The addition of capnography to standard monitoring reduces hypoxemic events during gastrointestinal endoscopic sedation: a systematic review and meta-analysis

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Background: The use of capnography monitoring devices has been shown to lower the rates of hypoxemia via early detection of respiratory depression, and facilitate more accurate titration of sedatives during procedures. The aim of the current meta-analysis was to compare the incidence of hypoxemia associated with standard monitoring alone during gastrointestinal endoscopy to that associated with standard monitoring with the addition of capnography.

Methods: The MEDLINE, EMBASE, and Cochrane Central Register of Controlled Trials scientific databases were searched to identify relevant studies. We performed a meta-analysis of randomized controlled trials undertaken up to January 2018 that met our predefined inclusion criteria. The study outcome measures were incidence of hypoxemia, severe hypoxemia, apnea, the use of assisted ventilation, the use of supplemental oxygen, and change in vital signs.

Results: We included nine trials assessing a total of 3,088 patients who underwent gastrointestinal procedural sedation. Meta-analysis of study outcome revealed that capnography significantly reduced the incidence of hypoxemia (odds ratio 0.61, 95% CI 0.49–0.77) and severe hypoxemia (odds ratio 0.53, 95% CI 0.35–0.81). However, there were no significant differences in other outcomes including incidence of apnea, assisted ventilation, supplemental oxygen, and changes in vital signs. Early procedure termination and patient satisfaction-related outcomes did not differ significantly in the capnography group and the standard monitoring group.

Conclusion: This study indicates that capnography monitoring is an important addition with regard to the detection of hypoxemia during gastrointestinal procedural sedation, and should be considered in routine monitoring during gastrointestinal endoscopy.

Keywords: endoscopy, additional monitoring, hypoxemia, apnea

Introduction

Sedation and analgesia are commonly used during gastrointestinal endoscopic procedures, and have been shown to increase the success rates of these procedures.1 However, the use of sedative agents can result in respiratory depression, airway obstruction, hypoxemia, and adverse cardiopulmonary events.2,3 Therefore, monitoring of vital signs including continuous pulse oximetry combined with a patient’s breathing pattern is advocated in many national guidelines on sedation for gastrointestinal endoscopy.4,5

Capnography measures the carbon dioxide (CO2) concentration throughout the respiratory cycle, which allows for real-time evaluation of a variety of respiratory factors including respiratory depression, apnea, and hypercapnia.6 Hypoventilation can...
occur in the presence of normal arterial oxygen saturation as determined via pulse oximetry (SpO₂), and inadequate ventilation can precede hypoxemia by several minutes.⁷,⁸ The use of a capnography monitoring device has been shown to lower the rates of hypoxemia via early detection of respiratory depression and facilitate more accurate titration of sedatives during procedures.⁹

The American Society of Anesthesiologists recommended that capnography monitoring should be considered for all patients receiving deep sedation.³ However, there are insufficient data to demonstrate that improved clinical outcomes or care quality derive from the use of capnography in patients undergoing targeted moderate sedation for upper endoscopy and colonoscopy.⁴,⁹,¹¹ Furthermore, it has been suggested that capnography may lead to unnecessary procedure interruption, delay, or termination, contributing to inefficiency.⁴ Thus, most guidelines for gastrointestinal sedation do not recommend routine use of capnography, but recommend it only in specific situations including high-risk patients, intended deep sedation, and long procedures such as endoscopic retrograde cholangiopancreatography (ERCP) and endoscopic ultrasonography (EUS).⁴,¹¹,¹²

The aim of the current meta-analysis was to determine whether the addition of capnography to standard monitoring including pulse oximetry reduces hypoxemic events during gastrointestinal endoscopic sedation.

Methods
In this systematic review, determination of the criteria for the selection of studies, extraction of relevant data, assessment of study quality, and statistical analyses were conducted and reported in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analyses statement.¹³

Literature search
We conducted a systematic literature search using MEDLINE, EMBASE, and the Cochrane Central Register of Controlled Trials updated from 1979 to January 2018. The following Medical Subject Heading terms were used: “capnography”, “hypoxia”, “monitoring”, “physiologic”, “endoscopy”, “endoscopic”, “gastrointestinal”, “gastroscopy”, “gastroscopes”, “colonoscopy”, and “colonoscopes”. All of these items were then explored to include secondary headings, and the reports considered were limited to those involving human subjects. Two investigators (SHK and YSC) independently reviewed all potentially relevant material to determine whether or not it met the inclusion criteria.

