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To cite this article: Andrea Ciardulli, Gabriele Saccone, Daniele Di Mascio, Claudia Caissutti & Vincenzo Berghella (2018) Chewing gum improves postoperative recovery of gastrointestinal function after cesarean delivery: a systematic review and meta-analysis of randomized trials, *The Journal of Maternal-Fetal & Neonatal Medicine*, 31:14, 1924-1932, DOI: [10.1080/14767058.2017.1330883](https://doi.org/10.1080/14767058.2017.1330883)

To link to this article: <https://doi.org/10.1080/14767058.2017.1330883>



Accepted author version posted online: 14 May 2017.  
Published online: 06 Jun 2017.



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REVIEW ARTICLE



## Chewing gum improves postoperative recovery of gastrointestinal function after cesarean delivery: a systematic review and meta-analysis of randomized trials

Andrea Ciardulli<sup>a</sup>, Gabriele Saccone<sup>b</sup> , Daniele Di Mascio<sup>c</sup> , Claudia Caissutti<sup>d</sup>  and Vincenzo Berghella<sup>e</sup> 

<sup>a</sup>Department of Obstetrics and Gynecology, Catholic University of Sacred Heart, Rome, Italy; <sup>b</sup>Department of Neuroscience, Reproductive Sciences and Dentistry, School of Medicine, University of Naples Federico II, Naples, Italy; <sup>c</sup>Department of Obstetrics and Gynecology, University of Sapienza, Rome, Italy; <sup>d</sup>Department of Experimental Clinical and Medical Science, DISM, Clinic of Obstetrics and Gynecology, University of Udine, Udine, Italy; <sup>e</sup>Division of Maternal-Fetal Medicine, Department of Obstetrics and Gynecology, Sidney Kimmel Medical College of Thomas Jefferson University, Philadelphia, PA, USA

### ABSTRACT

**Objective:** To examine whether chewing gum hastens the return of gastrointestinal function after a cesarean delivery.

**Methods:** All randomized controlled trials comparing the use of chewing gum in the immediate postoperative recovery period (i.e. intervention group) with a control group were included in the meta-analysis. The primary outcome was the time to first flatus in hours. Meta-analysis was performed using the random effects model of DerSimonian and Laird, to produce summary treatment effects in terms of mean difference (MD) or relative risk (RR) with 95% confidence interval (CI).

**Results:** Seventeen trials, including 3041 women, were analyzed. Trials were of moderate to low quality with different inclusion criteria. In most of the included trials chewing gum was given right after delivery, three times a day for 30 min each and until the first flatus. Women who were randomized to the chewing gum group had a significantly lower mean time to first flatus (MD – 6.49 h, 95%CI –8.65 to –4.33), to first bowel sounds (MD – 8.48 h, 95%CI –9.04 to –7.92), less duration of stay (MD – 0.39 days, 95%CI –0.78 to –0.18), lower time to first feces (MD – 9.57 h, 95% CI –10.28 to 8.87) and to the first feeling of hunger (MD – 2.89 h, 95%CI –4.93 to –0.85), less number of episodes of nausea or vomiting (RR 0.33, 95%CI 0.12 to 0.87), less incidence of ileus (RR 0.39, 95%CI 0.19 to 0.80) and significantly higher satisfaction.

**Conclusions:** Gum chewing starting right after cesarean delivery three times a day for about 30 min until the first flatus is associated with early recovery of bowel motility. As this is a simple, generally inexpensive intervention, providers should consider implementing cesarean postoperative care with gum chewing.

### ARTICLE HISTORY

Received 27 April 2017

Revised 11 May 2017

Accepted 11 May 2017

### KEYWORDS

Labor; delivery; vaginal delivery; operative delivery; cesarean section

## Introduction

Cesarean delivery is the most common major surgical operation in the United States, with about one million done annually for an overall rate of about 30% in 2015 [1]. Postoperative ileus is an impaired condition of gastrointestinal motility defined as an abnormal interval from surgery until the passage of flatus or stool and the tolerance of an oral diet, that should occur within day 4 postoperatively [2]. It can be characterized by nausea, vomiting, loss of appetite, cramps, and abdominal pain and distention, and complicates up to 20% of cesarean delivery [2]. Moreover, it can also be associated with an enhanced length of stay in the hospital [2,3].

