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Long-term Impact of Mode of Delivery on Stress Urinary Incontinence and Urgency Urinary Incontinence: A Systematic Review and Meta-analysis

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Abstract

Context: Stress urinary incontinence (SUI) and urgency urinary incontinence (UUI) are associated with physical and psychological morbidity, and large societal costs. The long-term effects of delivery modes on each kind of incontinence remain uncertain.

Objective: To investigate the long-term impact of delivery mode on SUI and UUI.

Evidence acquisition: We searched Medline, Scopus, CINAHL, and relevant major conference abstracts up to October 31, 2014, including any observational study with adjusted analyses or any randomized trial addressing the association between delivery mode and SUI or UUI ≥1 yr after delivery. Two reviewers extracted data, including incidence/prevalence of SUI and UUI by delivery modes, and assessed risk of bias.

Evidence synthesis: Pooled estimates from 15 eligible studies demonstrated an increased risk of SUI after vaginal delivery versus cesarean section (adjusted odds ratio [aOR]: 1.85; 95% confidence interval [CI], 1.56–2.19; I² = 57%; risk difference: 8.2%).

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1. Introduction

Stress urinary incontinence (SUI) is defined as the involuntary loss of urine on effort or physical exertion, or on sneezing or coughing. Urgency urinary incontinence (UUI) is defined as involuntary loss of urine associated with a sudden and compelling desire to pass urine [1]. Both from the population perspective and from an individual perspective, SUI and UUI are the most burdensome and bothersome of all urinary symptoms in women [2]. SUI and UUI are associated with substantial physical and psychological morbidity, and large societal costs [3,4]. Established risk factors for urinary incontinence include age and body mass index (BMI) [5]: the prevalence of these conditions is therefore likely to increase with future demographic changes.

Although advances in treatment during the last two decades have decreased morbidity, primary prevention of long-term SUI and UUI remains highly desirable. Mode of delivery is one potentially modifiable risk factor. Vaginal childbirth is known to have major impacts on the pelvic floor, weakening bladder neck support [6] and compromising innervation [7]. Cesarean delivery, particularly prelabor cesarean, is believed to offer substantial protection against such pelvic floor trauma; in contrast, assisted vaginal delivery, with vacuum or forceps, is believed to carry increased risks of trauma. The World Health Organization statement on caesarean section rates recommends that the ideal rate for cesarean sections is between 10% and 15% [8]. Observed rates, however, vary widely between countries. Although rates are <10% in most low-income countries [9], middle- and high-income countries have seen substantial increases since the 1970s. In 2011 rates were 24% in the United Kingdom [10], 33% in the United States [11], and 54% in Brazil [12]. The increasing use of cesarean section has substantial negative public health consequences, including peripartum infection, bleeding, and thrombosis, and it has an impact on future pregnancies [8]. Any positive consequences from the increased use of cesarean have not been well quantified.

An extensive body of evidence from the first year after delivery demonstrates that in this initial postpartum period, rates of SUI are higher in women delivering vaginally than those delivering by cesarean [13,14]. The long-term effects of delivery mode, however, are more important to patients than transient postpartum incontinence. Therefore to reach a better understanding of the association between individual delivery modes and the long-term risk of SUI and UUI, we conducted a systematic review and meta-analysis.

Conclusions: Compared with cesarean section, vaginal delivery is associated with an almost twofold increase in the risk of long-term SUI, with an absolute increase of 8%, and an effect that is largest in younger women. There is also an increased risk of UUI, with an absolute increase of approximately 3%.

2. Evidence acquisition

We registered the protocol (PROSPERO 2013: CRD42013006213) and followed the Preferred Reporting Items for Systematic Reviews and Meta-analysis guidance [15].

2.1. Search strategy

An experienced research librarian (M.A.) collaborated in planning the search strategy, performed on October 31, 2014, in Medline (1946 to present), Scopus (1995 to present), and CINAHL (1960 to present). We also searched abstracts published from the annual meetings of the International Continence Society and the International Urogynecological Association (1999–2014). The searches were conducted without language restrictions and adapted for each electronic database. The details of searches are available in Supplement 1.

