

Effect of spontaneous pushing versus Valsalva pushing in the second stage of labour on mother and fetus: a systematic review of randomised trials

M Prins, J Boxem, C Lucas, E Hutton

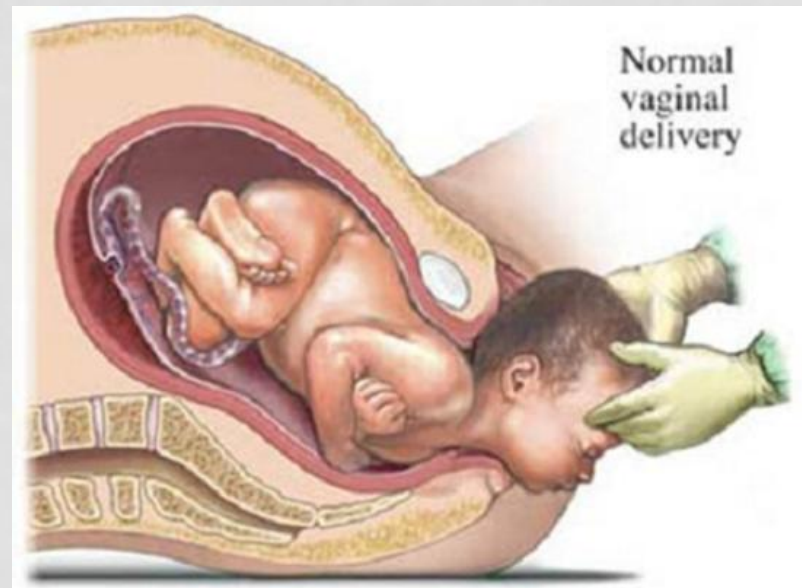
BJOG. 2011 MAY;118(6):662-70. doi:
10.1111/j.1471-0528.2011.02910.X.

EPUB 2011 Mar 10.

引言人:斯莉婷 日期:103.07.01

臨床現況

- 產婦於第二產程時開始閉氣用力，使胎頭下降至骨盆腔。
 - 方法-每次子宮收縮開始，深吸氣後憋氣(Valsalva pushing)，往下用力持續10秒，吐氣後再重覆一次
 - 可能產生的問題-
 - 臉紅脖子粗
 - 臉部微血管破裂
 - 骨盆底肌肉受傷
 -



Valsalva maneuver

- 持續閉氣用力(Valsalva maneuver)是日常生活中常見的動作，例如：搬起或運送移動重物、解使用力、咳嗽、嘔吐、噴嚏...等等。深呼吸之後閉氣用力，這個動作會造成會厭軟骨關閉、胸腹腔內的壓力上昇、進而出現一系列心臟血管的血液動力變化，包括下列的各個時期：

時期	動作	心臟的血液輸出量	血壓	機制
I	用力初期	不變	上昇	胸腹壓力向周邊血管傳導
II	持續用力	下降	下降	靜脈回流減少
III	用力剛解除	下降	下降	血液會積蓄在肺部中
IV	恢復期	正常或上昇	正常或上昇	1.靜脈回流增加 2.副交感神經活性上升

一定要閉氣用力嗎？

- Valsalva pushing vs. spontaneous pushing 對於母親及新生兒的利弊



Valsalva pushing

[HTTPS://WWW.YOUTUBE.COM/WATCH?V=GMZWU6YRYC0](https://www.youtube.com/watch?v=GMZWU6YRYC0)



spontaneous pushing

<https://www.youtube.com/watch?v=yreizEBmwHs>

SELECTED ARTICLE

DOI: 10.1111/j.1471-0528.2011.02910.x
www.bjog.org

Systematic review

Effect of spontaneous pushing versus Valsalva pushing in the second stage of labour on mother and fetus: a systematic review of randomised trials

