

1  
2  
3       **The prevalence of urinary incontinence 20 years after child birth:**

4       **a national cohort study in one-para women after vaginal or cesarean delivery**

5  
6  
7  
8       Maria Gyhagen, Maria Bullarbo, Thorkild F. Nielsen, Ian Milsom

9  
10  
11  
12       The Department of Obstetrics and Gynecology, Sahlgrenska Academy at Gothenburg  
13       University, Sahlgrenska University Hospital, Gothenburg and the Department of Obstetrics  
14       and Gynecology, Södra Älvborgs Hospital, Borås, Sweden

15       **Running title:**

16       Urinary incontinence 20 years after child birth

17  
18       **Author for Correspondence:**

19  
20       Ian Milsom, M.D., Ph.D.  
21       Department of Obstetrics and Gynecology  
22       Sahlgrenska University Hospital  
23       SE-416 85 Gothenburg  
24       Sweden  
25       Tel: +46 31 3434179  
26       Fax: + 46 31 192940  
27       E-mail: [ian.milsom@gu.se](mailto:ian.milsom@gu.se)

30 **Abstract**

31 **Objective** To investigate the prevalence and risk factors for urinary incontinence (UI) 20  
32 years after one vaginal delivery (VD) or one cesarean section (CS).

33 **Design** Registry-based national cohort study.

34 **Setting** Women who returned postal questionnaires (response rate 65.2%) in 2008.

35 **Population** Primipara with one, single birth in 1985-1988 and no further births (n = 5 236).

36 **Methods** The SWEPOP (Swedish pregnancy, obesity and pelvic floor) study linked Medical  
37 Birth Register (MBR) data to a questionnaire about UI.

38 **Main Outcome Measures** Prevalence of UI and UI for more than 10 years (UI>10 years)  
39 were assessed 20 years after childbirth.

40 **Results** The prevalence of UI (40.3% vs. 28.8%, OR 1.67; 95% CI 1.45-1.92) and UI>10  
41 years (10.1% vs. 3.9%, OR 2.75; 95% CI 2.02-3.75) was higher in women after VD than after  
42 CS. There was no difference in the prevalence of UI or UI>10 years after an acute CS or an  
43 elective CS. We found an 8% increased risk of UI per current BMI unit and age at delivery  
44 increased UI risk by 3% annually.

45 **Conclusions** Two decades after one birth, VD was associated with a 67% increased risk of UI  
46 and UI>10 years increased by 275% compared to CS. Our data indicated that it is necessary to  
47 perform 8-9 cesarean sections to avoid one case of UI. Weight control is an important  
48 prophylactic measure to reduce UI. Current BMI was the most important BMI-determinant for  
49 UI, which is important, as BMI is modifiable.

50 **Keywords** Urinary incontinence; vaginal delivery; cesarean section; body mass index; risk  
51 factor; epidemiology.

52

53

54

55 **Introduction**

56 In modern societies, women live the major part of their lives after giving birth to one or two  
57 children. Urinary incontinence (UI) is a common condition affecting adult women of all ages  
58 which may have a negative influence on quality of life.<sup>1</sup> Pregnancy and in particular vaginal  
59 delivery have been implicated in the etiology of UI.<sup>1,2</sup> An increasing number of women  
60 request cesarean section for non-medical indications and for some their demand appears to be  
61 motivated by a desire to prevent pelvic floor damage, including UI.

62

63 The etiology of UI is known to be multifactorial but obesity and ageing as well as obstetric  
64 trauma during childbirth are known to be three of the most important risk factors.<sup>1,2</sup> Although  
65 several studies have demonstrated an association between UI and vaginal delivery in the  
66 short- and medium long-term the long-term effects of childbirth on the risk of UI remain  
67 controversial.<sup>3-6</sup> The assessment of the influence of childbirth on urinary incontinence later in  
68 life has been hampered by the heterogeneity of study populations. Women of different ages  
69 and varying body weights have been included after a variable number of pregnancies often  
70 with different modes of delivery. The aim of this study was therefore to compare the  
71 prevalence of UI 20 years after delivery in a cohort of women who had given birth to only one  
72 child after vaginal delivery (VD) or cesarean section (CS).

73

74 **Methods**

75 A national survey of pelvic floor dysfunction, the SWEPOP (Swedish pregnancy, obesity and  
76 pelvic floor) study was conducted in 2008. The population studied and their obstetric data was  
77 obtained from the Swedish Medical Birth Register (MBR). The MBR, which was started in  
78 1973, is a national register that includes more than 98% of all births in Sweden. Data from all  
79 antenatal clinics and all obstetric units are sent to the MBR at the National Board of Health and

80 Welfare. Obstetrical parameters from the delivery were obtained from the MBR. Cesarean  
81 done before the onset of labor was denoted as an elective cesarean section (ECS) and cesarean  
82 done during labor was denoted as an acute cesarean section (ACS). The weight and height of  
83 pregnant women had been measured by a midwife at the antenatal clinic, usually at 8-10  
84 weeks of gestation and was also obtained from the MBR. Maternal weight at delivery and the  
85 weight gain during pregnancy were recorded at the delivery unit and were also obtained from  
86 the MBR. When individual data was initially examined it was noted that the maximum-  
87 recorded body weight from the MBR was 99 kg. Due to lack of data storage capacity in the  
88 1980s the MBR had decided to restrict registration of "heavy women" by recording up to two  
89 digits only. We therefore reviewed the patient records of the 300 women recorded as having a  
90 body weight of 99 kg to obtain the correct weights of these women.

