

Drug Induced Sleep Endoscopy for Obstructive Sleep Apnea Syndrome: An Initial Experience in a Tertiary Level Centre

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Submitted : November 5, 2019

Accepted : December 7, 2019

ABSTRACT

Drug induced sleep endoscopy (DISE) is a technique of performing endoscopy of the upper airway after inducing sleep by the use of anaesthetic agents in patients suffering from obstructive sleep apnea syndrome (OSAS). The main purpose of DISE is to detect the obstruction or collapse at one or more location in the upper airway, both for the successful diagnosis and management of OSAS. Here we discuss a case of OSAS who underwent DISE under novel anaesthetic agents midazolam and dexmedetomidine.

Keywords: *Dexmedetomidine, drug induced sleep endoscopy, obstructive sleep apnea syndrome, propofol*

INTRODUCTION

Obstructive sleep apnea syndrome (OSAS) is a condition in which upper airway collapse or gets obstructed during sleep time. The condition can be due to enlarged nasal turbinates, enlarged tonsils, soft tissue enlargement at pharynx or larynx, large tongue size, or collapse of airway during sleep at single or multiple levels.¹ The treatment modality includes weight reduction, exercise of pharyngeal muscles, application of Continuous positive airway pressure (CPAP) or the surgery of the airway at one or multiple settings.

CPAP is the gold standard therapy for OSAS² but the main problem is compliance with CPAP therapy. Surgical approach is good alternative but the outcome is variable. The outcome can be improved by the upper airway endoscopic study done at the sleep time as it represents the dynamic pathophysiology.³ Thus drug induced sleep endoscope (DISE) is performed by the use of the novel anaesthetic agents under monitoring in operating room. The common agents used for DISE are midazolam, propofol or dexmedetomidine.⁴

CASE PRESENTATION

Twenty six year old male (weight 75kg, BMI 24 kg/m²), otherwise healthy, presented with history of snoring, mouth breathing and few apneic episodes during sleep for one year. All his symptoms were gradually increasing. Upper airway examination was grossly normal. Apnea hypopnea index (AHI) was 72 and lowest oxygen saturation was 75% in polysomnography.

After adequate fasting on the day of DISE, intravenous cannula was secured, and Ringer's lactate was used as maintenance fluid. ECG, non-invasive blood pressure (NIBP) and pulse oximeter were monitored. Nasal cavity was anaesthetized using 10% lignocaine spray. To facilitate sedation, lights were dimmed, beeping of monitors was silenced and alarms were turned off, and noise was minimized by educating the staffs prior to the procedure and displaying the warning signs like "DISE, Keep silence" on the doors. A comfortable pillow was kept under the head before sedation. Baseline saturation was 98%, NIBP 140/70 mmHg and heart rate 80/minute. After confirmation of

backup required for anaesthetic management and adequate preparation for DISE, sedation was initiated with IV dexmedetomidine infusion at 1mcg/kg for 10 minutes and midazolam 1 mg IV concurrently. After 5 minutes of infusion, there was onset of snoring. A classical hypopnea and apnea cycle with fall in saturation upto 75% was seen with no change in ECG, heart rate and blood pressure. Nasal endoscope was inserted through right nostril and the dynamic airway anatomy during the period was recorded. Total duration was 8 minutes. After conclusion, dexmedetomidine infusion was stopped. The patient became conscious after 10 minutes. The patient was discharged home after the discharge criteria were met.

DISCUSSION

In OSAS there is dynamic obstruction at one or multiple levels during the sleep time. Polysomnography or Overnight Sleep study helps to diagnose and grade OSAS. OSAS is graded according to AHI (Mild 5-15, moderate 15-30 and severe >30).⁵ Our patient had severe OSAS with AHI of 72 and lowest oxygen saturation level of 75%.

Patient with OSAS produces loud snoring during sleep and has frequent arousals leading to cognitive dysfunction and day time somnolence.⁶ There can be difficult airway and other multi systemic problems including cardio-respiratory problems, which can be progressive over time.⁷ Our patient had day time somnolence. He snored loudly and had few apneic episodes, which were progressive.

Nasoendoscopic evaluation of upper airway of such patient in the awake state would give anatomical idea of the obstruction however true picture cannot be ensured. Thus sleep endoscopy would be the investigation of choice to figure out the dynamic picture of airway.⁸ These patients easily fall asleep any time of the day if comfortable environment is provided. However the duration of sleep and tolerance of nasal endoscopy is questionable. In our case, we had made our operation room dark, minimized the noise as much as possible including the beep of monitors and had offered a comfortable pillow for the patient.

Natural sleep of the patient can be simulated by using anesthetic drugs and drug induced sleep endoscopy can be done to evaluate the patients airway. An ideal anaesthetic agent for DISE should have rapid onset of action, short duration of action, has amnesic property, can easily be titrated to sedation level and has low residual effect for early discharge.⁵ Anaesthetic agents like midazolam,

propofol or dexmedetomidine have the above properties so can be used alone or in combination. Such patients are sensitive with the opioids and inhalational agents and hence not recommended for DISE.

Propofol is the most common inducing agent for DISE however it can cause pharyngeal collapse and central apnea, thus increasing the incidence of hypoxia.⁹ Besides, it can also cause haemodynamic instability at the dose. Midazolam has the property of amnesia, rapid onset of action and minimal or low residual sedative effect and can be used for the sedation of such patient however sole midazolam can lead to respiratory obstruction.¹⁰ Dexmedetomidine, a centrally acting alpha-2 agonist, has sedative, amnesic, analgesic, sympatholytics and minimal respiratory depressive property. The sleep induced by this drug resembles to that of the natural one.¹¹ Its use has been validated in many day care surgeries as anaesthetic adjunct.

We performed the case with midazolam and dexmedetomidine. Patient went to deep sleep after 5 minutes of infusion. There were no features of respiratory depression as minimum saturation was 75% and respiratory rate was 12-16 per minute. His blood pressure and heart rate were normal throughout.

There are no severe adverse effects associated with DISE in literature. However in DISE, sleep apnea cycle (SAC) is induced before endoscopy in OSA patients. Airway obstruction may be severe than expected and significant desaturation may occur necessitating advanced airway management. It is hence performed in operation theatre with backup preparation and difficult airway gadgets.⁵ Nasal bleeding due to trauma by endoscope, laryngospasm, pulmonary aspiration, cardiac dysrhythmia, and hypertension may also occur during the procedure.¹² No adverse events occurred in our patient.

The patient underwent septoplasty with bilateral inferior turbinate reduction with tonsillectomy with modified uvulopalatopharyngoplasty for the definite management on a later date.

CONCLUSION

A case of OSAS was evaluated with DISE using dexmedetomidine and midazolam smoothly.

ACKNOWLEDGEMENTS

We would like to acknowledge the contribution of the whole surgery team, anaesthesia team, nursing staff and operation theatre staff involved in the procedure.

CONFLICT OF INTEREST

None declared.

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