2.2. Study selection

We included any randomized trial, cross-sectional, or cohort study that recorded the delivery mode as well as SUI and/or UUI outcome beyond 1 yr after delivery among primip- and multiparous women and provided an analysis comparing at least two delivery modes with SUI and/or UUI. Because previous studies have established prognostic factors for SUI and UUI [16–20], we included only cross-sectional or cohort studies with an analysis that adjusted/matched for at least one of the following: age [3,16], BMI [17,19,20], or parity [18].

Because SUI and UUI have different etiologies [21–23], we excluded studies that reported on incontinence but did not report specifically on either SUI or UUI (eg, “any urinary incontinence”). We also excluded studies that only reported surrogate measures, such as urodynamic testing, cough stress test, or treatment rates. Reasoning that small studies...
are likely to be published only if they show anomalous results, we excluded studies with <100 participants. We accepted the definition of SUI and UUI used in each study, recognizing there would be heterogeneity in definitions, provided definitions captured the women’s own perception of incontinence.

2.3. Data extraction and risk of bias assessment

We used standard methods for screening and data extraction of systematic reviews (details in Supplement 2). For the risk of bias assessments, we evaluated each study according to six criteria: sampling and representativeness of population, assessment of the exposure, assessment of the outcome, presence of the outcome at the start of study, adjustment for confounding, and missing data (Supplementary Table 1). For each criterion, we judged studies to have either a high or low risk of bias. Studies with a high risk of bias for two or more criteria were classified as high risk of bias overall.

2.4. Data synthesis and analysis

For our primary analyses, we examined the association between mode of delivery and SUI or UUI. We calculated pooled estimates of adjusted estimates using the DerSimonian-Laird random-effects inverse variance method, and the I² statistic [24] and Cochran’s Q as indices of heterogeneity. We used prespecified hypotheses to examine heterogeneity using metaregression analysis weighted by the inverse of the variance in a random-effects model. We examined the following variables as potential sources of heterogeneity: age (as continuous variable), parity (as continuous variable), risk of bias (low vs high), composition of vaginal delivery group (including women delivering only ever by vaginal routes vs including women with both vaginal and cesarean deliveries), and the case definition of SUI or UUI (inclusive mild incontinence definitions vs restrictive severe definitions). We had prespecified hypotheses that effect sizes would be smaller for samples of older women, samples with higher parity, low risk of bias studies, mixed modes of delivery, and studies using a lower threshold (less severe symptoms) in their case definitions. We conducted a single sensitivity analysis including a randomized trial omitted from the primary analysis because of large crossover and concerns regarding applicability (all breech presentations).

To calculate the absolute risk increase of moderate or severe SUI or UUI with vaginal birth, we estimated the absolute risk of SUI or UUI after cesarean section using two large population-based studies [25,26]: 12.2% for moderate or severe SUI and 10.1% for moderate or severe UUI after any cesarean section, and 5.0% for SUI after elective cesarean section, and then used the odds ratio (OR) to calculate the absolute risk increase with vaginal delivery [27].

When primary papers had missing confidence interval (CI) information (i.e., providing ORs and p values but not CIs), we calculated the CIs (further information on data analysis in Supplement 3). Meta-analyses were performed using metan [28] and metareg in Stata v12.1 (StataCorp, College Station, TX, USA). We used the Harbord test to detect publication bias.

3. Evidence synthesis

3.1. Literature search and study characteristics

Our search yielded 3487 potentially relevant reports. After screening titles and abstracts, we retrieved 179 reports for full-text screening, of which 18 proved eligible. Of these 18, we did not include two studies in meta-analyses: a large-scale multicenter randomized trial of planned cesarean versus planned vaginal delivery due to low generalizability because it only included women with fetuses with breech presentations [29–31]; and a small cohort study [32] that combined spontaneous vaginal delivery and cesarean during labor and compared with vacuum or forceps. Of 16 studies included [5,25,26,33–47], 8 addressed the impact of delivery mode on SUI, 7 on both SUI and UUI, and 1 on UUI (Tables 1 and 2; Fig. 1). In these studies we identified 11 different comparisons between delivery modes assessing risk of SUI and 5 different comparisons assessing risk of UUI (Supplementary Fig. 1). The most common comparison was any vaginal delivery (including studies with spontaneous vaginal delivery only, vaginal delivery only, or at least one vaginal delivery) versus cesarean section (15 studies with 45 659 women for SUI and 8 studies with 49 623 women for UUI) for both SUI and UUI, followed by instrumental delivery versus spontaneous vaginal delivery for SUI (4 studies with 7417 women) (Figs. 2–4; Supplementary Fig. 1 and 2).

