

AWAKE CRANIOTOMY USING DEXMEDETOMIDINE INFUSION AND SCALP BLOCK: OUR EXPERIENCE IN SERIES OF CASES

MSSV Prasad P¹, Allu Padmaja², B. Krishna Chaitanya³

¹2nd Year Post Graduate, Department of Anaesthesiology, Andhra Pradesh, India.

²Associate Professor, Department of Anaesthesiology, Andhra Pradesh, India.

³Senior Resident, Department of Anaesthesiology, Andhra Pradesh, India.

ABSTRACT: BACKGROUND: Awake craniotomy for removal of intracranial tumors is most challenging procedure. The critical aspect of awake craniotomy is to maintain adequate analgesia and sedation, hemodynamic stability, airway safety, while keeping the patient immobile for duration of surgery, cooperative for neurological testing.

AIM OF THE STUDY: Dexmedetomidine is good analgesic, sedative and has anaesthetic-sparing properties without causing significant respiratory depression.^[1] We are reporting cases series of awake craniotomy under monitored anesthesia care using dexmedetomidine infusion as an adjuvant to scalp block, titrating the sedation level by BIS monitoring.

MATERIALS AND METHODS: after careful patient selection and psychological preparation Monitored Anesthesia care(MAC) was provided by continuous infusion of Dexmedetomidine at a rate of 0.2-0.5 mcg/kg/min titrating sedation level to a BIS value of 70-90%. Bilateral scalp block was administered using 0.5% bupivacaine. For dura mater incision, a pad with 2% lidocaine was applied for 3 minutes. The tumor removal was complete with no neurological deficiency. All the patients were discharged on 5th postoperative day without complications and with full patient satisfaction.

CONCLUSION: We conclude that monitored anesthesia care with dexmedetomidine infusion and scalp block for awake craniotomy is a safe and efficacious. Absence of complications and high patient satisfaction score makes this technique close to an ideal technique for awake craniotomy.

KEYWORDS: Awake craniotomy; Dexmedetomidine; Scalp block; BIS index.

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INTRODUCTION: Awake craniotomy for epilepsy surgery has been routine for many years. It is now also used for the resection of tumours located in the eloquent cortex.^[2,3] The challenges for anesthesiologists are performing sedation-analgesia and assuring cardiorespiratory stability without interfering with electrophysiologic monitoring and cognitive tests. Selection of drugs with rapid onset of action, easy titration, with minimal effects on cardiovascular and respiratory systems, which do not cause nausea or vomiting or interfere with the neurological assessment, is crucial. Moreover, careful selection of patients, high levels of motivation, and psychological and emotional preparation are key elements for a successful procedure.^[4]

Bispectral analysis is increasing in popularity for the monitoring of conscious level during anesthesia and sedation. The bispectral index correlates with hypnotic component of anesthesia and was used as a guide to the administration of dexmedetomidine infusion in our cases.^[5]

The aim of this cases report was to present a sedation technique with dexmedetomidine infusion combined with bupivacaine for scalp block for intracranial tumor resection under awake craniotomy. This technique allowed the main surgical steps without the occurrence of major complications, such as psychomotor agitation, hemodynamic changes, and over sedation, without airway manipulation and, mainly, it did not affect the patients' cognitive evaluation.

MATERIAL AND METHODS: After obtaining institutional ethics board approval and written patient consent five patients of intracranial tumors were enrolled were awake craniotomy. The patients' details were shown in the table. The routine blood investigations were normal; the patients did not have any other systemic illnesses

All members of the team were briefed in advance so that a calm atmosphere prevails in the operating room. On arrival into operation theatre two 18G IV cannulae were inserted and pantocid 40mg as an acid prophylaxis, midazolam 1mg for anxiolysis, ondansetron 4mg antiemetic, fentanyl 100mcg for analgesia and levetiracetam 500mg for seizure prophylaxis were administered intravenously. Oxygen 3L/min was administered through nasal prongs.

The patient was continuously monitored for ECG, oxygen saturation, noninvasive blood pressure, end tidal CO₂ and level of sedation was monitored by bispectral index). The bladder was catheterised because of prolonged

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Corresponding Author:

MSSV Prasad P, # 49-44-6/32, Flat No. 501,

Laxmi Residency, Near Chinnur Maszid,

Akkipalem, Vizag.

E-mail: jsp_4a@yahoo.co.in

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duration of procedure and as there is need for mannitol infusion.

Before placing the patient into operative position bolus dose of dexmedetomidine infusion 1mcg/kg administered over 20minutes, followed by continuous infusion at a rate of 0.2-0.5mcg/minute (titrated BIS index to 70-90). The patient was placed in supine position with head slightly rotated operative side facing up. Care was taken for adequate visibility of the patient to anaesthesiologist during intraoperative testing. Fentanyl 20mcg IV was administered intermittently to provide additional analgesia and patient comfort.

Supraorbital, supratrochlear, zygomaticofacial, auriculotemporal, greater auricular, greater and lesser occipital nerves were blocked using 20 ml of 0.5% bupivacaine. We have limited total dose to 3mg/kg body weight. In Addition 1% lignocaine was infiltrated at the three head holder pin sites and along skin incision line. Adrenaline (5ug/ mL) was added to minimizes acute rises in plasma anaesthetic concentration and maximise the duration of the block.^[5] 20gms mannitol was infused about 15 min prior to completion of craniotomy. A pad soaked in 1% lignocaine was applied over dura to provide analgesia during dural opening. Once the dura was opened, brain was found to be relaxed and pulsating. For localization of primary motor cortex, our patients were asked to "move your toes/fingers, squeeze the ball" and they are well educated and trained about these tasks in preoperative visits. The procedures have lasted for about 3-4 hr. At the end of the surgery, all the patients except one were fully awake and communicating. One patient was converted to general anesthesia due uncontrolled seizures. All the patients were shifted to the intensive care unit for further observation. The patients' haemodynamic parameters were stable throughout the procedure. The post-operative course was uneventful and discharged after 5 days.

