

Preoperative Fasting Times for Patients Undergoing Caesarean Delivery: Before and After a Patient Educational Initiative

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Abstract

Objective: Prolonged preoperative fasting may lead to dehydration, hypoglycaemia, ketoacidosis and delayed recovery. We hypothesised that a patient educational initiative would decrease our preoperative fasting periods for elective caesarean delivery.

Methods: This was an observational quality improvement impact study. Elective caesarean patients who delivered during our study period were included in the study, 40 patients in the pre-intervention and 40 patients in the post-intervention groups. Only English-speaking patients were included. We developed a patient educational pamphlet outlining preoperative fasting and analgesic expectations for caesarean delivery that was given to every patient at her preoperative anaesthesia consultation. The pamphlet included the American Society of Anesthesiologists' preoperative fasting and enhanced recovery carbohydrate drink recommendations. The primary outcome measure was intended fasting duration for liquids (defined as time from last reported liquid consumption to scheduled caesarean delivery) before and after the patient educational initiative. Secondary outcomes included solid fasting time, types of liquids and solids consumed.

Results: The intended median (interquartile range) fasting time for liquids decreased from 10 (8.9-12) h to 3.5 (2.5-10) h (p<0.001). The fasting period for solids was not significantly different: 12.5 (10.5-14) h pre- versus 12.4 (10.6-14) h post-pamphlet introduction (p=0.384). Despite the recommendation, only 22.5% consumed a carbohydrate-containing drink with a modest decrease in water consumption (87.5% before and 67.5% after; p=0.009).

Conclusion: A patient educational pamphlet significantly reduced fasting time for clear liquids. Future studies are needed to determine what barriers limited adherence to the recommended carbohydrate-containing drink consumption.

Keywords: Caesarean delivery, education, enhanced recovery, fasting, quality assurance

Introduction

The American Society of Anesthesiologists' (ASA) 2016 Practice Guidelines for Obstetric Anaesthesia (1, 2) recommends that patients restrict clear fluid consumption for at least 2 h and solid foods for 6-8 h before induction of general or neuraxial anaesthesia. Preoperative fasting reduces the risk of emesis; however, prolonged fasting may lead to hypoglycaemia, dehydration, ketoacidosis, delayed recovery and reduced patient satisfaction (3-5). Pregnant women who are hypovolaemic may be at higher risk of hypotension after spinal anaesthesia.

Institutional and national guidelines to withhold food and drink overnight may lead to excessive periods of fasting prior to caesarean delivery. Enhanced recovery after surgery (ERAS) protocols encourage drinking fluids, preferably carbohydrate-containing drinks, up to 2 h before surgery (5-8). Limiting the duration of fasting may enhance recovery after surgery and decrease length of stay (5, 6). To minimise fasting periods prior to caesarean delivery, implementation of a patient educational initiative was undertaken at our institution. The aim of this quality improvement project analysis was to determine fasting times for patients undergoing scheduled caesarean delivery before and after

the introduction of this initiative. We hypothesised that providing patients with an educational pamphlet outlining the recommended fasting times the day before their scheduled elective caesarean delivery would reduce preoperative liquid fasting time and increase consumption of carbohydrate-containing drinks.

Methods

Stanford University Institutional Review Board exemption was obtained (7 July 2016) prior to the initiation of this quality improvement project. The Institutional Review Board assessed that the project did not meet the federal definition of research or clinical investigation. No written informed consent was deemed required or obtained during this quality improvement project. We approached women undergoing elective caesarean delivery to determine fasting practices for liquids and solids before and after developing and initiating a patient-directed educational pamphlet (Appendix 1. See appendix https://doi.org/10.5152/TJAR.2019.95770) outlining anaesthesia for caesarean delivery. Prior to the implementation of this educational initiative, all women undergoing elective caesarean delivery underwent a preoperative anaesthesia consultation where they were instructed about the scheduled time of their caesarean delivery, as well as fasting recommendations and the risks and benefits of anaesthesia. Prior to this initiative, patients relied on verbal instruction from anaesthesia providers and nurses, and were not provided with specific instructional handouts from the anaesthesia service to take home following their consultation. We developed an educational pamphlet detailing information regarding anaesthesia for caesarean delivery. The educational pamphlet was written in a sixth-grade reading comprehension level and developed only in English. This pamphlet was implemented into our standard workflow and given to every patient at her preoperative anaesthesia consultation that occurred the day prior to her scheduled caesarean delivery at our labour and delivery unit. Patients received the same information during their preoperative consultation as before we implemented the educational initiative; however, they are now provided with this more standardised educational pamphlet to take home and read.

