the disc decompression is accomplished by the physical removal of disc material rather than its ablation. Traditionally, discectomy is performed manually through an open incision, using cutting forceps to remove nuclear material from within the disc annulus. This technique has been modified by automated devices that involve placement of a probe within the intervertebral disc and aspiration of disc material using a suction cutting device. Endoscopic techniques may be intradiscal or may involve the extraction of non-contained and sequestered disc fragments from inside the spinal canal using an interlaminar or transforaminal approach. Following insertion of the endoscope, the decompression is performed under visual control.

Regulatory Status

The Dekompressor® Percutaneous Discectomy Probe (Stryker), Herniatome Percutaneous Discectomy Device (Gallini Medical Devices), and the Nucleotome® (Clarus Medical) are examples of percutaneous discectomy devices that have been cleared for marketing by the U.S. Food and Drug Administration (FDA) through the 510(k) process. The FDA indication for these

products is for “aspiration of disc material during percutaneous discectomies in the lumbar, thoracic and cervical regions of the spine.” FDA product code: HRX.

A variety of endoscopes and associated surgical instruments have also been cleared for marketing by the FDA through the 510(k) process.

Rationale

Medical policies assess the clinical evidence to determine whether the use of a technology improves the net health outcome. Broadly defined, health outcomes are length of life, quality of life, and ability to function—including benefits and harms. Every clinical condition has specific outcomes that are important to patients and to managing the course of that condition. Validated outcome measures are necessary to ascertain whether a condition improves or worsens, and whether the magnitude of that change is clinically significant. The net health outcome is a balance of benefits and harms.

To assess whether the evidence is sufficient to draw conclusions about the net health outcome of a technology, 2 domains are examined: the relevance and the quality and credibility. To be relevant, studies must represent one or more intended clinical use of the technology in the intended population and compare an effective and appropriate alternative at a comparable intensity. For some conditions, the alternative will be supportive care or surveillance. The quality and credibility of the evidence depend on study design and conduct, minimizing bias and confounding that can generate incorrect findings. The randomized controlled trial (RCT) is preferred to assess efficacy, however, in some circumstances, nonrandomized studies may be adequate. RCTs are rarely large enough or long enough to capture less common adverse events and long-term effects. Other types of studies can be used for these purposes and to assess generalizability to broader clinical populations and settings of clinical practice.

Automated Percutaneous Discectomy

Clinical Context and Therapy Purpose

The purpose of automated percutaneous discectomy is to provide a treatment option that is an alternative to or an improvement of existing therapies for individuals with herniated intervertebral disc(s).

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with herniated intervertebral disc(s).

Interventions

The therapy being considered is automated percutaneous discectomy.

Comparators

The following therapies and practices are currently being used to treat herniated intervertebral disc(s): conservative therapy and open discectomy or microdiscectomy.

Outcomes

The general outcomes of interest are symptoms, functional outcomes, quality of life, and treatment-related morbidity. Specific outcomes measured by specific instruments include improvements in functional outcomes assessed on the Oswestry Disability Index (ODI), reductions in pain using a visual analog scale (VAS), improvements in quality of life measured on the 36-Item Short-Form Health Survey (SF-36) and Euro-QOL-5D, and treatment-related morbidity including surgical success/failure and complications. To assess outcomes, follow-up at 1 year is considered appropriate.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess longer term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

Systematic reviews have assessed automated percutaneous discectomy compared to other interventions; however, the majority of these reviews contained observational studies published more than a decade ago with generally small patient populations and inconsistent results. Lewis et al. (2015) published the most recent systematic review and network meta-analysis comparing trials of 21 different treatment strategies for sciatica. (1) Examples of the 21 treatment strategies included in the analysis are conservative care, disc surgery, intraoperative interventions, epidural injections, biologic agents, and percutaneous discectomy. Under the category of “percutaneous discectomy,” reviewers combined automated percutaneous discectomy, percutaneous automated nucleotomy, nucleoplasty, and laser discectomy. They searched 28 databases and trial registries through December 2009. Ninety studies were included and 10 involved the percutaneous discectomy category as an intervention. Of the 10, 4 are relevant to this medical policy: 2 case-control studies of percutaneous endoscopic discectomy (2006, 2007), 1 RCT of percutaneous endoscopic discectomy (1993), and 1 RCT of automated percutaneous discectomy (1995). The remaining studies were published in a foreign language or involved other comparators (nucleolysis and chemonucleolysis). The global effects odds ratio for the category of percutaneous discectomy compared with inactive control was 0.82 (95% confidence interval [CI], 0.39 to 1.72), which was inferior to disc surgery, epidural injections, and intraoperative interventions. The pain intensity weighted mean difference for the category of percutaneous discectomy compared with inactive control was 11.5 (95% CI, -18.6 to 41.6). Reviewers concluded that there was no support for the effectiveness of percutaneous discectomy for the treatment of sciatica. Due to the inclusion of additional

interventions into the broad category of percutaneous discectomy in this review, the relevance of these results to this medical policy is limited.

Randomized Controlled Trials

The 2002 Lumbar Automated Percutaneous Discectomy Outcomes Group (LAPDOG) trial was an RCT that compared automated percutaneous discectomy with open discectomy in patients with lumbar disc herniation. (2) No additional RCTs have been identified since the 2002 LAPDOG trial. The trial was designed to recruit 330 patients but enrolled 36 patients for reasons not readily apparent. Twenty-seven patients were available at follow-up, with efficacy reported by 41% of those undergoing automated percutaneous discectomy and by 40% of those undergoing conventional discectomy. The trialists concluded that “It is difficult to understand the remarkable persistence of percutaneous discectomy in the face of a virtually complete lack of scientific support for its effectiveness in treated lumbar disc herniation.” The tables below more fully describe key characteristics, results, and limitations of the LAPDOG trial.

Table 1. Characteristics of the LAPDOG Trial

Study	Countries	Sites	Dates	Participants	Interventions
Haines et al. (2002) (2)	US, Canada	10	NR	Patients with predominantly unilateral leg pain or paresthesia with no previous treatment for lumbar spinal disease, at least 2 of 4 objective signs, and an imaging study confirming disc herniation at the appropriate level	Automated percutaneous discectomy vs. conventional discectomy

LAPDOG: Lumbar Automated Percutaneous Discectomy Outcomes Group; NR: not reported; US: United States.

