

Determining gastric contents during general anaesthesia: evaluation of two methods

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Two methods used to measure the volume of gastric contents were evaluated in 24 supine anaesthetized adults. Methods compared were: (1) aspiration of stomach contents through a large, vented, multi-orificed gastric tube, and (2) indirect determination by a dye dilution method using polyethylene glycol (PEG) as the marker. The volumes determined by these methods (V_{asp} and V_{peg} respectively) were compared to the total volume (V_{tot}) present in the stomach, determined by direct inspection of the gastric pouch by the surgeon at the beginning of surgery. The results show that the volume of aspirated gastric fluid, using this type of tube, is a very good estimate of the total volume of gastric residue. The PEG dilution method yields similar results. However, correlation between V_{peg} and V_{tot} was not as close-fitting as the correlation between V_{asp} and V_{tot} . PEG dilution is more complicated, time-consuming and offers no advantage over aspiration.

Key words

GASTRO-INTESTINAL TRACT: stomach, contents;
MEASUREMENT TECHNIQUES: polyethylene glycol.

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Presented in part at the Canadian Anaesthetists' Society annual meeting in Montreal, June 1986.

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This work was supported by a grant from the University of Montreal.

Accurate measurement of the volume of gastric contents is of importance to help identify patient populations at increased risk of regurgitation and subsequent aspiration during general anaesthesia. Despite numerous studies on the volume and pH of gastric fluid of different patient populations coming for elective surgery, few authors have attempted to validate their method of quantifying gastric volume. Two methods are readily available in clinical practice to measure this volume: (1) gastric intubation and presumed complete aspiration of fluid contained in the stomach, and (2) indirect determination by a dye dilution method.

Both methods have their drawbacks. Aspiration of fluid may be incomplete, especially with conventional Levin tubes.¹ Dye dilution has been thoroughly investigated in the gastroenterology laboratory.²⁻³ The method has been shown to be accurate when patients were kept sitting, when the position of the stomach tube used for dye injection and sampling was checked radiologically, and when the previously determined optimal concentrations of indicator dye were used. However, the method has never been validated in the operating room environment. Furthermore, the functional division of the stomach into antral and fundal sacs in supine patients⁴ may interfere with both aspiration and dilution. Thus, we believed that a re-evaluation of these experimental methods was necessary. This study was designed to prospectively validate both gastric fluid measurement techniques in the supine, anaesthetized, patient.

Methods

The study was approved by the Ethics Committee and informed consent was obtained for all patients. Twenty-four ASA physical status I-III adult patients undergoing elective laparotomy, excluding surgery of the oesophagus and stomach, were studied. The anaesthetics were standardized. Induction consisted of a defasciculating dose of d-tubocurarine (DTC $50 \mu\text{g}\cdot\text{kg}^{-1}$), fentanyl $6 \mu\text{g}\cdot\text{kg}^{-1}$, pre-oxygenation with 100 per cent oxygen by face mask for three minutes, followed by thiopentone $4 \text{mg}\cdot\text{kg}^{-1}$ and succinylcholine $1.5 \text{mg}\cdot\text{kg}^{-1}$. Mask ventilation was avoided to prevent gastric insufflation. The trachea was

intubated with a cuffed endotracheal tube. Anaesthesia was maintained with nitrous oxide 60 per cent and isoflurane as needed, in oxygen. Patients were paralysed with pancuronium IV.

After the skin incision, but before exploration of the abdominal cavity, an 18 Fr Salem Sump tube (Argyle, St Louis, Mo.) was passed orally into the stomach. Gastric contents were aspirated as completely as possible (aspirated volume: V_{asp}), using a 50-ml syringe. The surgeon was then asked to inspect the stomach and ascertain complete gastric emptying. This direct inspection could lead to aspiration of additional gastric fluid upon mobilisation of the gastric pouch (additional volume: V_{add}). The stomach was then returned to its original position in the abdominal cavity.

In 15 patients, the total volume ($V_{tot} = V_{asp} + V_{add}$) less a 2-ml sample used for later determination of gastric pH was re-injected into the stomach and then diluted with 100 ml of a 0.4 per cent polyethylene glycol (PEG) solution. PEG is an indicator currently used in gastroenterology to measure the rate of gastric emptying. Mixing of the dye with gastric fluid was achieved by withdrawing the gastric contents into a 50 ml syringe and then reinjecting the fluid. This was repeated five times. The mixture was then re-aspirated as completely as possible (V_{re-asp}). PEG concentrations were determined by turbidimetry, in two 3 ml samples. Knowledge of PEG concentration enabled calculation of the volume diluting the indicator (V_{PEG} ; Figure 1). The biochemist in charge of determining V_{PEG} was blinded to V_{tot} .

In the first nine patients studied, dilution of gastric fluid with only 30 ml of indicator solution led to completely erratic determinations of V_{PEG} , which were excluded (see Discussion). After consultation with a gastroenterologist, the method described above was used to determine V_{PEG} in the remaining 15 patients.

Statistical analysis was performed by the Department of Mathematics and Statistics. Student's *t* tests for paired data and linear regression analysis were used. A $p < 0.05$ was considered significant.

