



Single and Two-Level Anterior Cervical Discectomy and Fusion (ACDF) with Stand-Alone Cage-Plate: Single Centre Experience

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ABSTRACT

Introduction: After almost a decade of employing the concept of intervertebral disc removal complemented by vertebral fusion as a therapeutic technique for herniated cervical discs, Anterior Cervical Discectomy and Fusion (ACDF) became the standard gold treatment of a spectrum of cervical diseases such as compressive myelopathy/radiculopathy, disc herniation, and trauma. There is no discrepancy regarding the technique of ACDF in terms of using a standalone cage with an integrated spacer system or plating. This study reports 20 cases that underwent ACDF with standalone cage-plate and their radiological outcomes. **Methods:** A total of 20 patients who underwent single or two-level ACDF with standalone cage-plate for radiculopathy or myelopathy between May 2017 and February 2019 at a tertiary University Hospital have retrospectively reported in this study. The patient's demographics and radiological outcomes, including disc height, segmental lordotic angle, and global lordotic angle, were reported for each case pre-operatively, immediately postoperatively, and at 12-month follow-up. As per protocol, all patients were on Calcium and Vitamin D for nine months after the procedure. **Results:** All cases had an excellent fusion rate at 12 months of follow-up. None of the cases we reported has faced subsidence. According to Odom's criteria, the surgical outcome has been reported as excellent for all patients at 12 months of follow-up. **Conclusion:** ACDF with a standalone cage-plate system has shown excellent fusion, clinical and radiological outcomes with no subsidence and pseudoarthrosis in short term.

Keywords: Anterior Cervical Discectomy and Fusion (ACDF), Myelopathy, Radiculopathy, Standalone cage plate

Abbreviations: ACDF: Anterior Cervical Discectomy and Fusion, PEEK: Poly Ether Ether Ketone, IRB: Institutional Review Board, KSU: King Saud University, SPSS: Statistical Package for the Social Science, GLA: Global Lordotic Angle, SLA: Segmental Lordotic Angle, RLNP: Recurrent Laryngeal Nerve Palsy

INTRODUCTION

Adverse drug reaction can be defined as, "An appreciably harmful or unpleasant reaction, resulting from an intervention. After almost a decade of employing the concept of intervertebral disc removal complemented by vertebral fusion as a therapeutic technique for herniated cervical discs, Anterior Cervical Discectomy and Fusion (ACDF) became the gold standard treatment of a spectrum of cervical diseases such as compressive myelopathy/radiculopathy, disc herniation, and trauma. They were initially innovated in the 1950s by Smith and Robinson [1]. There has been a consensus in the literature regarding the technique of ACDF in terms of using a standalone cage with an integrated spacer system or plating [2].

There have been multiple types of cages and systems reported in the literature for the ACDF technique with its pros and cons. Those various systems vary from metal, plastic, and ceramic cages. Even though the lack of data supporting the choice of cage materials, plastic cages made from Polyetheretherketone (PEEK) have become a popular method for stabilizing the disc space after ACDF [3].

Recently there has been an increase in the usage of standalone Polyetheretherketone (PEEK) cages [4]. It is due to its low modulus of elasticity which closely resembles bone. Furthermore, they are radiolucent and help to observe the fusion across the instrumented level more accurately. Jun Cho, et al. reported no clinical outcome differences between

those operated with standalone PEEK cages and zero profile devices [3].

There are many conflicts in the literature regarding single and multilevel ACDF with plating and without plating. Although, most of the authors report no differences in the clinical outcomes between the two procedures [5,6]. While, some were concerned that ACDF with standalone cage was associated with subsidence, pseudoarthrosis, and misalignment specifically [5,7,8]. Other studies have shown that cage-assisted ACDF without plating was associated with a lower complication rate and shorter hospital stay [9,10].

Our study aims to report the effectiveness of ACDF with standalone cage-plate, their radiological and clinical outcomes in terms of improved disc height, global lordotic angle, and segmental lordotic angle.

MATERIAL AND METHODS

Study Duration

The study duration is two years and it is conducted from May 2017 till May 2019.

Study Method

It's a retrospective study.

Study Sample (N)

Twenty patients were included who fulfilled the inclusion criteria.