The pamphlet included the ASA recommendations for fasting guideline of 2 h for clear fluids and 8 h for solids combined with ERAS recommendation for consumption of carbohydrate-containing drinks up to 2 h prior to surgery (5, 7, 9). Patients were encouraged to drink up to 12 oz of a carbohydrate-containing drink; however, the drink was not provided to them at the time of the preoperative anaesthesia consultation.

We interviewed a convenient sample of 40 women undergoing scheduled caesarean delivery on fasting practices for

liquids and solids prior to pamphlet implementation and interviewed a sample of 40 women post-pamphlet implementation. All women who undergo elective caesarean delivery at our institution are admitted to the preoperative unit 60-90 min prior to their scheduled caesarean delivery surgery time. During this time, the patients are prepared for their caesarean delivery, for example, a peripheral intravenous line is inserted, and consent and laboratory results are reviewed. We approached the women in the preoperative area on the day of surgery, after they had been admitted for their caesarean delivery, to collect their responses regarding current fasting status. Pre-pamphlet data were collected from June 2016 to July 2016, and post-pamphlet data were collected from January 2017 to June 2017. The following questions were asked in our pre-pamphlet group: (1) when was your last fluid intake prior to surgery and what did you drink? (2) when was your last solid meal prior to surgery and what did you eat? (3) were you hungry prior to your caesarean section? (4) if you were given the option of drinking a liquid energy drink or protein shake prior to your surgery, do you think that would have improved your experience? and (5) were you told ahead of time by a nurse or doctor about fasting prior to surgery? The following questions were asked in our post-pamphlet group: (1) did you receive a copy of the pamphlet? (2) did you read the pamphlet? (3) was the pamphlet helpful? (4) what is your understanding of how many hours before your caesarean that you can drink? (5) when was your last fluid intake? (6) what did you drink? (7) what is your understanding of how many hours before your caesarean that you can eat? (8) when was your last solid intake? (9) what did you eat? (10) are you hungry? and (11) are you thirsty? No demographic, obstetric or patient-identifying data were collected as part of this quality improvement project.

The primary outcome measure was the intended fasting duration for liquids, defined as the time duration between patient last reported liquid consumption and the scheduled caesarean delivery time. Secondary outcomes included the intended fasting duration for solids, the actual fasting duration for liquids and solids (defined as the time duration between patient last reported liquid or food consumption and neuraxial block placement), patient self-reported thirst and hunger, types of drinks consumed (water vs. other liquid) and solids consumed (heavy, defined as a meal containing high fat content vs. light, defined as a meal containing low fat content).

Statistical analysis

Data were analysed using IBM Statistical Package for the Social Sciences Statistics for Windows, version 23 (IBM SPSS Corp.; Armonk, NY, USA). Data are presented as mean±standard deviation, median (interquartile range; range) and count (percentage), as appropriate. Statistical tests for primary and secondary outcome measures included between

groups comparisons using the Student's t-test for normally distributed variables and Mann-Whitney U test for non-parametric comparisons. Data were graphed, and QQ test was performed to assess the normality of data distribution. Categorical variables were investigated using Pearson's chi-square test. A p-value of <0.05 was considered statistically significant.

Results

We had complete data from 33 out of 40 women in the pre-pamphlet group and 40 out of 40 women in the post-pamphlet implementation group. Missing data were a result of unrecorded scheduled caesarean delivery times and/or unrecorded block placement times in the pre-pamphlet group. We had a 97.5% (39 out of 40) implementation rate of women receiving the pamphlet at their preoperative anaesthesia visit. A self-reported 92.3% (36 out of 39) of those who received the pamphlet at their anaesthesia consult visit reported reading the pamphlet. All patients (36 out of 36) who read the pamphlet reported finding the information helpful.

