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# Primary care

## Effects of acupuncture and stabilising exercises as adjunct to standard treatment in pregnant women with pelvic girdle pain: randomised single blind controlled trial

Helen Elden, Lars Ladfors, Monika Fagevik Olsen, Hans-Christian Ostgaard, Henrik Hagberg

### Abstract

**Objectives** To compare the efficacy of standard treatment, standard treatment plus acupuncture, and standard treatment plus stabilising exercises for pelvic girdle pain during pregnancy.

**Design** Randomised single blind controlled trial.

**Settings** East Hospital, Gothenburg, and 27 maternity care centres in Sweden.

**Participants** 386 pregnant women with pelvic girdle pain.

**Interventions** Treatment for six weeks with standard treatment (n = 130), standard treatment plus acupuncture (n = 125), or standard treatment plus stabilising exercises (n = 131).

**Main outcome measures** Primary outcome measure was pain (visual analogue scale); secondary outcome measure was assessment of severity of pelvic girdle pain by an independent examiner before and after treatment.

**Results** After treatment the stabilising exercise group had less pain than the standard group in the morning (median difference = 9, 95% confidence interval 1.7 to 12.8; P = 0.0312) and in the evening (13, 2.7 to 17.5; P = 0.0245). The acupuncture group, in turn, had less pain in the evening than the stabilising exercise group (-14, -18.1 to -3.3; P = 0.0130). Furthermore, the acupuncture group had less pain than the standard treatment group in the morning (12, 5.9 to 17.3; P < 0.001) and in the evening (27, 13.3 to 29.5; P < 0.001).

Attenuation of pelvic girdle pain as assessed by the independent examiner was greatest in the acupuncture group.

**Conclusion** Acupuncture and stabilising exercises constitute efficient complements to standard treatment for the management of pelvic girdle pain during pregnancy.

Acupuncture was superior to stabilising exercises in this study.

### Introduction

Pelvic girdle pain is a common complaint among pregnant women worldwide,<sup>1</sup> and it causes severe pain in one third of affected women.<sup>2-3</sup> Strenuous work, previous low back pain, and previous pelvic girdle pain are known risk factors.<sup>3,4</sup>

Pelvic girdle pain generally arises in relation to pregnancy, trauma, or reactive arthritis. Pain is experienced between the posterior iliac crest and the gluteal fold, particularly in the vicinity of the sacroiliac joints. The pain may radiate in the posterior thigh and can occur in conjunction or separately in the symphysis. The endurance capacity for standing, walking, and sitting is diminished.<sup>5</sup> After pregnancy problems remain serious in about

7% of women with pelvic girdle pain, causing severe discomfort and reducing ability to work.<sup>3,6</sup>

Most studies do not distinguish between lumbar back pain and pelvic girdle pain.<sup>6</sup> Unspecified diagnosis is a problem, as the two conditions differ with respect to prognosis and treatment.<sup>2,5</sup> If a patient with pelvic girdle pain is treated for low back pain, the symptoms may be aggravated.<sup>2</sup> Pelvic girdle pain must be reproducible by specific pain provocation tests.<sup>3,5,7,8</sup> The posterior pelvic pain provocation test and Patrick's fabere test have the best sensitivity if pain is evident in the sacroiliac joints.<sup>5,8</sup> Modified Trendelenburg's test and palpation of the symphysis have better sensitivity if pain is evident in the symphysis pubis.<sup>8</sup> These tests have high intertester reliability.<sup>5,8</sup>

Standard treatment may consist of a pelvic belt, a home exercise programme, and patient education. A systematic review has shown that the efficacy of these interventions remains questionable.<sup>9</sup> Current treatment increasingly includes stabilising exercises and acupuncture.<sup>10-12</sup> However, insufficient evidence is available to give strong recommendations for or against any particular treatment modality for pelvic girdle pain.<sup>13</sup> We compared the efficacy of acupuncture or stabilising exercises as an adjunct to standard treatment with standard treatment alone for the treatment of pelvic girdle pain in pregnant women.

### Methods

The study was a randomised single blind trial done at East Hospital, Sahlgrenska Academy, and at 27 maternity care centres in the hospital's reference area in Gothenburg, Sweden, from 2000 to 2002.