Study Settings

The study is being conducted at King Saud University, Orthopedic surgery department.

Inclusion Criteria

Our inclusion criteria were cervical disc degeneration along with radiculopathy or myelopathy or a combination of both, and all of them must have undergone ACDF with a standalone cage plate system.

Exclusion Criteria

We excluded all those operated on for other cervical diseases such as infection, trauma, tumor, or deformity.

Institutional Review Board (IRB) Approval

The Institutional Review Board (IRB) approved the study at King Khalid University Hospital, King Saud University, Saudi Arabia, with the IRB number KSU-IRB 017E.

We reviewed retrospectively thirty-nine patients who underwent cervical spine surgery during May 2017 and May 2019 and included them in the study. Twenty patients met the criteria for our research, which was single or double-level ACDF with standalone cage-plate.

All involved patients were treated and followed up at King Khalid University Hospital. The data concerning each patient included age, gender, smoking status, the indication of surgery, levels involved, intra-operative time, estimated blood loss, length of hospital stay, and radiological outcomes including disc height, global lordotic angle, and segmental lordotic angle on standing lateral X-ray pre-operatively, immediate postoperatively, and at 12 months follow up. As per protocol, all patients were on Calcium and Vitamin D for nine months after the procedure. The flexion and extension imaging was performed at the final follow-up.

Disc height was measured by the length between the midpoints of two adjacent vertebrae; global and segmental lordotic angle was measured by the Cobb method [11]. Before treatment, any patient who initially had kyphosis had recorded global and lordotic angles with negative (-). Screw loosening was marked as a radiolucent line surrounding the implant >1 mm in width known as a halo sign [12]. Fusion was applied as a standard definition by observing bridging bone between adjacent vertebral endplates of the involved segments [13-18].

Surgical Technique

All surgeries were performed under general anesthesia. All the surgeries were performed through a left-sided skin

incision in line with a skin crease at the desired level. After identification of the carotid pulsation, blunt dissection was performed to approach the prevertebral fascia. After the prevertebral fascia's penetration, the appropriate level was confirmed under the image intensifier, followed by subperiosteal elevation of longus colli muscles and standard discectomy. After preparing the endplates, the suitable size cage is selected using a trail cage, and then the final cage was inserted and secured. Before closure, hemostasis was secured. No drains were used for any patient. The wound was thoroughly irrigated with 0.9% normal saline. Wound closure was performed with vicryl and monocryl. Finally, the antiseptic dressing was applied. Postoperatively all the patients were given rigid cervical collars until six weeks. For all the patients, there were not any intraoperative complications. Postoperative cervical spine X-rays were done (Figure 1).

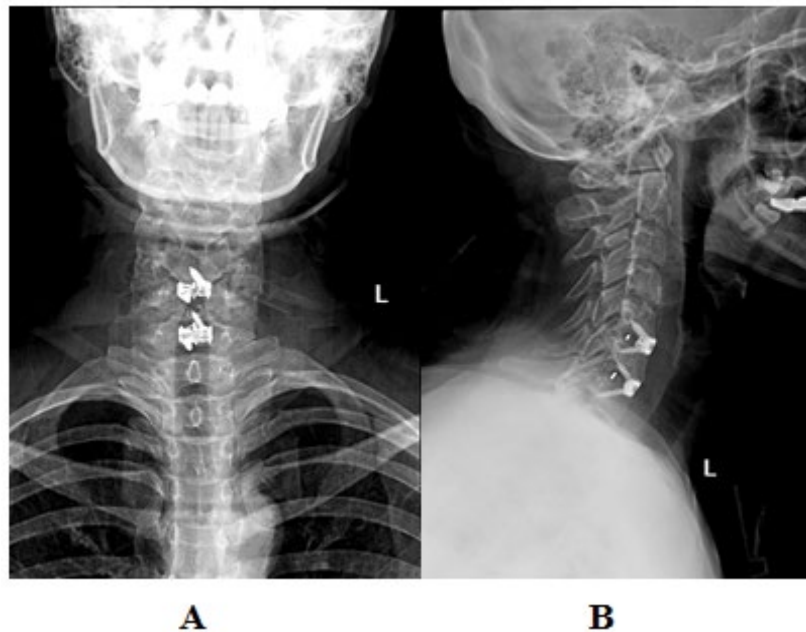


Figure 1 Immediate postoperative radiographs in AP and Lateral view in figures A and B respectively

All the patients were seen in our clinic at three months, and one year after the surgery. Cervical spine X-rays were done at three months (Figure 2) and one year (Figure 3) and calculations were made for disc height to record subsidence.

