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Nurse-administered propofol sedation for gastrointestinal endoscopic procedures: first Nordic results from implementation of a structured training program

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Pages 1503-1509 | Received 03 Jun 2011, Accepted 25 Aug 2011, Published online: 04 Nov 2011

 Cite this article  <https://doi.org/10.3109/00365521.2011.619274>

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Abstract

Introduction. Proper training to improve safety of NAPS (nurse-administered propofol sedation) is essential. **Objective.** To communicate our experience with a training program of NAPS. **Materials and methods.** In 2007, a training program was introduced for endoscopists and endoscopy nurses in collaboration with the Department of Anaesthesiology. During a 2.5-year period, eight nurses were trained. Propofol was given as monotherapy. The training program for nurses consisted of a 6-week course including theoretical and practical training whereas the training program for endoscopists consisted of 2.5 h of theory. Patients were selected based on strict criteria including patients in ASA (American Society of Anesthesiologists) group I-III. **Results.**

2527 patients undergoing 2.656 gastrointestinal endoscopic procedures were included. The patients were ASA group I, II and III in 34.7%, 56% and 9,3%, respectively. Median dose of propofol was 300 mg. No mortality was noted. 119 of 2527 patients developed short lasting hypoxia (4.7%); 61 (2.4%) needed suction; 22 (0.9%) required bag-mask ventilation and 8 (0.3%) procedures had to be discontinued. In 11 patients (0.4%), anesthetic assistance was called due to short lasting desaturation. 34 patients (1.3%) experienced a change in blood pressure greater than 30%. Conclusion. NAPS provided by properly trained nurses according to the present protocol is safe and only associated with a minor risk (short lasting hypoxia 4.7%). National or international structured training programs are at present few or non-existing. The present training program has documented its value and is suggested as the basis for the current development of guidelines.

Key Words::

[endoscopy-general](#)

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Declaration of interest: The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

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