Assessment of Different Regiments of Early Oral Feeding in Comparison to Delayed Feeding After Elective Uncomplicated Cesarean Section: A Randomized Control Trial

Adel Atef¹, Rania Elsayed², Akmal El-Mazny¹ and Sarah Hassan¹

¹Department of Obstetrics and Gynecology, Faculty of Medicine, Cairo University, Egypt
²Resident of Obstetrics and Gynecology at Damietta General Hospital, Damietta, Egypt

ABSTRACT

Aim: Starting oral feeding after caesarian section enhances bowel function return, patient's ambulation, and patient recovery. We aimed to evaluate effectiveness of two different approaches of early oral feeding following section on gastrointestinal function, and patient satisfaction.

Materials and Methods: In this randomized control study, 300 pregnant women with elective uncomplicated cesarean section done from October 2019 to March 2020 at Cairo University teaching hospital with were randomly allocated into three feeding groups.

Results: Significant differences were noticed in patients' satisfaction and time interval to return of gastrointestinal function, ambulation, regular diet, and hospital stay between the three groups and between the two groups of early feeding (P value < 0.05 for all) with highest satisfaction and shortest intervals noticed at introducing soft foods early irrespective of return of intestinal sounds (The first early feeding group). Non-significant differences noted concerning postoperative complications between the three groups or between the two groups of early feeding apart from vomiting found to be higher in the early feeding groups, but the cases were mild and easily treatable.

Conclusion: Early oral feeding reduces the time needed for normal bowel function return and increases satisfaction of the patients with no detrimental significant effects on the gastrointestinal complication.

Key Words: Cesarean, early feeding, gastrointestinal symptoms, satisfaction.

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Corresponding Author: Adel Atef, ¹Department of Obstetrics and Gynecology, Faculty of Medicine, Cairo University, Kasr Alainy Street, Cairo 11562, Egypt, Tel.: +20 1115241250, E-mail: dr.adel90@gmail.com

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INTRODUCTION

In the industrialized world Caesarean section (CS) is considered one of the major surgical procedures performed commonly in hospitals⁴¹. Rates in Egypt have been steadily increasing to about 52% in 2014⁴². With the changing attitudes of surgeons, early oral feeding after abdominal surgery, especially cesarean section, has been considered⁴³.

Postoperative ileus has become a public health problem because of its role in postoperative morbidity and increased hospital stay⁴⁴. Commonly, after abdominal surgery and until return of bowel function (evidenced by bowel sounds, flatus or stool passage, or hunger feeling) no fluids or food are given to patients. This practice is done for prevention of postoperative complications as nausea, vomiting, distention, and others⁴⁵. Hence limitation of bowel mobility was believed to follow CS, there was a fear that postoperative ileus will be of common incidence. So that the belief of bowl functions affection after oral feeding was prevalent among both the public and medical staff⁴⁶.

From a surgical point, bowel manipulation during cesarean delivery never takes place and peritoneal irritation occurs significantly less often as compared to another laparotomy procedure⁴⁷. Interventions as ambulation, probiotics, chewing gum after surgery aimed to accelerate the gastrointestinal motility recovery, and many systematic reviews and clinical trials proved the benefits of early over delayed oral feeding⁴⁸. Early oral feeding may help in reducing rate of body protein depletion, improvement of wound healing and psychological status, and help to reduce the incidence of nosocomial infections, hospital stay time and treatment costs⁴⁹.

The current study aimed to evaluate the effect of two different approaches of early oral feeding after cesarean section on gastrointestinal function and patient satisfaction.
METHODOLOGY

Study design and setting

We conducted a randomized controlled study employing a purposeful convenient sample from October 2019 to March 2020. The research was carried out at Labor and Delivery unit at obstetrics and gynaecology department in Kasr Alainy Teaching hospital, located in the metropolitan area of Cairo and serves as the largest tertiary referral center in Egypt and the region of Middle East[10]. The study has been approved by the research ethics committee, Faculty of Medicine, Cairo University prior to commencement of the study. The current research was prospectively registered in ClinicalTrials.gov registry website with clinical trial identifier (NCT04338737). Informed written consent was obtained from all potential participants following comprehensive explanation of the purpose and potential benefits of the study.

Sample size was calculated using G*Power software version 3.1.2 for MS Windows, with an error probability of 0.05, 80% for the power and 1 was set for intervention groups ratio resulting in a minimum sample of 100 in each group.

Study population

All pregnant women, primigravida or had previous one C/S, aged 20-40 years, with viable singleton pregnancy who planned to undergo elective c/s for obstetric indications were eligible to participate in the study. Patients were excluded from the study if they had chronic medical disorders such as diabetes, liver and kidney diseases, history of major abdominal surgery other than C/S and intraoperative intestinal surgery. Those who experienced obstructed labor with features suggestive of peritonitis and who declined to participate were also excluded from the study.