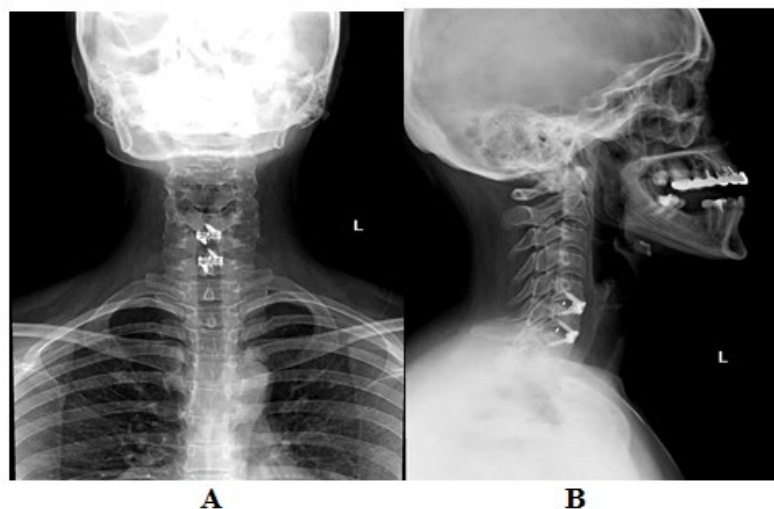


Figure 2 Radiographs taken at three months follow up in AP and Lateral view in figures A and B respectively

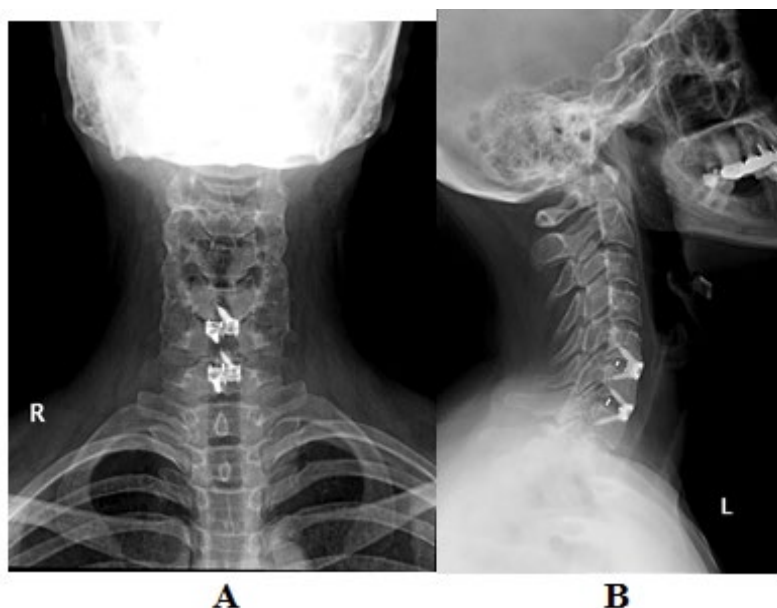


Figure 3 Radiographs taken at one year follow up in AP and Lateral view in figures A and B respectively

RESULTS

The analysis was conducted using Statistical Package for the Social Science (SPSS) software, version 23 (SPSS Inc., Chicago, Illinois, USA) is used for data entry and analysis.

Table 1 list all the patient's demographic characteristics, while Table 2 shows the indications, radiological outcomes at pre-op, immediate post-op, and 12 months post-op for each case.