#### Participants

Doctors and midwives at the 27 maternity care centres preselected consecutive patients. Participants filled in a previously validated questionnaire<sup>2</sup> and a diary for baseline information for one week before the inclusion visit. An independent specially trained physiotherapist then assessed patients who were eligible and willing to participate in the study. This assessment included a detailed standardised physical examination and collection of baseline data. The tests used were the posterior pelvic pain provocation test, Patrick's fabere test, a modified Trendelenburg's test, Lasegue test, and palpation of the symphysis pubis.<sup>5,8</sup> The main inclusion criteria were healthy women at 12-31 completed gestational weeks, well integrated in the Swedish language, with singleton fetuses and defined pregnancy related pelvic girdle pain. We excluded patients with other pain conditions, systemic disorders, or contraindications to treatment. The participants gave informed consent.

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### Randomisation

A research assistant used a computer generated random table to determine the allocation sequence before the study. Groups were coded, and the allocation was transferred to a series of presealed opaque envelopes. The independent examiner randomised the patients individually after doing the baseline assessment.

### Treatment protocols

The study comprised a one week baseline period, six weeks of treatment, and follow up one week after the last treatment. Patients were asked to avoid other treatments during the intervention period. Three experienced physiotherapists gave standard treatment, two experienced medical acupuncturists did acupuncture, and two experienced physiotherapists gave the stabilising exercises. All possible adverse events were recorded.

**Standard treatment group**—Standard treatment consisted of general information about the condition and anatomy of the back and pelvis. Adequate advice was given about activities of daily living. The physiotherapist made sure that the patient understood and respected the relation between impairment, load demand, and actual loading capacity as well as the importance of necessary rest. The purpose of this information was to reduce fear and to enable patients to become active in their own treatment. The patients were given a pelvic belt (Puff Igång AB, Sweden) and a home exercise programme designed to increase strength in the abdominal and gluteal muscles.

**Acupuncture**—Patients received the same treatment as in the standard group but in addition had acupuncture. Local acupuncture points were selected individually after diagnostic palpation to identify sensitive spots.<sup>14</sup> A total of 10 segmental points and seven extrasegmental points were used (table 1). The needles (Hegu AB, Landsbro, Sweden) were made of stainless steel (Ø 0.30) and inserted intramuscularly to a depth of 15-70 mm to evoke needle sensation (De Qi), described as tension, numbness, and often a radiating sensation from the point of insertion, reflecting activation of muscle-nerve afferents. The needles were left in situ for 30 minutes and manually stimulated every 10 minutes. Treatment was given twice a week over six weeks. Fetal heart rate and maternal heart rate and blood pressure were monitored before and after all treatments.

**Stabilising exercises**—Patients received the same treatment as in the standard group but in addition did stabilising exercises modified because of the pregnancy (box).<sup>15 16</sup> The training programme started by emphasising activation and control of local deep lumbopelvic muscles. Training of more superficial muscles in dynamic exercises to improve mobility, strength, and endurance capacity was gradually included. Patients received treatments individually for a total of six hours during six weeks. They were told to integrate the exercises in daily activities and to exercise in short sessions on several occasions during the day.

### Outcome measures

**Primary outcome measure**—Patients scored their current intensity of pelvic pain related to motion on a 100 point visual analogue scale every morning and every evening in the diaries.

**Secondary outcome measures**—The independent examiner assessed recovery from symptoms. Patients were asked not to reveal any information about their treatment during assessment.

### Statistical analysis

When planning the study we assumed that the mean pain score related to motion at baseline would be 60 mm (visual analogue scale) in all three groups. We did not expect any treatment effect in the standard group. We assumed that the mean pain score after treatment would be 60 mm in the standard group, 50 mm

