

Posterior Instrumentation for Occipitocervical Fusion

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Abstract: Since 1995, 29 consecutive patients with craniocervical spine instability due to several pathologies were managed with posterior occipitocervical instrumentation and fusion. Laminectomy was additionally performed in nineteen patients. The patients were divided in two groups: Group A which included patients managed with screw-rod instrumentation, and Group B which included patients managed with hook-and-screw-rod instrumentation. The patients were evaluated clinically and radiographically using the following parameters: spine anatomy and reconstruction, sagittal profile, neurologic status, functional level, pain relief, complications and status of arthrodesis. The follow-up was performed immediately postoperatively and at 2, 6, 12 months after surgery, and thereafter once a year. Fusion was achieved in all but one patient. One case of infection was the only surgery related complication. Neurological improvement and considerable pain relief occurred in the majority of patients postoperatively. There were neither intraoperative complications nor surgery related deaths. However, the overall death rate was 37.5% in group A, and 7.7% in group B. There were no instrument related failures. The reduction level was acceptable and was maintained until the latest follow-up in all of the patients. No statistical difference between the outcomes of screw-rod and hook-and-screw-rod instrumentation was detected. Laminectomy did not influence the outcome in either group. Screw-rod and hook-and-screw-rod occipitocervical fusion instrumentations are both considered as safe and effective methods of treatment of craniocervical instability.

Keywords: Occipitocervical fusion, craniocervical instability, spine, reconstruction, posterior procedures.

INTRODUCTION

The occipitocervical joint consists of the occiput, atlas, and axis and contributes more than 50% of the rotation and flexion-extension of the head and neck [1]. A wide variety of abnormalities affect the occipitocervical junction causing instability of the occipitocervical junction, which finally leads to neural compression and/or craniospinal, or spinal instability [2-8]. Whatever the pathological process might be, it demands especially if it is symptomatic and progressive, immediate correction of subluxation or malalignment, decompression, and stabilization.

The majority of occipitocervical lesions are anterior and anterolateral. Posterior surgical decompression remains invaluable for the treatment of these lesions in select cases, as well as the posteriorly situated lesions. Reduction surgery and internal fixation are chosen when positioning or traction alone are adequate for the realigning of the column, but irreducible deformities need additional decompressive surgery using an anterior/transoral or posterior approach depending on the site of compression [7,9,10].

Occipitoatlantoaxial instability due to RA, trauma, neoplasms, dens dysplasias or os odontoideum, Down syndrome [3, 5-9, 11], and occasionally a prior failed attempt at C1-2 fusion or a complex cervical deformity [12], are the commonest indications for occipitocervical fusions.

The purpose of this study is to report the use of occipitocervical fusion instrumentations in the craniocervical joint of 29 patients suffering from several diseases in order to investigate the versatility and the usefulness of the systems. Specifically, the operated patients were examined for neurologic status improvement, fusion rate, pain relief, and complications after using the fixation techniques.

MATERIALS AND METHODOLOGY

Twenty-nine consecutive patients with acute and chronic spinal lesions (Table 1) located in the region from C1 to C5 were managed by using either a hook-and-screw-rod system [posterior cervical Compact Cotrel Dubousset system (CCD) (Medtronic Sofamor Danek, USA)] or a screw-rod system [PCR Summit (DePuy Orthopaedics, Inc, USA) or the similar Vertex (Medtronic Sofamor Danek, USA)] for occipitocervical fusion in two affiliated academic spine units between 1995 and 2003, using the same surgical technique and evaluation protocol approved by the scientific committees with which the principal investigators are affiliated. The

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Table 1. Patients Profile Pre- and Postoperatively**GROUP A
Subgroup A1**