Table 1 Characteristics of patients included in this study

No.	Age	Gender	Smoking Status	Segments	Intraoperative Time (Min)	Estimated Blood Loss (ml)	Length Of Hospital Stay (Days)
1	42	Female	Non-Smoker	C5/6, C6/7	165	>150	5
2	68	Male	Non-Smoker	C5/6, C6/7	209	8	1
3	42	Male	Smoker	C5/6	96	7	2
4	56	Male	Smoker	C5/6, C6/7	129	50	4
5	66	Male	Smoker	C4/5, C5/6	195	200	3
6	54	Female	Non-Smoker	C5/6	178	6	4
7	72	Male	Smoker	C4/5	146	100	4
8	56	Female	Non-Smoker	C5/6	107	50	3
9	71	Male	Smoker	C4-5	102	15	8
10	52	Female	Non-Smoker	C5-C6-C6-C7	159	15	4
11	34	Male	Non-Smoker	C5-6	77	5	2
12	40	Male	Non-Smoker	C5-6	74	7	5
13	39	Female	Non-Smoker	C5-6	72	5	2
14	59	Male	Non-Smoker	C3-4	74	6	3
15	54	Female	Non-Smoker	C5-6	76	7	2
16	29	Male	Non-Smoker	C5-6/C6-7	84	11	2
17	44	Female	Non-Smoker	C5-6/C6-7	82	9	2
18	43	Female	Non-Smoker	C5-6	73	8	2
19	53	Male	Non-Smoker	C5-6	78	14	3
20	58	Male	Non-Smoker	C5-6	69	6	2

Table 2 Indication, Radiological outcome pre-op, post-op, and 12 months post-op for each case

Sr. No.	Indication of Surgery	Preop Radiology/Clinical outcome			Postop Radiology/Clinical outcome			12 months postop Radiology/Clinical outcome		
		Disc height (mm)	GLA (1) (degrees)	SLA (2) (degrees)	Disc height (mm)	GLA (degrees)	SLA (degrees)	Disc height (mm)	GLA (degrees)	SLA (degrees)
1	Cervical stenosis+radiculopathy	C5-6: 4.33 C6-7: 4.62	7.1°	0.8°	C5-6: 6.77 C6-7: 7.02	14.6°	16.2°	C5-6: 6.93 C6-7: 7.02	20.8°	15.4°
2	Cervical stenosis and myelopathy	C5-6: 2.37 C6-7: 4.69	10.6°	6.4°	C5-6: 5.56 C6-7: 6.11	18.4°	11.4°	C5-6: 2.83 C6-7: 5.55	11.3°	5.2°
3	Degenerative disc+myelomalacia	C5-6: 5.59	31°	1°	C5-6: 6.98	25.5°	4.1°	C5-6: 5.48	37°	7.5°
4	Cervical spinal+Myelopathy	C5-6: 3.88 C6-7: 3.77	17.7°	1.6°	C5-6: 5.97 C6-7: 6.94	21.6°	8.9°	C5-6: 5.65 C6-7: 6.56	19.03°	8.4°
5	Cervical stenosis+radiculopathy+myelopathy	C4-5: 3.79 C5-6: 3.48	9.7°	3.2°	C4-5: 6.63 C5-6: 6.98	23.5°	16.7°	C4-5: 6.58 C5-6: 6.93	24.6°	12.9°
6	Cervical stenosis+radiculopathy	C5-6: 3.6	2.4°	7.9°	C5-6: 6.68	3.9°	12°	C5-6: 6.68	15°	12°
7	Cervical stenosis+radiculopathy	C4-5: 5.94	10.2°	3°	C4-5: 7.82	11.7°	12°	C4-5: 5.51	19°	9.3°
8	Cervical myelopathy	C5-6: 5.93	10.2°	7.5°	C5-6: 5.85	6.2°	3.9°	C5-6: 5.68	3°	5.7°
9	Cervical stenosis (brown-Sequard syndrome)+myelomalacia	C4-5: 7.9	20.9°	5.4°	C4-5: 7.58	16.5°	5.1°	C4-5: 7.58	25.8°	5°
10	Cervical stenosis+myelopathy	C5-6: 1.81 C6-7: 3.64	4.1°	0.6°	C5-6: 6.57 C6-7: 6.57	9.4°	9°	C5-6: 6.58 C6-7: 2.65	8.6°	2.1°
11	Cervical stenosis+myelomalacia (radiculopathy)	C5-6: 3.9	3.6°	1.6°	C5-6: 5.9	11.7°	8.5°	C5-6: 5.9	12.4°	8.5°
12	Radiculopathy+myelopathy	C5-6: 3.7	7°	0.5°	C5-6: 7	20°	14.1°	C5-6: 7	25°	14.1°
13	Cervical myelopathy	C4-5: 2.9	-1°	-2.1°	C4-5: 4.9	11°	9°	C4-5: 4.9	12.1°	9°
14	Cervical myelopathy	C3-4: 3.9	14.9°	-3.6°	C3-4: 6.8	15.4°	-1.4°	C3-4: 6.8	18.9°	-1.4°
15	Cervical radiculopathy	C5-6: 3.8	20.5°	0.9°	C5-6: 5.9	28.4°	11.1°	C5-6: 5.6	28.1°	11°
16	Cervical radiculopathy	C5-6: 3.7 C6-7: 4.5	25.1°	16°	C5-6: 5.9 C6-7: 6.9	12.8°	23°	C5-6: 5.9 C6-7: 6.9	26.2°	23.1°
17	Cervical radiculopathy	C5-6: 2.9 C6-7: 3.2	7.1°	4°	C5-6: 6.1 C6-7: 6.1	16.2°	19.9°	C5-6: 6.1 C6-7: 6.1	19.7°	19.9°