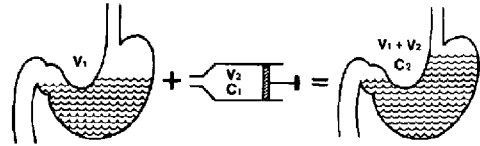
Results

Aspiration of gastric contents was performed in all 24 patients. The volume of fluid retrieved from the stomach on initial aspiration (V_{asp}) was 31.1 ± 28.8 ml (mean SD)

TABLE Summary of results (mean \pm SD)

	V_{asp} (ml)	V_{add} (ml)	V_{tot} (ml)	V_{PEG} (ml)
Initial 9 Patients	31 ± 32.2	4.7 ± 3.5	35.7 ± 32.6	*
Subsequent 15 Patients	31.2 ± 27.7	4.2 ± 4.3	35.4 ± 28.1	26.2 ± 28.8
Total (n = 24)	31.1 ± 28.8	4.4 ± 3.9	35.5 ± 29.1	*

*First nine determinations of V_{PEG} were excluded (see text).



$$V_1 = \frac{V_2 (C_1 - C_2)}{C_2}$$

FIGURE 1 Principle of the dye dilution technique. V_1 (or V_{PEG}) is volume of fluid initially present in stomach. V_2 is volume of indicator dye. C_1 is initial concentration of indicator dye. C_2 is final concentration of marker.

and ranged from 0 to 110 ml. The additional volume (V_{add}) aspirated following direct inspection by the surgeon was 4.4 ± 3.9 ml (range: 0 to 13 ml). The total volume (V_{tot}) present in the stomach at the beginning of surgery was 35.5 ± 29.1 ml (range: 1.5 to 118 ml). V_{tot} was significantly larger than V_{asp} , ($p < 0.0001$). The volume determined by indicator dilution (V_{PEG}) in 15 patients was 26.2 ± 28.8 ml. In these 15 patients who had both determinations of gastric contents, V_{PEG} was not statistically different ($p = 0.157$) from V_{asp} (31.2 ± 27.7 ml). Complete results are presented in the Table.

The volume re-injected into the stomach ($V_{tot} - 2$ ml for pH determination + 100 ml PEG solution) was 134.1 ± 28.2 ml. This volume was significantly different from V_{re-asp} (122.1 ± 34.3 ml; $p < 0.02$).

There was a statistically significant ($p < 0.001$) correlation between V_{asp} and V_{tot} ($r = 0.99$; Figure 2). V_{PEG} was significantly ($p < 0.001$) correlated to V_{tot} ($R = 0.89$, Figure 3). V_{re-asp} also significantly ($p < 0.001$) correlated with the volume re-injected in the stomach ($r = 0.87$).

Discussion

This study compared two methods of measuring the residual gastric fluid volume in anaesthetized adult patients and validated these methods in relation to actual volumes measured following direct stomach manipulation during laparotomy. The results indicate that volumes determined by both the aspiration and dye dilution methods are satisfactory estimates of the volume of fluid

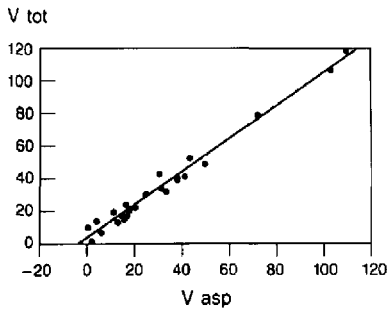


FIGURE 2 Correlation between V_{asp} and V_{tot} . $V_{tot} = V_{asp} + 4.3$ ml by linear regression analysis (24 data points; $r = 0.99$).

present in the stomach. The correlation with V_{tot} determined by aspiration ($r = 0.99$) is slightly better than that determined by PEG dilution ($r = 0.89$). These data are important since significant medical recommendations have been based on such measurements despite the lack of prior validation.

The principal disadvantages of the aspiration method are that it requires meticulous attention to detail and that there is always a small residue (about 4 ml) left in the stomach after aspiration. This should not be of major clinical significance.

V_{re-asp} was significantly smaller than the volume re-injected into the stomach to determine V_{PEG} . This difference may be explained in two ways: either the aspiration technique itself was at fault or, alternatively, fluid was lost through the pylorus during the syringing procedure. We believe the latter explanation is more likely since: (1) the additional volume was very small

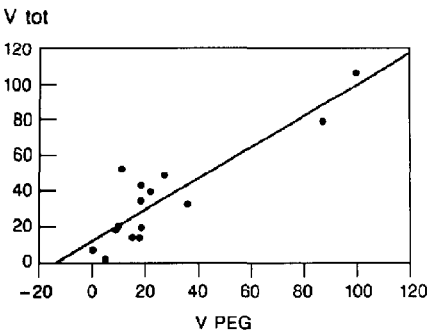


FIGURE 3 Correlation between V_{PEG} and V_{tot} . $V_{tot} = 0.87 V_{PEG} + 12.6$ ml by linear regression analysis (15 data points; $r = 0.89$).

initially, and (2) V_{re-asp} still significantly correlated with the re-injected volume.