Inclusion and exclusion criteria and definitions
The inclusion criteria were as follows: 1) study population, patients undergoing gastrointestinal endoscopic sedation; 2) intervention, capnography monitoring; control, standard monitoring; 3) outcome measure, the incidence of oxygen desaturation; and 4) study design, prospective and randomized controlled trials (RCTs). Duplicate publications, review articles, case reports, studies without raw data available for retrieval, and studies only available as abstracts from conferences were excluded.

Evaluation criteria for endpoints
The primary endpoint of this study was the weighted summary OR of the incidence of hypoxemia (SpO₂ of <90% to <95%) with a 95% CI. The secondary endpoints were 1) incidence of severe hypoxemia defined as SpO₂ of <85%; 2) incidence of apnea; 3) early detection of apnea; 4) use of assisted ventilation; 5) use of supplemental oxygen; 6) bradycardia; 7) hypotension; 8) incidence of early procedure termination; 9) patient satisfaction; and 10) patient cooperation with regard to the endoscopic procedure.

Study selection and data extraction
After removing duplicate studies, two investigators (SHK and YSC) independently perused the titles and abstracts of all publications retrieved and assessed their eligibility for inclusion in the meta-analysis. If available, the full texts of selected studies were screened with reference to the inclusion and exclusion criteria. Selected full-text articles were critically appraised for relevance and validity. When any disagreement occurred, this was resolved by discussion, together with clinical expert consultation. The following information was extracted from the articles included: first author, year of publication, endoscopic procedure, monitoring intervention, patients’ baseline characteristics, sedative agents, depth of sedation, duration of procedures, and primary and secondary outcomes. To avoid bias in the data extraction process, two investigators (SHK and YSC) independently evaluated each study quality and compared results with one another. In case of disagreement, the third investigator (EK) made a determination decision.

Assessment of methodological quality
The Cochrane Collaboration’s risk of bias tool was used to assess the risk of bias in all included studies. The following items were assessed and recorded: random sequence generation (selection bias), allocation concealment (selection bias),
blinding of participants and personnel (performance bias),
blinding of outcome assessment (detection bias), incom-
plete outcome data (attrition bias), and selective reporting
(reporting bias).

**GRADE assessment and summary of findings**

We used the Grading of Recommendations Assessment,
Development, and Evaluation (GRADE) approach to assess
the quality of the evidence associated with specific outcomes
(hypoxemia, severe hypoxemia, incidence of apnea, and
detection of apnea) in our review and to construct a “sum-
mmary of findings” table. The GRADE approach considers
study risk of bias (methodological quality), directness of
the evidence, heterogeneity of the data, precision of effect
estimates, and risk of publication bias.

**Statistical analysis**

All analyses were performed using Meta V.4.9–0 and
Meta for V.2.0–0 in R packages (R language version 3.4.3;
We calculated the OR as a summary statistic with a 95% CI
for dichotomous outcomes and the standardized mean differ-
ce with a 95% CI for continuous outcomes. Heterogeneity
among the studies was measured using the $I^2$ statistic and the
$P$-value was derived from the Cochran’s $Q$ test. Significant
between-study heterogeneity was deemed to be present when
there was an $I^2$ value of more than 50% or a Cochran’s $Q$ test
$P$-value of <0.1. $^{14}$ The presence of publication bias was
evaluated via Egger’s test with a funnel plot.$^{15}$

**Results**

**Identification of studies**

Nine studies derived from eight RCTs$^{6,7,9,16-20}$ in which a
total of 3,088 patients who underwent gastrointestinal pro-
cedural sedation were enrolled were ultimately included in
the final analysis (Figure 1). Overall, 1,542 patients were
assigned to the capnography group, and the remaining 1,546
were assigned to the standard monitoring group.

**Study characteristics and patient populations**

Of the eight studies, four were performed in Germany, three
were performed in the USA, and one was performed in
Denmark. Endoscopic procedures included upper endoscopy,
colonoscopy, EUS, ERCP, and other interventional pro-
cedures. One study, published by Mehta et al,$^{6}$ provided sepa-
rate data on colonoscopy and esophagogastroduodenoscopy.