18	Cervical radiculopathy	C5-6: 1.4	5.6°	-1.5°	C5-6: 5.9	8.2°	7.5°	C5-6: 5.9	15.8°	7.8°
19	Cord compression+radiculopathy	C5-6: 2.9	17.3°	-2.6°	C5-6: 6.9	25.3°	10.3°	C5-6: 6.9	27.1°	10.3°
20	Cervical stenosis+radiculopathy	C5-6: 4.1	16.6°	2.1°	C5-6: 7.1	21.4°	11.5°	C5-6: 7.1	24.2°	11.3 °

Legend: 1-Global Lordotic Angle, 2-Segmental Lordotic Angle.

The median pre-operative disc height was 3.8 with an interquartile range of 1.7, and the 12-months postoperative final disc height was 6 with an interquartile range of 1.3. The median pre-operative global lordotic angle was 10.2 with an interquartile range of 12, and the 12-months postoperative final global lordotic angle was 20.25 with an interquartile range of 8.89. The median pre-operative segmental lordotic angle was 1.85 with an interquartile range of 4.9, and the median 12-month postoperative final segmental lordotic angle was 9.8 with an interquartile range of 6.15 (Table 3, Table 4, and Table 5).

Table 3 Median and Interquartile range of pre-operative, immediate post-operative, and 12-months post-operative disc height

	Pre-operative disc height	Immediate post-op disc height	12 months post-op disc height
Median	3.8	6.6	6
IQR	1.7	1	1.3

Table 4 Median and Interquartile range of pre-operative, immediate post-operative, and 12-months post-operative global lordotic angle

	Pre-op Global Lordotic Angle	Post-op Global Lordotic Angle	12-months post-op Global Lordotic Angle
Median	10.2	6.35	20.25
IQR	12	10.95	11.55

Table 5 Median and Interquartile range of pre-operative, immediate post-operative, and 12-months post-operative segmental lordotic angle

	Pre-op Segmental Lordotic Angle	Post-op Segmental Lordotic Angle	12-months post-op Segmental Lordotic Angle
Median	1.85	11.25	9.8
IQR	4.9	6.37	6.15

We want to report further that none of our patients developed hoarseness of voice after surgery. Although that wasn't the variable under research, that was an additional finding.

DISCUSSION

Even though ACDF has been associated with high osseous fusion rates, pseudarthrosis is still a significant complication. Given that there is a lack of standard criteria for determining fusion, pseudarthrosis rates after ACDF vary widely [19]. Nevertheless, has reported the percentage of pseudarthrosis to be 2.6%, with a reduction of pseudarthrosis rate when using autograft fusion compared to allograft fusion [19]. It has also been found that patients who underwent ACDF with a standalone cage had a higher pseudarthrosis rate than the patients who underwent the same procedure in addition to plate fixation [20].