Table 2. Results of the LAPDOG Trial

Study	Treatment success ^a (at 6 months)	Treatment failure ^b (at 6 months)	SF-36 Physical Functioning Subscore	SF-36 General Health Subscore	Modified Roland Score
Haines et al. (2002) (2)					
N	27	27	NR	NR	NR
Automated percutaneous discectomy	7 (41%)	10 (59%)	Pre- vs. postoperative mean difference: 35.7	Pre- vs. postoperative mean difference: 5.0	Pre- vs. postoperative mean difference: 9.7
Conventional discectomy	4 (40%)	6 (60%)	Pre- vs. postoperative mean	Pre- vs. postoperative mean	Pre- vs. postoperative mean

			difference: 36.1	difference: 8.0	difference: 10.6
p	.95	.95	.96	.58	.74

LAPDOG: Lumbar Automated Percutaneous Discectomy Outcomes Group; NR: not reported; SF-36: 36-Item Short-Form Health Survey.

^a Success was defined as either an excellent or good result as defined by an outcome matrix.

^b Failure was defined as not achieving success or requiring a second procedure during the follow-up period.

Table 3. Study Relevance Limitations of the LAPDOG Trial

Study	Population ^a	Intervention ^b	Comparator ^c	Outcomes ^d	Duration of Follow-up ^e
Haines et al. (2002) (2)	3. Investigators believed that study inclusion criteria reflected an existing population with lumbar disc disease; however, results from only 27 patients were eventually analyzed from a planned enrollment of 330 patients			4. Primary outcomes of "success" or "failure" largely subjective in nature; investigators admit that the outcome measurement tool used cannot be precisely reproduced	1, 2. Outcomes reported only for 6 months of follow-up; 12 month follow-up was achieved for only 19 patients and the study did not report any of these results

LAPDOG: Lumbar Automated Percutaneous Discectomy Outcomes Group.

The study limitations stated in this table are those notable in the current literature review; this is not a comprehensive gaps assessment.

^a Population key: 1. Intended use population unclear; 2. Clinical context is unclear; 3. Study population is unclear; 4. Study population not representative of intended use; 5. Other.

^b Intervention key: 1. Not clearly defined; 2. Version used unclear; 3. Delivery not similar intensity as comparator; 4. Not the intervention of interest; 5. Other.

^c Comparator key: 1. Not clearly defined; 2. Not standard or optimal; 3. Delivery not similar intensity as intervention; 4. Not delivered effectively; 5. Other.

^d Outcomes key: 1. Key health outcomes not addressed; 2. Physiologic measures, not validated

surrogates; 3. No CONSORT reporting of harms; 4. Not establish and validated measurements; 5. Clinical significant difference not prespecified; 6. Clinical significant difference not supported; 7. Other.

^e Follow-Up key: 1. Not sufficient duration for benefit; 2. Not sufficient duration for harms; 3. Other.

Table 4. Study Design and Conduct Limitations of the LAPDOG Trial

Study	Allocation ^a	Blinding ^b	Selective Reporting ^c	Data Completeness ^d	Power ^e	Statistical ^f
Haines et al. (2002) (2)		1, 2. Blinding did not appear to occur		1. Of 34 initially randomized patients, 9 were lost to follow-up, 6 month follow-up data was obtained on only 27 patients, and 12 month follow-up data was obtained for only 19 patients	3. Power estimates led the investigators to plan enrollment of 330 patients in order to reliably identify a difference in success rate of 15% or greater; results were analyzed on 27 patients	1. Beyond the cursory discussion of lack of power, a discussion of the statistical analyses is nonexistent

The study limitations stated in this table are those notable in the current literature review; this is not a comprehensive gaps assessment.

^a Allocation key: 1. Participants not randomly allocated; 2. Allocation not concealed; 3. Allocation concealment unclear; 4. Inadequate control for selection bias.

^b Blinding key: 1. Not blinded to treatment assignment; 2. Not blinded outcome assessment; 3. Outcome assessed by treating physician.

^c Selective Reporting key: 1. Not registered; 2. Evidence of selective reporting; 3. Evidence of selective publication.

^d Data Completeness key: 1. High loss to follow-up or missing data; 2. Inadequate handling of missing data; 3. High number of crossovers; 4. Inadequate handling of crossovers; 5. Inappropriate exclusions; 6. Not intent to treat analysis (per protocol for noninferiority trials).

^e Power key: 1. Power calculations not reported; 2. Power not calculated for primary outcome; 3. Power not based on clinically important difference.

^f Statistical key: 1. Analysis is not appropriate for outcome type: (a) continuous; (b) binary; (c) time to event; 2. Analysis is not appropriate for multiple observations per patient; 3. Confidence intervals and/or p values not reported; 4. Comparative treatment effects not calculated.

All published trials have focused on lumbar disc herniation. There were no RCTs of automated percutaneous discectomy for cervical or thoracic disc herniation. A review of the evidence from the American Society of Interventional Pain Physicians (ASIPP) (2013) noted that “even though

Dekompressor [disc removal system] may be considered a new interventional modality, the early studies were published approximately 8 years ago. Consequently, one would expect that the technique's continued use would be supported by more recent, high-quality evaluations.”
(3)

Section Summary: Automated Percutaneous Discectomy

The evidence for automated percutaneous discectomy in individuals who have herniated intervertebral disc(s) includes small RCTs and systematic reviews. Evidence from small RCTs does not support the use of this procedure. Well-designed and executed RCTs are needed to determine the benefits and risks of this procedure.

Percutaneous Endoscopic Discectomy

Clinical Context and Therapy Purpose

The purpose of percutaneous endoscopic discectomy is to provide a treatment option that is an alternative to or an improvement on existing therapies for individuals with herniated intervertebral disc(s).

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Populations

The relevant population of interest is individuals with herniated intervertebral disc(s).

Interventions

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Comparators

The following therapies and practices are currently being used to treat herniated intervertebral disc(s): conservative therapy and open discectomy or microdiscectomy.

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The general outcomes of interest are symptoms, functional outcomes, quality of life, and treatment-related morbidity. Specific outcomes measured by specific instruments include improvements in functional outcomes assessed on the ODI, reductions in pain using a VAS, improvements in quality of life measured on the SF-36 and Euro-QOL-5D, and treatment-related morbidity including surgical success/failure and complications. To assess outcomes, follow-up at 1 year is considered appropriate.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.