No	Age and Sex	Pathology	Operation	Pain		Complications		Frankel		F-Up	Fusion
				Pre-Op	Post-Op	Surgery Related	Not Surgery Related	Pre-Op	Post-Op		
1	52 (F)	Rheumatoid Arthritis	CCD (O-C5) + C1-C2 Fusion Wire (C1-C2) + Laminectomy	7	0	—	—	C	E	40m	Yes
2	60 (M)	Tumor	CCD (O-C4) + Laminectomy	8	4		Meningitis (5 th post-op day) Death 3 rd month	D	E	3m	No
3	75 (M)	Ankylosing Spondylitis	CCD (O-C5) + Laminectomy	10	2	—	Death 7 th month	D	E	12m	Yes
4	67 (M)	Rheumatoid Arthritis	CCD (O-A5)+ C1-C2 Fusion Wire (C1-C2) + Laminectomy	9	0	—	—	D	E	30m	Yes
5	74 (F)	Tumor	CCD (O-C5) + Laminectomy	9	0	—	Death 28 th month	D	E	28m	Yes
6	68 (M)	Dens Pseudarthrosis	CCD (O-C7) + Laminectomy	8	0	—	—	D	D	9 y	Yes
7	80 (F)	Tumor	CCD (O-C7) + Laminectomy	9	0	—	Death 13 th month	E	E	12m	Yes
8	58 (F)	Rheumatoid Arthritis	CCD (O-C4) + Laminectomy	8	1	—	—	E	E	36m	Yes
9	63 (F)	Rheumatoid Arthritis	CCD (O-C5) + Laminectomy	8	1	—	—	E	E	5y	Yes
10	61 (F)	Rheumatoid Arthritis	CCD (O-C7) + Laminectomy	9	2	—	—	C	D	5 y	Yes
11	70 (M)	Tumor	CCD (O-C7) + Laminectomy	8	0	—	Death 28 th month	E	E	28m	Yes

Subgroup A2

No	Age and Sex	Pathology	Operation	Pain		Complications		Frankel		F-Up	Fusion
				Pre-Op	Post-Op	Surgery Related	Not Surgery Related	Pre-Op	Post-Op		
1	40 (F)	Trauma	CCD (O-C4)	9	0	—	—	E	E	36m	Yes
2	70 (M)	Ankylosing Spondylitis	CCD (O-C5)	10	2	—	—	D	E	38m	Yes
3	52 (F)	Rheumatoid Arthritis	CCD (O-C6)	6	1	—	—	E	E	36m	Yes
4	71 (F)	Cervical Spondylosis	CCD (O-C4)	10	1	—	—	D	E	28m	Yes
5	57 (M)	Tumor	CCD (O-C6) + Cage + ORION plate + Anterior procedure	9	8	Posterior Fusion Infection	Death 18 th month	D	D	18m	No

(Table 1) contd.....

GROUP B
Subgroup B1

No	Age and Sex	Pathology	Operation	Pain		Complications		Frankel		F-Up	Fusion
				Pre-Op	Post-Op	Surgery Related	Not Surgery Related	Pre-Op	Post-Op		
1	77 (M)	Trauma	PCR Summit (O-C4) + C1-C2 Fusion Wire (C1-C2) + Laminectomy	9	1	—	—	E	E	36m	Yes
2	69 (F)	Rheumatoid Arthritis	VERTEX (O-C4) + Laminectomy	9	0	—	—	D	E	30m	Yes
3	75 (F)	Tumor	VERTEX (O-C5) + Laminectomy	9	0	—	—	E	E	24m	Yes
4	70 (F)	Rheumatoid Arthritis	VERTEX (O-C7) + C1-C2 Fusion Wire (C1-C2) + Laminectomy	7	1	—	—	D	D	18m	Yes
5	64 (F)	Dens Pseudarthrosis	VERTEX (O-C4) + Laminectomy	8	0	—	—	E	E	15m	Yes
6	65 (M)	Trauma	VERTEX (O-C6) + Laminectomy	8	1	—	—	A	A	12m	Yes
7	78 (M)	Tumor	VERTEX (O-C6) + Laminectomy	9	0	—	—	E	E	9m	Yes
8	75 (M)	Tumor	VERTEX (O-C6) + Laminectomy + Anterior procedure	9	0	—	—	D	E	6m	Yes

Subgroup B2

No	Age and Sex	Pathology	Operation	Pain		Complications		Frankel		F-Up	Fusion
				Pre-Op	Post-Op	Surgery Related	Not Surgery Related	Pre-Op	Post-Op		
1	59 (M)	Trauma	PCR Summit (O-C4)	8	0	—	—	E	E	36m	Yes
2	80 (F)	Tumor	PCR Summit (O-C7)	8	0	—	—	C	D	36m	Yes
3	30 (M)	Tumor	PCR Summit (O-C5)	7	0	—	—	E	E	24m	Yes
4	70 (M)	A1-A2 Infection	PCR Summit (O-C5) + Anterior Procedure	10	0	—	—	B	D	6m	Yes
5	66 (F)	Tumor	VERTEX (O-T2) + VARIGRIP + Kyphoplasty	9	0	—	Death 12 th month	D	E	12m	Yes