Three-hundred women participated in this work were randomly and evenly assigned to early oral feeding (EOF) or late oral feeding (LOF) groups. Patients allocated to EOF were subsequently divided into 2 groups, A and B. Group A comprised of 100 women received water and clear fluid approximately 2 hours after surgery, followed by soft food and regular diet 4 hours later irrespective to intestinal sounds, flatus, or stool. In the same context, 2 hours post C/S, water and clear fluid were initiated in group B till the return of intestinal sounds, then soft food, and regular diet later. Group C participants were made NPO and received 2-3 litres intravenous fluid till the return of bowel functions. Thereafter, water and clear fluids were given followed by soft food and, eventually a regular diet.

Study measurements

All eligible patients participating in the study were subjected to spinal anaesthesia using heavy Marcaine injected by a spinal needle no 25. Preoperative prophylactic dose of 3rd generation cephalosporin was administered to all women preceded by skin sensitivity test. All C/S deliveries were performed according to the local hospital protocol by a well-trained obstetrician. We adopted double-layer closure of the uterine incision with gentle manipulation of the tissue[11]. Postoperatively, all patients were examined hourly for the first 6 hours then four hourly till the time of patients discharge. First audible intestinal sounds and passage of first flatus, postoperatively, were recorded in hours for all study population. Assessment of possible postoperative complications and/or discomfort such as nausea, vomiting, abdominal distension, and manifestations suggestive of ileus was done. The patient satisfaction was evaluated by all participants using visual analogue scale prior to discharge.

Statistical analysis

Statistical package for the Social Sciences (SPSS) version 26 (IBM Corp., Armonk, NY, USA) was used for coding and entering data. Summarization of data was done using mean and standard deviation for quantitative variables and frequencies (number of cases) and relative frequencies (percentages) for categorical variables. Analysis of variance (ANOVA) was used for group comparison and post hoc test for multiple comparisons. Chi square (χ²) test was used for categorical data comparison. Statistical significance is considered with P-values less than 0.05.

RESULTS

In our study 525 women were allocated for the study where 225 women were excluded as 119 women did not fit the inclusion criteria and 86 women met the exclusion criteria and 20 women refused to participate. The 300 women participated in this work were randomly allocated into three equal groups 100 in each.

The mean age in the participant women was 26.88±3.9 years, the mean BMI was 29.04±0.9415kg/M², the mean parity was 1.84±0.819, and the mean operative duration was 44.32±6.59 minutes. (Table 1) showed basic demographic characteristics of included participants with no significant difference between the three groups. There was no significant difference between the three groups as regarding presence of adhesions (p value 0.517).

Clear fluid was allowed for Group A and B 2 hours after surgery and only allowed for the third group after open bowel. Semisolid and regular diet were allowed for group A 4 hours after surgery regardless return of intestinal function while for group B semisolid and regular diet were allowed after return of intestinal sound. For group C semisolid and regular diet were allowed after open bowel after somewhat from clear fluid. So, there was statistically highly significant difference between the three
groups as regarding post-operative interval to return of bowel sounds (\(p \text{ value } 0.0001\)), return of bowel movement (\(p \text{ value } 0.0001\)), return of regular diet (\(p \text{ value } 0.0001\)), starting ambulation (\(p \text{ value } 0.0001\)), time to be eligible for discharge (\(p \text{ value } 0.0001\)), and as regarding satisfaction assessed using the VAS tool at discharge (\(p \text{ value } 0.0001\)) as shown in (Table 2). These results showing that group A had the best outcome as regarding bowel function and satisfaction (Figure 1) and has earlier ambulation and shorter hospital stay (Figure 2) followed by group B and then group C.

![Fig. 1: Participants satisfaction among different groups](image1)

![Fig. 2: The hospital stay length among three groups.](image2)

### Table 1: Demographic data of the studied groups

<table>
<thead>
<tr>
<th>Group</th>
<th>Age (years)</th>
<th>BMI</th>
<th>Parity</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>26.47</td>
<td>28.96</td>
<td>1.82</td>
</tr>
<tr>
<td>B</td>
<td>27.42</td>
<td>29.18</td>
<td>1.95</td>
</tr>
<tr>
<td>C</td>
<td>26.75</td>
<td>29.01</td>
<td>1.75</td>
</tr>
<tr>
<td><strong>P value</strong></td>
<td>0.141</td>
<td>0.204</td>
<td>0.215</td>
</tr>
</tbody>
</table>