- To assess longer term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

A number of systematic reviews have evaluated the efficacy and safety of percutaneous endoscopic discectomy compared to open discectomy or microendoscopic discectomy. A comparison of the trials included in more recent systematic reviews (2017 to present) is shown in Table 5. Characteristics and results of these reviews are summarized in Tables 6 and 7A/B.

Table 5. Trials Included in Systematic Reviews of Percutaneous Endoscopic Discectomy Versus Other Discectomy Procedures

Trials	Systematic Reviews							
	Phan et al. (2017) (4)	Shi et al. (2019) (5)	Yu et al. (2019) (6)	Zhou et al. (2020) (7)	Xu et al. (2020) (8)	Bao et al. (2021) (9)	Gadjradj et al. (2021) (10)	Zhao et al. (2022) (11)
Ma et al. (2022) (12)								●
Wang et al. (2021) (13)								●
Rajamani et al. (2021) (14)								●
Jing et al. (2021) (15)								●
Jarebi et al. (2021) (16)								●
Meyer et al. (2020) (17)								●
Chen et al. (2020) (18)								●
Kim et al. (2019) (19)								●
Ahn et al. (2019) (20)								●
Liu et al. (2018) (21)								●
Sun et al. (2017) (22)								●

Jeong et al. (2006) (23)								●
Akcakaya et al. (2016) (24)							●	
Choi et al. (2018) (25)							●	
Dai et al. (2020) (26)							●	
Krappel et al. (2001) (27)							●	
Tacconi et al. (2019) (28)							●	
Tacconi et al. (2020) (29)							●	
Tao et al. (2018) (30)							●	
Wang et al. (2017) (31)							●	
Xu et al. (2020) (32)							●	
Ahn et al. (2016) (33)						●		●
Chang et al. (2018) (34)						●	●	●
Liu et al. (2017) (35)						●		●
Pan et al. (2016) (36)						●	●	●
Yao et al. (2017) (37)						●		
Yao et al. (2017) (38)						●		●
Gibson et al. (2017) (39)				●			●	●

Hsu et al. (2013) (40)				●				
Kim et al. (2007) (41)				●		●		●
Qu et al. (2017) (42)				●				●
Wang et al. (2013) (43)				●				●
Zhao et al. (2012) (44)				●				
Yoon et al. (2012) (45)	●	●	●		●			
Li et al. (2015) (46)	●				●			●
Sinkemani et al. (2015) (47)	●	●	●		●			●
Song et al. (2017) (48)		●	●		●			●
Tu et al. (2017) (49)					●			
Liu et al. (2018) (21)		●	●	●	●	●		
Li et al. (2018) (50)		●	●	●	●			●
Abdurexiti et al. (2018) (51)		●	●		●			
Chen et al. (2018) (52)		●	●	●	●	●		●
Liu et al. (2012) (53)			●					
Wu et al. (2009) (54)		●						
Yang et al. (2015) (55)		●		●				
Duan et al. (2016) (56)		●						
Zhao et al. (2016) (57)		●						
Ding et al. (2017) (58)		●						●

Li et al. (2017) (59)		●						
Liu et al. (2017) (60)		●						
Luo et al. (2017) (61)		●						
Qu et al. (2017) (62)		●						
Chen et al. (2018) (63)		●						
Wu et al. (2018) (64)		●						
Belykh et al. (2016) (65)		●						
Chen et al. (2015) (66)	●							●
Choi et al. (2016) (67)	●			●				●
Garg et al. (2011) (68)	●							
Hermantin et al. (1999) (69)	●						●	
Huang et al. (2005) (70)	●							
Hussein et al. (2014) (71)	●							
Kleinpeter et al. (1995) (72)	●							
Lee et al. (2009) (73)	●					●		●
Martin-Laez et al. (2012) (74)	●							
Mayer et al. (1993) (75)	●			●		●	●	●
Ohya et al. (2016) (76)	●							

Pan et al. (2014) (77)	●							●
Righesso et al. (2007) (78)	●							
Ruetten et al. (2008) (79)	●							
Ruetten et al. (2009) (80)	●					●		
Sasaoka et al. (2006) (81)	●							
Schizas et al. (2005) (82)	●							
Teli et al. (2010) (83)	●							
Ruetten et al. (2007) (84)	●							
Ruetten et al. (2008) (85)						●		
Lee et al. (2006) (86)						●		

Table 6. Summary of Systematic Reviews of Percutaneous Endoscopic Discectomy Versus Other Discectomy Procedures

Study	Dates	Trials	Participants	N (Range)	Design	Duration
Zhao et al. (2022) (11)	To May 2022	33	Patients with lumbar disc herniation who underwent PTED, MED or other surgical procedures	6467 (20-1856)	7 RCTs; 26 non-randomized controlled retrospective studies	Not reported
Bai et al. (2021) (9)	To February 2018	14	Patients with lumbar disc herniation who underwent PELD or	2528 (74-902)	4 RCTs; 10 cohort studies	Not reported

			other surgical procedures			
Gadjradj et al. (2021) (10)	To April 2020	14	Patients with lumbar disc herniation who underwent PTED or open microdiscectomy	1465 (30-462)	9 RCTs; 5 prospective nonrandomized comparative studies	Follow-up: 3 to 12 months
Xu et al. (2020) (8)	Search dates not stated; included trials from 2012 to 2018	9	Patients with single-level lumbar disc herniation who underwent PELD or MED for treatment	984 (51-216)	1 Prospective RCTs; 8 Retrospective nonrandomized comparative studies	Follow-up: 1 to > 6 years
Zhou et al. (2020) (7)	To October 2018	12	Patients with lumbar disc herniation who underwent PELD or MED for treatment	2400 (40-915)	4 RCTs; 8 Retrospective nonrandomized comparative studies	Follow-up: 3 to 46 months
Yu et al. (2019) (6)	To August 31, 2018	8	Patients with lumbar disc herniation who underwent PTED or MED procedures and were followed for at least 6 months	805 (51-216)	1 Prospective RCTs; 7 Observational studies	Follow-up: 6 months to 5 years
Shi et al. (2019) (5)	To July 2018	18	Patients with single-level lumbar disc herniation with sciatica who underwent PELD or MED for treatment	2161 (51-273)	8 Prospective studies; 10 Retrospective studies	Follow-up: 3 months to >6 years
Phan et al. (2017) (4)	To February 2016	23	Patients who underwent either an endoscopic or open approach for disc herniation; the endoscopic approach consisted of patients who	24,487 (20-26,612)	10 Prospective RCTs; 4 Prospective observational studies; 9 Retrospective observational studies	Follow-up: 3 to 104 months

			underwent either FED or MED while the open approach included those who underwent open discectomy or micro-discectomy			
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FED: full-endoscopic technique discectomy; MED: microendoscopic discectomy; PELD: percutaneous endoscopic lumbar discectomy; PTED: percutaneous transforaminal endoscopic discectomy; RCT: randomized controlled trial