**Table 1** Acupuncture points, their anatomical position, and their innervation

Points	Segmental innervation	Muscle localisation
GV 20	Nn trigeminus (V), occipitalis minor (C2), occipitalis major (C2-3)	Aponeurosis epicrani tissue
LI 4 bilateral	Nn ulnaris medianus (C8, Th 1)	Mm interosseus dorsalis I, lumbricalis II, adductor pollicis
BL 26 bilateral	Nn thoracodorsalis (C6-8), toracicus (Th 9-12), lumbalis (L1-3)	Nn thoracolumbalis, m erector spinae
BL 32 bilateral	Nn thoracodorsalis (C6-8), toracicus (Th 9-12), lumbalis (L1-3)	Fascia thoracolumbalis, m erector spinae
BL 33 bilateral	Nn thoracodorsalis (C6-8), toracicus (Th 9-12), lumbalis (L1-3)	Fascia thoracolumbalis, m erector spinae
BL 54 bilateral	Nn gluteus inferior (L5, S1-2)	M gluteus maximus
KI 11 bilateral	Nn thoracicus (Th 6-12), subcostalis	Vagina m recti abdominis
BL 60 bilateral	Nn suralis (S2)	Fibrotic tissue
EX 21 bilateral	Nn lumbalis, sacralis (L4-5, S1-2)	Fascia thoracolumbalis, m erector spinae
GB 30 bilateral	Nn gluteus inferior (L5, S1), obturatorius internus (L4-5, S1)	Mm gluteus maximus, gemellus superior, piriformis
SP 12 bilateral	Nn thoracicus, lumbalis (Th 7-12, L1)	Aponeurosis mm obliquus externus, abdominus internus
ST 36 bilateral	Nn peroneus profundus (L4-5)	M tibialis anterior

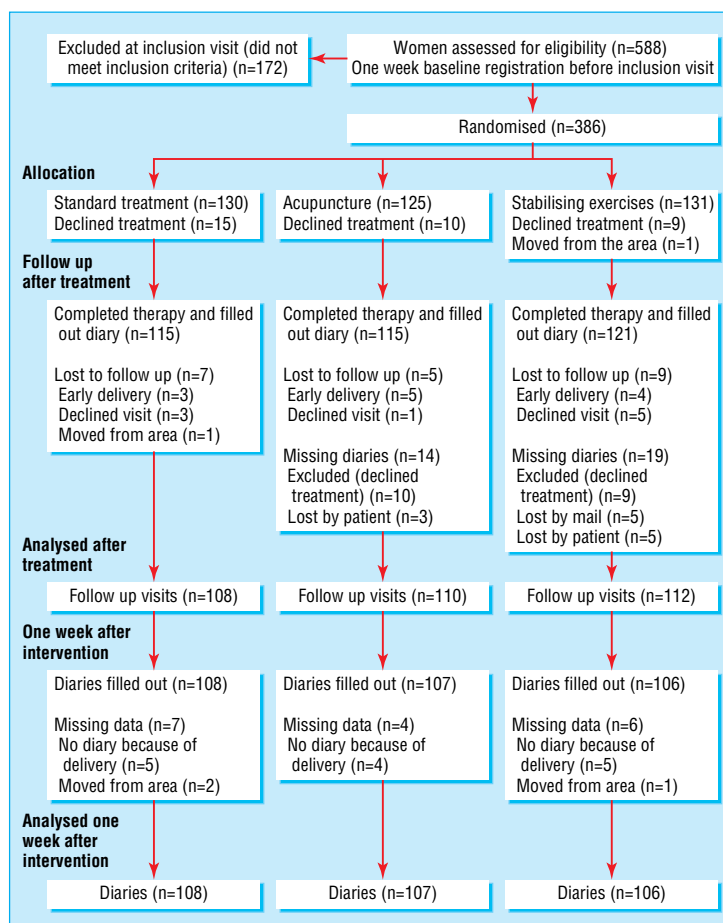
BL=bladder channel; EX=extra channel; GB=gall bladder channel; GV=governor vessel channel; KI=kidney channel; LI=large intestine channel; SP=spleen channel; ST=stomach channel.

in one treatment group, and 40 mm in the other. To achieve a 90% power of detecting a significance (at the two sided 5% level), with an assumed standard deviation of 40, we needed 103 patients for each study group. To compensate for an anticipated loss to follow up of 20%, we needed 386 patients. When analysing the study we decided to apply the Mann-Whitney U test for comparing changes in pain scores, with Bonferroni's correction (P values multiplied by three). The power achieved with that method was 86% for comparison of the two most extreme groups, provided that  $\Delta\mu/\sigma = 0.5$ .