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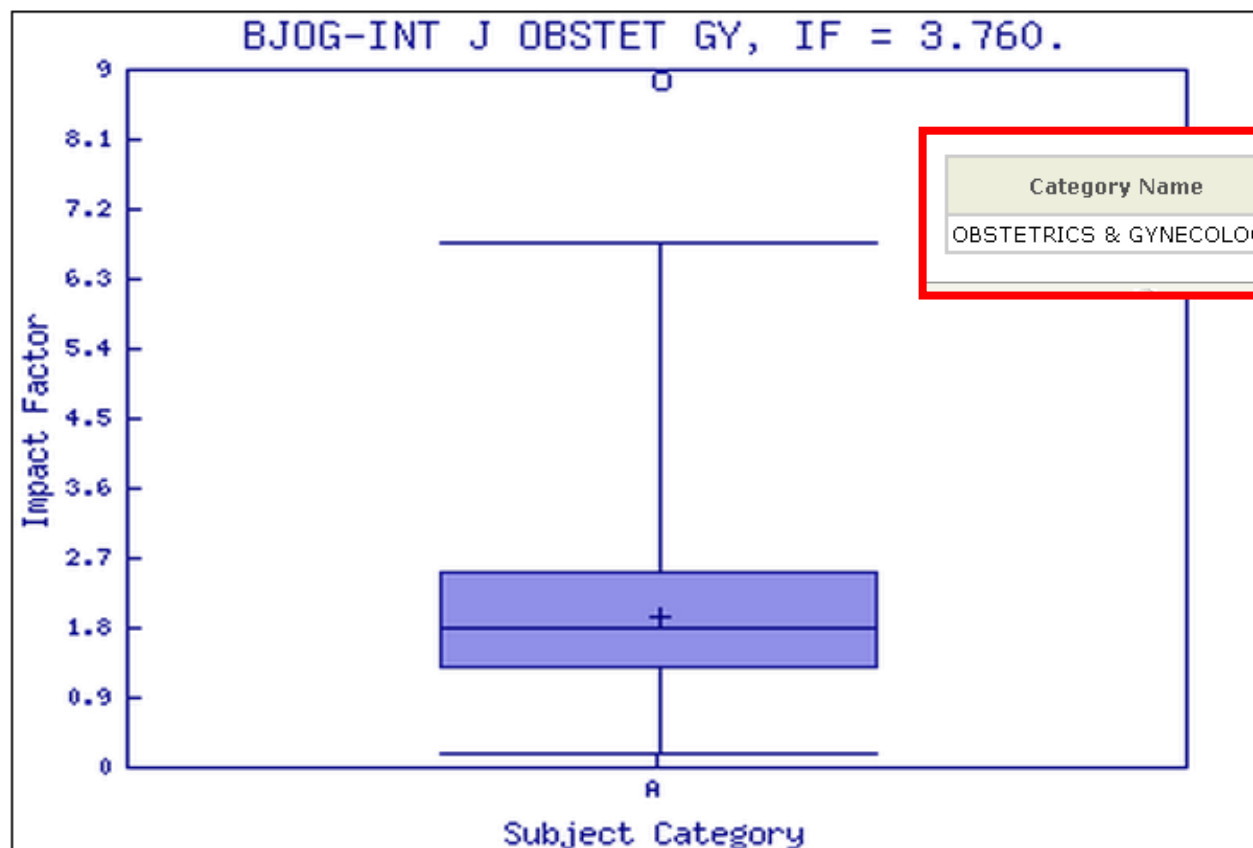
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Accepted 16 December 2010. Published Online 10 March 2011.

BJOG: AN INTERNATIONAL JOURNAL OF OBSTETRICS AND GYNAECOLOGY



Category Name	Total Journals in Category	Journal Rank in Category	Quartile in Category
OBSTETRICS & GYNECOLOGY	78	7	Q1