There was a significant difference in liquid fasting times between the pre- and post-pamphlet group. Median (interquartile range) liquid fasting times were 10 (8.9-12) h in the pre- and 3.5 (2.5-10) h in the post-pamphlet introduction groups (p<0.001; Figure 1). Women's understanding of liquid fasting duration was 2 (2-2) h, consistent with the recommen-

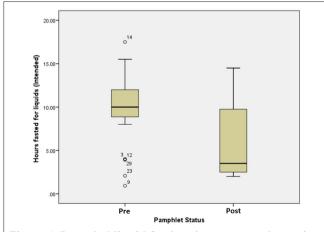


Figure 1. Intended liquid fasting times pre- and post-intervention

dation in the pamphlet. Actual liquid fasting times between pre- and post-pamphlet groups were also statistically different (p<0.001; Table 1). The median (interquartile range) delay for surgical start time was 0.5 (0.1-0.8) h. Delay in surgery caused a difference between intended and actual fasting times for both liquids (p=0.007) and solids (p<0.001), respectively. Before intervention, 87.5% (35 out of 40) of women consumed water, and after pamphlet introduction, 67.5% (27 out of 40) of women consumed water (p=0.009). Of the 40 patients, 0% and 22.5% consumed a carbohydrate-containing drink in the pre- and post-pamphlet groups, respectively.

Before intervention, 53.8% (21 out of 39) of women reported eating a heavy meal the night prior to caesarean delivery, and after pamphlet introduction, 50% (20 out of 40) consumed a heavy meal the night prior (p=0.732). Pamphlet introduction did not impact patients' perception of hunger, and 62.5% (25 out of 40) reported hunger before vs. 75% (30 out of 40) reported hunger after the pamphlet implementation (p=0.228). Subjective perception of thirst was only asked in the post-pamphlet group. Although women reported shorter liquid fasting times, a high proportion, 82.5% of women, still reported thirst.

Discussion

This quality improvement project demonstrates that a patient educational pamphlet is effective at decreasing preoperative liquid fasting times in patients undergoing elective caesarean delivery. Successful implementation of a patient educational pamphlet requires integration into the workflow of preoperative anaesthesia consultation for elective caesarean patients. ERAS protocols utilised in other surgical subspecialties, such as colorectal, urology and gynaecology-oncology, recommend pre-admission education and counselling (9-12). Prior to this educational initiative, patients were verbally instructed on the current fasting practice during their preoperative anaesthesia consultation; however, median (interquartile range) liquid fasting time was 10 (8.9-12) h, suggesting that not all women understood nor retained the information relayed to them during their consultation. We had a very high (97.5%) pamphlet distribution rate, and all the patients who read the pamphlet reported finding the information helpful. Our patients also demonstrated understanding of the preoperative

Table 1. Liquid and solid fasting times pre-and post-intervention			
	Pre- Intervention n=33	Post- Intervention n=40	p
Intended Fasting Time Liquids	10 [8.9-12]	3.5 [2.5-10]	< 0.001
Actual Fasting Time Liquids	10.4 [9.1-12.4]	3.9 [3.0-10.1]	< 0.001
Intended Fasting Time Solids	12.5 [10.5-14.0]	12.4 [10.6-14.0]	0.384
Actual Fasting Time Solids	13.6 [10.8-14.5]	12.9 [10.8-14.5]	0.907
Values are median [interquartile range] in ho	urs		

fasting recommendation with a median (interquartile range) for their understanding of liquid fasting time of $2\ (2-2)\ h$. However, the actual post-intervention liquid fasting times were higher $(3.5\ h)$ than the $2\ h$ recommended, and the interquartile range of $2.5-9.9\ h$ suggests variability in patient application of this knowledge.

The ASA 2016 Guidelines for Obstetric Anaesthesia (1, 2) recommends preoperative fasting to 2 h for clear liquids. Prolonged fasting (i.e. nil per os after midnight) has not been shown to lower the amount of gastric contents nor decrease gastric acidity (10, 13, 14). Perioperative fasting is associated with increased surgical stress, increased insulin resistance and impaired gastrointestinal function (6, 7, 15). Impaired gastrointestinal function, such as postoperative ileus, has been demonstrated to significantly prolong post-caesarean delivery length of stay (16). Measures that can improve patient's nutritional status in the perioperative period could potentially lead to improved clinical outcome in the caesarean delivery patient population. ERAS protocols utilised in colorectal surgery have shortened hospital length of stay and decreased complication rates by up to 30-50% (9, 11, 15). Although there are no national guidelines for enhanced recovery fasting recommendations or guidelines for caesarean delivery, Wrench et al. (17) have successfully demonstrated adoption of ERAS protocol to the obstetric unit and its impact on decreased length of stay for elective caesarean deliveries. Adoption of an ERAS protocol and associated reduction in fasting duration before elective caesarean may have other benefits, such as decreased complication rates and decreased postoperative ileus that have been observed in other surgical subspecialties.