Table 1 provides a description of the 16 studies. Table 2 provides authors’ definitions of SUI and UUI. Three of the studies included only primiparous women [26,37,46]. SUI prevalence estimates varied from 9% to 68%; UUI from 8% to 27%. Six (37.5%) of the 16 authors confirmed the accuracy of

Table 1. Characteristics of included studies

<table>
<thead>
<tr>
<th>Study</th>
<th>Mode of Delivery</th>
<th>Sample Size</th>
<th>Source</th>
<th>Case Definition</th>
<th>Risk of Bias</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study 1</td>
<td>Vaginal (n)</td>
<td>1000</td>
<td>Hospital</td>
<td>SUI, UUI</td>
<td>Low</td>
</tr>
<tr>
<td>Study 2</td>
<td>Cesarean</td>
<td>2000</td>
<td>Clinic</td>
<td>Moderate SUI</td>
<td>High</td>
</tr>
<tr>
<td>Study 3</td>
<td>Spontaneous Vaginal</td>
<td>1500</td>
<td>Community</td>
<td>UUI</td>
<td>Moderate</td>
</tr>
</tbody>
</table>

Fig. 1 – Flowchart outlining the literature search and article evaluation process.
SUI = stress urinary incontinence; UUI = urgency urinary incontinence.
<table>
<thead>
<tr>
<th>Study</th>
<th>Analyzed participants, n</th>
<th>Year(s) data collected</th>
<th>Sampling frame</th>
<th>Type of survey</th>
<th>Specific inclusion criteria</th>
<th>Age, yr, mean (range)</th>
<th>Follow-up time postpartum</th>
</tr>
</thead>
<tbody>
<tr>
<td>El-Azab et al [36]</td>
<td>1652</td>
<td>Not reported</td>
<td>Living in upper Egypt</td>
<td>In person</td>
<td>Women ≥20 yr of age</td>
<td>Not reported</td>
<td>Unclear</td>
</tr>
<tr>
<td>Fritel et al [37]</td>
<td>307</td>
<td>2000</td>
<td>2 hospitals in France</td>
<td>Mailed questionnaire</td>
<td>Primiparous women, with singleton, vertex, nonpremature birth 1996</td>
<td>33 (21–51)</td>
<td>4 yr</td>
</tr>
<tr>
<td>Fritel et al [38]</td>
<td>2625</td>
<td>1990–1996</td>
<td>Employed by the French national power company</td>
<td>Mailed questionnaire</td>
<td>Women with first birth (singleton, nonpremature) 5–10 yr before enrollment</td>
<td>40 (23–54)</td>
<td>5–10 yr</td>
</tr>
<tr>
<td>Goldberg et al [39]</td>
<td>733</td>
<td>2001</td>
<td>Participants of the National Organization of Mothers of Twins Clubs, USA</td>
<td>Given questionnaire</td>
<td>Mothers of multiples</td>
<td>37 (22–75)</td>
<td>Unclear</td>
</tr>
<tr>
<td>Handa et al [33,34]</td>
<td>1011/449</td>
<td>2008–2013</td>
<td>1 US hospital</td>
<td>Given questionnaire</td>
<td>Women with first birth (singleton, nonpremature) 5–10 yr before enrollment</td>
<td>38 (not reported)</td>
<td>5–12 yr</td>
</tr>
<tr>
<td>Handa et al [40]</td>
<td>1481</td>
<td>2008–2013</td>
<td>1 US hospital</td>
<td>Given questionnaire</td>
<td>Women with singleton, nonpremature delivery</td>
<td>40 (unclear)</td>
<td>Unclear</td>
</tr>
<tr>
<td>Kepenekci et al [41]</td>
<td>4002</td>
<td>2005–2007</td>
<td>6 different family medicine centers in Turkey</td>
<td>Questionnaire administered by trained staff</td>
<td>Women accompanying or supporting a patient</td>
<td>41 (15–86)</td>
<td>Unclear</td>
</tr>
<tr>
<td>Song et al [45]</td>
<td>5392</td>
<td>2002</td>
<td>Population-based study in one community in China</td>
<td>Mailed questionnaire</td>
<td>Women attending obstetrics/gynecology clinic</td>
<td>40 (18–87)</td>
<td>Unclear</td>
</tr>
<tr>
<td>Zhu et al [47]</td>
<td>19 024</td>
<td>2006</td>
<td>Population-based study in six provinces in China</td>
<td>Questionnaire administered by a doctor</td>
<td>Women attending obstetrics/gynecology clinic</td>
<td>45 (20–99)</td>
<td>Unclear</td>
</tr>
</tbody>
</table>

SUI = stress urinary incontinence; UUI = urgency urinary incontinence.