Personnel from an independent institution coded all results from the study and entered them into a database. Analyses were done by intention to treat. The statistician who did the analysis was blinded to group and treatment. In the analysis of the pain diaries we defined the median visual analogue scale baseline levels in the mornings and in the evenings for each patient by cal-

#### Treatment protocol for the stabilising exercise group

- Additional information about anatomy and the genesis of pelvic pain and discussion about how the basic written regimen could be integrated at home and at work
- Exercises for stabilising the pelvis and back—that is, training of the transversus abdominis and the multifides facilitated by contractions of the pelvic floor muscles, according to Richardson and Jull but modified because of the pregnancy.<sup>16</sup> While the standard positions (prone and supine) were not suitable the pressure biofeedback unit was not used. The exercises were done with the patient lying on her side, four point kneeling, sitting, and standing. Arm and leg movements were added when the basic movement was correct
- Exercises for increasing circulation in hip rotator muscles. The exercises were done with many repetitions during low force and in a limited range of motion in a side lying position with a pillow between the legs or sitting without foot support
- Massage, effleurage, and petrissage of hip extensors and rotators
- Stretching of hip external rotators and extensors in the sitting position: 20 sec/stretch



Participants' progress through trial and withdrawals

culating the median for the days before treatment (five to seven days). The same calculations of median pain were done for the first week after the end of treatment.

We calculated the medians, quartiles, means, and standard deviations when possible. We used the Mann-Whitney U test to compare differences between the groups for continuous variables and  $\chi^2$  for categorical variables. Adjustments (multiplication by three) of the P values due to multiple comparisons were done by Bonferroni's method. We considered an adjusted P value  $< 0.05$  to be statistically significant. We calculated median differences and confidence intervals for the differences between medians on the basis of the Mann-Whitney U test. We also did an analysis of treatment effects in patients divided into four subgroups<sup>17</sup>: one sided sacroiliac pain, double sided sacroiliac pain, one sided sacroiliac pain plus symphysis pubis pain, and pelvic girdle syndrome (double sided sacroiliac pain plus symphysis pubis pain). The results were analysed with the SAS software package, version V8.

## Results

### Randomisation and progress through the trial

Of 558 women referred for the first assessment, 172 did not meet the inclusion criteria; 386 women were included in the trial. Baseline characteristics were similar in the three treatment groups (table 2). The figure show the progress of patients through the trial and withdrawals from the study.

### Primary outcome measure

Table 3 shows improvements in pain scores. The reduction in pain was most pronounced in the evening in the acupuncture group one week after the end of treatment, compared with the other treatment groups.

**Table 2** Characteristics of 386 pregnant women with pelvic girdle pain included in trial. Values are numbers (percentages) unless stated otherwise

Characteristic	Standard group (n=130)	Acupuncture group (n=125)	Stabilising exercise group (n=131)
Mean (SD) maternal age (years)	30.8 (4.8)	30.6 (4)	30.0 (4)
Gestation weeks (+ days) at inclusion	24 (+3)	24 (+3)	24 (+3)
First pregnancy	33 (25)	34 (27)	36 (27)
Full time work	71 (55)	72 (58)	76 (58)
Smoker	12 (9)	11 (9)	13 (10)
Previous low back pain	90 (69)	89 (71)	84 (64)
Physical activity during leisure before pregnancy:			
Not at all	29 (22)	36 (29)	30 (23)
Once a week	52 (40)	37 (30)	50 (38)
More than twice a week	50 (38)	51 (41)	50 (38)
Lifting heavy objects >10 times a day	59 (45)	50 (40)	53 (40)
No or rare ability to take rest breaks	76 (58)	72 (58)	68 (52)

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**Table 3** Primary outcome measure: pain on visual analogue scale related to motion one week after treatment compared with baseline measurements