Key

A - OBSTETRICS & GYNECOLOGY

步驟 1：系統性文獻回顧探討的問題為何？

研究族群/問題	low-risk, healthy, nulliparous women with uncomplicated pregnancy 36 weeks
介入措施(Intervention)	spontaneous pushing
比較(Comparison)	Valsalva pushing
結果(Outcome)	<p>primary outcome</p> <ul style="list-style-type: none">• Instrumental/operative delivery <p>Other outcomes</p> <ul style="list-style-type: none">• Length of labour, Caesarean section, Episiotomy• Perineal/vaginal/anal sphincter laceration• Bladder function• Maternal satisfaction• Postpartum haemorrhage >1000ml <p>Infant outcomes</p> <ul style="list-style-type: none">• Low Apgar score <7 after 5 minutes• Umbilical arterial pH <7.2, Need for intubation,• Admission to neonatal intensive care unit,• Serious neonatal morbidity

步驟 2：系統性文獻回顧的品質如何？ (FAITH)

F - 研究是否找到 (Find) 所有的相關證據？

最好的狀況是？

良好的文獻搜尋至少應包括二個主要的資料庫(如：Medline, Cochrane考科藍實證醫學資料庫, EMBASE 等)，並且加上文獻引用檢索(參考文獻中相關研究、Web of Science, Scopus或 Google Scholar)、試驗登錄資料等。文獻搜尋應不只限於英文，並且應同時使用 MeSH字串及一般檢索詞彙(text words)。

- Included **published and unpublished randomised controlled trials**. We **had no language restriction**.
- Search in the **Cochrane library**, **MEDLINE** (1950 to May 2010), **EMBASE** (1980 to May 2010) and **CINAHL** (1982 to May 2010). We searched the **registers of ongoing clinical trials**, and performed an internet hand search to find **relevant master's theses and dissertations**. We **cross-checked** the reference lists of all relevant papers.
- The search terms included **Mesh headings and key words** linked with **Booleans OR, AND**.

評讀結果：■是 □否 □不清楚

步驟 2：系統性文獻回顧的品質如何？ (FAITH)

A - 文獻是否經過嚴格評讀 (Appraisal)？

最好的狀況是？應根據不同臨床問題的文章類型，選擇適合的評讀工具，並說明每篇研究的品質(如針對治療型的臨床問題，選用隨機分配、盲法、及完整追蹤的研究類型)

- **Quality was assessed** for randomisation (**sequence, blinding**), **loss to follow up**, **number of participants**, description of **intervention treatment and control treatment**, outcome variables, **blinding of outcome measurements**, **intention-to-treat analysis** and **blinding of analysis**.
- **Two reviewers** independently assessed the eligibility of the randomised controlled trials for inclusion; **any disagreement of assessment was discussed and resolved by consensus**.

評讀結果：■是 □否 □不清楚

步驟 2：系統性文獻回顧的品質如何？ (FAITH)