A key component of ERAS protocols is minimising fasting period and increasing nutritional support. Surgery induces a catabolic state; the recommendation of a preoperative carbohydrate-containing drink 2 h prior to caesarean delivery aims to keep the patient metabolically balanced and decrease insulin resistance (5, 6, 8, 12, 15). Our educational pamphlet recommends that women drink a carbohydrate-containing drink as their clear liquid 2 h before their scheduled elective caesarean delivery. Before the implementation of this educational initiative, patients were verbally instructed on the types of liquids that met the criteria for clear liquids but were not consistently instructed to drink a carbohydrate-containing drink as their clear liquid. Our pamphlet initiative standardised our clear liquid recommendations, such that all women were instructed to consume a carbohydrate-containing drink as their clear liquid. Despite this, our recommendation had a modest impact with only 22.5% of patients consuming a carbohydrate-containing drink, such as Gatorade, after the educational intervention, and water was still consumed by most women (87.5% before and 67.5% after introduction of the pamphlet). Self-reported reasons for not choosing a carbohydrate-containing drink include not readily available at home, preference for unsweetened drink and preference for water. Giving patients a carbohydrate-containing drink on the day of their preoperative anaesthesia consultation visit, as is often done for colorectal and other ERAS protocols, may help to improve adherence with this recommendation.

Our pamphlet initiative did not significantly change the intended and actual fasting times for solids for our pre- and post-pamphlet groups. Therefore, patient self-report of hunger was not significantly different between the two groups. Our educational pamphlet did not make recommendations regarding the types of solid food to consume; consequently, the percentage of women who ate a heavy meal in the pre- and post-pamphlet groups was not significantly different. Recommendations for type of solids consumption could be considered when implementing patient fasting educational initiatives; however, we did not find a difference in reported hunger between women consuming or not consuming a 'heavy meal'.

We found a delay between actual and scheduled caesarean delivery times. On average, this increased fasting times for both liquids and solids by only 0.5 h. During this quality improvement analysis period, wrongful adherence to the fasting guidelines did not contribute to surgical delay. Scheduled elective caesarean deliveries are sometimes delayed to accommodate urgent cases on the obstetric suite, and delays will vary amongst institutions with different workload demands or staffing availability. A prolonged delay can significantly impact a woman's fasting duration and contribute to hypovolaemia and potentially greater haemodynamic instability after neuraxial anaesthesia. Offering clear fluids to women awaiting caesarean delivery should be considered if significant delays are expected or unexpectedly occur.

Our quality assurance study has several limitations. The goals of our analysis were to examine the feasibility of implementing new fasting recommendations and to observe any change in behaviour with regard to fasting practices. We did not examine barriers to adherence of our recommendations or of the significance the change in liquid fasting time would have on patient outcomes. This patient educational pamphlet was initially developed in English and given to English-speaking patients; therefore, we cannot generalise its effect to patients whose primary language is not English.

Conclusion

This quality assurance analysis demonstrates that a patient educational pamphlet given at the time of preoperative anaesthesia consultation can significantly reduce fasting time for clear liquids. However, the consumption of carbohydrate-containing drinks was limited despite the pamphlet's recommendation for these liquids rather than water to enhance recovery. Future studies are needed to examine barriers against carbohydrate-containing drink consumption and if different initiatives can increase carbohydrate-containing drink utilisation. The impact of shorter liquid fasting duration and heavy versus light preoperative meals on maternal and foetal outcomes after caesarean delivery need further investigation.

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Ethics Committee Approval: Notice of Determination of Human Subject Research was obtained 7 July 2016. The Stanford University Institutional Review Board assessed that this quality improvement project did not meet the federal definition of research or clinical investigation, and Institutional Review Board exemption was obtained prior to the initiation of this quality improvement project.

Informed Consent: No written informed consent was deemed required by the Stanford University Institutional Review Board for this quality improvement project.

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Conflict of Interest: The authors have no conflicts of interest to declare.

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Appendix 1.

Recovery

You will stay in recovery for about one hour.

As the numbness wears off, you may feel discomfort at the surgery site. You will be offered medicine to help treat this.

You may be itchy, feel nauseous or shiver after delivery. If you do, we have medicines to help treat these symptoms.