Several approaches have emerged in an attempt to hasten the return of gastrointestinal motility after cesarean delivery, including early hydration and ambulation. Chewing gum may offer an efficacious intervention for improving postoperative gastrointestinal function recovery after cesarean delivery [4–15], since it has been already proven to improve gastrointestinal function in non-obstetric abdominal surgery [3]. Considering the number of people who undergo cesarean delivery each year globally, this could have implications for health care costs and recovery [16]. It is therefore essential that benefits and costs are carefully evaluated.

Thus, the aim of this systematic review and meta-analysis of randomized controlled trials (RCTs) was to

examine whether chewing gum after cesarean delivery hastens the return of gastrointestinal function.

## Materials and methods

### Search strategy

This review was performed according to a protocol designed *a priori* and recommended for systematic review [17]. Electronic databases (i.e. MEDLINE, Scopus, ClinicalTrials.gov, EMBASE, Sciencedirect, the Cochrane Library at the CENTRAL Register of Controlled Trials, Scielo) were searched from their inception until November 2016. Search terms used were the following text words: “gum,” “cesarean,” “caesarean,” “delivery,” “labor,” “labour,” “chewing,” “sham feeding,” “general anesthesia,” “morbidity,” “mortality,” “meta-analysis,” “metaanalysis,” “review,” “randomized,” “post-operative,” “clinical trial,” “randomised,” “effectiveness,” “guidelines,” “cost,” “ileus,” and “clinical trial.” No restrictions for language or geographic location were applied. In addition, the reference lists of all identified articles were examined to identify studies not captured by electronic searches. The electronic search and the eligibility of the studies were independently assessed by two authors (AC, GS). Differences were discussed with a third reviewer (VB).

### Study selection

We included all RCTs comparing the use of chewing gum in the immediate postoperative recovery period (i.e. intervention group) with a control group for comparison. Studies in which the gum contained an active therapeutic agent were not included. Studies in which the intervention consisted of gum in combination with another intervention were also excluded. Quasi RCTs (i.e. trials in which allocation was done on the basis of a pseudo-random sequence, e.g. odd/even hospital number or date of birth, alternation) were not included.

### Risk of bias

The risk of bias in each included study was assessed by using the criteria outlined in the *Cochrane Handbook for Systematic Reviews of Interventions*. Seven domains related to risk of bias were assessed in each included trial since there is evidence that these issues are associated with biased estimates of treatment effect: (1) random sequence generation; (2) allocation concealment; (3) blinding of participants and personnel; (4) blinding of outcome assessment; (5) incomplete outcome data; (6) selective reporting;

and (7) other bias. Review authors' judgments were categorized as “low risk,” “high risk,” or “unclear risk” of bias [17].

Two authors (AC, GS) independently assessed inclusion criteria, risk of bias, and data extraction. Disagreements were resolved by discussion with a third reviewer (VB).

### Outcomes

All analyses were done using an intention-to-treat approach, evaluating women according to the treatment group to which they were randomly allocated in the original trials. Primary and secondary outcomes were defined before data extraction.

The primary outcome was time to first flatus in hours after cesarean delivery. The secondary outcomes were time to first bowel sounds in hours, length of hospital stay in days, time to first feces in hours, maternal satisfaction, assessed by self-reported patient satisfaction survey, first feeling of hunger in hours, number of episodes of nausea or vomiting after cesarean, need for additional analgesics or antiemetics, and incidence of paralytic ileus, defined as symptoms or signs of gastrointestinal disturbance such as nausea, vomiting, abdominal cramping, or abdominal distension within the first 72 h after the cesarean delivery or as defined by the original trial.

We planned to assess the primary outcome (i.e. time to first flatus in hours) in subgroup analysis according to the type of cesarean delivery.

### Statistical analysis

The data analysis was completed independently by two authors (AC, GS) using Review Manager v. 5.3 (The Nordic Cochrane Centre, Cochrane Collaboration, 2014, Copenhagen, Denmark). The completed analyses were then compared, and any difference was resolved by discussion with a third reviewer (VB).

Data from each eligible study were extracted with-out modification of original data onto custom-made data collection forms. For continuous outcomes means  $\pm$  standard deviation were extracted and imported into Review Manager v. 5.3.

Meta-analysis was performed using the random effects model of DerSimonian and Laird, to produce summary treatment effects in terms of mean difference (MD) or relative risk (RR) with 95% confidence interval (CI). Heterogeneity was measured using I-squared (Higgins  $I^2$ ).