<table>
<thead>
<tr>
<th>Reports identified from literature search (n=547)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duplication (n=362)</td>
</tr>
<tr>
<td>Articles screened on basis of title and abstract (n=185)</td>
</tr>
<tr>
<td>Title and abstract excluded by inclusion and exclusion criteria (n=174)</td>
</tr>
<tr>
<td>Manuscript review and application of inclusion criteria (n=11)</td>
</tr>
<tr>
<td>Unrelated to study aims (n=3)</td>
</tr>
<tr>
<td>RCT included in this meta-analysis (n=8)</td>
</tr>
</tbody>
</table>

Five studies used propofol as a sedative agent$^{7,9,18-20}$ and
other studies used either fentanyl, midazolam, meperidine,
or ketamine, or combination of these.$^{6,16,17}$ The main charac-
teristics of the studies are presented in Table 1.

Apnea was defined as the absence of respiratory activity
denoted by a flat line on the capnograph in 7 studies,$^{6,7,9,16,17,19,20}$
and hypoventilation was defined as a respiratory rate of less
than or equal to 8/min in two studies.$^{6,19}$ In another two
studies,$^{6,17}$ greater than a 75% reduction in the amplitude
of the capnograph waveform compared with baseline
for 5–10 seconds was defined as abnormal or disordered
ventilation.

In five of the eight studies, the endoscopy team (physi-
cians and nurses) was not blinded to the assignment of
patients. Capnographic data were available for the capnogra-
phy group, but not for the standard monitoring group. In the
other three studies,$^{6,16,17}$ the endoscopy team was not aware of
the capnographic findings, and capnographic data were only
available to an independent observer. In these three studies,
interventions were initiated in both groups based on capnog-
raphy, albeit at different intervals (5–10 seconds for abnormal
ventilation in the capnography group and >30 seconds for
apnea in the standard monitoring group).

The studies included in this meta-analysis utilized four
types of capnography. The Capnostream 20 (Oridion Medical,
Needham, MA, USA) was used in four studies, and the other
four studies used Capnostream 20 (Covidien, Mansfield, MA,
USA), Philips M4 with Microstream CO$_2$ (Philips Medical
Table 1 The main characteristics of the studies included in the meta-analysis

<table>
<thead>
<tr>
<th>Study (publication year)</th>
<th>Number (cap, control)</th>
<th>Population and procedures</th>
<th>Sedatives, depth of sedation</th>
<th>Duration of procedures (cap vs control, minute)</th>
<th>Definition of hypoxemia</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lighthale et al (2006)</td>
<td>163 (83, 80)</td>
<td>ASA 1–2, children (0–19 years), endoscopy and colonoscopy</td>
<td>Fentanyl and midazolam, moderate</td>
<td>Endoscopy: 10 (0–24) vs 10 (4–25), colonoscopy: 39 (34–67) vs 40 (15–69)</td>
<td>SaO₂ &lt;95% for 5 seconds</td>
</tr>
<tr>
<td>Qadeer et al (2009)</td>
<td>247 (124, 123)</td>
<td>ASA 1–3, adults, ERCP or EUS</td>
<td>Midazolam in combination with meperidene or fentanyl, N/A</td>
<td>37.2 (16.1) vs 34.4 (12.5)</td>
<td>SaO₂ &lt;90% for 15 seconds</td>
</tr>
<tr>
<td>Beitz et al (2012)</td>
<td>757 (383, 374)</td>
<td>ASA 1–3, adults, colonoscopy</td>
<td>Propofol, adequate</td>
<td>35.9 (22.0) vs 33.8 (20.6)</td>
<td>Fall in SaO₂ ≥5% or &lt;90%</td>
</tr>
<tr>
<td>Slegelse et al (2013)</td>
<td>540 (263, 277)</td>
<td>ASA 1–3, adults, endoscopy or colonoscopy</td>
<td>Propofol, N/A</td>
<td>23.6 (12–30) vs 24 (13–31)</td>
<td>SaO₂ &lt;92%</td>
</tr>
<tr>
<td>Friedrich-Rust et al (2014)</td>
<td>533 (267, 266)</td>
<td>ASA 1–3, adults, colonoscopy alone or in combination with EGD</td>
<td>Propofol in combination with ketamine, N/A</td>
<td>38 (16) vs 38 (17)</td>
<td>SaO₂ &lt;90% for 15 seconds</td>
</tr>
<tr>
<td>Klare et al (2016)</td>
<td>238 (108, 115)</td>
<td>ASA 1–4, adults, ERCP</td>
<td>Propofol and midazolam, deep</td>
<td>38 (6–165) vs 38 (5–164)</td>
<td>SaO₂ &lt;90%</td>
</tr>
<tr>
<td>Mehta et al (2016)</td>
<td>209 (101, 108)</td>
<td>ASA 1–2, adults, endoscopy</td>
<td>Fentanyl or midazolam or meperidene, moderate</td>
<td>5.6 (2.6) vs 5.6 (2.6)</td>
<td>SaO₂ &lt;90% for 10 seconds</td>
</tr>
<tr>
<td>Mehta et al (2016)</td>
<td>231 (117, 114)</td>
<td>ASA 1–2, adults, colonoscopy</td>
<td>Fentanyl or midazolam or meperidene, moderate</td>
<td>17.3 (7.3) vs 17.4 (7.5)</td>
<td>SaO₂ &lt;90% for 10 seconds</td>
</tr>
<tr>
<td>Riphaus et al (2017)</td>
<td>170 (83, 87)</td>
<td>ASA 1–3, adults, interventional endoscopy (EUS or other interventional procedures)</td>
<td>Midazolam and propofol, adequate</td>
<td>25 vs 26</td>
<td>SaO₂ &lt;90%</td>
</tr>
</tbody>
</table>