91 The quality of this national database has been shown to be good and suitable for population  
92 studies of this type. A description of the MBR in english can be found at  
93 (<http://www.socialstyrelsen.se/register/halsodataregister/medicinskafoedelseregistret/inenglish>) and  
94 an evaluation of the MBR has been performed by Cnattingius et al.<sup>7</sup> as well as by the  
95 National Board of Health and Welfare and is available at  
96 <http://www.socialstyrelsen.se/publikationer2002/2002-112-4>. Inclusion criteria for  
97 participation in this study were *primiparae* with *one, single birth* 1985-1988 and *no further*  
98 *births*. Multifetal pregnancies were excluded. Ethical approval was obtained from the  
99 Regional and the National Ethic Review Boards (the Ethics Committee at Sahlgrenska  
100 Academy, Gothenburg University, and the National Board of Health and Welfare).

101 The results of this study have been reported according to the STROBE statement.

102

103 The questionnaire was sent to 9 423 women (Fig. 1) who were asked to provide written,  
104 informed consent to participate and to complete a questionnaire. Subjects were excluded from

105 the study, based on the answers in the questionnaire, if they affirmed multiparity  
106 (misdiagnosis of ‘parity’ is predominantly related to immigration, the first birth in Sweden is  
107 sometimes misdiagnosed as the first ever), multifetal or ongoing pregnancy (Fig. 1). The  
108 results regarding anal incontinence and genital prolapse will be reported separately.

109  
110 The 31-item questionnaire included questions about current height and weight, urinary or anal  
111 incontinence and genital prolapse, menstrual status, hysterectomy, the menopause, hormone  
112 treatment etc. Urinary incontinence (UI) was defined according to the International  
113 Continence Society and by the question “Do you have involuntary loss of urine?”.<sup>8</sup>  
114 Participants reporting UI were grouped according to the duration of UI (UI< 5 years, 5–10  
115 years, or >10 years). The severity of UI (frequency, leakage amount) was assessed using the  
116 Sandvik score.<sup>9</sup> After three mailing cycles during a four month period the questionnaire was  
117 returned by 6148 women (65.2 %).

118  
119 Maternal BMI was categorized as normal (<25), overweight ( $\geq 25$ -29.9) and obese ( $\geq 30$ )  
120 according to the WHO classification<sup>10</sup> and was calculated for each woman according to  
121 weight and height measurements in early pregnancy at week 8-10, (BMI-Early Pregnancy), at  
122 delivery (BMI-Delivery) and 20 years after delivery (Current BMI).

123  
124 ***Characteristics of the sample population and the non-responders***  
125 The proportion of missing data varied between 0% (age) and 15.9% for hysterectomy in the  
126 population cohort. There was little difference difference in the proportions of missing data  
127 between groups, eg. the proportion of missing data for hysterectomy which had the greatest  
128 proportion of missing data was 15.5% (620/3995) in the VD-group and 17% (205/1204) in the  
129 CS-group. The non-responders were 1.6 years younger (49.6 yrs  $\pm$  5.9 vs. 51.2 yrs  $\pm$  5.9;

130 p<0.001), and they were more often overweight or obese (37% vs. 27%; p<0.001 and had an  
131 infant birth weight <4000 g (43% vs. 48%; p<0.003) compared to responders.

132

133 **Statistical analysis**

134 Statistical analysis was performed with SAS 9.1 (SAS Institute Inc, Cary, NC, USA). For  
135 cohort characteristics  $\chi^2$  test was used to compare categorical variables and the Students t-test  
136 for continuous variables. A *p*-value less than 0.05 was considered statistically significant.  
137 Adjusted frequencies and odds ratios for UI were calculated using a covariance analysis  
138 model to obtain effect measures. A logistic regression model was used to assess risk factors  
139 for UI while controlling for potential confounders. Potential risk factors used in the analysis  
140 were mode of delivery, maternal age at delivery, maternal BMI (at delivery, and current),  
141 hysterectomy, hormone replacement therapy, gestational age, infant birth weight and head  
142 circumference. Odds ratio (OR) and their 95% confidence intervals (CI) were calculated from  
143 the model. Prevalence figures permitted the calculation of the number of cesarean sections  
144 needed to avoid one case of UI using the number needed to treat principle (NNT). The NNT  
145 was calculated as the inverse of the absolute risk reduction, where risk reduction was the  
146 difference of adjusted UI prevalence between VD and CS.

147 **Results**

148 Basic characteristics of the women grouped according to mode of delivery are shown in  
149 Tables 1 and 2. The mean follow-up time after delivery was 21.5 yrs. (SD 1.5) in VD group  
150 and 21.8 yrs (SD 1.1) in the CS group. Women delivered by CS were older (current age 53.7  
151 yrs. (SD 6.3) compared to 50.4 yrs. (SD 5.6) in the VD group p<0.001) and gave birth to an  
152 infant with a lower birth weight (p<0.001) at a lower gestational week (p<0.001) compared to  
153 women who were delivered vaginally (Table 1). The proportion of women aged  $\geq 35$  years at  
154

155 delivery was higher ( $p<0.001$ ) in the CS group whereas the proportion of infants with a birth  
156 weight  $\geq 3500$  g was lower ( $p<0.001$ ) in the CS group compared to the VD group.

157

158 The prevalence of UI (Table 3) was 67% higher (OR 1.67 CI; 1.45-1.92) after a vaginal  
159 delivery (40.3%) compared to women who had been delivered by cesarean section (28.8%).  
160 From the prevalence data available on UI it was possible to calculate using the NNT principle  
161 that it is necessary to perform 8-9 cesarean sections to avoid one case of UI. Furthermore, the  
162 prevalence and risk increase of UI for more than 10 years almost tripled after VD compared  
163 to after CS. Prevalence of UI for >10 years after VD was 10.1% compared to 3.9% after CS  
164 (OR 2.75 CI: 2.02-3.75). There was however no significant differences in the prevalence of UI  
165 (27.1% vs. 24.4%, OR 1.15 CI: 0.88-1.51) or UI for more than 10 years (6.5% vs. 5.1%, OR  
166 1.30 CI: 0.79-2.14) between women delivered by acute cesarean section (ACS) or elective  
167 cesarean section (ECS) respectively.