Of the 16 studies included, 8 addressed the impact of delivery mode on SUI [5,25,33,34,37–40,45,46], 7 on both SUI and UUI [26,35,36,41–44], and 1 on UUI [47].

* From the same study, two eligible articles [33,34] using the baseline data and one article [40] using the prospective data have been published.

* Median age.

* Median age at study enrollment.

* Pregnant women, 6 mo postpartum, and women with cognitive disorders or neurologic diseases, a history of previous gastrointestinal, anorectal, or gynecologic surgery, or staying at nursing homes were excluded.
Table 2 – Stress urinary incontinence and urgency urinary incontinence assessment in the 16 eligible studies

<table>
<thead>
<tr>
<th>Study</th>
<th>Validated questionnaire used</th>
<th>Type of urinary incontinence assessed</th>
<th>Specific SUI question used</th>
<th>Specific UUI question used</th>
<th>Response options/Definition of normal-abnormal</th>
</tr>
</thead>
<tbody>
<tr>
<td>Altman et al [35]</td>
<td>Modeled CCI Score</td>
<td>Stress and urgency</td>
<td>Do you experience involuntary loss of urine at physical activities?</td>
<td>Do you experience sudden urges to void urine that are followed by involuntary loss of urine?</td>
<td>No/Less than once/week-More than once/week-Daily</td>
</tr>
<tr>
<td>El-Azab et al [36]</td>
<td>UDI-6, Arabic version</td>
<td>Stress and urgency</td>
<td>Do you experience, and, if so, how much are you bothered by urine leakage related to physical activity, coughing, or sneezing?</td>
<td>Do you experience, and, if so, how much are you bothered by urine leakage related to the feeling of urgency?</td>
<td>Not at all/Slightly-Moderately-Greatly</td>
</tr>
<tr>
<td>Fritel et al [37]</td>
<td>BFLUTS, French version</td>
<td>Stress and urgency</td>
<td>Does urine leak, when you are physically active, cough, or sneeze?</td>
<td></td>
<td>Never/Occasionally-Sometimes-Occasionally-All the time</td>
</tr>
<tr>
<td>Fritel et al [38]</td>
<td>BFLUTS, French version</td>
<td>Stress</td>
<td>Does urine leak, when you are physically active, cough, or sneeze?</td>
<td></td>
<td>Never/Occasionally-Sometimes-Occasionally-All the time</td>
</tr>
<tr>
<td>Goldberg et al [39]</td>
<td>PFDI and IIQ, UDI</td>
<td>Stress and urgency</td>
<td>Do you leak urine with coughing, straining, laughing, physical activity, or exercise?</td>
<td></td>
<td>Not at all/Slightly-Moderately-Greatly</td>
</tr>
<tr>
<td>Goldberg et al [5]</td>
<td>PFDI and IIQ</td>
<td>Stress and urgency</td>
<td>Do you leak urine with coughing, straining, laughing, physical activity, or exercise?</td>
<td></td>
<td>Not at all/Slightly-Moderately-Greatly</td>
</tr>
<tr>
<td>Gyhagen et al [26]</td>
<td>Sandvik questionnaire</td>
<td>Stress and urgency</td>
<td>Do you have involuntary loss of urine in connection with coughing, sneezing, laughing, lifting heavy items?</td>
<td>Do you have involuntary loss of urine in connection with sudden and strong urge to void?</td>
<td>Not at all/Slight-Moderate-Severity</td>
</tr>
<tr>
<td>Handa et al [33,34,40]</td>
<td>EPIQ</td>
<td>Stress and urgency</td>
<td>Do you experience urine leakage related to activity, coughing, or sneezing?</td>
<td></td>
<td>No/Yes</td>
</tr>
<tr>
<td>Kepenekci et al [41]</td>
<td>UDI</td>
<td>Stress and urgency</td>
<td>Do you experience, and if so, how much are you bothered by leakage related to physical activity, coughing, or sneezing?</td>
<td>Do you experience, and if so, how much are you bothered by leakage related to feeling of urgency?</td>
<td>Not at all/Slightly-Moderately-Greatly</td>
</tr>
<tr>
<td>Lukacz et al [25]</td>
<td>EPIQ</td>
<td>Stress and urgency</td>
<td>Do you experience urine leakage related to activity, coughing, or sneezing?</td>
<td></td>
<td>No/Yes</td>
</tr>
<tr>
<td>Manonai et al [42]</td>
<td>Stress and urgency</td>
<td>Not reported</td>
<td></td>
<td></td>
<td>Not reported</td>
</tr>
<tr>
<td>Rortveit et al [43]</td>
<td>Sandvik questionnaire</td>
<td>Stress and urgency</td>
<td>Do you leak when coughing, sneezing, laughing, lifting heavy items?</td>
<td>Is leakage accompanied with a sudden and strong urge to void?</td>
<td>Not at all/Slight-Moderate-Severity</td>
</tr>
<tr>
<td>Singh et al [44]</td>
<td>Stress and urgency</td>
<td>Screening question: “Do you have complaint of urinary leakage?” was used. Subsequently, specific incontinence questions were used but remain unclear</td>
<td>Screening question: “Do you have complaint of urinary leakage?” was used. Subsequently, specific incontinence questions were used but remain unclear</td>
<td></td>
<td>No/Yes</td>
</tr>
<tr>
<td>Song et al [45]</td>
<td>Stress</td>
<td>Do you experience urine leakage related to activity, coughing, or sneezing?</td>
<td></td>
<td></td>
<td>No/Yes/ Severity: 0–5</td>
</tr>
<tr>
<td>Yang et al [46]</td>
<td>Modified from BFLUTS, Chinese version</td>
<td>Stress and urgency</td>
<td>Do you experience urine leakage related to activity, coughing, or sneezing?</td>
<td></td>
<td>Never/Occasionally-Sometimes-Occasionally-All the time</td>
</tr>
<tr>
<td>Zhu et al [47]</td>
<td>BFLUTS, Chinese version</td>
<td>Urgency and stress</td>
<td>Does urine leak before you can get to the toilet?</td>
<td></td>
<td>Never/Occasionally-Sometimes-Occasionally-All the time</td>
</tr>
</tbody>
</table>