Notes: Duration of procedures are presented as mean (SD) or median (interquartile range).
Abbreviations: cap, capnography; ASA, American Society of Anesthesiologists; ERCP, endoscopic retrograde cholangiopancreatography; EUS, endoscopic ultrasonography; EGD, esophagogastroduodenoscopy; N/A, not applicable; SaO₂, oxygen saturation.

**Systems**: Andover, MA, USA; Oridion), Microcap (Oridion), and Philips MP 20 monitor, respectively.

**Quality of the included studies**
The risks of performances and outcome assessment biases were substantial due to lack of blinding. In five studies, neither the patients nor the endoscopy team was blinded, and there were no independent observers for the outcome assessment. The overall quality of the studies was moderate to low using the GRADE approach. We have included the risk of bias and a summary of findings in Tables S1 and S2, respectively.

**Hypoxemia**
All studies reported hypoxemia, with the definitions varying from SpO₂ of <90% to <95%. The study by Lighthale et al was performed in pediatric patients, and the definition of hypoxemia was SpO₂ <95%, which was higher than that of the other studies. In another studies, SpO₂ of <90%–<92% was the criterion for hypoxemia. Publication bias was assessed by Egger’s test with a funnel tunnel plot (P=0.691).

The analysis revealed that capnography significantly reduced the incidence of hypoxemia (OR 0.61, 95% CI 0.49–0.77) (Figure 2). Six studies reported severe hypoxemia, defined as SpO₂ of <85%. The analysis showed that the addition of capnography to standard monitoring has benefit in reducing the incidence of severe hypoxemia (OR 0.53, 95% CI 0.35–0.81) (Figure 3).

**Apnea**
Apnea was assessed with capnography in four studies, and the incidence of apnea was comparable between the two groups (OR 0.74, 95% CI 0.39–1.43) (Figure 4). In these four studies, capnography was used to detect apnea in both groups. In three studies, the endoscopy team was unaware of the capnographic findings, and apnea was assessed by an independent observer. In the remaining study, the endoscopy team was not blinded to the capnographic findings, and the incidence of apnea was retrospectively analyzed in the standard monitoring group.