168

169 The prevalence of urinary incontinence was higher after VD compared to CS for each current  
170 BMI class ( $BMI<25$ ,  $BMI\geq 25-29.9$ , and  $BMI\geq 30$ ) with differences ranging from 11 to 14%  
171 (Table 4). Again using the NNT principle ,we calculated the number of cesarean section that  
172 would need to be performed to prevent one case of UI for the different BMI groups (9 for  
173  $BMI <25$ ; 7 for  $BMI 25-29.9$  and 8 for  $BMI \geq 30$  ). The combined effect of BMI and mode of  
174 delivery was substantial, for example the adjusted frequency of UI after CS with a current  
175  $BMI <25$  was 24.7% whereas the frequency more than doubled to 54.8% after VD with a  
176 current  $BMI \geq 30$  (Table 4). When using “normal BMI” as reference the risk of UI increased  
177 significantly for both overweight and obese women after both modes of delivery. The risk  
178 increase of UI in obese women more than doubled in comparison to women with a normal  
179 BMI after VD and more than tripled after CS (Table 5). In the logistic regression analyses we

180 found an 8% (range 6-10%) increased risk of UI per BMI unit increase and the increased rate  
181 of UI was apparent for both modes of delivery (Table 6).

182

183 Due to a interaction term between IBW and mode of delivery ( $p<0.01$ ) we made separate  
184 analyses of IBW for the CS group and VD group. The prevalence of UI following VD was  
185 higher than after CS in all infant birth weight groups except for weights  $<3000$  g. For women  
186 who delivered vaginally rates of incontinence increased with increasing infant birth weight  
187 but this was not observed after CS (Table 4). Logistic regression analysis in the total cohort  
188 failed to demonstrate a significant increase of UI risk for infant birth weights  $\geq 4500$  g (Table  
189 5).

190

191 The multivariable analysis (Table 6) did not demonstrate any significant increased risk of UI  
192 associated with infant head circumference. However, there was an increased risk of UI after  
193 VD compared to CS regardless of fetal head circumference. The risk increase associated with  
194 VD in comparison to CS was stronger for fetal head circumference  $\geq 36$  cm than for head  
195 circumference less than 36 cm, OR 2.46 (1.66–3.63) vs. 1.64 (1.40–1.91) (Table 4). Nor were  
196 there any differences in UI prevalence in the women grouped according to fetal head  
197 circumference (Table 5) after both modes of delivery.

198

199 The prevalence of UI was 10% higher in women  $\geq 35$  years at delivery compared to women  
200  $<23$  years who had undergone CS and 7% higher in women of the same ages who had a VD  
201 (Table 4). In the logistic regression analysis a higher maternal age was associated with an  
202 increased risk of UI (OR 1.03, 1.02–1.04), which corresponds to an annual risk increase of 3%  
203 per year.

204

205 **Discussion**

206 The risk of developing UI was found to be 67-71% higher after VD than after CS and the  
207 prevalence of UI for more than 10 years almost tripled after VD compared to CS. We found  
208 no difference in the prevalence of UI or UI for more than 10 years between women delivered  
209 by ACS or ECS, indicating that it is during the later stages of delivery, when the fetus passes  
210 through the pelvic floor, that leads to the increased risk of UI. Maternal weight was also an  
211 important risk factor and in the logistic regression analyses we found an 8% increased risk of  
212 UI per BMI unit increase and the rate of UI was apparent for both modes of delivery. Current  
213 BMI was the most important BMI-determinant for UI and this finding is important, as BMI is  
214 modifiable. For women who delivered vaginally rates of incontinence increased with  
215 increasing infant birth weight but this was not observed after CS. The prevalence of UI  
216 increased with maternal age and there was an annual increase in UI prevalence of 3% per  
217 year.

218

219 The main strengths of this study are the use of a large national, population-based cohort of I-  
220 para women and the high response rate. There are advantages of studying I-para women as the  
221 first delivery is considered to exert the greatest risk increase for UI even if subsequent  
222 deliveries contribute to a further increase in the risk of UI.<sup>1,11</sup> Including multiparous women  
223 would disrupt obstetric homogeneity and since most risk factors also covariate with time/age  
224 also this would confound effect measures of the analysis. The inclusion of I-para women  
225 regardless of maternal health status, maternal and fetal complications is considered a strength  
226 as it allows for a greater generalisation of results and therefore a better basis for consultation  
227 about elective cesarean section on request. The weight and height data during pregnancy were  
228 objectively measured at the antenatal clinics and a validated questionnaire was used.<sup>9,12</sup>

229

230 Some limitations of the present investigation must also be considered. First, women with  
231 incontinence may be more predisposed to participate in studies and therefore UI might be  
232 over-estimated. Secondly, the symptoms of UI were self-reported. However, several studies  
233 have shown that self-reported symptoms are consistent and valid when assessing current UI  
234 and changes in incontinence severity over time which applies to our study.<sup>11,13</sup> This study  
235 also lacks information on whether UI was present or not before or/and during pregnancy or  
236 started after delivery. However there is little evidence to suggest any difference in UI  
237 prevalence before the first pregnancy or during pregnancy in women grouped according to  
238 mode of delivery. It was not possible to assess the importance of the length of the second  
239 stage of delivery, as this is unfortunately not documented in the MBR. Obstetric techniques  
240 and parameters have varied over time (fewer episiotomies, increasing number of vacuum  
241 extractions and severe lacerations, older mothers, higher BMI and heavier children) which  
242 may also influence the clinical interpretation of our results. It may also in some respects seem  
243 unrepresentative to study the consequences of giving birth to only one child. However, United  
244 Nations data show that total fertility rates (TFR) study<sup>14</sup> are rapidly declining globally and the  
245 predicted TFR in the middle of this century is predicted to be less than 2.0 children/woman  
246 and in many developed countries the TFRs is already between 1.0 and 1.5. Analyses of the  
247 non-responder group suggest a small selection bias on our results acting in both directions  
248 (younger women and smaller children leading to overestimation of results; overweight/obesity  
249 to the opposite).