Table 7A. Results of Systematic Reviews of Trials of Percutaneous Endoscopic Discectomy Versus Other Discectomy Procedures

Study	Length of stay	Leg pain VAS	Lower back pain VAS
Zhao et al. (2022) (11)			
Total (N)	1231	1487	1372
Pooled effect (95% CI); p value	MD -2.42 (-3.21 to -1.63); .0001	MD -0.23(-0.61 to 0.15); .60	MD -0.49 (-0.84 to -0.14); .006
I ² (p)	95%; .00001	51%; .03	90%; .00001
Bai et al. (2021) (9)			
Total (N)	NR	NR	NR
Pooled effect (95% CI); p value	MD -2.59 (-3.87 to -1.31); <.001	MD 0.00 (-0.10 to 0.10); .991	MD -0.17 (-0.55 to 0.21); .384
I ² (p)	72.1%; .001	0.0%; .996	88.3%; <.001
Gadjradj et al. (2021) (10)			
Total (N)		621 and 152	
Pooled effect (95% CI); p value		3 to 6 month MD 0.05 (-0.10 to 0.21) 12 month MD 0.11 (-0.30 to 0.53)	
I ² (p)		30%; .23	
Xu et al. (2020) (8)			
Total (N)	NR	NR	NR
Pooled effect (95% CI); p value	OR -1.041 (-1.493 to -0.583); .000	6 months to 2 years OR -0.138 (-0.384 to 0.108); .270 2 years OR 0.020 (-0.193 to 0.233); .855	6 months to 2 years -0.456 (-0.947 to 0.034); .068 2 years OR -0.856 (-1.488 to -0.224); .008
I ² (p)		53.8%; .090; 6 months to 2 years 4.4%; .351; 2 years	88%; .000; 6 months to 2 years 86.7%; .001; 2 years
Zhou et al. (2020) (7)			

Total (N)			
Pooled effect (95% CI); p value			
I ² (p)			
Yu et al. (2019) (6)			
Total (N)	707	NR	NR
Pooled effect (95% CI); p value	MD -1.92 (-2.90 to -0.94); <.001	1 year postop or last follow-up: MD -0.07 (-0.22 to 0.08); .38	1 year postop or last follow-up: MD -0.41 (-0.76 to -0.06); .02
I ² (p)	88%		
Shi et al. (2019) (5)			
Total (N)	1717	742	742
Pooled effect (95% CI); p value	MD -2.29 (3.03 to -1.55); <.00001	At last follow-up: MD -0.18 (-0.45 to 0.09); .19	At last follow-up: MD -0.77 (-1.31 to -0.24); .005
I ² (p)	96%; <.00001	88%; <.00001	95%; <.00001
Phan et al. (2017) (4)			
Total (N)	685	390	
Pooled effect (95% CI); p value	MD -4.79 (-6.52 to -3.07); <.00001	MD -0.04 (-0.37 to 0.30); .84	
I ² (p)	99%; <.00001	70%; .003	

CI: confidence interval; MD: mean difference; NR: not reported; ODI: Oswestry Disability Index; OR: odds ratio; RR: risk ratio; VAS: visual analogue scale; WMD: weighted mean difference

Table 7B. Results of Systematic Reviews of Trials of Percutaneous Endoscopic Discectomy Versus Other Discectomy Procedures

Study	ODI	Overall complication rate	Reoperation	Recurrence or residue
Zhao et al. (2022) (11)				
Total (N)	1687	2,372	2,226	2,621
Pooled effect (95% CI); p value	MD -2.21 (-4.17 to -0.25); .03	OR 0.94 (0.67 to 1.32); .71	OR 1.67 (1.17 to 2.36); .004	OR 1.55 (1.07 to 2.24); .02
I ² (p)	88%; .00001	0%; .65	0%; .89	0%; .93
Bai et al. (2021) (9)				
Total (N)	NR	NR		NR
Pooled effect (95% CI); p value	MD -0.29 (-1.00 to 0.43); .434	relative risk 0.86 (0.63 to 1.18); .361		relative risk 1.65 (1.08 to 2.52); .021
I ² (p)	0.0%; .996	51.5%; .024		26.1%; .220
Gadraj et al. (2021) (10)				
Total (N)	621 and 152			

Pooled effect (95% CI); p value	3-to-6-month MD -0.09 (-0.24 to 0.07) 12-month MD -0.11 (-0.45 to 0.24)			
I ² (p)	9%; .83			
Xu et al. (2020) (8)				
Total (N)	NR	NR	NR	NR
Pooled effect (95% CI); p value	6 months to 2 years -0.077 (-0.370 to 0.215); .604 2 years OR -0.425 (-0.724 to -0.127); .005	OR 0.972 (0.635 to 1.488); .896	OR 1.136 (0.415 to 3.108); .805	OR 1.306 (0.664 to 2.566); .439
I ² (p)	75.3%; .000; 6 months to 2 years 52.7%; .121; 2 years			
Zhou et al. (2020) (7)				
Total (N)			787	972
Pooled effect (95% CI); p value			OR 1.77 (1.18 to 2.64); .006	OR 1.60 (1.01 to 2.53); .05
I ² (p)			0%; .97	0%; .94
Yu et al. (2019) (6)				
Total (N)	NR	659		443
Pooled effect (95% CI); p value	1 year postop or last follow-up: MD -0.27 (-1.71 to 1.16); .71	MD 1.01 (0.60 to 1.69); .98		MD 1.31 (0.54 to 3.17); .54
I ² (p)		0%		0%
Shi et al. (2019) (5)				
Total (N)	1337	1527	805	928
Pooled effect (95% CI); p value	At last follow-up: MD -0.30 (-1.02 to 0.42); .41	OR 0.96 (0.65 to 1.43); .85	OR 2.67 (1.07 to 6.67); .04	OR 2.22 (1.02 to 4.83); .05
I ² (p)	55%; .01	0%; .90	0%; .79	0%; .86
Phan et al. (2017) (4)				
Total (N)	303	27,699	995	1081