In another two studies, apnea was more frequently detected in the capnography group than in the control group (OR 44.76, CI 25.35–79.04) (Figure 5). In these two studies, capnography was only used in the capnography group and was not used to detect apnea in the standard monitoring group.
Capnography monitoring during gastrointestinal procedural sedation assisted ventilation and supplemental oxygen

Five studies reported on the use of assisted ventilation, and there was no evidence of heterogeneity ($I^2=0\%$). The fixed effect model demonstrated that the need to provide assisted ventilation was comparable between the two groups (OR 0.56, 95% CI 0.25–1.22). The need to provide supplemental oxygen was reported in five studies, and analysis indicated that the provision of supplemental oxygen was comparable in the two groups (OR 0.74, 95% CI 0.52–1.03).

Other side effects

Seven studies reported bradycardia, with definitions varying from heart rates of <50 beats/min to <60 beats/min. Hypotension (systolic blood pressure varying from <100 mmHg to <80 mmHg) was reported in eight studies. The incidences of bradycardia (OR 1.18, 95% CI 0.84–1.65) and hypotension (OR 0.92, 95% CI 0.68–1.25) did not differ significantly between the two groups. Four studies reported the incidence of procedure termination secondary to concerns for patient safety, and there were no significant differences in early procedure termination (OR 4.96, 95% CI 0.24–104.39). Three studies reported patient satisfaction outcome and patient cooperation. Neither patient satisfaction (standard mean difference (SMD) $-0.02$, 95% CI $-0.14$ to 0.10) nor patient cooperation (SMD 0.00, 95% CI $-0.12$ to 0.11) differed significantly between the two groups.

Discussion

Recently, the use of sedatives has improved patient comfort and their acceptance of gastrointestinal endoscopy. Intervventional endoscopic procedures under endoscopic sedation via a sedative agent such as propofol are now used widely. During endoscopic sedation, cardiorespiratory complications such as hypoxemia and apnea may occur, and require technical monitoring including pulse oximetry. However, this method only provides an indirect measure of respiratory function, and the detection of cardiorespiratory depression may potentially be delayed.

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Discussion

Recently, the use of sedatives has improved patient comfort and their acceptance of gastrointestinal endoscopy. Intervventional endoscopic procedures under endoscopic sedation via a sedative agent such as propofol are now used widely. During endoscopic sedation, cardiorespiratory complications such as hypoxemia and apnea may occur, and require technical monitoring including pulse oximetry. However, this method only provides an indirect measure of respiratory function, and the detection of cardiorespiratory depression may potentially be delayed.
Capnography facilitates continuous, real-time, non-invasive measurement of expiratory $\text{CO}_2$. Capnography monitoring entails the measurement of end-tidal $\text{CO}_2$ concentrations using an infrared sensor attached to a nasal cannula or face mask. This facilitates the early detection of arterial oxygen desaturation via near real-time graphic assessment of respiratory activity. Many studies of surgical operations and non-surgical procedures have shown that the use of capnography monitoring is associated with a reduction in the incidence of hypoxic events. Nevertheless, the use of capnography during gastrointestinal endoscopy is not an established recognized standard for monitoring respiratory function.

In a recent meta-analysis, there was a lack of convincing evidence that the addition of capnography to standard monitoring in emergency departments reduced the rate of adverse events including hypoxemia. However, most of the procedures included in that analysis were orthopedic manipulation, abscess incision and drainage, or cardioversion, which differ from elective procedures. In another meta-analysis including gynecologic procedures, abortion, and emergency department procedures, capnography reduced the rate of hypoxemia, which is consistent with the results of the current meta-analysis. Notably, there was no consensus on whether capnography could reduce the need for assisted ventilation in those two studies. Further, there was no previously published meta-analysis of the use of capnography during gastrointestinal endoscopic sedation.

In previous sedation guidelines for non-anesthesiologists, continuous monitoring via visual observation of breathing and its frequency was recommended, but capnography monitoring was not mandatory. However, recent standards advocated by the American Society of Anesthesiologists suggest that the adequacy of ventilation should be evaluated via continuous capnography monitoring.

The results of our meta-analysis show that capnography reduced the rate of hypoxemia during gastrointestinal sedation but was not significantly associated with incomplete endoscopic procedures or patient satisfaction. Many factors can influence respiratory depression, such as the degree and depth of sedation, the sedative agents used, the patient’s baseline disease status, age, and the duration of the procedure during endoscopy. The nine studies assessed in the current investigation evaluated hypoxemia, and the median procedure times of four studies were >30 minutes. The duration of the procedure is an important risk factor for oxygen desaturation. Therefore, it is expected that the role of capnography will increase in future in conjunction with the ongoing development of interventional endoscopy procedures with long procedure times, and the widespread use of sedatives associated with those procedures.