250

251 In this study, CS was often used as reference for comparison with VD to quantify the effect of  
252 vaginal birth on UI. The baseline outcome after CS can then be interpreted as representing the  
253 risk of pregnancy *per se* and the risk of VD represented the risk of pregnancy plus VD and

254 hence the difference between VD and CS is therefore a measure (in terms of UI prevalence  
255 and risk) of vaginal birth trauma. Even if the nulliparous pelvis represents the best available  
256 clinical model of normal function, the prevalence of UI in nulliparous women of childbearing  
257 age has been reported to be 10-15% study.<sup>2,15</sup> Urinary leakage preceding pregnancy in  
258 nulliparous women has been shown to be a strong precursor for increased prevalence of UI 4-  
259 12 years post partum.<sup>1,15</sup> Pregnancy *per se*, independent of labour and delivery practice, has  
260 been reported to be a risk factor for postpartum UI<sup>4,16</sup> especially if the incontinence started  
261 during the first trimester.<sup>17</sup> Several studies have demonstrated that postpartum UI is a risk  
262 factor for UI after varying terms of follow-up.<sup>3-5</sup>

263

264 There is still no general agreement whether or not the long-term maternal effects of the two  
265 delivery modes differ with regard to prevalence of UI. The prospective multicenter study of  
266 McKinnie et al.<sup>18</sup> did not show a significant difference of risk for bothersome UI between  
267 women delivered by one or more VD compared to one or more CS. Also The Omnibus  
268 Survey of MacLennan et al.<sup>19</sup> could not demonstrate an increased risk for any type of UI after  
269 VD when compared to CS. In these studies however the CS groups were relatively small and  
270 heterogeneous with respect to parity. On the other hand the EPINCONT study demonstrated a  
271 1.7-fold increased age-adjusted risk of UI after one or more VD compared with one or more  
272 CS. The age-standardized prevalence rate of UI was 15.9% for the CS group and 21.0% for  
273 the VD group.<sup>2</sup> The study population was younger and the follow-up time shorter in the  
274 EPINCONT-study compared to the cohorts and the follow up time of our study. Other later  
275 studies have also indicated an increased risk of UI following VD compared to CS.<sup>20-22</sup>

276

277 Several studies have reported that a higher BMI is a risk factor for UI<sup>1</sup> and cross-sectional  
278 studies have confirmed this association in middle-aged women.<sup>23,24</sup> We found an increased

279 risk of UI of  $\approx$  8 % per unit BMI. Our findings correspond with those of others who showed a  
280 risk increase varying between 2-10% per unit increase of BMI.<sup>25,26</sup> The overall assessment of  
281 the relation between BMI and prevalence of UI in this study indicated that there was a dose-  
282 response relationship between BMI and UI whereas the effect of mode of delivery (i.e. VD or  
283 CS) appeared to be constant regardless of maternal BMI status.

284 The results of this study indicated that current BMI was the most important determinant for  
285 UI and this finding is important, as BMI is modifiable. Resolution of UI has been  
286 demonstrated after weight loss.<sup>27</sup> Intervention by non-surgical means or laparoscopic gastric  
287 bypass surgery indicate that there is a dose-response association between prevalence of UI  
288 and the magnitude of weight reduction.<sup>28</sup> The strong association between the prevalence of UI  
289 and current BMI is encouraging as it means that it is never too late to achieve improvement of  
290 UI through weight reduction and weight control.

291  
292 The negative effect of VD on urinary continence is consistent with results of several clinical  
293 studies that have demonstrated poor urethral support and increased urethral mobility after VD,  
294 leading to UI.<sup>29,30</sup> Impaired urethral function could also be shown after VD<sup>29</sup> but this was not  
295 observed after CS.<sup>31</sup>

296  
297 In conclusion, the risk of developing UI and UI for more than 10 years was higher 20 years  
298 after a VD compared to a CS. Prevalence did not differ between women delivered by ACS or  
299 ECS, indicating the importance of the later stages of delivery during the passage of the fetus  
300 through the pelvic floor for the occurrence of UI in later life. Weight control was also shown  
301 to be an important preventive measure to reduce UI. Our data also provide a quantification of  
302 the importance of mode of delivery and body weight for the risk of future UI. The results of  
303 this study indicate that one has to perform 8-9 cesarean sections to avoid one case of UI.

304 However there may be other advantages regarding the possible protective effective of CS on  
305 future pelvic floor function, such as a reduced prevalence of vaginal prolapse which could be  
306 included in the decision of whether or not CS is advantageous. Vaginal delivery and BMI  
307 have been shown to be important risk factors for UI but operative delivery by CS also  
308 involves a degree of risk for morbidity and mortality over and above that of VD.<sup>32</sup>

309 **Acknowledgements**

310 We thank Ms Marianne Sahlén and Ms Anja Andersson for help with data registration and  
311 Björn Areskoug MSc for expertise in statistical programming.

312

313 **Disclosure of interests**

314 We declare that we have no conflict of interests.

315

316 **Contribution to Authorship**

317 All authors were involved in the conception and design of the study, acquisition of data and  
318 interpretation of the results as well as the writing of the manuscript. All authors approved the  
319 final version of the submitted manuscript. MG and IM take full responsibility for the integrity  
320 of the data and the accuracy of the data analysis.