BFLUTS = Bristol Female Lower Urinary Tract Symptoms Questionnaire; CCI Score = Cleveland Clinic Incontinence Score; EPIQ = Epidemiology of Prolapse and Incontinence Questionnaire; International Consultation on Incontinence Questionnaire-Female Lower Urinary Tract Symptoms; IIQ = Incontinence Impact Questionnaire; PFDI = Pelvic Floor Distress Inventory; SUI = stress urinary incontinence; UDI = Urinary Distress Inventory; UUI = urgency urinary incontinence.

a Cut-off point (threshold) used for normal versus abnormal symptom occurrence. Response options classified as abnormal are shown in boldface type. All studies used the same response options for both SUI and UUI.

b Specific question not provided in publication.

c Information provided by the author, not in the published reference.

d Information regarding UUI was also measured, but effect estimates were not reported.

* Information regarding SUI was also measured, but effect estimates were not reported.
our consensus data extraction [25,26,34,37,42,47]; four (25%) corrected some errors or provided additional information [37,38,41,43]; six (37.5%) were unable to assist with our requests for data checks and clarifications [5,35,39,44–46].

We identified only two prospective studies [30,40]. One randomized trial [29–31] included only women with fetuses presenting in breech position. In this study there was a significant crossover between groups: for women randomized to planned cesarean section group, 941 (90.4%) delivered by cesarean section, but for those randomized to the planned vaginal delivery group, only 591 (56.7%) were delivered vaginally. The study assessed rates of SUI for the “previous 7 d” at 3 mo, and for the “previous 3–6 mo” at 2 yr, which likely explains the observed higher rates of SUI at 2 yr compared with 3 mo. At 3 mo postpartum, the authors noted a lower rate of SUI among women assigned to

Fig. 2 – Forest plot showing risk of stress urinary incontinence between vaginal delivery and cesarean section. CI = confidence interval; OR = odds ratio.