The results of the current investigation showed that capnography has a beneficial effect by reducing the incidence of hypoxemia and severe hypoxemia during gastrointestinal endoscopic sedation. Because the target level of sedation was equal in both the standard monitoring and capnography

<table>
<thead>
<tr>
<th>Study</th>
<th>Capnography events</th>
<th>Total</th>
<th>Control events</th>
<th>Total</th>
<th>Odds ratio</th>
<th>OR (95% CI)</th>
<th>Weight (fixed)</th>
<th>Weight (random)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Qadeer et al (2009) ¹</td>
<td>51</td>
<td>124</td>
<td>77</td>
<td>123</td>
<td>0.42</td>
<td>(0.25–0.70)</td>
<td>28.5%</td>
<td>25.6%</td>
</tr>
<tr>
<td>Mehta (EGD) et al (2016)ª</td>
<td>59</td>
<td>101</td>
<td>46</td>
<td>108</td>
<td>1.89</td>
<td>(1.09–3.28)</td>
<td>24.7%</td>
<td>25.0%</td>
</tr>
<tr>
<td>Mehta (CFS) et al (2016)ª</td>
<td>65</td>
<td>117</td>
<td>72</td>
<td>114</td>
<td>0.73</td>
<td>(0.43–1.24)</td>
<td>26.9%</td>
<td>25.4%</td>
</tr>
<tr>
<td>Riphaus et al (2017) ²</td>
<td>31</td>
<td>83</td>
<td>46</td>
<td>87</td>
<td>0.53</td>
<td>(0.29–0.98)</td>
<td>19.9%</td>
<td>24.0%</td>
</tr>
</tbody>
</table>

**Fixed effect model**

- 425
- 432
- 0.74 (0.56–0.97)
- 0.74 (0.39–1.43)

**Random effects model**

- Heterogeneity: $I^2$=83%, $t^2$=0.3691, $P$<0.01

**Figure 4** Forest plot showing the odds ratios and 95% CIs of each study for the incidence of apnea.

**Abbreviations:** EGD, esophagogastroduodenoscopy; CFS, colonoscopy.

**Figure 5** Forest plot showing the odds ratios and 95% CIs of each study for the detection of apnea.
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Capnography monitoring during gastrointestinal procedural sedation

groups; the incidence of apnea was not different between the two groups; however, in two studies evaluating the detection of apnea, there was no evidence of heterogeneity and the results showed that apnea was more often detected with the addition of capnography than it was in the standard monitoring group. Whereas most episodes of apnea might remain undetected using standard monitoring with visual assessment of the patient, the use of capnography can facilitate early interventions such as patient stimulation, withholding or reducing sedative medications, and using chin-lift or jaw-thrust maneuvers. Easier detection of apnea via capnography may explain the difference in the incidence of hypoxemia between the two groups, and these results demonstrate the value of adding capnography monitoring during sedation for gastrointestinal procedure.

The target level of sedation in the trials included in this study ranged from moderate to deep sedation. In moderate sedation, spontaneous ventilation is adequate and maintenance of airway patency does not require interventions. However, it is not always possible to predict how a patient will respond to specific sedative and analgesic medications. Thus, the level of sedation could become deeper than initially intended. In this meta-analysis, the addition of capnography monitoring was effective in reducing hypoxemia, and it did not increase the rate of early procedure termination. Therefore, we suggest capnography monitoring should be considered for routine monitoring during gastrointestinal endoscopic procedures performed under moderate sedation in the future.

The present meta-analysis had several limitations that should be taken into account when interpreting the results. It included studies that varied with respect to the sedative agents and corresponding doses used. This may account for some of the heterogeneity in the results of the analysis. Second, because our analysis was based on only eight RCTs, we did not perform subgroup analysis based on the type of endoscopic procedure. Thus, it cannot be decisively concluded that diagnostic and therapeutic endoscopy have similar safety with regard to sedation. We also analyzed both upper endoscopy through the oral cavity and colonoscopy together. Further studies with larger numbers of patients are warranted to clarify the beneficial effects of capnography during gastrointestinal endoscopic sedation.