Pain	Standard group		Acupuncture group		Stabilising exercise group		Group comparisons		
	No	Median (25th-75th centile)	No	Median (25th-75th centile)	No	Median (25th-75th centile)	Comparison	Differences of medians (95% CI)	*P values
Morning:									
Baseline	131	23 (13-41)	125	23 (15-44)	130	22 (13-43)			
One week after treatment	108	27 (12-58)	107	15 (7-29)	106	18 (9-37)	S-ACU:	12 (5.9 to 17.3)	<0.001
							S-SE:	9 (1.7 to 12.8)	0.0312
							ACU-SE:	-3 (-7.8 to 0.3)	NS
Evening:									
Baseline	131	63 (49-75)	125	65 (47-76)	130	60 (4-73)			
One week after treatment	108	58 (40-74)	107	31 (12-58)	106	45 (21-68)	S-ACU:	27 (13.3 to 29.5)	<0.001
							S-SE:	13 (2.7 to 17.5)	0.0245
							ACU-SE:	-14 (-18.1 to -3.3)	0.0130

ACU=acupuncture; S=standard; SE=stabilising exercise.

\*P values from Mann-Whitney U test. All original two tailed P values were multiplied by three (Bonferroni's correction); NS=not significant.

**Secondary outcome measures**

Table 4 shows that attenuation of pelvic girdle pain assessed by the independent examiner was greatest in the acupuncture group. Three of four subgroups of pelvic girdle pain improved after acupuncture compared with standard treatment and one of four subgroups improved compared with the stabilising exercise group.

**Side effects**

No serious complication occurred during treatments or during the follow up period after any of the treatments.

**Discussion**

The main finding of this study was that acupuncture or stabilising exercises as an adjunct to standard treatment offer clear clinical advantages over standard treatment alone for reduction of pain in pregnant women with pelvic girdle pain. This is supported by the patients' own estimates and by independent examiners. Our results are also supported by earlier findings that showed beneficial effects of stabilising exercises for women with pelvic girdle pain after pregnancy.<sup>10</sup> The training was aimed at affecting dysfunction of the muscle-tendon-fascia system that controls force closure of the pelvis.<sup>18</sup> Exactly how the exercises influence this system is unknown, but research has shown that

contraction of the transversus abdominis decreases the laxity of the sacroiliac joint.<sup>15</sup> The exercises were intended to affect mainly the local stability system, but whether an addition of global stabilising muscle exercises could have provided the same effect is not known. No major differences exist between current recommendations about stabilisation training and the exercises given in our study.<sup>15</sup>

The stabilisation group also got additional treatment of stretching exercises of specific muscles plus massage. All treatment was more or less multifactorial, and massage as well as stretching may have had some contributory effect. The main training, however, was the stabilisation exercises that were done on several occasions during the day, in contrast to the stretching or massage that was done only at the visits.

In the standard group, pain remained constant during treatment. This is in line with earlier findings that indicated lack of evidence for the effect of standard treatment for pelvic girdle pain.<sup>9</sup>

Previous studies of acupuncture for low back pain in pregnancy reported pain relief.<sup>11 12</sup> However, these studies had methodological shortcomings, as the type of back pain was not clearly defined. Furthermore, the acupuncture stimulation given previously must be considered weak compared with that given in studies of men and non-pregnant women. One of the studies

**Table 4** Secondary outcome measure: assessment of severity of pelvic girdle pain by an independent examiner before intervention and at follow up after last treatment. Values are numbers (percentages) unless stated otherwise

Tests for assessment of pelvic girdle pain	Standard group		Acupuncture group		Stabilising exercise group		Group comparisons after treatment	
	Inclusion (n=130)	Follow up (n=108)	Inclusion (n=125)	Follow up (n=110)	Inclusion (n=131)	Follow up (n=112)	Comparison	P value*
Positive pain drawing	130 (100)	100 (93)	125 (100)	94 (85)	131 (100)	97 (87)		NS
Posterior pelvic pain provocation test	130 (100)	92 (85)	125 (100)	72 (65)	131 (100)	95 (85)	ACU-S	0.0021
							ACU-SE	0.0024
Pain when turning in bed	130 (100)	95 (88)	125 (100)	73 (66)	131 (100)	80 (71)	ACU-S	<0.001
							SE-S	0.0072
Palpation of pubic symphysis	47 (36)	50 (46)	51 (41)	32 (29)	62 (47)	39 (35)	ACU-S	0.0261
Patrick's fabere test	65 (50)	57 (53)	69 (55)	36 (33)	74 (56)	47 (42)	ACU-S	0.0084
Trendelenburg's test	51 (39)	43 (40)	52 (42)	30 (27)	45 (34)	30 (27)		NS
<b>Subgroups of pelvic girdle pain</b>								
Pelvic girdle syndrome	34 (26)	33 (31)	43 (34)	20 (18)	49 (37)	25 (22)		NS
Double sided sacroiliac pain	47 (36)	50 (46)	51 (41)	32 (29)	62 (47)	39 (35)	ACU-S	0.0261
One sided sacroiliac pain + symphysis pubis pain	49 (38)	45 (42)	51 (41)	23 (21)	63 (48)	36 (32)	ACU-S	0.0027
One sided sacroiliac pain	130 (100)	92 (85)	125 (100)	72 (65)	131 (100)	95 (85)	ACU-S	0.0021
							ACU-SE	0.0024