### Table 2: Effect of different approaches on patient's outcome

<table>
<thead>
<tr>
<th></th>
<th>Group A</th>
<th>Group B</th>
<th>Group C</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st oral fluid feeding (hours)</td>
<td>2.00</td>
<td>2.00</td>
<td>10.14</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Semisolid diet intake (hours)</td>
<td>4.00</td>
<td>5.59</td>
<td>4.16</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Regular diet (hours)</td>
<td>4.00</td>
<td>8.30</td>
<td>10.14</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>1st intestinal sound (hours)</td>
<td>5.35</td>
<td>5.59</td>
<td>7.07</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Open bowel (hours)</td>
<td>6.08</td>
<td>8.30</td>
<td>10.14</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Satisfaction</td>
<td>6.80</td>
<td>6.01</td>
<td>4.16</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Ambulation (hours)</td>
<td>5.68</td>
<td>6.62</td>
<td>7.51</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Admission time (hours)</td>
<td>9.30</td>
<td>12.65</td>
<td>14.69</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

Significant \(P < 0.05\)
Table 3 showed the effect of each feeding approach on unpleasant outcome as nausea where there was no significant difference between the three approaches (P value 0.27), while significant difference was observed between different groups on vomiting (P value 0.014), where delayed feeding group has the least number and patients experienced vomiting were received IV metoclopramide and none of them had second attack. As regarding distension, ileus and readmission, there was no statistical difference between the studied groups.

Table 3: Postoperative complications in different groups

<table>
<thead>
<tr>
<th>Group</th>
<th>Count</th>
<th>%</th>
<th>Count</th>
<th>%</th>
<th>Count</th>
<th>%</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nausea</td>
<td>Yes</td>
<td>16</td>
<td>15</td>
<td>15.0%</td>
<td>23</td>
<td>23.0%</td>
<td>0.276</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>84</td>
<td>85</td>
<td>85.0%</td>
<td>77</td>
<td>77.0%</td>
<td></td>
</tr>
<tr>
<td>Vomiting</td>
<td>Yes</td>
<td>13</td>
<td>8</td>
<td>8.0%</td>
<td>2</td>
<td>2.0%</td>
<td>0.014</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>87</td>
<td>92</td>
<td>92.0%</td>
<td>98</td>
<td>98.0%</td>
<td></td>
</tr>
<tr>
<td>Distension</td>
<td>Yes</td>
<td>0</td>
<td>1</td>
<td>1.0%</td>
<td>0</td>
<td>0.0%</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>100</td>
<td>99</td>
<td>99.0%</td>
<td>100</td>
<td>100.0%</td>
<td></td>
</tr>
<tr>
<td>Ileus</td>
<td>Yes</td>
<td>0</td>
<td>1</td>
<td>1.0%</td>
<td>0</td>
<td>0.0%</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>100</td>
<td>99</td>
<td>99.0%</td>
<td>100</td>
<td>100.0%</td>
<td></td>
</tr>
<tr>
<td>Re admission</td>
<td>Yes</td>
<td>0</td>
<td>0</td>
<td>0.0%</td>
<td>0</td>
<td>0.0%</td>
<td>--------</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>100</td>
<td>100</td>
<td>100.0%</td>
<td>100</td>
<td>100.0%</td>
<td></td>
</tr>
</tbody>
</table>

**DISCUSSION**

In the current study, information about the return of bowl function following C-section under spinal anaesthesia in both the EOF (groups A&B) and LOF (group C) was evaluated. Evaluation of this comparison revealed that EOF especially group A is superior to LOF as EOF provoked earlier bowel sound after surgery. Also, no significant difference was found in GI complications between both groups apart from mild vomiting. Finally, the maternal satisfaction increased among EOF.

In this study 2 h after surgery was chosen for the early feeding groups because of possible complications that may occur during this period such as hemorrhage. Jalilian, et al 2013, and Ahmed et al, 2018, also chose the 2h post-operative time for the early feeding group.[22,23]

The early feeding groups especially group A had a statistically significant shorter mean time to first bowel sound (P value 0.0001). This finding was similar with other related studies.[9,14-17] The possible variations reported may be related to the difference in both nature of diet in different countries and the time length before initiation of oral feeding.

Furthermore, women in the early feeding groups had a statistically significant shorter mean time to first passage of flatus (P value 0.0001). This was also reported by Teoh 2007 with different mean time to first passage of flatus in each study and this may be due to the different approaches of feeding we used.[19] Both Ahmed et al 2018, and Kovavisarach, 2005 found the time until the passage of flatus in the early hydration group was shorter than the delayed hydration group but the difference was statistically insignificant (P value >0.05).[13,19]. Kovavisarach, 2005 was including general and spinal anaesthesia in their study and compared their effect and this may be the cause why the difference is insignificant in his study[19].