Conclusion
In the current meta-analysis, capnography monitoring was associated with reduced incidence of hypoxemia during gastrointestinal procedural sedation, and there was no evidence of an association with procedural interruption. Capnography monitoring should be considered in routine monitoring in the near future.

Author contributions
All authors contributed toward data analysis, drafting and critically revising the paper and agree to be accountable for all aspects of the work.

Disclosure
The authors report no conflicts of interest in this work.

References


Supplementary materials

Table S1 Risk of bias summary

<table>
<thead>
<tr>
<th>Study (publication year)</th>
<th>Random sequence generation</th>
<th>Allocation concealment</th>
<th>Blinding of participants and personnel</th>
<th>Blinding of outcome assessment</th>
<th>Incomplete outcome data</th>
<th>Selective reporting</th>
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<tbody>
<tr>
<td>Beitz et al (2012)</td>
<td>Low</td>
<td>High</td>
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<td>Slagelse et al (2013)</td>
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<td>High</td>
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<td>Low</td>
<td>High</td>
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<td>Klare et al (2016)</td>
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<td>High</td>
<td>Low</td>
<td>Low</td>
</tr>
</tbody>
</table>

Table S2 GRADE approach

Date:

Question: Capnography monitoring compared to standard monitoring for gastrointestinal procedural sedation

Setting:

Bibliography:

Certainty assessment

<table>
<thead>
<tr>
<th>No of studies</th>
<th>Study design</th>
<th>Risk of bias</th>
<th>Inconsistency</th>
<th>Indirectness</th>
<th>Imprecision</th>
<th>Other considerations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypoxemia (assessed with: oxygen saturation)</td>
<td>9</td>
<td>Randomized trials</td>
<td>Serious</td>
<td>Not serious</td>
<td>Not serious</td>
<td>None</td>
</tr>
<tr>
<td>Severe hypoxemia (assessed with: oxygen saturation)</td>
<td>6</td>
<td>Randomized trials</td>
<td>Serious</td>
<td>Not serious</td>
<td>Not serious</td>
<td>None</td>
</tr>
<tr>
<td>Incidence of apnea (assessed with: capnography)</td>
<td>4</td>
<td>Randomized trials</td>
<td>Not serious</td>
<td>Not serious</td>
<td>Not serious</td>
<td>None</td>
</tr>
<tr>
<td>Detection of apnea (assessed with: capnography or visual inspection)</td>
<td>2</td>
<td>Randomized trials</td>
<td>Serious</td>
<td>Not serious</td>
<td>Not serious</td>
<td>Not serious</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>No of patients</th>
<th>Effect</th>
<th>Certainty</th>
<th>Importance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Capnography monitoring</td>
<td>standard monitoring</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Relative (95% CI)</td>
<td>Absolute (95% CI)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>463/1,529 (30.3%)</td>
<td>OR 0.61 (0.49–0.77)</td>
<td>Moderate</td>
<td>Important</td>
</tr>
<tr>
<td>622/1,544 (40.3%)</td>
<td>111 fewer per 1,000 (from 61 fewer to 154 fewer)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>91/1,100 (8.3%)</td>
<td>OR 0.53 (0.35–0.81)</td>
<td>Low</td>
<td>Important</td>
</tr>
<tr>
<td>158/1,100 (14.4%)</td>
<td>62 fewer per 1,000 (from 24 fewer to 88 fewer)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>206/425 (48.5%)</td>
<td>OR 0.74 (0.39–1.43)</td>
<td>Moderate</td>
<td>Important</td>
</tr>
<tr>
<td>241/432 (55.8%)</td>
<td>75 fewer per 1,000 (from 86 more to 228 fewer)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>283/491 (57.6%)</td>
<td>OR 44.76 (25.35–79.04)</td>
<td>Moderate</td>
<td>Important</td>
</tr>
<tr>
<td>14/489 (2.9%)</td>
<td>540 more per 1,000 (from 399 more to 671 more)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Notes: *In five studies, endoscopy team was not blinded to assignment of patients because of organizational reasons. *In three studies, endoscopy team was not blinded to assignment of patients. *In two studies, endoscopy team was not blinded to assignment of patients.