choice of instrumentation either screw-rod or screw-and-hook-rod was randomized. The occipitocervical fusion was combined with laminectomy in nineteen cases, with Brooks-Gallie technique of fusion in two cases, and an anterior procedure in three cases. Grafts (Allomatrix) were applied in most of the cases. Bone cement was applied in some cases with very poor bone stock, especially in tumors and severe rheumatoid arthritis. The occipital screws were placed at the centre of the posterior inferior occipital bone which is thick and mainly consists of dense cortical bone. Computer assisted navigation was not used in our series. Postoperatively the stabilization was in every case (so that the study would be unbiased as to the postoperative use of orthoses) secured with a Halo-jacket for at least 3 months and a Philadelphia collar for an additional 3-month period in order to augment fusion which was to our opinion especially necessary in cases of tumor or very poor bone stock.

The indications for surgery were acute or chronic instability, presence of neurologic deficit, and resistant pain. The pathology of each case is shown in Table 1. There were 14 men and 15 women, and their average age was 65 years (range, 30-80 years). Eleven patients suffered from chronic cervical spine diseases (rheumatoid arthritis, ankylosing spondylitis, cervical degeneration) (Fig. 1), 11 patients of tumor of the cervical spine (Fig. 2), 3 of fresh traumatic lesion of the cervical spine, and two of dens pseudarthrosis. In all of the patients, fusion involved the occipitocervical junction. The patients were clinically and radiologically re-evaluated immediately postoperatively and also at 2, 6 and 12 months after surgery, and thereafter once a year. Clinical evaluation consisted of the neurological and functional status, and also pain improvement. The neurological and functional statuses were estimated using the Frankel classification; patients preoperatively classified as A (whose status was unlikely to improve) or B-C-D (whose status could either deteriorate or improve or remain unchanged) or E (whose status could either deteriorate or remain unchanged) were studied separately. Pain was subjectively estimated by the following method: the patient chose a number between 0 and 10 that best fitted to the pain status considering 0 as no pain at all, 1 as the mildest possible pain, and 10 as the worst possible pain. Radiological evaluation for the diagnosis of the underlying cause and the follow-up comprised plain and flexion/extension roentgenograms and CT scans; since 1999 MRI was also performed routinely in every patient for the diagnosis and the follow-up.

Patients were divided into two groups: Group A which included patients treated with CCD system, and Group B which included patients treated with PCR or Vertex systems. Groups A and B were each subdivided into two subgroups: A1/A2 and B1/B2 respectively; groups A1 and B1 included patients who had undergone laminectomy, and groups A2 and B2 included patients who had not undergone laminectomy (Table 1). The data collection of the two groups was completed at 12 months follow up and it was statistically evaluated.

Student "t-test" was used to evaluate any statistical difference as to the postoperative improvement/deterioration of pain and neurological status, whereas χ^2 with Yates correction was used to estimate any statistical difference as to the reoperation rate and the fusion rate. Statistical significance was set at $p < 0.05$.

Surgical Procedure

In all patients with dislocation, reduction was achieved by halo traction. After intubation, patients were placed prone on a frame and a horseshoe-type head holder or halo traction device. Correct cervical alignment was established under fluoroscopy. The head was taped to the head holder, and the shoulders were pulled caudal by a heavy bandage. During these maneuvers the cervical spine was strictly maintained in a neutral position. With a median skin incision, the paravertebral muscles were dissected laterally to expose the occipitocervical junction, the laminae and the articular masses in the upper cervical vertebrae. The skull was subperiosteally prepared up to the occipital notch. Then the occipitocervical instrumentation was applied. In all patients potentials were checked prior to anaesthesia, after anaesthesia and prior to the rolling of the patient, and after the rolling of the patient. Intraoperative monitoring was performed routinely since 2000; before 2000, a wake-up test was performed. Unless impaired neurologically or by their general condition the patients were encouraged to ambulate or sit up on the bed the day after surgery.