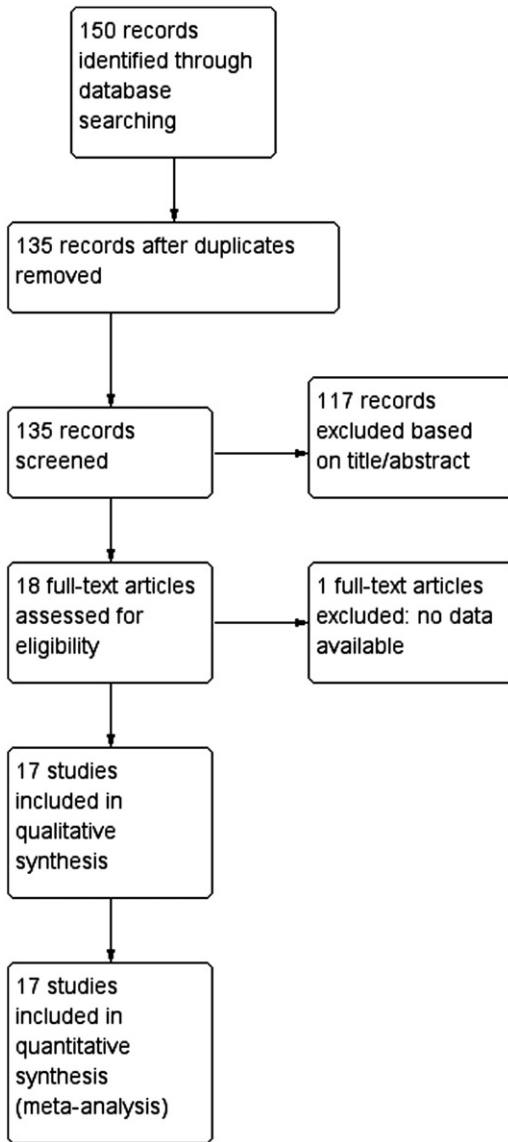
Potential publication biases were assessed statistically by using Begg's and Egger's tests. The meta-

analysis was reported following the Preferred Reporting Item for Systematic Reviews and Meta-analyses (PRISMA) statement [18]. Before data extraction, the review was registered with the PROSPERO International Prospective Register of Systematic Reviews (CRD 42017056270).

**Results**

**Study selection and study characteristics**

The flow of study identification is shown in Figure 1. 18 trials were assessed for eligibility [4–15,19–24]. One was excluded since no data were available [24]. Therefore, 17 trials, including 3041 women, were analyzed



**Figure 1.** Flow diagram of studies identified in the systematic review. Prisma template (Preferred Reporting Item for Systematic Reviews and Meta-analyses).

[4–15,19–23]. No quasi-randomized trials were included. Publication bias, assessed using Begg’s and Egger’s tests, was not significant ( $p = .75$  and  $.84$ , respectively). The quality of the included trials in general was low and most of the trials had high or unclear risk of bias in most of the seven Cochrane domains related to the risk of bias (Figure 2).

|                     | Random sequence generation (selection bias) | Allocation concealment (selection bias) | Blinding of participants and personnel (performance bias) | Blinding of outcome assessment (detection bias): Subjective | Blinding of outcome assessment (detection bias): Objective | Incomplete outcome data (attrition bias) | Selective reporting (reporting bias) | Other bias |
|---------------------|---|---|---|---|--|--|--------------------------------------|------------|
| Ledari 2013         | ?   | ?                                       | +   | +   | ?  | -  | -                                    | ?          |
| Abd-El-Maeboud 2009 | +   | ?                                       | +   | +   | ?  | -  | -                                    | +          |
| Ajuzieogu 2014      | +   | ?                                       | +   | +   | -  | -  | -                                    | ?          |
| Akhlaghi 2008       | ?   | ?                                       | +   | +   | ?  | -  | -                                    | ?          |
| Dehcheshmeh 2011    | ?   | ?                                       | +   | +   | ?  | -  | ?                                    | ?          |
| Garshasbi 2010      | ?   | ?                                       | +   | +   | ?  | -  | ?                                    | ?          |
| Jakkaew 2013        | +   | +                                       | +   | +   | ?  | ?  | +                                    | +          |
| Kafali 2010         | +   | ?                                       | +   | +   | ?  | ?  | -                                    | ?          |
| Ledari 2012         | +   | ?                                       | +   | +   | ?  | ?  | +                                    | +          |
| Liang 2007          | ?   | ?                                       | +   | +   | ?  | -  | +                                    | ?          |
| Lu 2010             | ?   | ?                                       | +   | +   | ?  | -  | ?                                    | ?          |
| Luo 2010            | ?   | ?                                       | +   | +   | ?  | -  | -                                    | ?          |
| Rashad 2013         | ?   | ?                                       | +   | +   | ?  | -  | -                                    | +          |
| Satij 2006          | ?   | ?                                       | +   | +   | ?  | -  | ?                                    | ?          |
| Shang 2010          | +   | +                                       | +   | +   | -  | -  | -                                    | ?          |
| Wang 2011           | ?   | ?                                       | +   | +   | ?  | +  | -                                    | ?          |
| Zamora 2012         | ?   | ?                                       | +   | +   | ?  | -  | -                                    | ?          |