The use of a large more rigid, vented and multi-orificed tube appears to be much more effective than the use of a conventional Levin tube, and this may well explain the discrepancy between this and older¹ studies. A larger and more rigid tube is easier to position correctly in the stomach, but may be more difficult to pass through the nose. The distal openings of a vented tube will not so readily be obstructed by gastric mucosa. Finally, multiple distal openings spaced on a sufficient length of tubing will ensure that nearly all the gastric pouch is properly drained. The technique is simple, fast and inexpensive. It does not entail the use of ancillary equipment or personnel. This method could also be suitable to empty the stomach of its liquid contents prior to anaesthesia.

The gastric volume determined by dye (PEG) dilution is a satisfactory predictor of V_{tot} , but V_{PEG} will underestimate V_{tot} by about 12 ml. The method is delicate, complicated and time consuming. Thus, even as a research tool, the technique is not superior to aspiration. It requires homogeneous mixing of gastric fluid and indicator, and a precise determination of PEG concentration. Homogeneous mixing requires a properly positioned tube and an adequate exchange of fluid. The gastric tubes used in this study were most often appropriately positioned in the body² of the stomach, as determined at the time of surgery. Aspiration and re-insertion of five 50-ml syringes is a procedure that should ensure adequate mixing (mixing for 30 seconds or more has been found to be adequate²). The marker used must be non-toxic, water-soluble, non-absorbable and must not be affected by gastric pH.

The validity of PEG as an indicator in the human stomach has been demonstrated.⁶ It is readily measured by a turbidimetric method but this requires the collaboration of experienced laboratory personnel. However, as for other indicators, the behaviour of PEG during initial instillation into the stomach (as compared to subsequent, repeated instillations) raises the possibility that the luminal distribution of indicators may differ from other solutes, such as hydrogen ion, and hence give biased data (spuriously high concentrations).⁶ These falsely elevated PEG concentrations will lead to an underestimation of the volume present in the stomach (Figure 1). The contribution of this phenomenon to our results is not known.

Another possible source of error when determining V_{PEG} is the loss of stomach fluid through the pylorus during the syringing procedure. This may have occurred, V_{re-asp} being smaller than the volume re-injected into the stomach. The exact composition of the lost fluid (gastric fluid to indicator solution ratio) will determine the magnitude and direction of the discrepancy between V_{tot}

and V_{PEG} , but the importance of this possible source of error appears to be minimal when relatively large volumes of indicator solution are used. Conversely, the use of a small volume of PEG solution is less likely to promote adequate mixing and any loss of indicator (or gastric fluid) will result in important variations of V_{PEG} . This could explain the erratic results observed in our first nine attempts to determine V_{PEG} .

Division of the stomach into antral and fundal sacs⁴ was shown radiologically to occur in the gas-distended stomach of one female patient. Experiments using an isolated pig's stomach divided by a 35 mm diameter cylinder to "mimic the effect of the bulge of the vertebral column into a patient's abdomen" showed that a 360° rotation was needed to ensure adequate mixing of antacid and gastric contents.⁴ While application of these findings to the conditions of the present study is uncertain, we did not encounter any difficulty with aspiration or mixing of gastric fluid and indicator, secondary to this possible functional division of the stomach.

A previous study⁵ showed the presence of larger volumes of gastric fluid, as determined by a PEG dilution method, when compared to the volumes estimated by aspiration. These results, at conflict with the present report, may be explained by either an incomplete aspiration of gastric fluid using conventional Levin tubes, or by an over-estimation of gastric contents using a PEG dilution method that had not been validated in the anaesthetic setting.

We conclude that the volume of aspirated gastric fluid through a large, vented, multi-orificed tube is a very good estimate of the volume present in the stomach at the time of induction in the anaesthetized supine adult patient. The technique is fast, simple and inexpensive. The PEG dilution method does not yield superior results. Furthermore, it is time-consuming, relatively complicated, and requires the use of ancillary equipment and personnel.

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Résumé

Cette étude avait pour but de valider, chez l'adulte anesthésié en décubitus dorsal, deux méthodes de mesure du volume du résidu gastrique couramment disponibles en clinique. Il s'agit de 1) l'aspiration directe du contenu gastrique avec une sonde de gros calibre, ventilée et pourvue d'orifices multiples, et de 2) la mesure indirecte du volume gastrique par dilution de l'indicateur polyéthylène glycol (PEG). Les volumes mesurés par ces deux méthodes étaient comparés au volume absolu présent dans l'estomac déterminé par inspection directe au début de la laparotomie. Nos résultats démontrent que le volume mesuré par aspiration directe, lorsque l'on utilise ce type de sonde, est une bonne évaluation du volume du contenu liquide de l'estomac, le sous-estimant d'environ 4 ml. La méthode utilisant la dilution du PEG fournit également des résultats satisfaisants. La technique est cependant plus compliquée, nécessite la collaboration du laboratoire, et n'offre aucun avantage sur l'aspiration directe.