RESULTS

Group A: There was no intraoperative complication. Infection in one patient was the only surgery related postoperative complication (infection rate 6.1%). Among the sixteen patients, six died during the follow-up (death rate 37.5%). None of the deaths was surgery related: four patients died due to a neoplasm, one patient with ankylosing spondylitis died to a heart attack 7 months postoperatively, and, finally, one death happened at the early postoperative period (<3 months) due to a brain stroke that complicated meningitis related to the neurosurgical operation that was performed prior to the orthopaedic operation (intracanal tumor resection) that was performed by a neurosurgeon just prior to the orthopaedic stabilization performed by an orthopaedic surgeon.

Among the fifteen patients that survived more than 3 months a solid fusion was achieved in all but one patient (fusion was not achieved in this patient because an infection occurred) (fusion rate: 93.75%). The pain, on a scale of 0-10, improved in all patients treated with CCD (including the one in which an infection occurred) by an average of 7.4 grades (range, 1-9), but excluding the patient with infection and the patient who survived less than 3 months the pain improved by an average of 7.9 grades (range, 5-9). The neurological status improved in the vast majority of patients with a neurological deficit preoperatively. There was no patient preoperatively classified as Frankel A or B. Eight out of ten preoperatively Frankel C-D patients improved

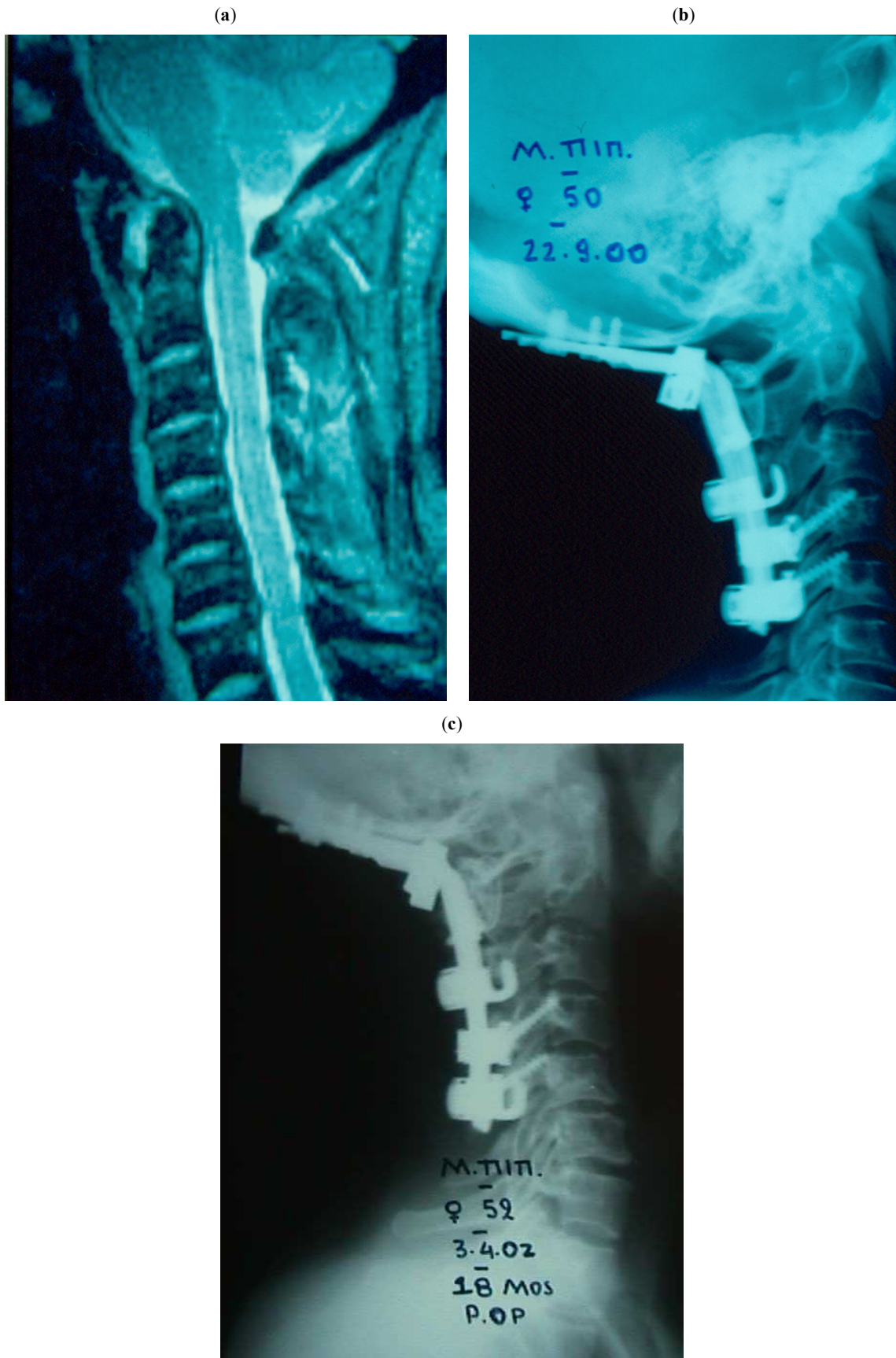


Fig. (1). (a-c) A patient with rheumatoid arthritis (case 4): after the application of a posterior CCD system, the sagittal profile of the cervical spine and the C1-C2 stability was restored and maintained during an 18 months follow-up. (a) MRI scan: C1-2 instability, vasilar invagination, and cord compression are obvious, (b) 2nd post-op day, (c) 18th post-op month.



Fig. (2). (a-c) A patient with a metastasis in the cervical spine. (a): MRI showing cord compression. (b): CT scan showing severe osteolysis resulting in C1-2 instability. (c): Laminectomy and application of PCR Summit instrumentation for occipitocervical fusion.

Table 2. Postoperative Reduction of the Sagittal Profile

Instrumentation	Parameters	Preoperatively Mean (Range)	Postoperatively Mean (Range)	Follow-Up Mean (Range)
Screw-rod system (Group A)	Occiput-C7 lordosis (degrees)	-31.1 (-13 to -40)	-40.7 (-37 to -46)	-40.3 (-34 to -45)
	Vertebral translation (mm)	1.4 (0 to 2.5)	0.5 (0 to 1.5)	0.5 (0 to 1.5)
Hook-and-screw-rod system (Group B)	Occiput-C7 lordosis (degrees)	-31.4 (-11 to -43)	-41 (-36 to -45)	-40.6 (-34 to -44)
	Vertebral translation (mm)	1.3 (0 to 2.5)	0.4 (0 to 2)	0.4 (0 to 2)