I - 是否只納入
(included) 具良好效度
的文章？

最好的狀況是？

僅進行文獻判讀是不足夠，系統性
文獻回顧只納入至少要有一項研究
結果是極小偏誤的試驗。

**The quality assessment data
sheet is available on request.**

評讀結果： ☐是 ☐否 ☒不清楚

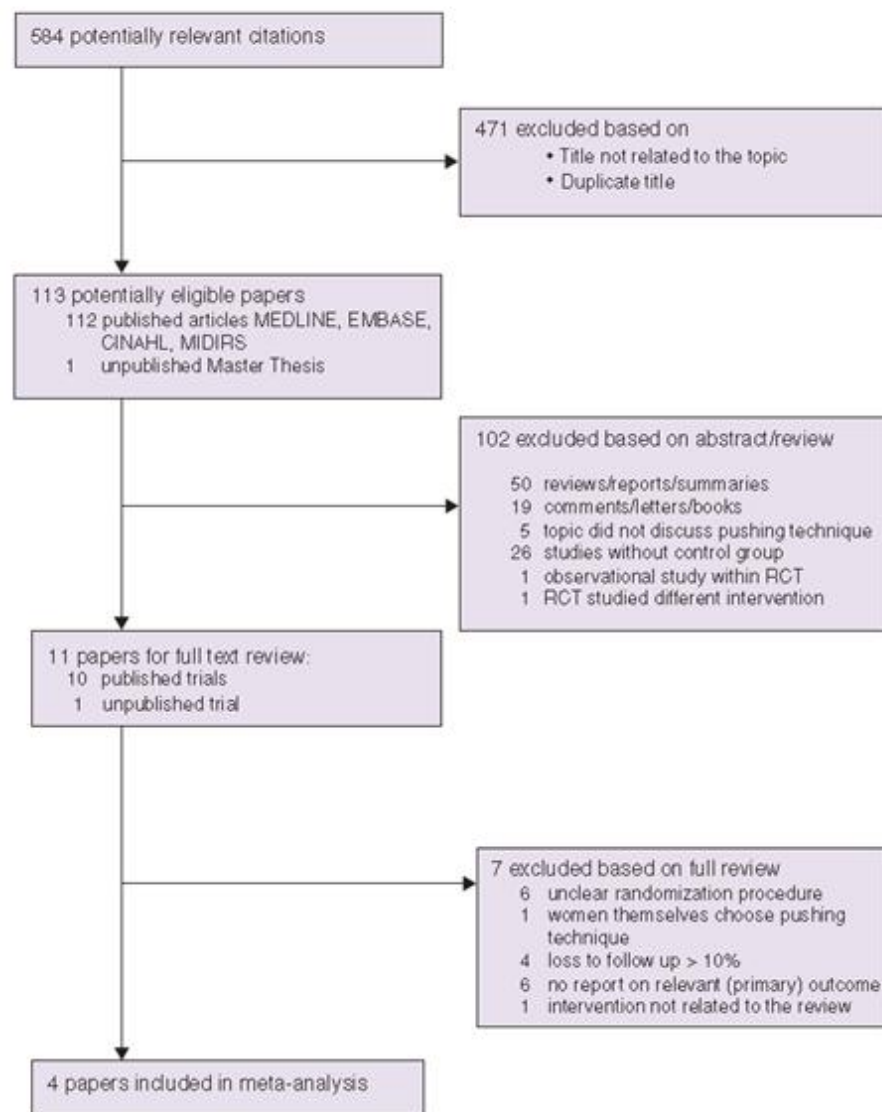


Figure 1. Results of literature review.

步驟 2：系統性文獻回顧的品質如何？ (FAITH)

T - 作者是否以表格和圖表「總結」 (total up) 試驗結果？

最好的狀況是？應該用至少 1 個摘要表格呈現所納入的試驗結果。若結果相近，可針對結果進行統合分析(meta-analysis)，並以「森林圖」(forest plot)呈現研究結果，最好再加上異質性分析。

3 studies, 425women
[RR] 0.70
95% CI 0.34–1.43

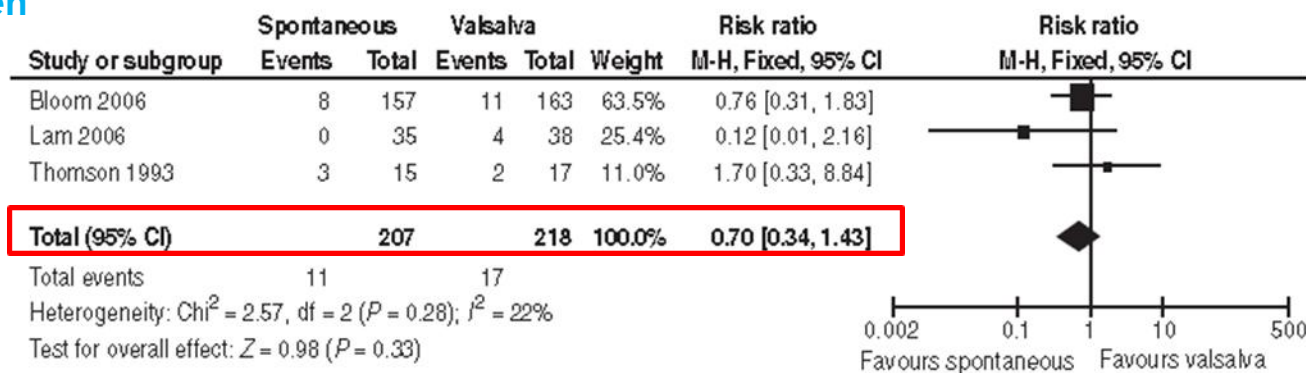


Figure 2. Meta-analysis results: instrumental/operative delivery.