321

322 **Details of ethics approval**

323 Ethical approval for the SWEPOP-study was obtained from the Regional and the National  
324 Ethic Review Boards (the Ethics Committee at Sahlgrenska Academy, Gothenburg  
325 University, Ref No 381-07, August 13<sup>th</sup> 2007 and the National Board of Health and Welfare,  
326 Ref No 34-9148/2007, October 26<sup>th</sup> 2007).

327 **Funding**

328 The study was supported by a National LUA/ALF grant nr 11315 and the Region of Västra  
329 Götaland, grants from The Göteborg Medical Society and Hjalmar Svenssons Fund, The  
330 funding source had no role in the study design, data analysis, data interpretation or writing of  
331 the report. MG and IM had full access to all study data and had final responsibility for the  
332 decision to submit for publication.

333

334

335 **References**

336 1. Milsom I, Altman D, Herbison P, Lapitan MC, Nelson R, Sillén U, Thom D.  
337 Epidemiology of urinary (UI) and faecal (FI) Incontinence and pelvic organ prolapse  
338 (POP). *In: Incontinence*, Editors Abrams, Cardozo, Kourhry and Wein. Health  
339 Publications Ltd, Paris 2009.

340

341 2. Rortveit G, Daltveit AK, Hannestad YS, Hunskaar S. Urinary incontinence after vaginal  
342 delivery or cesarean section. *N Engl J Med* 2003; 348: 900-07.

343

344 3. Viktrup L, Lose G, Rolff M, Barfoed K. The symptom of stress incontinence caused by  
345 pregnancy or delivery in primiparas. *Obstet Gynecol* 1992; 79: 945-49.

346

347 4. MacArthur C, Glazener CM, Wilson PD, Lancashire RJ, Herbison GP, Grant AM.  
348 Persistent urinary incontinence and delivery mode history: a six-year longitudinal study.  
349 *BJOG* 2006; 113: 218-24.

350

351 5. Press JZ, Klein MC, Kaczorowski J, Liston RM, von Dadelszen P. Does cesarean  
352 section reduce postpartum urinary incontinence? A systematic review. *Birth* 2007; 34:  
353 228-37.

354

355 6. MacArthur C, Glazener C, Lancashire R, Herbison P, Wilson D. Exclusive caesarean  
356 section delivery and subsequent urinary and faecal incontinence: a 12-year longitudinal  
357 study. *BJOG* 2011;118: 1001-7.

358 7. Cnattingius S, Ericson A, Gunnarskog J, Källén B. A quality study of a medical birth  
359 registry. *Scand J Soc Med* 1990; 2:143-8.

360

361 8. Abrams P, Cardozo L, Fall M, et al. The standardisation of terminology in lower urinary  
362 tract function: report from the standardisation sub-committee of the International  
363 Continence Society. *Urology* 2003; 61: 37-49.

364

365 9. Sandvik H, Hunskaar S, Seim A, Hermstad R, Vanvik A, Bratt H. Validation of a  
366 severity index in female urinary incontinence and its implementation in an  
367 epidemiological survey. *J Epidemiol Community Health* 1993; 47: 497-9.

368

369 10. World Health Organization. BMI classification. 2006 [updated 02/02/201102/02/2011];  
370 Available from: [http://apps.who.int/bmi/index.jsp?introPage=intro\\_3.html](http://apps.who.int/bmi/index.jsp?introPage=intro_3.html).

371

372 11. Milsom I, Ekelund P, Molander U, Arvidsson L, Areskoug B. The influence of age,  
373 parity, oral contraception, hysterectomy and menopause on the prevalence of urinary  
374 incontinence in women. *J Urol* 1993; 149: 1459-62.

375

376 12. Sandvik H, Seim A, Vanvik A, Hunskaar S. A severity index for epidemiological  
377 surveys of female urinary incontinence: comparison with 48-hour pad-weighing tests.  
378 *Neurourol Urodyn* 2000; 19: 137-45.

379

380 13. Alling Moller L, Lose G, Jorgensen T. Risk factors for lower urinary tract symptoms in  
381 women 40 to 60 years of age. *Obstet Gynecol* 2000; 96: 446-51.

382

383 14. United Nations Children's Fund. Total fertility rate (TFR). 2008 [updated  
384 07/201003/02/2011]; Available from:  
385 <http://data.un.org/Data.aspx?d=SOWC&f=inID%3A127>.  
386

387 15. Brown SJ, Donath S, MacArthur C, McDonald EA, Krastev AH. Urinary incontinence  
388 in nulliparous women before and during pregnancy: prevalence, incidence, and  
389 associated risk factors. *Int Urogynecol J Pelvic Floor Dysfunct* 2010; 21: 193-202.  
390

391 16. Thom DH, Brown JS. Reproductive and hormonal risk factors for urinary incontinence  
392 in later life: a review of the clinical and epidemiologic literature. *J Am Geriatr Soc*  
393 1998; 46: 1411-17.  
394

395 17. Eason E, Labrecque M, Marcoux S, Mondor M. Effects of carrying a pregnancy and of  
396 method of delivery on urinary incontinence: a prospective cohort study. *BMC*  
397 *Pregnancy Childbirth* 2004; 4: 4.  
398

399 18. McKinnie V, Swift SE, Wang W, et al. The effect of pregnancy and mode of delivery  
400 on the prevalence of urinary and fecal incontinence. *Am J Obstet Gynecol* 2005; 193:  
401 512-17; discussion 517-8.  
402

403 19. MacLennan AH, Taylor AW, Wilson DH, Wilson D. The prevalence of pelvic floor  
404 disorders and their relationship to gender, age, parity and mode of delivery. *BJOG* 2000;  
405 107: 1460-70.  
406

407 20. Lukacz ES, Lawrence JM, Contreras R, Nager CW, Luber KM. Parity, mode of  
408 delivery, and pelvic floor disorders. *Obstet Gynecol* 2006; 107: 1253-60.

