The prevalence, implications, and clinical course of pregnancy-related pelvic girdle pain

by

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Scientific Environment

Scientific environment

The work presented in this thesis was carried out in association with the Norwegian Centre for Movement Disorders and the Department of Obstetrics and Gynaecology at Stavanger University Hospital, Stavanger, Norway. The Research Department at Stavanger University Hospital has organized the necessary office facilities at Forskningens Hus.

My supervisors have been Jan Petter Larsen MD PhD, neurologist and professor at the University of Stavanger, and Kolbjørn Brønnick, PhD, psychologist and professor at Faculty of Health Sciences, University of Stavanger. My co-supervisor has been Inger Økland, MD PhD, obstetrician, Associate Professor II at the University of Stavanger, and Head of research at Stavanger University Hospital. The statistic work in all three papers has been guided, supervised, and conducted by Kolbjørn Brønnick.

Knut Andersen, chiropractor, PhD, has been co-author of all three papers, Inger Kjaermann, chiropractor, MSc, has been co-writing paper

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I and paper II, and Anne-Mari Gausel, chiropractor, PhD, was cowriting paper III.

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Background

Pelvic girdle pain (PGP) during pregnancy is common and, indeed, has always been considered normal. It is commonly associated with moderate to severe pain that impairs everyday activities such as getting up from a chair, bending, walking, working in the home and caring for children, as well as, of course, paid employment. Also, PGP is a frequent cause of sick leave during pregnancy. The aetiology of PGP is poorly understood and there is no official nomenclature, no effective evidence-based preventive measures or treatment, known risk factors or detailed knowledge of the clinical course of the various subgroups of this condition.

Objectives

The objectives for this project were to determine the prevalence of PGP during pregnancy in a random population of women, detect factors associated with the development of this condition, explore what influences taking sick leave due to PGP, and examine whether pregnant women with PGP, who have been sub-grouped on the basis

of two clinical tests, differ with regards to demographic characteristics and/or the clinical course of PGP during the second half of their pregnancy.

Methods

The thesis consists of three papers, based on two separate data collections at Stavanger University Hospital. Paper I and II originate from a retrospective cohort study conducted in 2009, in which women giving birth at Stavanger University hospital in a 4-month period were asked to fill in a questionnaire on demographic features, pain, disability, PGP, pain-related activities of daily living, sick leave in general and for PGP, frequency of exercising before and during pregnancy, and Oswestry Disability Index.

Inclusion criteria were singleton pregnancy of at least 36 weeks and competence in the Norwegian language. Drawings of the pelvic and low back area were used for the localization of pain. PGP intensity was then rated retrospectively on a numerical rating scale. Non-parametric tests, multinomial logistic regression and sequential linear regression analysis were used in the statistical analysis.

Paper III originate from a prospective longitudinal cohort study carried out in 2010. Inclusion criteria were the as for the retrospective data collection and took place at the second-trimester routine ultrasound examination. All eligible women (n=503) filled in questionnaires and answered a weekly SMS question during pregnancy until delivery. Women with pain in the pelvic area underwent a clinical examination following a test procedure recommended in the European guidelines for the diagnosis and treatment of PGP.

Results

Paper I report that nearly 50% of the women experienced moderate and severe PGP during pregnancy. Approximately half of them had PGP syndrome, whereas the other half experienced lumbopelvic pain. Ten percent of the women experienced moderate and severe LBP alone. These pain syndromes increased sick leave and impaired general level of function during pregnancy. Approximately 50% of women with PGP had pain in the area of the symphysis pubis. The analysis of risk factors did not present a unidirectional and clear picture.

In Paper II PGP is reported to be a frequent and major cause of sick leave during pregnancy among Norwegian women, which is also reflected in activities of daily living as measured with scores on all Oswestry disability index items. In the multivariate analysis of factors related to sick leave and PGP were work satisfaction, problems with lifting and sleeping, and pain intensity risk factors for sick leave. Also, women with longer education, higher work satisfaction and fewer problems with sitting, walking, and standing, were less likely to take sick leave in pregnancy, despite the same pain intensity as women being on sick leave. In Paper III, 42% (212/503) reported pain in the lumbopelvic region and 39% (196/503) fulfilled the criteria for a probable PGP diagnosis. 27% (137/503) reported both the posterior pelvic pain provocation (P4) and the active straight leg raise (ASLR) tests positive at baseline in week 18, revealing 7.55 (95% CI 5.54 to 10.29) times higher mean number of days with bothersome pelvic pain compared with women with both tests negative. They presented the highest scores for workload, depressed mood, pain level, body mass index, Oswestry Disability Index and the number of previous pregnancies. Exercising regularly before and during pregnancy was more common in women with negative tests.