no significant difference in instrumental/operative deliveries

步驟 2：系統性文獻回顧的品質如何？ (FAITH)

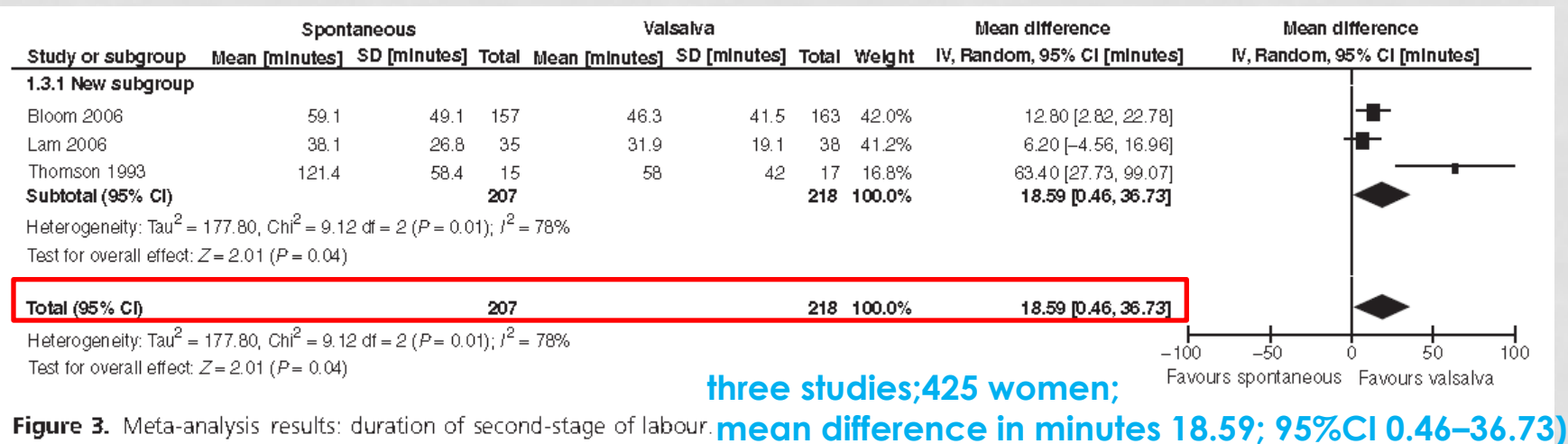


Figure 3. Meta-analysis results: duration of second-stage of labour. **mean difference in minutes 18.59; 95%CI 0.46-36.73**

- Because the I^2 was 78% a sensitivity analysis was carried out by consecutively removing the studies and rerunning the analysis.
- By excluding the study of Thomson the mean difference **was 9.75 minutes; 95% CI 2.43-17.06**

評讀結果：■是 □否 □不清楚

Table 1. Characteristics of the studies included comparing spontaneous pushing in the second stage of labour with the Valsalva pushing technique