Fig. 3 – Forest plot showing risk of urgency urinary incontinence between vaginal delivery and cesarean section. CI = confidence interval; OR = odds ratio.

Fig. 4 – Relative and absolute risk of stress urinary incontinence between vaginal delivery and cesarean section by age group. CI = confidence interval; CS = cesarean section; OR = odds ratio; VD = vaginal delivery.
planned cesarean section compared with planned vaginal delivery (4.5% vs 7.3%; \( p = 0.02 \)). At 2 yr, the difference between groups was no longer significant (17.8% vs 21.8%; \( p = 0.14 \)). Inclusion of this trial in a sensitivity analysis did not materially change the results (adjusted odds ratio [aOR]: 1.78; 95% CI, 1.51–2.10; risk difference: 7.6%).

The other prospective study [40] reported longitudinal changes of pelvic floor disorders for parous women with and without a history of vaginal delivery. This study recruited women at 5–10 yr after a first birth to annual follow-up over 5 yr. At the baseline assessment (no difference in maternal age or in time from first delivery to study enrollment between groups), the prevalence of SUI was 54% in vaginal versus 20% in cesarean delivery; for UUI, prevalence was 17% in vaginal and 7% in cesarean delivery. We used cross-sectional analyses of these baseline data [33,34] in our meta-analyses. The longitudinal data collection demonstrated that differences between vaginal and cesarean section diminished over time from delivery.

### 3.2. Risk of bias

In all studies, women undergoing different delivery modes were drawn from the same database, over the same time frame, and we judged the assessment of mode of delivery exposure as accurate. Eight studies nevertheless met criteria for high risk of bias (Supplement 4 and Supplementary Fig. 3). Thirteen studies (81%) had little missing data or used self-reported validated questionnaires or another method with demonstrated validity. Twelve studies (75%) adjusted/matched for all the most important confounders (age, BMI, parity). No study collected information regarding SUI or UUI before delivery.

### 3.3. Impact of delivery mode on stress and urgency urinary incontinence

In the pooled analysis (15 studies: 7 low and 8 high risk of bias), the odds of reporting SUI was almost double after any vaginal delivery (spontaneous or assisted) (aOR: 1.85; 95% CI, 1.56–2.19; heterogeneity: \( p = 0.003; I^2 = 57\% \); risk difference: 8.2%) compared with any cesarean section. All studies but one [36] suggested an increased risk, and in 12 of 15 studies the CI excluded no effect (Fig. 2). When comparing elective cesarean with the decision made before the onset of labor only, two (both high risk of bias) studies [20,28] reported a risk of SUI over three times higher with vaginal delivery (aOR: 3.53; 95% CI, 2.55–4.90; heterogeneity: \( p = 0.84; I^2 = 0\% \); risk difference: 10.7%).

The pooled analysis (four studies, two low and two high risk of bias) demonstrated no significant difference in SUI between instrumental delivery, including vacuum and forceps, and spontaneous vaginal delivery (aOR: 1.11; 95% CI, 0.84–1.45; heterogeneity: \( p = 0.11; I^2 = 50\% \)) (Supplementary Fig. 2). The results were similar when comparing vacuum with spontaneous vaginal delivery (two studies, both high risk of bias; aOR: 1.10; 95% CI, 0.80–1.51; heterogeneity: \( p = 0.60; I^2 = 0\% \)) [31,38] or forceps to spontaneous vaginal delivery (three studies, two low and one high risk of bias; aOR: 1.16; 95% CI, 0.71–1.89; heterogeneity: \( p = 0.06; I^2 = 65\% \)) [5,34,38].

The pooled analysis (eight studies, three low and five high risk of bias) demonstrated that the risk of UUI was modestly increased after vaginal delivery when compared with cesarean delivery (aOR: 1.30; 95% CI, 1.02–1.65; heterogeneity: \( p = 0.14; I^2 = 37\% \); risk difference: 2.6%) (Fig. 3). No study reported the impact of elective cesarean only versus vaginal delivery on UUI.