postoperatively by at least one Frankel grade and two patients remained at the same Frankel level (improvement rate 80%). The overall improvement of the neurological Frankel status among the preoperatively Frankel C-D patients averaged 0.9 grades. The neurological status did not deteriorate in the six patients with Frankel E level preoperatively.

There was neither hardware failure nor screw loosening until the latest follow-up. The sagittal profile of the occipitocervical junction as shown on the lateral radiographs was reduced after the operation and remained so until the final follow-up in all of the cases (Table 2).

Group B: There was no surgery related (intraoperative or postoperative) complication. The infection rate was 0%. Among the thirteen patients, one died during the follow-up (death rate: 7.7%); this death was related to a neoplasm. A solid fusion was achieved in all patients (fusion rate: 100%). The pain, on a scale of 0-10, improved in all patients by an average of 8.1 grades (range, 6-10).

The neurological status improved in the vast majority of patients with a neurological deficit preoperatively. One patient, who preoperatively was classified as Frankel A, did not improve postoperatively. Five out of six preoperatively Frankel B-D patients improved postoperatively by at least one Frankel grade, and one patient remained at the same Frankel level (neurological improvement rate 83.5%). The overall improvement of the neurological Frankel status among the preoperatively Frankel B-D patients averaged 1 grade. The neurological status did not deteriorate in the six patients with Frankel E level preoperatively.

There was neither hardware failure nor screw loosening until the latest follow-up. The sagittal profile of the occipitocervical junction as shown on the lateral radiographs

was reduced after the operation and remained so until the final follow-up in all of the cases (Table 2).

The comparison between the results of the application of hook-and-screw-rod (Group A) and screw-rod (Group B) instrumentation in our study did not reveal any statistically significant difference as to surgery related death rate, infection rate, pain improvement, fusion rate, neurological status improvement, and reduction of sagittal profile.

Laminectomy was not found to influence the outcome of the application of either the hook-and-screw-rod instrumentation (CCD) or the screw-rod instrumentation (PCR or Vertex); no statistically significant difference was detected between groups A1 and A2 or between B1 and B2 or between all patients with laminectomy (A1 and B1 subgroups) and all patients without laminectomy (A2 and B2 subgroups) (Tables 3 and 4) as to surgery related death rate, infection rate, pain improvement, fusion rate, neurological status improvement, and reduction of sagittal profile.