DISCUSSION: There is an increase in number of indications in intracranial surgery for the patient to be awake during some or all of the operation. To achieve a goal of increased lesion removal, improved survival benefit.^[1] shorter hospitalization time, reduced cost and a decreased incidence of postoperative complications, a sound anatomical knowledge of the nerve blocks and the knowledge to predict intraoperative events is extremely rewarding for the neuroanaesthetist.^[6,7]

One of the most important considerations for awake craniotomy is careful patient selection. The patient should cooperate in an unfamiliar and stressful environment for an extended period of time. Infact, the only absolute contraindication for awake technique is an uncooperative patient. A Our anesthesia team in preoperative visit detailed about expected discomforts and level of co-operation expected, potential and tasks that will be performed for motor testing.

There is considerable variation in the anesthetic management of the awake craniotomy in different institutions. Propofol sedation, commonly in combination

with a shorter acting opioid such as fentanyl, or remifentanyl, is an effective and popular technique during awake craniotomy, achieving a high degree of patient satisfaction and acceptance. a blinded, prospective, randomized study while comparing the efficacy of dexmedetomidine versus propofol-remifentanyl based sedation in patients undergoing awake craniotomy for resection of tumors, hypothesised that the efficacy of performing intra-operative brain mapping is identical between dexmedetomidine and the propofol-remifentanyl based sedation.^[8]

In our patients, combination of the dexmedetomidine infusion and the scalp block with bupivacaine provided effective sedation, analgesia and allowed full excision of the tumor, without damage or deficits for the patient who had early mobilization and was discharged uneventfully.^[9]

Dexmedetomidine (highly specific α_2 -adrenoreceptor agonist) has potential application during awake craniotomy for tumor resection because they provide analgesia and sedation that is easily reversed with verbal stimulation. There is no risk of respiratory depression.^[9] The primary action is excitation inhibition in the central nervous system. It is a cerebral vasoconstrictor by stimulating α_2b receptors in the cerebral blood vessels with no effect on CMRO₂. Dexmedetomidine inhibits the cerebral vasodilatation induced by hypercapnia, thus avoiding increased intracranial pressure and brain bulging. Dexmedetomidine also has anticonvulsant effects that might be helpful during epilepsy surgery or tumor resection. The analgesic effect of dexmedetomidine consistently reduces opioid administration.^[10] In a case report by Basavaraj G Kallapur and Raghavendra Bhosale, by administration of dexmedetomidine 1–0.5 mcg/kg/h and propofol 60 mg/h, achieved adequate sedation, analgesia and fully co-operative patient to perform cognitive tests successfully once the infusions were stopped 10 min prior.^[10]

In our case series the level of sedation during Dexmedetomidine infusion was monitored by BIS index, maintained BIS score between 70-90 by increasing level of sedating during positioning and head pin fixation and reducing it during testing for functional area intraoperatively.

Patient tolerance of an awake craniotomy relies on effective analgesia of the surgical field, and cannot rely on sedation or anaesthesia alone. In our case scalp block combined with local infiltration provided adequate analgesia, haemodynamic stability and decreased the stress response to painful stimuli.

Complications of awake craniotomy include seizures, cerebral edema, nausea and vomiting, decreased level of consciousness, neurological deficit, pain, and loss of patient cooperation.^[1,11] Airway management is uneventful during awake craniotomy under sedation in our reported cases. However, over sedation inevitably runs the risk of apnoea and airway obstruction. Airway obstruction (incidence 0% to 20%) with oxygen desaturation (0-28%) may result in brain swelling because of elevated levels of Paco₂.^[12]

Equipment for emergency airway control should be available throughout awake craniotomy.

Seizures may occur unexpectedly (0% to 24%) due to decreased levels of anticonvulsants or local anesthetic toxicity. Most of the seizures can be resolved by irrigation of the surgical field with cold saline or administration of propofol.^[13] An antiepileptic prophylaxis may be helpful to prevent intraoperative seizures. However, one of reported cases developed uncontrolled seizures in spite of prophylactic measures.

During the procedure precautions were taken to prevent shivering by using warm infusions and blanket.^[1] Dexmedetomidine can cause hypotension and bradycardia. However the patients in our case series were hemodynamically stable.

CONCLUSIONS: In our case series, infusion of dexmedetomidine 0.2-0.5 mcg/kg/h and scalp block, has achieved adequate sedation and analgesia during awake craniotomy. By monitoring depth of anesthesia with BIS, we could achieve adequate sedation without airway compromise. Our patients were fully co-operative for cognitive testing. We conclude that dexmedetomidine as an adjunct to scalp block is useful during awake craniotomy for tumour resection.

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Case serial No	Age (yrs)	Sex	Weight (kg)	Presenting symptom	Diagnosis	Duration of surgery (hours)
1	45	Male	72	Seizures, right upper limb weakness	Right frontal glioma	3
2	31	Male	65	Partial seizures	Left premotor and motor strip glioma	3.5
3	22	Female	55	Headache, seizures	Left parietal opecular glioma	4
4	33	Female	67	Seizures of four months duration	Left parietal glioblastoma multiform	3.0
5	26	Female	60	seizures	Right frontal glioma	Converted to general anesthesia due to uncontrolled seizures
Table 1: Patient details of case series						



Case 1: BIS Monitoring (Spacelab workstation with inbuilt BIS module)



Case 2: Positioning for awake craniotomy