**Figure 2.** Assessment of risk of bias. Summary of risk of bias for each trial; Plus sign: low risk of bias; minus sign: high risk of bias; question mark: unclear risk of bias.

**Table 1.** Characteristics of the included studies.

| Study location      | Simple size | Chewing gum Start    | Chewing gum Times per day        | Chewing gum Duration | Chewing gum End             | Postoperative diet   | Out of bed policy           | Need for intestinal enema                     | Prior cesarean delivery | Prior abdominal surgery | Type of anesthesia                       | Primary outcome <sup>a</sup>   |
|---------------------|-------------|----------------------|----------------------------------|----------------------|-----------------------------|--|-----------------------------|---|-------------------------|-------------------------|--|--|
| Satij [19]          | USA         | 32 (15 versus 17)    | Three                            | 30 min               | First flatus or defecation  | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Spinal anesthesia, or general anesthesia | Evaluate the effect of the gum chewing on the return of bowel function in cesarean delivery patients<br>Time to first flatus, to first bowel movement, complications, tolerance of gum |
| Liang [23]          | China       | 120 (60 versus 60)   | Three                            | 15 min               | First flatus                | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Not reported                             | To investigate the effect of gum chewing as false nutrition on the bowel movement and prevention of post cesarean ileus  |
| Akhalghai [15]      | Iran        | 400 (200 versus 200) | Three                            | 45 min               | First flatus                | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Not reported                             | To test the hypothesis that gum chewing would enhance rapid return of bowel motility after elective cesarean section   |
| Abd-El-Maeboud [14] | Egypt       | 200 (93 versus 107)  | Every 2 h during daytime         | 15 min               | First flatus                | Oral intake of clear fluids and soft foods began after passage of first flatus   | Not reported                | Not reported                                  | Not reported            | 3/93 versus 11/107      | General anesthesia                       | To test the hypothesis that gum chewing would enhance rapid return of bowel motility after elective cesarean section   |
| Luo [21]            | China       | 300 (150 versus 150) | Four                             | 10–15 min            | 3 days after CS             | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Not reported                             | Time to first bowel flatus, first bowel movement, first bowel sound and complications  |
| Lu [22]             | China       | 97 (47 versus 50)    | Every 2 h (at least 6 h chewing) | 30–40 min            | First flatus                | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Not reported                             | Time to first flatus, time to first bowel movement   |
| Kafali [11]         | Turkey      | 150 (74 versus 76)   | Three                            | 1° 15 min; then 1 h  | First flatus                | Oral fluids 6 h after surgery, irrespective of return of bowel sound. Oral food after 24h on detection of bowel sounds on auscultation | Early ambulation encouraged | If no flatus in the first 48h postoperatively | 28/74 versus 22/76      | 22/74 versus 20/76      | Spinal anesthesia, or general anesthesia | To assess the effects of gum chewing on post-operative bowel function after cesarean section   |
| Shang [13]          | China       | 386 (195 versus 191) | Three                            | 30 min               | Defecation of discharge     | Oral intake of clear fluids and soft foods after first flatus  | Early ambulation encouraged | Not reported                                  | 34/195 versus 30/191    | Not reported            | Spinal anesthesia                        | Time to first operative passage of flatus  |
| Garshasbi [12]      | Iran        | 500 (238 versus 262) | Three                            | At least 30 min      | Until start of regular diet | Not reported   | Not reported                | Not reported                                  | Not reported            | Not reported            | Not reported                             | To determine whether gum chewing in the immediate postoperative period facilitated recovery from ileus following cesarean section  |

(continued)