In our study, time for return to regular diet was statistically significant shorter among group A (P value 0.0001). The same was reported by LR, 2017, and Devi et al, 2015 with different mean time to return to regular diet in each study and this may be due to the different approaches of feeding we used.[20,21]

Ahmed et al 2018 found the time until the return to regular diet in the early hydration group was shorter than the delayed hydration group but the difference was statistically insignificant (P value 0.296) [13]. This may be due to difference in the meaning of regular diet between our studies (They considered the regular diet as the 2nd tolerated solid meal). Our findings regarding return of bowel function may be explained by that gastrointestinal hormonal secretion could be stimulated by food ingestion. Moreover, some reflexes may be stimulated which produces coordination of propulsive activity resulting in beneficial effect on bowel motility[22].

Also, in the early feeding groups especially group A had a statistically significant shorter mean time to ambulation (P value 0.0001). This was also reported by Mohamed A 2018, and Nantasupha et al. 2016 and this may be because prohibiting oral feeding gave the participant the attitude that they are sick and should be in bed (made them play the role of the patient)[23,24].

In our study, group A had a statistically significant shorter mean time to be eligible for discharge (P value
0.0001) than other groups. The same was observed by Al-Ghareeb 2013[23]. Jalilian et al 2013 found that the time was shorter in the early feeding group than the delayed feeding group, but the difference was insignificant, and they explained this as their hospital protocols did not allow post CS discharge before 48 hrs. till achieving other criteria[12].

Statistical comparison between the three groups using the VAS tool at discharge showed that women in the early feeding group especially group A had a significantly higher satisfaction than those in group B & C (P value 0.0001). The same was observed from previous studies[17,10]. Nantasupha et al. 2016, found that patients’ satisfaction scores were similar among the three groups (P value 0.110) [24]. This may be due to the small sample size they included (40 participant for each group). Assessment of maternal satisfaction is an essential outcome as it affects postpartum blues, future pregnancies decisions, and reputation of both doctor and hospital.

Most of previous studies together with our study confirmed no difference in GI complications (except for vomiting) associated with EOF groups versus LOF group. Nausea was experienced by 16%,15% and 23% among groups A, B and C respectively but with no significant difference either between the three groups (P value >0.276) or between the two groups of early feeding (P value 0.500). These results were parallel to the results of Mawson et al 2019, and Ahmed et al 2018[13,15]. Teoh 2007 showed more nausea in the early feeding group than the delayed feeding group with no statistical significance (P value 0.3). This may be explained as Teoh 2007 introduced feeding very early (0.5 h after surgery)[10].

Vomiting was observed in 13%,8% and 2% of groups A, B and C respectively. With significant difference between the three groups (P value 0.014) having more vomiting in the early feeding groups but no significant difference between the two groups of early feeding (P value 0.178) but the cases was treated easily and didn’t affect maternal satisfaction, prevent progress in the stepping diet, or delay their eligibility from discharge later than the time experienced in other participants. Vomiting in those participants may be due to the effect of pregnancy hormones in delaying the gastric emptying time or may be due to their lower vomiting threshold (personal variation). Unlike that, studies of all, Al-Ghareeb 2013, Mawson et al 2019 and Teoh 2007 showed no significant difference between the early and delayed feeding groups as regarding vomiting with less vomiting in the delayed feeding groups[13,18,23].

In our study, there was no cases of distension, paralytic ileus, or readmission in the three groups. Other studies as LR 2017, and Mohamed A 2018 found that there was no significant difference between the early and delayed feeding groups as regarding distension and ileus (P value >0.05) with less cases in the early feeding groups and the ileus when occurred was found to be mild with no cases of severe ileus[20,23]. All the above-mentioned studies had no cases of readmission except one case in one study which was readmitted for a septic wound and not for a cause related to the study.

CONCLUSION

In conclusion, early feeding within 2 hours after C-section and introduction of regular diet regardless to open bowel was found to be more convenient with best maternal satisfaction and improved bowel function with accepted mild complication and shorter hospital stay and found to be superior to other early feeding modality waiting return of bowel sound for regular diet and by far more favorable than the standard delayed feeding regimen. Further studies including emergency cases and other cesarian section categories should be considered.

AUTHORS' CONTRIBUTIONS

Dr. AA: supervision of the practical work, writing and final revision of the paper. Prof. AM: final revision of the paper. Dr. SH: assistance in writing and final revision. Dr. RE: responsible for the practical part of the research. All authors read and approved the final manuscript.

CONFLICT OF INTERESTS

There are no conflicts of Interest.

REFERENCES