You are welcome to continue skin-to-skin and nursing in recovery. We encourage sips of water and ice chips.

After your Cesarean

You will be asked to rate your pain from 0 (no pain) to 10 (worst pain imaginable). Pain after Cesarean is usually very well controlled and we try to keep pain at 4 or less. Pain usually peaks on the second day after surgery (when the pain medicine placed in your spinal wears off and you start to move around).

You will receive pain medicine at scheduled times (without needing to ask) in order to keep you comfortable. If you are still uncomfortable, you can ask your nurse for more medicine. All of these medicines are safe for nursing mothers.

An anesthesiologist will come and visit you and check on how you are doing a day or two after your surgery. Your nurse can call an anesthesiologist back sooner if you have any questions or needs.

Commonly asked questions

I am very nervous about my Cesarean, can I be asleep for it instead?

General anesthesia has added risks for mom and baby compared to spinal anesthesia. Therefore we only put you to sleep in unique situations and emergencies.

Can my partner stay with me during my Cesarean? Yes. Your partner can stay with you from the moment you go into the operating room to the moment you go to recovery. If there is an emergency and you must go to sleep, your partner will be escorted out of the operating room so that we may focus solely on taking care of you.

Can I still do skin-to-skin with baby if I am having a Cesarean?

Yes. The doctors will examine the baby right after birth and if he/she is doing well, and it is an appropriate time during surgery, the baby will be brought to you for skin-to-skin.

Will I get back pain from a spinal or epidural? Spinals and epidurals do not cause long-term back pain. You may have mild bruising and tenderness where the spinal or epidural is placed that may last a few days.

Will a resident do my anesthetic?
Stanford offers the highest quality of care. An attending physician anesthesiologist who is an expert in obstetric anesthesia will closely supervise experienced residents and fellows, and participate in all key aspects of your care. Our team care model offers many advantages over isolated clinical practice.

Lucile Packard
Children's Hospital
Stanford

Anesthesia for Your Upcoming Cesarean Delivery

Congratulations

Below is some information to make your Cesarean delivery as safe and pleasant as possible.

Before your Cesarean

You may eat up to 8 hours before your scheduled cesarean delivery. We recommend eating a regular meal the night before surgery.

You may drink water or a carbohydrate-containing drink up to 2 hours before you come to the hospital for your surgery. Drinking water may help you feel less thirsty and hungry. A carbohydrate-containing drink (Gatorade, Powerade or Vitaminwater) may help speed your recovery. Therefore we encourage you to drink up to 12 Oz (or 1.5 cups) of Gatorade, Powerade.

Vitaminwater or water up to 2 hours before you come to the hospital for your surgery.

Please take your medicines as recommended by your obstetrician and anesthesiologist.

<u>When you arrive</u>

Your obstetrician and anesthesiologist will come and talk to you and answer any questions.

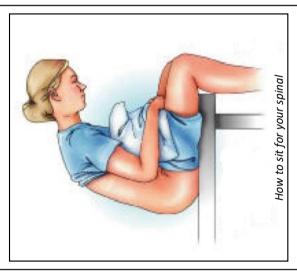
You will receive an IV and a warming blanket, and your nurse will prepare you for surgery.

To allow for the safest care possible, your surgery may be delayed if there are any emergencies on labor and delivery.

During your Cesarean

Your anesthesiologist will stay with you throughout the surgery, and will help keep you safe and comfortable.

Most scheduled Cesareans are done with a spinal or a combined spinal-epidural anesthetic. The anesthesia we use is extremely safe.



Your anesthesiologist will guide you through your spinal or combined spinal-epidural. You will be positioned as in the picture above. You may feel pressure during the procedure, but it should not be painful. We use medicine to numb where the anesthetic will be placed, and you are welcome to request more numbing medicine if you feel discomfort during the anesthetic procedure.

Within a few minutes of your spinal, your legs will become numb. The anesthesiologist will test several times to make sure you are numb. Surgery will not start until we are sure that you are adequately numb and will be comfortable during surgery.

The spinal will make you numb and heavy from the mid-chest down. You may feel some pressure and tugging during surgery but you should not feel surgical pain. If you feel pain or discomfort please let your anesthesiologist know. You will be awake during surgery. Sedation can be used, but we prefer to avoid giving any so you can enjoy and remember the birth.

Please feel free to ask any questions or express any needs that you have during your surgery.