Table 1. Continued

| Study location | Simple size          | Chewing gum Start                  | Chewing gum Times per day | Chewing gum Duration | Chewing gum End            | Postoperative diet  | Out of bed policy | Need for intestinal enema | Prior cesarean delivery | Prior abdominal surgery | Type of anesthesia                       | Primary outcome <sup>a</sup>  |
|----------------|----------------------|------------------------------------|---------------------------|----------------------|----------------------------|---|-------------------|---------------------------|-------------------------|-------------------------|--|---|
| Dehdeshmeh [4] | 120 (60 versus 60)   | From delivery                      | Four                      | -                    | First flatus or defecation | Not reported  | Not reported      | Not reported              | 0/60 versus 0/60        | Not reported            | Spinal anesthesia                        | To assess the effects of chewing of sugar free gum after elective cesarean delivery of return of bowel function in primiparous women        |
| Wang [20]      | 233 (116 versus 117) | From 2 h postoperatively           | Every 2 h during day time | 15 min               | First flatus               | Not reported  | Not reported      | Not reported              | Not reported            | Not reported            | Spinal anesthesia                        | Time to first flatus  |
| Ledari [9]     | 100 (50 versus 50)   | From 6 h postoperatively           | Three                     | At least 1 h         | Discharge                  | Not reported  | Not reported      | Not reported              | 50/50 versus 50/50      | Not reported            | Spinal anesthesia                        | To evaluate the effect of chewing gum on the recovery of bowel function after cesarean section  |
| Zamora [10]    | 53 (18 versus 35)    | From 12 h postoperatively          | -                         | 15 min               | First flatus               | Not reported  | Not reported      | Not reported              | Not reported            | Not reported            | Spinal anesthesia                        | To compare the effects of post-operative gum chewing with traditional feeding on the early return of bowel motility after cesarean delivery |
| Rashad [6]     | 60 (30 versus 30)    | From delivery                      | Three                     | 30 min               | Discharge                  | Not reported  | Not reported      | Not reported              | Not reported            | Not reported            | Spinal anesthesia, or general anesthesia | To identify the effect of sugarless chewing gum on intestinal movement after cesarean section   |
| Ledari [7]     | 60 (30 versus 30)    | From 6 h postoperatively           | Three                     | 1 h                  | Discharge                  | Not reported  | Not reported      | Not reported              | 0/30 versus 0/30        | 0/30 versus 0/30        | Spinal anesthesia                        | To investigate the effect of gum chewing on the return of intestinal function in women with cesarean section                                |
| Jakkaew [8]    | 50 (25 versus 25)    | From delivery                      | Four                      | 30 min               | First flatus               | Oral intake of clear fluids and soft foods after first flatus | Not reported      | Not reported              | 4/25 versus 8/25        | 0/25 versus 1/25        | Spinal anesthesia, or general anesthesia | To evaluate the effect of gum chewing on recovery of bowel function after cesarean section  |
| Ajuzteogu [5]  | 180 (90 versus 90)   | From the first day postoperatively | Three                     | 30 min               | 5 days                     | Not reported  | Not reported      | Not reported              | 0/90 versus 0/90        | 0/90 versus 0/90        | Spinal anesthesia                        | To identify the effect of chewing gum on duration of post operative ileus following cesarean section  |

Data are presented as total number (number in the intervention versus number in the control group).

<sup>a</sup>When the primary outcome was not specifically stated, all listed outcomes were included in this table.



**Table 2.** Inclusion and exclusion criteria.

|                     | Inclusion criteria   | Exclusion criteria   |
|---------------------|--|--|
| Satij [19]          | Women at term undergoing planned CD                            | Preterm, emergency CD  |
| Liang [23]          | Women undergoing CD  | Not reported   |
| Akhlaghi [15]       | Women at term undergoing planned CD                            | Preterm, emergency CD  |
| Abd-El-Maeboud [14] | Women at term undergoing planned CD under general anesthesia   | Preterm, emergency CD, spinal anesthesia, cesarean hysterectomy, prior abdominal surgery   |
| Luo [21]            | Women undergoing CD  | Not reported   |
| Lu [22]             | Women undergoing CD  | Not reported   |
| Kafali [11]         | Women undergoing planned or emergency CD                       | Chronic medical disorders, high risk pregnancy, antepartum hemorrhage, blood transfusion, postoperative admission to intensive care unit |
| Shang [13]          | Women at term undergoing planned or emergency CD               | Preterm, preexisting gastrointestinal disorders, blood transfusion   |
| Garshasbi [12]      | Women planned or emergency CD                                  | Not reported   |
| Dehcheshmeh [4]     | Primiparous women at term undergoing planned CD                | Preterm, emergency CD, multiparous   |
| Wang [20]           | Women undergoing CD  | Not reported   |
| Ledari [9]          | Women at term undergoing planned or emergency CD with prior CD | Preterm, primiparous, prior abdominal surgery  |
| Zamora [10]         | Women at term undergoing planned or emergency CD               | Preterm  |
| Rashad [6]          | Women at term undergoing planned or emergency CD               | Preterm  |
| Ledari [7]          | Primiparous women at term undergoing planned or emergency CD   | Preterm, multiparous, prior abdominal surgery  |
| Jakkaew [8]         | Women at term undergoing planned or emergency CD               | Preterm, cesarean hysterectomy, recent chemotherapy, postoperative admission to intensive care unit                                      |
| Ajuzieogu [5]       | Primiparous women at term undergoing planned CD                | Preterm, emergency CD, multiparous, prior abdominal surgery, diabetic, hypothyroid, women who were on opioids                            |