Pooled effect (95% CI); p value	MD -1.88 (-4.06 to 0.29); .09	OR 0.77 (0.45 to 1.31); .33	OR 1.46 (0.33 to 6.43); .61	OR 1.12 (0.60 to 2.09); .73
I ² (p)	67%; .03	60%; .004	66%; .004	0%; .97

CI: confidence interval; MD: mean difference; NR: not reported; ODI: Oswestry Disability Index; OR: odds ratio; RR: risk ratio; VAS: visual analogue scale; WMD: weighted mean difference.

Results from the systematic reviews were fairly consistent with a significantly reduced length of hospitalization observed with endoscopic discectomy and sometimes significant improvements in VAS or ODI, but only at specific time points. Overall, no consistently significant improvement in VAS, ODI, total complication rate, reoperation, or recurrence was observed with endoscopic discectomy versus other interventions. Authors of the systematic reviews noted multiple limitations including the innate flaws of included studies (i.e., observational designs, a limited number of studies meeting criteria for inclusion, small sample sizes, lack of allocation concealment and blinding), different methodologies contributing to heterogeneity in analyses, loss of usable and sufficient data resulting in difficulty performing accurate analysis of outcomes, and that a majority of the more recently completed studies were completed in China, which may affect the generalizability of the results to other populations.

Randomized Controlled Trials

More recent RCTs not included in any of the systematic reviews were also identified. (87-90) Results of these trials were similar to those seen in the more comprehensive systematic reviews - percutaneous endoscopic discectomy was associated with a significant reduction in length of stay with no consistent improvement in patient-reported outcome measures such as VAS and ODI. Two of the 4 RCTs evaluated treatment-related morbidities, and reported a reduced incidence of intraoperative and postoperative complications and repeat surgeries with percutaneous endoscopic discectomy. Key characteristics, results, and limitations of these RCTs are summarized in the following tables.

Table 8. Characteristics of RCTs of Percutaneous Endoscopic Discectomy

Study	Countries	Sites	Dates	Participants	Interventions
Liu et al. (2023) (90)	Korea	1	July 2016 to July 2021	Patients with L5-S1 lumbar disc herniation	Interlaminar endoscopic lumbar discectomy vs microscopic lumbar discectomy
Gadjradj et al. (2022) (87)	Netherlands	4	February 2016 to April 2019	Patients with sciatica caused by lumbar disc herniation	PTED vs microendoscopic discectomy
Ran et al. (2021) (88)	China	1	August 2016 to February 2020	Patients with highly migrated lumbar disc herniation	PELD with computerized tomography

					navigation vs open discectomy
Wang et al. (2019) (89)	China	1	July 2015 to July 2016	Patients with single-segment lumbar disc herniation with imaging results consistent with symptoms	PTED vs microendoscopic discectomy

PELD: percutaneous endoscopic lumbar discectomy; PTED: percutaneous transforaminal endoscopic discectomy; RCT: randomized controlled trials.

Table 9A. Results of RCTs of Percutaneous Endoscopic Discectomy

Study	Length or stay (days)	Leg pain VAS	Lower back pain VAS	ODI
Liu et al. (2023) (90)				
N	28	28		
Mean difference at 12 months (95% CI)		0.71 (-2.54 to 1.12)	0.08 (-2.25 to 2.42)	8.48 (-1.67 to 18.63)
Interlaminar endoscopic lumbar discectomy	3.69±1.60 days			
Microscopic lumbar discectomy	5.47±1.36 days			
p-value	.003			
Gadjradj et al. (2022) (87)				
N	420	413	413	413
Pooled effect at 12 months (95% CI)	Median (IQR) PTED: 0 (0 to 0) Microendoscopic discectomy: 1 (1 to 1)	MD 7.1 (2.8 to 11.3)	MD 6 (2 to 10)	MD 5.3 (3.0 to 7.7)
p-value				
Ran et al. (2021) (88)				
N		66		
PELD with computerized tomography navigation at 12 months		0.58 ± 0.90		

Open discectomy at 12 months		0.75 ± 0.84		
p-value		.58		
Wang et al. (2019) (89)				
N	90	90	90	90
Percutaneous transforaminal endoscopic discectomy	Postoperative: 3.01 ± 0.52	Preoperative mean score vs. 6 months after surgery: 7.21 vs. 1.05	Preoperative mean score vs 6 months after surgery: 6.40 vs. 1.36	Preoperative mean score vs 6 months after surgery: 58.21% vs. 17.05%
Microendoscopic discectomy	Postoperative: 6.68 ± 0.30	Preoperative mean score vs. 6 months after surgery: 7.09 vs. 0.98	Preoperative mean score vs 6 months after surgery: 6.34 vs. 1.65	Preoperative mean score vs 6 months after surgery: 57.17% vs. 16.98%
p	.001	.097	.523	.864

CI: confidence interval; IQR: interquartile range; MD: mean difference; ODI: Oswestry Disability Index; PELD: percutaneous endoscopic lumbar discectomy; PTED: percutaneous transforaminal endoscopic discectomy; RCT: randomized controlled trials; VAS: visual analogue scale.