One low risk of bias study [48] compared vacuum deliveries with a combination of spontaneous vaginal deliveries and forceps deliveries (aOR: 0.8; 95% CI, 0.6–1.0 for SUI; and aOR: 1.2; 95% CI, 0.7–2.2 for UUI), and forceps deliveries to a combination of spontaneous vaginal deliveries and vacuum deliveries (aOR: 0.9; 95% CI, 0.7–1.1 for SUI; and aOR: 0.8; 95% CI, 0.4–1.5 for UUI). One high risk of bias study [41] compared the risk of UUI between vacuum and spontaneous vaginal delivery (aOR: 1.03; 95% CI, 0.64–1.67 in our reanalysis adjusted for age and parity using data provided by authors).

One high risk of bias study [33] reported no significant differences between cesarean in the first stage of labor versus elective cesarean (aOR: 0.88; 95% CI, 0.40–1.91) or cesarean in the second stage of labor versus elective cesarean (aOR: 1.30; 95% CI, 0.57–2.95) but had more than a fourfold risk of SUI (aOR: 4.45; 95% CI, 2.14–9.27) in instrumental delivery versus elective cesarean.

### 3.4. Variability across studies

Mean age and parity of study populations, case definition of SUI and UUI, definition of vaginal delivery groups, risk of bias, and survey methods varied across studies (Tables 1 and 2). In the 15 studies addressing the association between vaginal versus cesarean delivery and SUI, in unpublished metaregressions we found that the mean sample age at ascertainment of outcome \( (p = 0.005) \) modified the effect of delivery mode on SUI (older age, smaller effect). Other hypothesized effect modifiers were nonsignificant in unpublished metaregression. Based on the results of the metaregression, we were able to calculate the estimated OR for the association between delivery mode and SUI, at various levels of mean sample age (Fig. 4). Ascertained at age 30, the OR associated with vaginal delivery was 2.51 (95% CI, 1.96–3.21); ascertained at age 60, the OR was 1.29 (95% CI, 0.97–1.72).

In addressing the risk of UUI between vaginal delivery and cesarean, the small number of studies limited the power of the metaregressions. We did not identify statistically significant sources of heterogeneity in effect size for these meta-analyses. There was no evidence of publication bias, either on visual inspection of funnel plots (Supplementary Fig. 4) or when applying the Harbord test.

### 3.5. Discussion

This systematic review examining the association between delivery mode and the presence of SUI and UUI \( > 1 \) yr after delivery identified 11 different comparisons between
delivery modes assessing risk of SUI and 5 different comparisons assessing risk of UUI. Meta-analysis of data from 15 cross-sectional studies demonstrated an almost twofold increase in the risk of developing long-term SUI, an absolute increase of approximately 8% in moderate or severe SUI when comparing any vaginal delivery with cesarean section. The impact was age dependent and decreased in cohorts of older women. Ascertained at age 30, the OR associated with vaginal delivery versus cesarean was 2.51; ascertained at age 60, the OR was 1.29. This difference in gradient reflects the increasing incidence of incontinence for reasons other than mode of delivery as women age.

When SUI was compared with specifically elective cesarean, the risk was over three times higher, an absolute increase of >10%. Meta-analysis also showed a small increased risk of UUI after vaginal delivery compared with cesarean, an absolute increase of approximately 3%. Results showed no difference in the risk of SUI when comparing instrumental vaginal delivery and spontaneous vaginal delivery.

Aside from one randomized trial [29–31] including only breech presentations, only one optimally adjusted longitudinal study addressed the question of interest [40]. In this study, symptoms related to SUI and UUI were more common and of greater severity after vaginal than cesarean birth. Consistent with results in our metagression of age on effect size, SUI symptom differences between these two groups decreased with increasing time from childbirth.

3.6. Strengths and limitations

The strengths of our study include the comprehensive search without language restrictions, the duplicate assessment of eligibility and data abstraction, the appraisal of risk of bias, and the contribution of authors of primary studies to confirmation and clarification of our data abstraction. We used appropriate statistical methods to generate pooled estimates and explored possible sources of heterogeneity, demonstrating that apparent effects of spontaneous vaginal delivery versus cesarean section on SUI decreased with increasing age of assessment of SUI. We have also separately quantified the larger benefit associated with elective prelabor cesarean, compared with any cesarean section (either before or during labor). Finally, we not only estimated relative effects but also provided absolute estimates.