Data are presented as total number (number in the intervention versus number in the control group).  
CD: cesarean delivery.

Table 1 shows the characteristics of the included clinical trials. All the studies used postoperatively sugar-free gum chewing as intervention. In most of the included trials, chewing gum was given right after or within 2 h of delivery (7/17, 41%, immediately after, and 5/17, 29%, 2 h after), three times a day (10/17, 59%) for 15–30 min each (7/17, 41%, 30 min, and 5/17, 29%, for 15 min) and until the first flatus (10/17, 59%). Women in both groups received routine post-operative diet. Most of the included women were undergoing planned CD at term (Table 2).

Eight studies described in details how the outcomes were assessed [5–9,11,13,15]. In Shang et al. and Rashad et al., every patient was checked for bowel sounds and/or flatus five times a day, and women were asked to tell to study investigators when they passed a bowel movement [6,13]. Ajuzieogu et al., Ledari 2012 et al., Ledari 2013 et al., and Jakkaew et al. reported that a research assistant who was not aware of the gum prescription and groups, visited the patients regularly, every 1 h and recorded the time of the first bowel sounds, passage of flatus, and defecation [5,7–9]. In the other two trials, women were checked five times every day [11,15].

### Synthesis of results

Table 3 shows the primary and secondary outcomes in the overall and in subgroup analyses. In the overall

analysis, the statistical heterogeneity ranged from 0% to 92%, with  $I^2=29%$  for the primary outcome.

Women who were randomized to the chewing gum group had a significantly lower mean of the first flatus time (MD – 6.49 h, 95%CI –8.65 to –4.33; Figure 3), first bowel sounds (Figure 4), less duration of stay, lower time to first fees and to first feeling hunger, less incidence of ileus, less episodes of nausea or vomiting, and significantly higher satisfaction (Table 3). Subgroup analyses concur with the overall analysis (Table 3).

## Discussion

### Main findings

This meta-analysis from 17 RCTs [4–15,19–23], provides evidence that gum chewing after CD is an intervention that enhances early recovery of bowel function. In most of the included trials, chewing gum was given right after delivery three times a day for 30 min each and until the first flatus. Our meta-analysis represented level 1 data and included only RCTs. Test of heterogeneity and sensitivity analyses all point to the efficacy of gum chewing as studied so far. However, the quality of the included trials is low.

### Comparison with existing literature

Our data support earlier findings by two prior Cochrane Reviews. Short et al. in a meta-analysis of 81