Table 9B. Results of RCTs of Percutaneous Endoscopic Discectomy

Study	SF-36 PCS	Complication rates	Repeat surgery within 1 year
Liu et al. (2023) (90)			
N		28	
Mean difference at 12 months (95% CI)			
Interlaminar endoscopic lumbar discectomy		Blood loss: 44±26.67 mL	
Microscopic lumbar discectomy		Blood loss: 20±20.99 mL	
p-value		.009	
Gadjradj et al. (2022) (87)			
N	413	420	420
Pooled effect at 12 months (95% CI)	MD -2.8 (-4.1 to -1.6)	PTED vs microendoscopic discectomy: Dural tears (n=0 vs 8)	PTED vs microendoscopic discectomy: n=9 (5%) vs 14 (6%)

		Nerve root injury (n=0 vs 1) Wound infection (n=3 vs 0) Cerebrospinal fluid leakage (n=1 vs 0)	
p-value			
Ran et al. (2021) (88)			
N		66	
PELD with computerized tomography navigation at 12 months		Infection, n=0 Recurrence, n=1	
Open discectomy at 12 months		Infection, n=1 Recurrence, n=0	
p-value		>.99	
Wang et al. (2019) (89)			
N			
Percutaneous transforaminal endoscopic discectomy			
Micro-endoscopic discectomy			
p			

CI: confidence interval; MD: mean difference; PELD: percutaneous endoscopic lumbar discectomy; PTED: percutaneous transforaminal endoscopic discectomy; RCT: randomized controlled trials; SF-36 PCS: Short-Form-36 Physical Component Score.

Table 10. Study Relevance Limitations of the RCTS of Percutaneous Endoscopic Discectomy

Study	Population ^a	Intervention ^b	Comparator ^c	Outcomes ^d	Duration of Follow-up ^e
Liu et al. (2023) (90)	4. Limited to participants from single site in Korea			1. Morbidity-related outcomes such as complications and reoperation were limited	

Gadjradj et al. (2022) (87)	4. Limited to participants from 3 sites in the Netherlands				
Ran et al. (2021) (88)	4. Limited to participants from single site in China	4. PELD was used with computerized tomography navigation		1. Morbidity-related outcomes such as complications were limited	
Wang et al. (2019) (89)	4. Study population similar to other trials with regard to age, sex; however, included patients from a single Chinese hospital			1. Morbidity-related outcomes such as complication and reoperation rates were not reported	1,2. Outcomes reported only for 6 months of follow-up

PELD: percutaneous endoscopic lumbar discectomy; RCT: randomized controlled trials.

The study limitations stated in this table are those notable in the current literature review; this is not a comprehensive gaps assessment; 4. Enrolled populations do not reflect relevant diversity; 5. Other.

^a Population key: 1. Intended use population unclear; 2. Study population is unclear; 3. Study population not representative of intended use.

^b Intervention key: 1. Not clearly defined; 2. Version used unclear; 3. Delivery not similar intensity as comparator; 4. Not the intervention of interest.

^c Comparator key: 1. Not clearly defined; 2. Not standard or optimal; 3. Delivery not similar intensity as intervention; 4. Not delivered effectively.

^d Outcomes key: 1. Key health outcomes not addressed; 2. Physiologic measures, not validated surrogates; 3. No CONSORT reporting of harms; 4. Not established and validated measurements; 5. Clinically significant difference not prespecified; 6. Clinically significant difference not supported.

^e Follow-Up key: 1. Not sufficient duration for benefit; 2. Not sufficient duration for harms.

Table 11. Study Design and Conduct Limitations of the RCTs of Percutaneous Endoscopic Discectomy

Study	Allocation ^a	Blinding ^b	Selective Reporting ^c	Data Completeness ^d	Power ^e	Statistical ^f
Liu et al. (2023) (90)		1,2. Blinding did not occur			1. No mention of power	

Gadjraj et al. (2022) (87)	4. A proportion of patients with a strong preference for PTED who were randomised to open microdiscectomy dropped out of the study after randomization	1,2. Blinding did not occur				
Ran et al. (2021) (88)	3. Allocation concealment unclear	1,2. Blinding did not occur			1. Power calculations not reported	
Wang et al. (2019) (89)	3. Allocation concealment unclear	1,2. Blinding did not appear to occur			1. Power calculations not reported	

PTED: percutaneous transforaminal endoscopic discectomy; RCT: randomized controlled trials.

The study limitations stated in this table are those notable in the current literature review; this is not a comprehensive gaps assessment.

^a Allocation key: 1. Participants not randomly allocated; 2. Allocation not concealed; 3. Allocation concealment unclear; 4. Inadequate control for selection bias.

^b Blinding key: 1. Not blinded to treatment assignment; 2. Not blinded outcome assessment; 3. Outcome assessed by treating physician.

^c Selective Reporting key: 1. Not registered; 2. Evidence of selective reporting; 3. Evidence of selective publication.

^d Data Completeness key: 1. High loss to follow-up or missing data; 2. Inadequate handling of missing data; 3. High number of crossovers; 4. Inadequate handling of crossovers; 5. Inappropriate exclusions; 6. Not intent to treat analysis (per protocol for noninferiority trials).

^e Power key: 1. Power calculations not reported; 2. Power not calculated for primary outcome; 3. Power not based on clinically important difference.

^f Statistical key: 1. Analysis is not appropriate for outcome type: (a) continuous; (b) binary; (c) time to event; 2. Analysis is not appropriate for multiple observations per patient; 3. Confidence intervals and/or p values not reported; 4. Comparative treatment effects not calculated.

Observational Studies

Comparative observational studies with at least a 2-year follow-up are summarized below.

Yang et al. (2024) published the results of a retrospective study that compared transforaminal endoscopic lumbar discectomy (n=89) to microdiscectomy (n=65) in patients with lumbar disc herniation. (91) The mean follow-up was 5.5 years. Postsurgical VAS scores of the leg and back reached their lowest point at 1 year and were maintained until the final follow-up. Oswestry

Disability Index scores continued to decrease until final follow-up in patients who underwent transforaminal endoscopic lumbar discectomy but remained fairly similar between 1 year and final follow-up among patients who underwent microdiscectomy. Recurrence occurred in 4.49% and 1.54% ($p=.31$) of patients in the transforaminal endoscopic lumbar discectomy and microdiscectomy groups, respectively.

Saghebdoost et al. (2023) published the results of a retrospective study in 434 patients with lumbar disc herniation who underwent transforaminal endoscopic lumbar discectomy or open microdiscectomy. (92) At the end of the 7-year follow-up period, records for 412 patients were evaluable. A similar proportion of patients in both groups had outcomes that were rated as excellent or good (about 88%) according to the modified MacNab criteria. Perioperative complications were similar between groups, but intraoperative blood loss ($p<.05$) and length of hospital stay ($p<.05$) were significantly less in the transforaminal endoscopic lumbar discectomy group. Recurrence that required reoperation occurred in 21 patients in the transforaminal endoscopic lumbar discectomy group and 9 patients in the open microdiscectomy group ($p<.05$).