The limitations of our review are largely the weaknesses of the eligible studies. Investigators have conducted only one randomized trial [30] and only one prospective cohort study [40] examining the impact of delivery mode on SUI and UUI. Although there were numerous comparisons between delivery modes assessing the risk of SUI and UUI, it was frequently impossible to compare data quantitatively. In particular, most primary studies combined all cesarean sections, irrespective of timing. We were able to conduct analyses specifically for prelabor cesarean compared with vaginal delivery, but we were not able to compare prelabor cesarean with cesarean after cervical dilatation; nor were we able to compare elective cesarean with planned vaginal delivery (ie, including both vaginal deliveries and cesarean after cervical dilatation). Furthermore, the effect estimates in the analysis comparing instrumental delivery and cesarean section, and elective cesarean and vaginal delivery were imprecise due to lack of statistical power. None of the studies collected information about SUI or UUI before delivery. In addition, 11 studies had unknown follow-up time [5,25,36,38,39,41,42,44–47]. However, the median of mean/median ages of the women included in these studies was >40 yr, implying that these studies also examined long-term impact of delivery mode on SUI and UUI.

As in all large-scale studies of incontinence, the included primary studies used self-report of incontinence, rather than diagnoses reached using urodynamics. We consider that self-report of SUI or UUI provides the most patient-relevant outcome. However, symptomatic incontinence may show limited correlation with urodynamic diagnosis, and thus these findings should not be generalized to diagnoses of urodynamic SUI or detrusor overactivity [49].

3.7. Relation to prior work

Although previous systematic reviews have demonstrated an increased risk of early postpartum incontinence after vaginal delivery compared with cesarean section [13,14], investigators have not previously conducted a rigorous review of long-term effects. One earlier systematic review focused entirely on the short-term postpartum period and included only studies with follow-up <1 yr [14], whereas the other one [13] included two appropriately adjusted cross-sectional studies of incontinence beyond the first postpartum year [37,43]. In contrast, our own search found 16 studies that could be included in meta-analyses. These much larger pooled analyses have provided more precise estimates of the impact of vaginal delivery compared with cesarean section on SUI.

The potential impact of delivery mode on UUI has received little consideration compared with the impact of delivery mode on SUI [21]. The only available prior review found no statistically significant difference between vaginal delivery and cesarean section [13]. In our much larger pooled analysis of eight studies (including 49 623 women), we were able to detect a modest effect size with tight CIs. For the first time, we have also been able to perform a quantitative synthesis of studies of instrumental delivery, finding no impact on SUI.

3.8. Implications of findings

Incontinence is very common among women irrespective of delivery history: prevalence estimates vary from 2.8% to 30.8% for SUI and from 0.7% to 19.9% for UUI [50,51]. Hence potentially increasing use of cesarean section may have beneficial public health consequences from the perspective of pelvic floor health, including decreased need for SUI and pelvic organ prolapse surgery [52,53]. Our results are consistent with those of a Swedish cohort study that reported vaginal deliveries increased surgical treatment...
for incontinence (hazard ratio: 2.9; 95% CI, 2.4–3.6) compared with women only having cesarean deliveries. The increased risk persisted for >3 decades [52]. The estimates provided here may be useful when counseling women about the risk and benefits of different delivery modes. Although we have quantified one benefit of planned cesarean, women and their caregivers must consider other consequences. Planned cesarean section confers an increased risk of neonatal intensive care admission for the baby and a substantially longer hospital stay for the mother [54]. A prior cesarean also carries risks in future pregnancies, including an increased risk of uterine rupture and abnormal placentaion [54]. In general, the medicalization of pregnancy associated with planned cesarean may also be undesirable from both individual and societal perspectives [55].

4. Conclusions

Our results demonstrate that vaginal delivery is associated with almost double the odds of long-term SUI, an absolute increase of approximately 8% when compared with cesarean section. The effect is largest in younger women but diminishes with age. The odds of UUI is also increased after vaginal delivery, but the pooled absolute difference is sufficiently small (3%) that cesarean section rates have only a small impact on UUI at a population level. The available evidence suggests no difference in the odds of SUI, if planned vaginal delivery results in instrumental delivery instead of spontaneous vaginal delivery. These data quantify one important aspect of cesarean section, to help women and their physicians make decisions regarding mode of delivery.

Author contributions: Kari A.O. Tikkinen had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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Appendix A. Supplementary data

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