409

410 21. Dolan LM, Hilton P. Obstetric risk factors and pelvic floor dysfunction 20 years after  
411 first delivery. *Int Urogynecol J* 2010; 21: 535-544.

412

413 22. Leijonhufvud A, Lundholm C, Cnattingius S, Granath F, Andolf E, Altman D. Risks of  
414 stress urinary incontinence and pelvic organ prolapse surgery in relation to mode of  
415 childbirth. *Am J Obstet Gynecol* 2011; 204: 70.e1-7.

416

417 23. Chiarelli P, Brown W, McElduff P. Leaking urine: prevalence and associated factors in  
418 Australian women. *Neurourol Urodyn* 1999; 18: 567-77.

419

420 24. Sampselle CM, Harlow SD, Skurnick J, Brubaker L, Bondarenko I. Urinary  
421 incontinence predictors and life impact in ethnically diverse perimenopausal women.  
422 *Obstet Gynecol* 2002; 100: 1230-38.

423

424 25. Mommsen S, Foldspang A. Body mass index and adult female urinary incontinence.  
425 *World J Urol* 1994; 12: 319-22.

426

427 26. Waetjen LE, Liao S, Johnson WO, et al. Factors associated with prevalent and incident  
428 urinary incontinence in a cohort of midlife women: a longitudinal analysis of data: study  
429 of women's health across the nation. *Am J Epidemiol* 2007; 165: 309-18.

430

431 27. Greer WJ, Richter HE, Bartolucci AA, Burgio KL. Obesity and pelvic floor disorders: a  
432 systematic review. *Obstet Gynecol* 2008; 112: 341-349.

433

434 28. Subak LL, Wing R, West DS, et al. Weight loss to treat urinary incontinence in  
435 overweight and obese women. *N Engl J Med* 2009; 360: 48-90.

436

437 29. Tapp A, Cardozo L, Versi E, Montgomery J, Studd J. The effect of vaginal delivery on  
438 the urethral sphincter. *Br J Obstet Gynaecol* 1988; 95: 142-46.

439

440 30. Peschers U, Schaer G, Anthuber C, Delancey JO, Schuessler B. Changes in vesical neck  
441 mobility following vaginal delivery. *Obstet Gynecol* 1996; 88: 1001-06.

442

443 31. van Geelen JM, Lemmens WA, Eskes TK, Martin CB, Jr. The urethral pressure profile  
444 in pregnancy and after delivery in healthy nulliparous women. *Am J Obstet Gynecol*  
445 1982; 144: 636-49.

446

447 32. National Institute of Health State of the Science conference statement. Cesarean  
448 delivery on maternal request. March 27-29, 2006. <http://consensus.nih.gov>.

449

450

451

452

453

454

455

456

457

458

459

460

461

462

463

464

465

466

467

468

469

470

471

472

473 **Legend to Figure**

474

475 **Figure 1.** Flowchart of the women who gave birth to one child 1985–1988 identified from the