Yu et al. (2021) published the results of a retrospective multicenter study that followed patients for 2 years after receipt of transforaminal percutaneous endoscopic discectomy ($n=632$) and microendoscopic discectomy ($n=421$) for lumbar disc herniation. (6) Mean blood loss ($p<.001$) and mean duration of hospital stay ($p=.018$) were significantly reduced with transforaminal percutaneous endoscopic lumbar discectomy compared to microendoscopic discectomy. Rates of complications, recurrence, and revisions were similar in both groups. The VAS pain scores did not differ between groups after the first postoperative day. At 1 month postoperatively, there was a significant difference in ODI scores between groups ($p=.016$) in favor of transforaminal percutaneous endoscopic discectomy, but there was no significant difference at other time points.

Song et al. (2021) published a retrospective single-center study that compared percutaneous endoscopic lumbar discectomy ($n=306$) and microendoscopic discectomy ($n=116$) in patients undergoing same day ambulatory surgery for lumbar disc herniation. (48) Mean blood loss and mean duration of hospital stay were significantly less with percutaneous endoscopic lumbar discectomy (both $p<.001$ compared to microendoscopic discectomy). After 3 years of follow-up, VAS pain scores for the back were also significantly lower in the percutaneous endoscopic lumbar discectomy group compared to the microendoscopic discectomy group ($p=.001$) but there was no difference between groups in pain scores for the legs ($p=.224$). Overall recurrence rates ($p=.201$) and ODI scores ($p=.220$) were also similar between groups.

A number of observational studies have also assessed the learning curve (93-95) and the need for longer follow-up for endoscopic discectomy. (96, 97) The largest and longest follow-up to date has been reported by Choi et al. (2015), who examined 10,228 patients at their institution who had had percutaneous endoscopic lumbar discectomy over a 12-year period. (98) They found that 4.3% of cases required reoperation in the first 6 weeks due to incomplete removal of

herniated discs (2.8%), recurrence (0.8%), persistent pain (0.4%), and approach-related pain (0.2%).

Section Summary: Percutaneous Endoscopic Discectomy

The evidence for percutaneous endoscopic discectomy in individuals who have herniated intervertebral disc(s) includes a number of RCTs, systematic reviews, and comparative observational studies with at least 2 years of follow-up. Many of the more recent RCTs were conducted at institutions within China. There are few reports from the United States. Overall, results from RCTs and systematic reviews reveal a significantly reduced length of hospitalization with endoscopic discectomy and occasionally significant improvements in VAS or ODI, but only at specific time points. No consistently significant improvement in VAS, ODI, total complication rate, reoperation, or recurrence was observed with percutaneous endoscopic discectomy versus other interventions.

Summary of Evidence

For individuals who have herniated intervertebral disc(s) who receive automated percutaneous discectomy, the evidence includes randomized controlled trials (RCTs) and systematic reviews of observational studies. Relevant outcomes are symptoms, functional outcomes, quality of life, and treatment-related morbidity. The published evidence from small RCTs is insufficient to evaluate the impact of automated percutaneous discectomy on the net health outcome. Well-designed and executed RCTs are needed to determine the benefits and risks of this procedure. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have herniated intervertebral disc(s) who receive percutaneous endoscopic discectomy, the evidence includes a number of RCTs, systematic reviews, and observational studies. Relevant outcomes are symptoms, functional outcomes, quality of life, and treatment-related morbidity. Many of the more recent RCTs were conducted at institutions within China. There are few reports from the United States. Results do not reveal a consistently significant improvement in patient-reported outcomes and treatment-related morbidity with percutaneous endoscopic discectomy in comparison to other discectomy interventions. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

Practice Guidelines and Position Statements

National Institute for Health and Care Excellence

The National Institute for Health and Care Excellence (NICE; 2005) published guidance on automated percutaneous mechanical lumbar discectomy, indicating that there was limited evidence of efficacy based on uncontrolled case series of heterogeneous groups of patients, and evidence from small RCTs showed conflicting results. (99) The guidance indicated that, in view of uncertainty about the efficacy of the procedure, it should not be done without special arrangements for consent and for audit or research. The guidance was considered for an update in 2009, but failed review criteria; the 2005 guidance is therefore considered current.

A NICE (2016) guidance on percutaneous transforaminal endoscopic lumbar discectomy for sciatica was published. (100) The guidance stated that current evidence is adequate to support the use of percutaneous transforaminal endoscopic lumbar discectomy for sciatica. Choice of operative procedure (open discectomy, microdiscectomy, or percutaneous endoscopic approaches) may be influenced by symptoms, and location and size of prolapsed disc.

A NICE (2016) guidance on percutaneous interlaminar endoscopic lumbar discectomy for sciatica was also published. (101) The guidance stated that current evidence is adequate to support the use of percutaneous interlaminar endoscopic lumbar discectomy for sciatica. Choice of operative procedure (open discectomy, microdiscectomy, or percutaneous endoscopic approaches) may be influenced by symptoms, location, and size of the prolapsed disc.

American Society of Interventional Pain Physicians

The guidelines from the American Society of Interventional Pain Physicians (2013) indicated that the evidence for percutaneous disc decompression with the Dekompessor was limited. (3) There were no recommended indications for the Dekompessor.

North American Spine Society

The North American Spine Society (NASS) (2014) published clinical guidelines on the diagnosis and treatment of lumbar disc herniation. (102) Table 12 summarizes recommendations specific to percutaneous endoscopic discectomy and automated percutaneous discectomy.

Table 12. Recommendations for Lumbar Disc Herniation With Radiculopathy

Recommendations	Grade or LOE^a
Endoscopic percutaneous discectomy is suggested for carefully selected patients to reduce early postoperative disability and reduce opioid use compared with open discectomy	B
There is insufficient evidence to make a recommendation for or against the use of automated percutaneous discectomy compared with open discectomy	I
Endoscopic percutaneous discectomy may be considered for treatment.	C
Automated percutaneous discectomy may be considered for treatment.	C
Patients undergoing percutaneous endoscopic discectomy experience better outcomes if <40 years and symptom duration <3 months.	II

LOE: level of evidence

^a Grade B: fair evidence (level II or III studies with consistent findings; Grade C: poor quality evidence (level IV or V studies). Level of evidence II: lesser quality randomized controlled trial (e.g., <80% follow-up, no blinding, or improper randomization), prospective comparative study, systematic review of level II studies or level I studies with inconsistent results; level of evidence III: case control, retrospective, systematic review of level III studies; level of evidence IV: case series; level of evidence V: expert opinion.