ACU=acupuncture; S=standard; SE=stabilising exercise.

\*P values from  $\chi^2$  test. All original P values were multiplied by three (Bonferroni's correction); NS=not significant.

**What is already known on this topic**

No cure exists for pelvic girdle pain during pregnancy

No studies have been published on the effects of acupuncture on well defined isolated pelvic girdle pain during pregnancy

**What this study adds**

This study shows large treatment effects on pain among pregnant women with well defined isolated pelvic girdle pain

Acupuncture was the treatment of choice for patients with one sided sacroiliac pain, one sided sacroiliac pain combined with symphysis pubis pain, and double sided sacroiliac pain

used almost exclusively auricular acupuncture<sup>11</sup>; the other study used segmental points, but no extrasegmental points were used and no further stimulation after insertion was given.<sup>12</sup> Our findings that acupuncture gives acceptable pain relief in women with pelvic girdle pain is supported by physiological models explaining the mechanisms in the treatment of both acute and chronic pain conditions.<sup>14 19</sup>

Pain may be inhibited at the segmental level, and for this reason the acupuncture points were selected in muscle segments according to the patient's pain drawing, including diagnostic palpation to identify sensitive spots. The aim was to establish control of pelvic pain to prevent dysfunction of muscles of the spine and pelvis. Speculatively, this was achieved through activation of both the segmental pain inhibitory system, involving the so called gate control mechanism, and the central pain inhibitory system, involving secretion of endogenous opioids.<sup>14</sup> Extrasegmental points to the lumbosacral area were used to strengthen and lengthen the effect of the central control systems. In addition, well known general pain relieving points were selected. Whether the choice of acupuncture points and the method of stimulation in this study are optimal remains to be elucidated.

Earlier research found that poor muscle function in the back and pelvis at the beginning of pregnancy is related to severe pain and disability throughout pregnancy.<sup>20</sup> This could be the case for the patients in the standard and stabilising exercise groups, as the acupuncture group estimated significantly lower visual analogue scale ratings after treatment. The acupuncture treatment succeeded in establishing control of the pain, and this may have been important in preventing dysfunction of muscles. Our results are supported by earlier research on acupuncture for low back pain.<sup>21 22</sup>

This study shows that methods other than structured physiotherapy may be effective in treating pelvic girdle pain in pregnancy and that acupuncture represents an effective alternative. A combination of several methods is probably even better. Each method needs to be evaluated individually, however, before combinations can be recommended for future research, and only after that should recommendations for treatment be made.

We conclude that acupuncture as well as stabilising exercises constitute effective complements to standard treatment for pregnant women with pelvic girdle pain. Acupuncture was superior to stabilising exercises in this study. The findings are of particular importance because no previous study has shown such marked treatment effects among pregnant women with well defined pelvic girdle pain.

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Contributors: HE initiated and coordinated the study, did most of the data collection, and contributed to study design and interpretation of results. LL advised on data collection and assisted in study design. MFO assisted in study design and gathered data. H-CO guided the scientific process, assisted in study design, provided advice on the epidemiology of pelvic girdle pain, and obtained funding. HH guided the scientific process and assisted in study design. All investigators contributed to data interpretation and preparation of the manuscript. H-CO and HH are the guarantors.

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Competing interests: None declared.

Ethical approval: The local ethics committee approved the study.

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