**Table 3.** Primary and secondary outcomes in overall and subgroup analyses.

| Overall analysis                              | Number of included studies   | Chewing gum group | Control group  | Number of included women | RR or MD (95%CI)           | I <sup>2</sup> |
|---|------------------------------|-------------------|----------------|--------------------------|----------------------------|----------------|
| First flatus (h)                              | 14 [4-7,9-11,13-15,20-23]    | 23.1 h            | 29.5 h         | 2459                     | -6.49 (-8.65 to -4.33)     | 95%            |
| First bowel sounds (h)                        | 10 [4-7], [9,11,13,14,20,21] | 13.5 h            | 18.3 h         | 1789                     | -4.63 h (-6.20 to -3.05)   | 92%            |
| Duration of stay (days)                       | 7 [4,5,10,11,13-15]          | 2.85 days         | 3.24 days      | 1489                     | -0.39 (-0.78 to -0.18)     | 50%            |
| First feces (h)                               | 5 [5,7,9,13,15]              | 33.9 h            | 43.2 h         | 2076                     | -9.57 h (-10.28 to -8.87)  | 0%             |
| Satisfaction (points)                         | 2 [5-8]                      | 8.25 points       | 6.8 points     | 230                      | 1.99 points (1.70 to 2.29) | 60%            |
| First feeling of hunger (h)                   | 3 [7-9]                      | 12.7 h            | 15.2 hours     | 210                      | -2.89 h (-4.93 to -0.85)   | 0%             |
| Nausea or vomiting                            | 4 [8,10,11,14]               | 5/210 (2.4%)      | 16/243 (6.6%)  | 453                      | 0.33 (0.12 to 0.87)        | 69%            |
| Need for additional analgesics or antiemetics | 3 [11,13,14]                 | 15/362 (4.1%)     | 25/364 (6.9%)  | 726                      | 0.50 (0.12 to 2.13)        | 39%            |
| Ileus <sup>a</sup>                            | 4 [10,12-14]                 | 28/544 (4.6%)     | 68/595 (11.4%) | 1139                     | 0.39 (0.19 to 0.80)        |                |
| First cesarean delivery                       |                              |                   |                |                          |                            |                |
| First flatus (h)                              | 4 [4,5,20,21]                | 28.6 h            | 36.4 h         | 833                      | -7.71 (-11.89 to -3.52)    | 97%            |
| Repeated cesarean delivery                    |                              |                   |                |                          |                            |                |
| First flatus (h)                              | 1 [14]                       | 93.2 h            | 107.2 h        | 200                      | -6.50 (-8.14 to -4.86)     | Not applicable |
| Planned cesarean delivery                     |                              |                   |                |                          |                            |                |
| First flatus (h)                              | 7 [4,5,8,9,14,15,20]         | 24.9 h            | 29.7 h         | 1293                     | -4.83 (-8.06 to 2.58)      | 90%            |
| Emergent cesarean delivery                    |                              |                   |                |                          |                            |                |
| First flatus (h)                              | 3 [4,10,13]                  | 19.8 h            | 25.5 h         | 499                      | -5.89 (-7.13 to -4.65)     | 3%             |

Data are presented as number (percentage) or as mean difference  $\pm$  standard deviation. Boldface data, statistically significant.

RR: relative risk; MD: mean difference; CI: confidence interval.

<sup>a</sup>Definitions of ileus were given in only two of the four RCTs which reported this outcome: "the delayed return of physiological coordinated bowel motility"[11]; "a group of manifestations persisting longer than 24h or requiring naso-gastric tube placement" [14].

studies, including 9072 participants who underwent abdominal surgery, found some evidence for the benefit of postoperative chewing gum in improving recovery of gastrointestinal function after obstetric and non-obstetric abdominal surgery [25]. Pereira Gomes Morais et al. showed a beneficial effect of chewing gum in women undergoing cesarean delivery [26]. However, both of these meta-analyses did not include all currently available RCTs on cesarean delivery. In 2014, Craciunas et al. also showed efficacy of chewing gum in reducing the incidence of postoperative ileus in 1462 patients (seven RCTs) who underwent cesarean delivery [27]. Conversely, Huang et al. in a meta-analysis of five RCTs found no benefit of postoperative gum chewing in women who underwent cesarean delivery [28]. Another review by Wen et al. was recently published, including 10 trials ( $n = 1659$  women). The authors concluded that gum chewing hastens the intestinal function recovery after cesarean delivery offering a safe and inexpensive option for postoperative care [29].

### Strengths and limitations

Our study has several strengths. Intent-to-treat analysis was used. In addition, publication bias was not apparent by statistical analysis. Limitations of our study are mostly inherent to the limitations of the included studies. The quality of evidence as well as the quality of the included trials was low. There were no reports of adverse effects related to gum chewing in the 17 included trials. However, since none of them specifically stated that this was a pre-specified outcome in their protocols or methods, we cannot be sure that gum chewing in the postoperative period of cesarean delivery is devoid of adverse effects. None of the trials assessed or reported adherence to gum chewing. None of the included trial was double blind. We used a random effect model in all analyses given the high statistical and clinical heterogeneity within the trials.

### Conclusions

In summary, this meta-analysis showed that gum chewing reduced the time to first passage of flatus and reduced the rate of ileus when given right after cesarean delivery, three times a day for 30 min each and until the first flatus. As simple, generally inexpensive intervention, providers should consider implementing cesarean delivery postoperative care with gum chewing.