American Pain Society

The clinical practice guidelines from the American Pain Society (2009) found insufficient evidence to evaluate alternative surgical methods to standard open discectomy and microdiscectomy, including laser or endoscopic-assisted techniques, various percutaneous techniques, coblation nucleoplasty, or the Dekompressor. (103)

American Society of Pain and Neuroscience

The American Society of Pain and Neuroscience (ASPN; 2022) published clinical guidance for interventional treatments for low back pain. (104) The guideline states that discectomy procedures (such as percutaneous and endoscopic disc procedures) have favorable safety and efficacy profiles for the treatment of lumbar disc herniation with persistent radicular symptoms; however, it is stated that further research is needed to evaluate complications rates in order for these procedures to supplant classic open microdiscectomy. Recommendations specific to percutaneous endoscopic discectomy are summarized in Table 13.

Table 13. Recommendations for Percutaneous and Endoscopic Procedures

Recommendation	Grade^a	Level of Evidence^b	Level of Certainty [Net Benefit]^c
Percutaneous Endoscopic Discectomy	B	I-a	High

^a Grade B: (The ASPN Back Group recommends the service. There is high certainty that the net benefit is moderate or there is moderate certainty that the net benefit is moderate to substantial.

^b Evidence Level: I-A: At least one controlled and randomized clinical trial, properly designed.

Ongoing and Unpublished Clinical Trials

Currently unpublished trials that might influence this policy are listed in Table 14.

Table 14. Summary of Key Trials

NCT Number	Trial Name	Planned Enrollment	Completion Date
<i>Unpublished</i>			
NCT02602093	Percutaneous Transforaminal Endoscopic Discectomy vs. Open Microdiscectomy for Lumbar Disc Herniation (PTED-study)	682	May 2024 (unknown status)
NCT01997086	Percutaneous Transforaminal Endoscopic Discectomy (PTED) vs. Microendoscopic Discectomy (MED) for the treatment of Lumbar Disc Herniation: A Prospective Randomized Controlled Study	125	Aug 2023 (unknown status)
NCT02742311	EuroPainClinics® Study V Prospective Observational Study (EPCSV)	500	Dec 2021 (completed)

NCT: national clinical trial.

Coding

Procedure codes on Medical Policy documents are included **only** as a general reference tool for each policy. **They may not be all-inclusive.**

The presence or absence of procedure, service, supply, or device codes in a Medical Policy document has no relevance for determination of benefit coverage for members or reimbursement for providers. **Only the written coverage position in a Medical Policy should be used for such determinations.**

Benefit coverage determinations based on written Medical Policy coverage positions must include review of the member's benefit contract or Summary Plan Description (SPD) for defined coverage vs. non-coverage, benefit exclusions, and benefit limitations such as dollar or duration caps.

CPT Codes	62287, 62380
HCPCS Codes	C2614

*Current Procedural Terminology (CPT®) ©2024 American Medical Association: Chicago, IL.

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Centers for Medicare and Medicaid Services (CMS)

The information contained in this section is for informational purposes only. HCSC makes no representation as to the accuracy of this information. It is not to be used for claims adjudication for HCSC Plans.

The Centers for Medicare and Medicaid Services (CMS) does not have a national Medicare coverage position. Coverage may be subject to local carrier discretion.

A national coverage position for Medicare may have been developed since this medical policy document was written. See Medicare's National Coverage at <<https://www.cms.hhs.gov>>.

Policy History/Revision	
Date	Description of Change
10/15/2025	Reviewed. No changes.
10/15/2024	Document updated with literature review. Coverage unchanged. Added references 90-92.
02/01/2024	Document updated with literature review. Coverage unchanged. Added references 11-23, 87, 88, and 103.
10/01/2022	Reviewed. No changes.
11/01/2021	Document updated with literature review. Coverage unchanged. The following references were added: 7-27, 29-41, 43-57, 60, 62-68, 71, 72, 74, 77-80, 93 and 94.
09/15/2020	Reviewed. No changes.
04/01/2019	Policy updated with literature review. Coverage unchanged. Reference 21 added, others updated.
06/15/2018	Reviewed. No changes.
02/15/2018	Document updated with literature review. Coverage has changed, the word percutaneous was added to the following coverage statement: Percutaneous endoscopic discectomy is considered experimental, investigational and/or unproven as a technique of intervertebral disc decompression in patients with back pain and/or radiculopathy related to disc herniation in the lumbar, thoracic, or cervical spine. The Title has changed from: Automated Percutaneous Discectomy and Endoscopic Discectomy.
09/15/2016	Document updated with literature review. Coverage unchanged.
07/01/2015	Reviewed. No changes.
08/01/2014	Medical document divided into: SUR712.023, SUR712.037 and SUR712.038. Document title changed from "Intervertebral Techniques to Treat Chronic Discogenic Back Pain". Document updated with literature review. The previous coverage statement was Percutaneous discectomy by any means is considered experimental, investigational and/or unproven as a technique of intervertebral disc decompression in patients with back pain related to disc herniation in the lumbar, thoracic or cervical spine. The following was added to coverage: Endoscopic discectomy is considered experimental, investigational and/or unproven as a technique of intervertebral disc decompression in patients with back pain and/or radiculopathy related to disc herniation in the lumbar, thoracic, or cervical spine.

06/01/2012	Coverage revised, the following was added; Axial lumbosacral interbody fusion (axial LIF) is considered experimental, investigational and unproven. Description, rationale, and references updated.
12/15/2011	Document updated with literature review. Document completely revised. All references to minimally invasive interbody fusion were removed. Title was changed from "Percutaneous Intervertebral Techniques to Treat Chronic Discogenic Back pain".
12/15/2009	Coverage revised
12/01/2008	Coverage Revised. Codes Revised/Added/Deleted
05/01/2008	Revised/Updated Entire Document
06/01/2006	Revised/Updated Entire Document
11/01/2004	New Medical Document