DISCUSSION

The cervical spine is unique because of its limited biomechanical strength, the paucity of some osseous structural elements, the neural and vascular structures contained within the spine, the frequent variation in anatomy among patients, and the complexity of the occipitocervical joint. Thus, any instrumentation used in this region should reach particular standards such as mechanical strength in multiple axes (flexion, extension, bilateral lateral bending, bilateral rotation, distraction, and axial loading), high fineness and appropriate dimensions so as to integrate into the cervical spine elements, and great flexibility to allow adjustment to several anatomical variations of this region [13].

Table 3. Results of Subgroups A1, A2, B1, B2, A1+B1, A2+B2. Subgroups A1 and B1 Represent Patients of Groups A and B with Laminectomy. Subgroups A2 and B2 Concern Patients of Groups A and B without Laminectomy

Subgroups	Pain Improvement Post-Op (Mean Degrees)	Neurological Status Frankel Classification (Mean) -Improvement rate: (%) / Grades of Level Improvement	Fusion Rate	Infection Rate
A1	7.6	85.7% / 1	100%	0%
A2	6.4	67% / 0.67	80%	20%
B1	8.22	67% / 0.67	100%	0%
B2	8.4	100% / 1.34	100%	0%
A1+B1	7.79	80% / 0.9	100%	0%
A1+B2	7.4	83.3% / 1	90%	10%

Table 4a-c. Comparative Analysis of the Postoperative Reduction of the Sagittal Profile Between Patients with and without Laminectomy**a) Comparison Between A1 and A2 Subgroups**

Instrumentation	Parameters	Preoperatively Mean (Range)	Postoperatively Mean (Range)	Follow-Up Mean (Range)
Subgroup A1	Occiput-C7 lordosis (degrees)	-30.7 (-13 to -37)	-40.3 (-37 to -44)	-40.4 (-35 to -45)
	Vertebral translation (mm)	1.2 (0 to 2.5)	0.5 (0 to 1.5)	0.5 (0 to 1.5)
Subgroup A2	Occiput-C7 lordosis (degrees)	-32 (-16 to -40)	-41.4 (-38 to -46)	-40 (-34 to -44)
	Vertebral translation (mm)	1.5 (0 to 2.5)	0.5 (0 to 1.5)	0.5 (0 to 1.5)

b) Comparison Between B1 and B2 Subgroups

Instrumentation	Parameters	Preoperatively Mean (Range)	Postoperatively Mean (Range)	Follow-Up Mean (Range)
Subgroup B1	Occiput-C7 lordosis (degrees)	-31.8 (-13 to -43)	-41.5 (-38 to -45)	-41 (-36 to -44)
	Vertebral translation (mm)	1.4 (0 to 2.5)	0.4 (0 to 2)	0.4 (0 to 2)
Subgroup B2	Occiput-C7 lordosis (degrees)	-30.8 (-11 to -40)	-40 (-36 to -43)	-39.5 (-34 to -43)
	Vertebral translation (mm)	1.1 (0 to 2.5)	0.5 (0 to 2)	0.5 (0 to 2)

c) Comparison Between Patients with Laminectomy (Subgroups A1 and B1) and Patients without Laminectomy (Subgroups A2 and B2)

Instrumentation	Parameters	Preoperatively Mean (Range)	Postoperatively Mean (Range)	Follow-Up Mean (Range)
Subgroups A1 and B1	Occiput-C7 lordosis (degrees)	-31.2 (-13 to -43)	-40.8 (-37 to -45)	-40.6 (-35 to -45)
	Vertebral translation (mm)	1.3 (0 to 2.5)	0.5 (0 to)	0.5 (0 to 2)
Subgroups A2 and B2	Occiput-C7 lordosis (degrees)	-31.4 (-11 to -40)	-40.8 (-36 to -46)	-39.8 (-34 to -44)
	Vertebral translation (mm)	1.4 (0 to 2.5)	0.5 (0 to 2)	0.5 (0 to 2)

The first preliminary report on successful use of a screw-rod stabilizing system posteriorly applied to the occipitocervical junction was in 1996 by Jeanneret *et al.* [14].

