Do OSA patients under deep sedation keep you awake at night?



Undergoing a deep sedation procedure places OSA patients at risk for upper airway obstruction. This increases the chance of hypoxemia and other pulmonary complications.

ASA guidelines include CPAP as a recommended therapy to minimize the risks associated with OSA patients. The guidelines suggest that supplemental oxygen should be given to all patients who are at an increased perioperative risk of OSA². They also state that CPAP should be administered preoperatively when OSA is severe or during sedation for patients who have been previously treated with CPAP².

SuperNO₂VA[™] nasal PAP ventilation system

The SuperNO₂VA nasal PAP ventilation system uses the flow from any oxygen source. It delivers both oxygen and nasal positive pressure ventilation to patients, which can maintain a patent upper airway.

This complete system allows for portability and continued use from preoperative oxygenation through postoperative care without any additional equipment.

Please contact your local representative.

vyaire.com/SuperNO2VA





REFERENCES

1. Franklin K. et al. Obstructive sleep apnea is a common disorder in the population—a review on the epidemiology of sleep apnea. J Thorac Dis 2015;7(8):1311-1322

2 Practice Guidelines for the Perioperative Management of Patients with Obstructive Sleep Apnea. Anesthesiology 2006; 104:1081–93.

GLOBAL HEADQUARTERS

Vyaire Medical, Inc. 26125 North Riverwoods Blvd Mettawa, IL 60045 USA CareFusion 22745 Savi Ranch Parkway Yorba Linda CA 92887, USA



ECREP Emergo Europe Prinsessegracht 20 2514 AP The Hague The Netherlands Vyaire Medical Pty Ltd Level 5, 7 Eden Park Drive Macquarie Park, NSW, 2113 Australia

vyaire.com/SuperNO2VA

For Africa, Australia, Canada, Europe and New Zealand distribution only.

Trademarks are the property of their respective owners. © 2019 Vyaire Medical, Inc. or one of its affiliates. All rights reserved. Vyaire, the Vyaire Medical logo and SuperNO2VA are trademarks or registered trademarks of Vyaire Medical, Inc or one of its affiliates. Medical devices Class IIa according to Medical Devices Directive 93/42/EEC. Please read the complete Instructions For Use that come with the devices or follow the instructions on the product labelling. VYR-INTL-1800029

SuperNO₂VA™ NASAL PAP VENTILATION SYSTEM