Author, Study period, Country, Population	Criteria for inclusion/exclusion	Primary outcomes instrumental/ operative delivery		Other outcomes	Comments
		Intervention Spontaneous pushing	Control Valsalva pushing		
Thomson ³ 1993 UK <i>n</i> = 32	<p>Inclusion</p> <ul style="list-style-type: none"> primiparous women aged over 18 singleton pregnancy >37 weeks cephalic presentation absence of maternal or fetal condition that would affect the management of the second stage of labour <p>Exclusion</p> <ul style="list-style-type: none"> conception achieved <i>in vitro</i> baby was to be adopted use of epidural analgesia 	12/15 normal delivery	15/17 normal delivery	<p>No significant difference in</p> <ul style="list-style-type: none"> perineal trauma rate estimated maternal blood loss umbilical venous blood gas levels fetal acid-base status <p>Significantly longer second stage in the spontaneous pushing group.</p>	<p>Randomisation using a table with random numbers, women at least 6 cm dilated. Researcher who was present at the time of randomisation.</p> <p>Pushing started as soon as the second stage of labour was diagnosed.</p> <p>After 90 minutes of pushing the midwife could adopt whatever clinical management she thought would fit. Mean duration of second stage in spontaneous group was 121.4 minutes.</p> <p>Outcome data all complete.</p> <p><u>Intention-to-treat analysis.</u></p>
Schaffer ^{21*} 2005 USA <i>n</i> = 128 Hispanic women	<p>Inclusion</p> <ul style="list-style-type: none"> nulliparous women singleton pregnancy 36–41 weeks cephalic presentation uncomplicated pregnancy cervical dilatation at least 4 cm <p>Exclusion</p> <ul style="list-style-type: none"> use of oxytocin use of epidural analgesia history of urinary/anal incontinence history of pelvic organ prolapse estimated fetal weight >4000 g any known complication of pregnancy 			<p>No significant difference in</p> <ul style="list-style-type: none"> maximum urethral closure functional urethral length positive Valsalva leak point pressure maximum flow rate ml/sec detrusor pressure at peak flow detrusor overactivity urodynamic stress incontinence <p>Significant difference</p> <ul style="list-style-type: none"> decreased bladder capacity in Valsalva group decreased first urge to void in Valsalva group 	<p>Allocation to study group adequate, see Bloom's study. Three months postpartum women were invited to undergo urodynamic tests. 61 from the intervention and 67 from the control group participated</p> <p>Urogynaecology nurse practitioners measured outcomes and were blinded to obstetric management</p>

Table 1. (Continued)

Author, Study period, Country, Population	Criteria for inclusion/exclusion	Primary outcomes instrumental/ operative delivery		Other outcomes	Comments
		Intervention Spontaneous pushing	Control Valsalva pushing		
Bloom ^{20*} 2006 USA n = 325 Hispanic women	<p>Inclusion</p> <ul style="list-style-type: none"> nulliparous women singleton pregnancy 36–41 weeks cephalic presentation uncomplicated pregnancy cervical dilatation at least 4 cm regular uterine contractions <p>Exclusion</p> <ul style="list-style-type: none"> use of oxytocin use of epidural analgesia estimated fetal weight >4000 g known complication of pregnancy history of urinary/anal incontinence history of pelvic organ prolapse 	7/157 instrumental 1/157 caesarean section	6/163 instrumental 5/163 caesarean section	<p>No significant difference in</p> <ul style="list-style-type: none"> umbilical arterial pH Apgar score <7 Admission to neonatal intensive care unit perineal lacerations <p>Significantly longer second stage in the spontaneous pushing group</p>	<p>Computer generated randomisation in blocks of ten women at the onset of the second stage</p> <p>Pushing started as soon as the second stage of labour was diagnosed.</p> <p>Data outcome 320/325 complete <u>Intention-to-treat analysis</u></p>
Lam ¹⁹ 2006 Hong Kong n = 78, Chinese women	<p>Inclusion</p> <ul style="list-style-type: none"> woman between 18 and 40 years experienced their first birth healthy singleton fetus ≥37 weeks cephalic presentation uncomplicated pregnancy <p>Exclusion</p> <ul style="list-style-type: none"> medical or obstetric complication which would affect the management of the second stage of labour use epidural analgesia 	0/35 instrumental (0/35 caesarean section)	4/38 instrumental (0/38 caesarean section)	<p>No significant difference in</p> <ul style="list-style-type: none"> length of the second stage of labour umbilical arterial pH Apgar score <7 admission to neonatal intensive care unit perineal lacerations <p>Significantly more instrumental deliveries in the Valsalva group</p>	<p>Block randomisation on balanced blocks of ten at the onset of the second stage</p> <p>After 60 minutes of pushing the woman was changed over to the clinical management policy of the unit</p> <p>Pushing started when the fetal head was at +1 level of the ischial spines</p> <p>Control group 3/41 incomplete forms. In the experimental group 2/37 incomplete forms <u>Intention-to-treat analysis</u></p>

*Bloom and Schaffer used the same data from the trial with 325 women but presented different variables. In the table and meta-analysis we counted these women once.