Future large, better designed, randomized trials will help to increase the quality of the evidence for this intervention. Future trials should also establish the



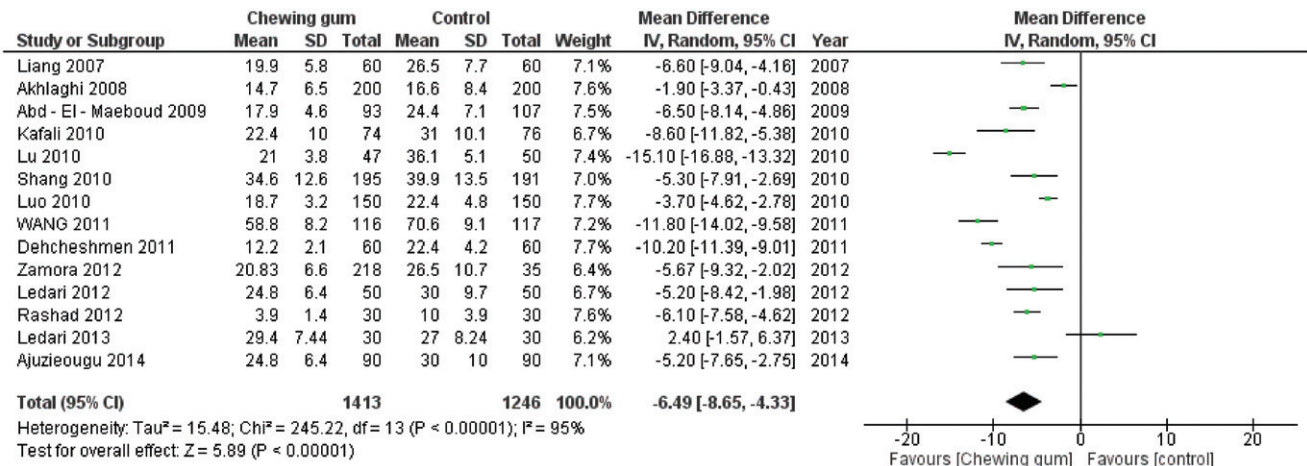


Figure 3. Forest plot for the mean of time to the first flatus in hours.

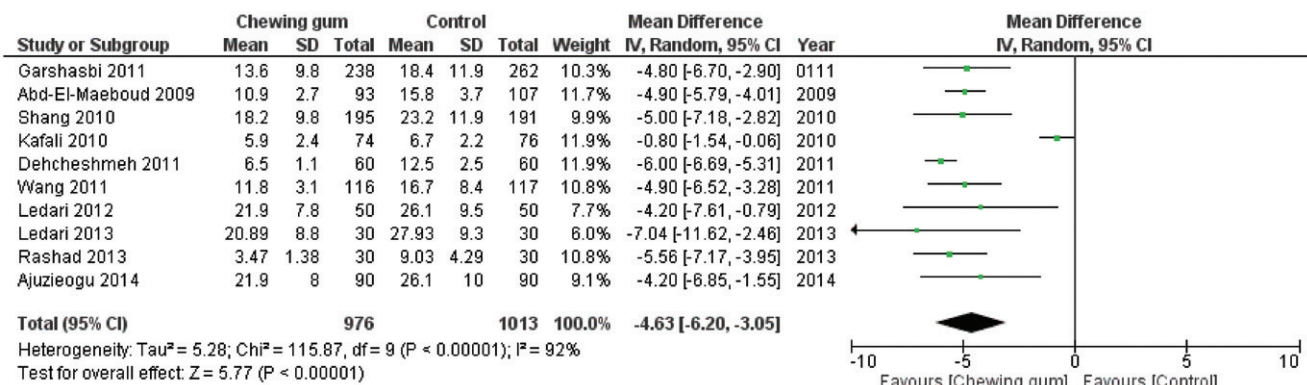


Figure 4. Forest plot for the mean of time to the first bowel sounds in hours.

optimal regimen of gum chewing (e.g. initiation, number, and duration of sessions per day) to enhance bowel function recovery after cesarean delivery, and assess how other beneficial interventions, such as early feeding [30,31], may give additional benefits during postpartum care for women who undergo gum chewing.

## Acknowledgements

We acknowledge Yadollah Omidi MD, Mohammad Rafi MD, and Jaleh Barar MD to provide assistance in the translation of the manuscripts.

## Disclosure statement

The authors declare that they have nothing to disclose.

## ORCID

Gabriele Saccone <http://orcid.org/0000-0003-0078-2113>  
Daniele Di Mascio <http://orcid.org/0000-0002-6560-3393>  
Claudia Caissutti <http://orcid.org/0000-0002-6535-4497>  
Vincenzo Berghella <http://orcid.org/0000-0003-2854-0239>

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