476 Swedish Medical Birth Register (MBR).

477

478

479

480

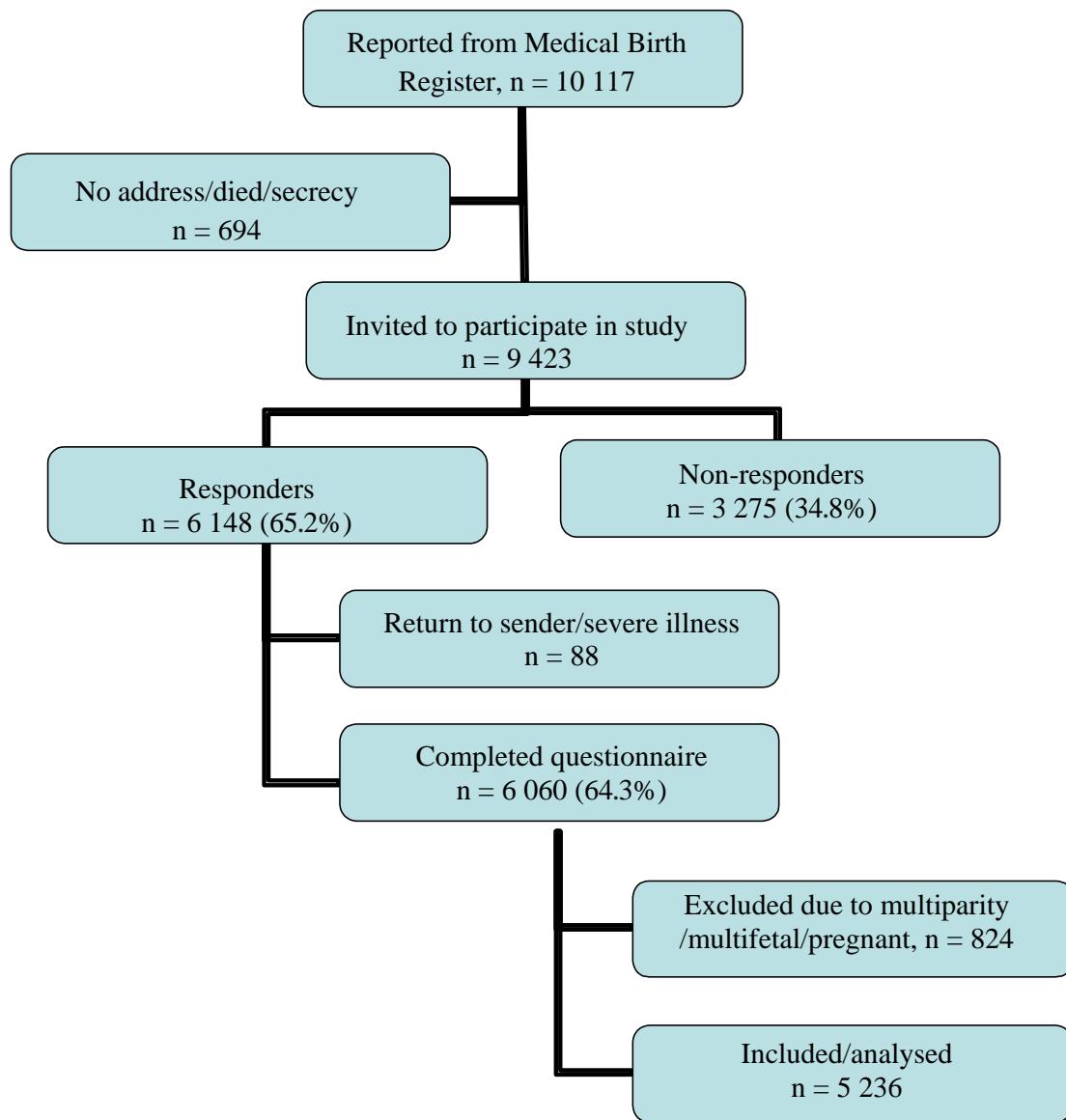
481

482

483

484

Fig. 1



The current addresses of 9423 of these women could be traced in the Swedish population address register and these women were invited to participate in the study. The difference, 694 women, was due to newly deceased women or women with unknown address or hidden personal identity. Of the 6 148 who returned questionnaires 6060 women were able to participate or gave their informed consent for participation in the study. At this stage however a further 824 women were excluded from the study population due to the fact that they had given birth abroad unknown to the MBR, were currently pregnant or had been wrongly categorized as having had only one child in the MBR. Excluded due to multiparity (n = 716); multifetal (n = 43); ongoing pregnancy (n = 6) and 59 missing data about parity in the questionnaire.

**Table 1.** The basic characteristics of the women grouped according to mode of delivery.

	<b>VD</b> mean (range)/%	<b>CS</b> (ECS and ACS) mean (range)/%	<b>Difference</b> CI (95%)	<b>p-value</b>
	n = 3 995	n = 1 204		
Age at delivery	29.0 (15–46)	31.9 (15–45)	2.9 (2.6–3.3)	<0.001
Age ≥35 years	17.4%	35.6%	18.2 (15.3–21.2)	<0.001
BMI early pregnancy	23.0 (15.0–45.6)	23.0 (15.2–41.7)	0.0 (-0.3–0.3)	=0.94
BMI at delivery	28.3 (17.2–50.3)	28.3 (18.3–47.2)	0.0 (-0.3–0.3)	=0.95
BMI at delivery ≥25	79.0%	78.1%	0.9 (-4–2)	=0.56
BMI current	26.1 (14.5–63.0)	26.3 (16.1–53.8)	0.2 (-0.1–0.5)	=0.16
BMI current ≥25	51.3%	51.2%	0.1 (-3–3)	=0.98
Infant birth weight (g)	3585 (850–5680)	3294 (820–5615)	291 (245–336)	<0.001
Infant birth weight ≥3500 g	55.8%	38.2%	17.6 (14.4–20.7) .	<0.001
Gestational age (weeks)	39.7 (24–45)	38.5 (27–43)	1.2 (1.1–1.4)	<0.001
Hysterectomy	7.9%	9.9%	2.0 (0.00–0.04)	=0.06
Estrogen therapy	7.1%	10.0%	2.9 (0.8–4.9)	<0.01

VD = vaginal delivery; CS = cesarean section; ECS = Elective cesarean section; ACS = Acute cesarean section; BMI = body mass index. Student's t-test was used for statistical comparison between groups.

**Table 2.** Cohort characteristics for the women who underwent elective or acute cesarean section.

	<b>ECS mean (range)/%</b>	<b>ACS mean (range)/%</b>	<b>Difference CI (95%)</b>	<b>p-value</b>
	n = 766	n = 438		
Age at delivery	32.5 (15–45)	30.9 (18–45)	-1.64(-2.32 – - 0.95)	<0.001
Age ≥ 35 years	40.7%	26.7%	-14.0 (-19.4 – -8.6)	<0.001
BMI early pregnancy	22.4 (15.2–41.7)	23.9 (15.4–41.7)	1.52 (1.00–2.04)	<0.001
BMI at delivery	27.6 (18.3–47.2)	29.3 (18.9–46.5)	1.75 (1.20–2.30)	<0.001
BMI at delivery ≥ 25	74.7%	83.6%	8.9 (3.8–14.0)	<0.001
BMI current	25.8 (18.9–50.8)	27.0 (18.1–53.8)	1.16 (0.55–1.77)	<0.001
BMI current ≥ 25	47.1%	58.2%	11.1 (5.2–17.0)	<0.001
Infant birth weight (g)	3157(820–5615)	3534 (950–5310)	377 (286–468)	<0.001
Infant birth weight ≥ 3500 g	27.2%	57.5%	30.3 (24.7–36.0)	<0.001
Gestational age (weeks)	38.0 (27–43)	39.4 (27–43)	1.44 (1.15–1.73)	<0.001
Hysterectomy	10.1%	9.5%	-0.01 (-0.04–0.03)	=0.75
Estrogen therapy	11.5%	7.4%	-0.04(-0.08–0.00)	<0.05

ECS = Elective cesarean section; ACS = Acute cesarean section; BMI = body mass index.

Student's t-test was used for statistical comparison between groups.

**Table 3.** Crude and adjusted\* prevalence and odds ratio of urinary incontinence and urinary incontinence for more than 10 years in relation to mode of delivery.