However, in Hwang SL, et al. study patients who underwent titanium cage assisted-ACDF had long-term stabilization, increased lordosis, increased segmental height, and increased foraminal size in comparison to patients who underwent

the same procedure in addition to plate fixation [9]. Moreover, the cage-only group patients had a lower complication rate and shorter hospital stays.

Furthermore, fusion rates were reported variably as well. In a study, both cage and plate use resulted in a higher fusion rate [20]. On the other hand, Kaiser MG, et al. reported fusion rates for one- and two-level ACDF with anterior fixation was 96% and 91%, respectively, compared to 90% and 72% for one- and two-level ACDF without anterior fixation [21].

The unsettlement of the adverse effects and reduced biomechanical stability of interbody cages was reported in the literature [22]. Goel first described a modification by discussing bicortical and tricortical screws integrated with the cage in 1997 [23]. The technique's idea was to position the screws in an oblique manner to engross the body's anterior cortex, traverse the cortices, and adjoin the disc space to enhance stabilization [24].

We, therefore, reported 20 cases of which an ACDF with coalition integrated plate and spacer system was the surgeon's method of choice, and we observed radiological outcomes in terms of disc height, global lordotic angle, and segmental height at three different time intervals which are pre-operatively, immediately postoperatively, and at 12 months post-operation as seen in Table 2 previously. According to the subsidence definition found in the literature, which is a decrease in disc height more than 3 mm from the immediate postoperative image until the 12 months postoperative image, none of the cases we reported has faced subsidence [25]. Similarly, a study with the same method used said 80% of their subjects had some degree of implant subsidence, yet none of their patients had subsidence >2 mm or collapse of any segment [26].

We operated on all the patients through the left-sided incision; there is enough data to support the left-sided approach based on its benefits, especially avoidance of the injury to the recurrent laryngeal nerve. We want to report further that none of our patients develop a change in voice after surgery. Although that wasn't the variable under research, that was an additional finding. The Left-sided approach followers believe that the recurrent laryngeal nerves follow an inconsistent course in the lower neck and increase the risk of Recurrent Laryngeal Nerve Palsy (RLNP) [27]. On the other hand, proponents of the right-sided approach confirmed that the right-sided method is technically easier for right-handed surgeons. There is a high incidence of right-sided RLNP because most of the data are from left-sided surgeries [28,29]. Recently, Johnson, et al. showed in their work that there is no increased risk associated with either approach [30]. In another series by Justin M. Haller, and his colleagues reported that superior to C7-T1, both RLNs had similar anatomic courses and received equal protection *via* soft-tissue [31]. So, they supposed there was not a side-to-side difference. The selection of side for ACDF is based chiefly on the surgeon's preference unless there is a specific, compelling indication to advocate one approach over another. All of our patients reported complete resolution of symptoms and were satisfied with the surgical outcome at 12 months follow-up. According to Odom's criteria of surgical outcomes, all patients were graded excellent [32].

Limitation

Shortcomings of this study may include a small sample size. We suggest that the method of ACDF and its sequelae needs to be observed on larger sample sizes to give a more definitive answer on the pros and cons of ACDF with a standalone cage-plate system.

CONCLUSION

In conclusion, ACDF with a standalone cage-plate system has shown excellent fusion, clinical and radiological outcomes with no subsidence and pseudoarthrosis in short term.

DECLARATIONS

Conflicts of Interest

The authors declared no potential conflicts of interest concerning the research, authorship, and/or publication of this article.

Ethics Approval and Consent to Participate

Ethical approval was granted by our institutional review board following the National Committee of Bio-Ethics (NCBE) guidelines with the IRB number of KSU-IRB 017E.

Availability of Data and Materials

The datasets used and analyzed during the current study are available from the corresponding author on request.

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Authors' Contributions

Waleed Awwad reviewed the final version of the manuscript. Rohail Mumtaz contributed to the study design and manuscript preparation and performed the literature review and statistical analysis. Khalid Alasaleh, Abdulaziz Almaawi, and Waleed Albishi contributed to the study design and manuscript preparation. Sarah Aljasser and Mana Almuhaideb contributed to the study design and data collection. Waleed Awwad contributed to the manuscript preparation and data collection. Rohail Mumtaz contributed to the manuscript preparation and data collection. All authors read and approved the final manuscript.

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