步驟 2：系統性文獻回顧的品質如何？FAITH

Table 2. Summary of the meta-analysis of maternal and fetal outcomes

Outcome or subgroup	Studies	Participants	Statistical method*	Effect estimate	P value	Heterogeneity I ² (%)
Operative or instrumental delivery	3	425	Risk Ratio (M-H, Fixed, 95% CI)	0.70 (0.34–1.43)	0.33	27
Duration of second stage defined from full dilatation (minutes)	3	425	Mean difference (IV, Random, 95% CI [minutes])	18.59 (0.46–36.73)	0.04	78
Episiotomy	1	320	Risk ratio (M-H, Fixed, 95% CI)	0.79 (0.53–1.19)	0.26	
pH arterial <7.20 mmol/l	1	320	Risk ratio (M-H, Fixed, 95% CI)	0.65 (0.22–1.94)	0.44	
Mean venous pH	1	32	Mean difference (IV, Fixed, 95% CI)	–0.05 (–0.12 to 0.02)	Not applicable	
Mean arterial pH	1	320	Mean difference (IV, Fixed, 95% CI)	0.00 (–0.22 to 0.22)	1.00	
Mean Apgar score after 5 minutes	1	73	Mean difference (IV, Fixed, 95% CI)	0.00 (–0.23 to 0.23)	1.00	
Apgar score <7 after 5 minutes	2	393	Risk ratio (M-H, Fixed, 95% CI)	0.35 (0.01–8.43)	0.51	Not applicable
Any perineal repair	2	352	Risk ratio (M-H, Random, 95% CI)	0.95 (0.64–1.40)	0.79	57
Third- or fourth-degree tear	1	320	Risk ratio (M-H, Fixed, 95% CI)	0.87 (0.45–1.66)	0.66	
Need for resuscitation	2	352	Risk ratio (M-H, Fixed, 95% CI)	0.83 (0.40–1.75)	0.63	0
Admission to neonatal intensive care unit	2	393	Risk ratio (M-H, Fixed, 95% CI)	1.08 (0.30–3.79)	0.91	0
Mean estimated blood loss	2	105	Mean difference (IV, Fixed, 95% CI)	9.72 (–37.84 to 57.29)	0.69	0
First urge to void	1	128	Mean difference (IV, Fixed, 95% CI)	41.50 (8.40–74.60)	0.01	
Bladder capacity	1	128	Mean difference (IV, Fixed, 95% CI)	54.60 (13.31–95.89)	0.010	

*IV, inverse variance; M-H, Mantel–Haenszel.

H - 試驗的結果是否相近
- 異質性
(Heterogeneity) ?

最好的狀況是？在理想情況下，各個試驗的結果應相近或具同質性，若具有異質性，作者應評估差異是否顯著(卡方檢定)。根據每篇個別研究中不同的PICO及研究方法，探討造成異質性的原因。

I² was 57% one study;
320 women; RR 1.25,
95% CI 0.76–2.06

評讀結果：■是 □否 □不清楚

結論

- 不支持常規使用Valsalva pushing
 - 降低產後3個月的膀胱容量，可能對骨盆底功能有負面影響(Schaffer, 2005))
- Spontaneous pushing
 - 鼓勵產婦選擇適合自己的用力方法
 - 支持產婦自己的身體感受，隨著往下的壓力感，持續用力
 - 護理人員提供明確的用力指導和協助

討論(1)

- 臨床上曾試過spontaneous pushing，但產婦用力的強度及效果不佳，產前就需要開始練習，很難短時間學會
- 生產後的解尿困難，與產程時間長、使用產鉗及吸引器生產、膀胱脹太久或會陰4度裂傷...等因素，可能會使膀胱神經鈍化，影響解尿功能，但本研究結果顯示，差異量不大

討論(2)

- 是否在第二產程時，建議產婦使用 Spontaneous pushing ?

- ✓ 同意7人
- ✓ 懷疑11人
- ✓ 不同意1人



THANK YOU!