In the present study, the clinical utility, safety, and effectiveness of hook-and-screw-rod and screw-and-rod instrumentation for posterior craniocervical fixation was demonstrated. The best technique for occipitocervical fusion in case of an unstable craniocervical junction (CCJ) remains controversial since the desired rigid immediate stabilization demands increasing surgical complexity and risk [15].

Current techniques of posterior occipitocervical fusion include:

Wire fixation-strut bone graft technique: A 98% fusion rate has been achieved with this technique, despite several biomechanical disadvantages reported [9, 7, 11, 16].

Onlay bone graft technique [17]: Despite its safety [18], the lack of immediate rigid postoperative fixation necessitates prolonged postoperative immobilization.

C1-C2 transarticular screw fixation [19, 20, 21]: A significant range of movement of the cervical spine is maintained when this construct is used, and this is considered to be another advantage in comparison his of to screw-rod or hook-rod occipitocervical fusion systems, since the latter systems lead to considerable cervical spine stiffness which remains even after the hardware

removal. However, this method demands high level of expertise and does not lack potential danger of vertebral artery injury and screw malposition [21-23].

Occipitocervical plates and screws: It has been reported to provide very good results [20, 22, 24, 25], even though it is associated with a considerable rate of screw malposition and the danger of vertebral artery injury [25-28].

Contoured rods are another alternative described with a high fusion rate [12, 29, 30-33]. Screw-rod systems appear to offer several advantages compared to screw-plate systems [34]. A study of screw-rod posterior system applied in 30 patients reported good results in all but one patient, and one neurologic deterioration [35]. The use of hook-and-screw-rod system for the treatment of occipitocervical instability is reported to provide stable bony fusion, very low surgery-related morbidity or mortality (0-3%) and no hardware failure with a 100% fusion rate [36-39]. In many cases of this series where CCD was applied, a multiple-level fusion was necessary even if the abnormality included one level, because CCD as a hook-and-screw-rod system requires the use of the “claw” configuration at the two lowermost levels of fixation, and the avoidance of levels with significant canal stenosis or prior laminectomy is desirable or even demanded. This need for longer instrumentation is the main drawback of CCD when compared with lateral mass screw and pedicular systems [25, 35].

The present study showed that posterior occipitocervical fusion (either with hook-and-screw-rod systems or screw-rod systems) applied in the craniovertebral spine for several abnormalities provided adequate reduction maintaining until the latest observation, and immediate and rigid stabilization of the spine even in cases of instability due to a primary neoplasm or metastasis. Indeed, the heterogeneity in the underlying cause of occipitocervical instability in the patients of our study might make the interpretation of the outcome of occipitocervical fusion difficult. However, our study demonstrated a very good outcome of posterior occipitocervical fusion using either screw-rod or hook-and-screw-rod instrumentation irrespective of the underlying cause; this indicates the versatility of the method which is very important especially in cases of neoplasms or severe degenerative disease.

There was no significant difference between the results of the application of the hook-and-screw-rod and screw-rod system in our series. The fusion rates of both groups (A and B) of our study are comparable to those of the literature. Relief of acute and chronic pain occurred in most of the patients of either group contributing to a considerable improvement to their quality of life. Neurologic improvement and good to excellent functional results in the vast majority of the operated patients of both groups combined with the clinical improvement of the pain confirms the successful use of the two methods.

Although, the death rate in Group A was high, this cannot be attributed to the technique or any other surgery related reason. Four of these deaths were related to the underlying disease which was a malignancy that directly caused the unwanted outcome despite the adequacy of the surgical technique. However, even in these patients surgery proved to be effective as it permitted a considerable pain relief and, therefore, better living conditions.

This finding combined with the very low rate of surgery related complications of this study also confirms the safety of screw-rod and hook-and-screw-rod instrumentation for occipitocervical fusion. Additionally, laminectomy did not influence the outcome of either surgical technique; this might indicate that the stability of occipitocervical fusion is not jeopardized by laminectomy.

In conclusion, occipitocervical fusion with hook-and-screw-rod or screw-rod instrumentation provided good clinical and radiological results in the vast majority of abnormalities of the occipitocervical junction without neurologic, vessel or other major complication. Even though a cohort with mixed diagnoses and constructs was evaluated in our series, it seems that these systems alone or in combination with anterior instrumentation may provide a rigid immediate fixation with very high fusion rate. It is a safe and effective alternative in the management of occipitocervical instability.

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Received: March 01, 2011

Revised: April 07, 2011

Accepted: April 11, 2011

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