	CS %	VD %	Crude OR (95% CI)	CS %	VD %	Adjusted* OR (95% CI)
Urinary incontinence	30.0	40.2	1.56 (1.36-1.80)	28.8	40.3	1.67 (1.45-1.92)
Urinary incontinence for >10 years	4.6	10.0	2.30 (1.73-3.08)	3.9	10.1	2.75 (2.02-3.75)

VD = vaginal delivery; CS = cesarean section; BMI = body mass index.

\*Adjusted for BMI-current, BMI at delivery, maternal age, gestational weeks, infant birth weight and head circumference.

**Table 4.** The results\* of logistic regression analysis of possible risk factors for urinary incontinence (odds ratio 95% CI).

<b>Risk factors/confounders</b>	<b>OR (95% CI)</b>
Vaginal delivery	1.71 (1.41–2.08)
BMI-current	1.08 (1.06–1.10)
BMI-at delivery	0.98 (0.96–1.00)
Maternal age at delivery	1.03 (1.02–1.04)
Gestational weeks	1.01 (0.96–1.06)
Infant birth weight (hg)	1.00 (0.98–1.02)
Infant head circumference ≥ 36 cm	1.06 (0.84–1.34)
Hysterectomy	1.21 (0.90–1.63)
Hormone replacement therapy yes vs. no	1.34 (0.98–1.82)

BMI = body mass index. \* 3488 women contributed to the model.

**Table 5.** Crude and adjusted\* prevalence and odds ratio of urinary incontinence in relation to mode of delivery stratified for each risk factor

Prevalence of urinary incontinence							
	CS %	VD %	Crude OR (95% CI)	CS %	VD %	Adjusted* OR (95% CI)	
<b>Infant birth weight (g)</b>							
< 3000	33.0	37.6	1.23 (0.91–1.64)	35.5	37.4	1.08 (0.81–1.49)	
3000–3499	28.6	38.6	1.57 (1.22–2.00)	27.3	39.1	1.71 (1.33–2.19)	
3500–3999	30.9	40.5	1.53 (1.14–2.05)	27.6	41.3	1.85 (1.36–2.50)	
4000–4499	27.8	41.5	1.85 (1.28–2.67)	25.3	41.3	2.08 (1.42–3.03)	
≥ 4500	23.1	50.4	3.39 (1.63–7.04)	21.2	48.8	3.56 (1.68–7.53)	
<b>Infant head circumference (cm)</b>							
< 36	29.8	39.9	1.56 (1.34–1.82)	29.0	40.1	1.64 (1.40–1.91)	
≥ 36	26.9	43.1	2.06 (1.41–3.01)	24.0	48.8	2.46 (1.66–3.63)	
<b>BMI early pregnancy</b>							
< 25	30.9	37.5	1.34 (1.12–1.61)	29.7	37.8	1.44 (1.20–1.74)	
25–29.9	29.2	44.5	1.94 (1.37–2.76)	27.6	44.9	2.14 (1.50–3.05)	
≥ 30	45.5	51.9	1.29 (0.67–2.50)	47.7	51.9	1.18 (0.61–2.28)	
<b>BMI at delivery</b>							
< 25	29.4	32.7	1.17 (0.84–1.63)	28.9	32.9	1.20 (0.86–1.68)	
25–29.9	30.8	39.3	1.45 (1.17–1.80)	30.2	39.5	1.51 (1.21–1.87)	
≥ 30	30.5	46.3	1.96 (1.49–2.59)	28.4	46.9	2.93 (1.68–2.95)	
<b>Current BMI</b>							
< 25	24.5	34.1	1.59 (1.29–1.97)	24.7	35.6	1.68 (1.36–2.08)	
25–29.9	29.3	41.7	1.73 (1.34–2.23)	28.1	42.4	1.88 (1.46–2.43)	
≥ 30	45.3	52.8	1.35 (1.01–1.81)	41.6	54.8	1.71 (1.27–2.29)	
<b>Maternal age at delivery (years)</b>							
< 23	22.8	35.6	1.87 (1.18–2.96)	22.8	36.0	1.90 (1.20–3.02)	
23–29	27.9	39.0	1.65 (1.26–2.17)	26.0	38.3	1.77 (1.34–2.33)	
30–34	33.8	42.9	1.47 (1.14–1.89)	33.2	43.2	1.53 (1.19–1.97)	
≥ 35	30.4	43.0	1.72 (1.33–2.23)	32.3	42.7	1.56 (1.21–2.01)	

VD = vaginal delivery; CS = cesarean section; BMI = body mass index.

\*Adjusted for BMI-current, BMI at delivery, maternal age, gestational weeks, infant weight and head circumference.

**Table 6.** Adjusted\* additional risks of UI in relation to stratified risk factors grouped according to mode of delivery.

	CS (95% CI)	VD (95% CI)
<b>BMI-current</b>		
<25	ref	ref
25–29.9	1.50 (1.11–2.03)	1.32 (1.14–1.53)
≥30	3.27 (2.34–4.59)	2.50 (2.10–2.98)
<b>Age at term</b>		
<23	ref	ref
23–29	1.29 (0.73–2.26)	1.22 (1.02–1.46)
30–34	1.84 (1.06–3.18)	1.49 (1.23–1.80)
≥35	1.66 (1.00–2.74)	1.49 (1.21–1.84)
<b>Infant birth weight</b>		
< 4500	ref	ref
≥ 4500	0.66 (0.33–1.29)	1.23 (0.87–1.76)
<b>Infant head circumference</b>		
< 36	ref	ref
≥ 36	0.86 (0.59–1.25)	1.07 (0.88–1.29)

VD = vaginal delivery; CS = cesarean section; BMI = body mass index.

\*Adjusted for maternal age, gestational length, BMI at term, current BMI, infant birth weight and infant head circumference.