Conclusions

Pelvic pain in pregnancy is a health care challenge in which moderate and severe pain develops rather early and has important implications for society. The observed associations between possible causative factors and moderate and severe LBP and PGP in the analysis of the retrospective data may, together with results from other studies, bring some valuable insights into their multifactorial influences and provide background information for future studies.

Some pregnant women with PGP show a higher pain tolerance, most likely dependant on education, associated with work situation and/or work posture, which decreases sick leave. These issues are recommended to be further examined in a prospective longitudinal study since they may have important implications for sick leave frequency during pregnancy.

If both P4 and ASLR tests were positive mid-pregnancy, a persistent bothersome pelvic pain of more than 5 days per week throughout the remainder of pregnancy could be predicted. Increased individual control over work situation and an active lifestyle, including regular exercise before and during pregnancy, may serve as a PGP prophylactic.

List of publications

List of publications

Paper I

Malmqvist S, Kjaermann I, Andersen K, Økland I, Brønnick K, Larsen JP. Prevalence of low back and pelvic pain during pregnancy in a Norwegian population. J Manipulative Physiol Ther. 2012;35:272-8.

Paper II

Malmqvist S, Kjaermann I, Andersen K, Økland I, Larsen JP, Brønnick K. The association between pelvic girdle pain and sick leave during pregnancy; a retrospective study of a Norwegian population.

BMC Pregnancy Childbirth. 2015;15:23

Paper III

Malmqvist S, Kjaermann I, Andersen K, Gausel AM, Økland I, Larsen JP, Brønnick K. Can a bothersome course of pelvic pain from midpregnancy to birth be predicted? A Norwegian prospective longitudinal SMS-track study. BMJ Open 2018;8:e021378.

Abbreviations

Abbreviations

Active straight leg raise test ASLR

Activities of daily living ADL

Analysis of variance ANOVA

Body Mass Index BMI

Long dorsal sacroiliac ligament test LDL

Low back pain LBP

Numerical rating scale NRS

Oswestry Disability Index ODI

Pelvic girdle pain PGP

Pelvic Girdle Questionnaire PGQ

Pelvic pain PP

Posterior pelvic pain provocation test P4

Roland-Morris Disability Questionnaire RMDQ

Sacroiliac joint SIJ

Short message service SMS

Transversus abdominis muscle TrA

Visual Analogue Scale VAS

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1 Introduction

Pelvic girdle pain (PGP) during pregnancy is common and, indeed, has always been considered normal. However, from the World Health Organization perspective, which defines health as "a state of complete physical, mental, and social well-being and not merely the absence of disease or infirmity", women who experience PGP are not healthy [1]. This condition is commonly associated with moderate to severe pain that impairs everyday activities such as getting up from a chair, bending, walking, working in the home and caring for children, as well as, of course, paid employment. The aetiology of PGP is poorly understood, and there is no official nomenclature [2, 3], no effective evidence-based preventive measures or treatment, known risk factors or detailed knowledge of the clinical course of the various subgroups of this condition.

1.1 Historical perspective

PGP during pregnancy was mentioned by Hippocrates as symphysis pubis dysfunction in his theory of "disjunctio pelvica" more than 2,000 years ago [4]. For centuries, research on this condition focused primarily on the laxity of the pelvic joints and its aetiology. However, in the second

half of the 1970s, this focus became more concerned with symptoms [5]. Questionnaires and illustrations allowed for a more detailed and accurate assessment of pain, which brought into question the assumption that joint relaxation is the main cause of this pain [6]. As a result of its potential negative impact on the woman's quality of life during pregnancy, as well as the cost of this condition on society, the medical profession has been paying more and more attention to PGP during the past 20 years [2].

1.2 Definition

PGP is defined as originating in the pelvic musculoskeletal system, excluding ailments of gynaecological and urological character. The PGP diagnosis is independent of pregnancy and sex, according to the European guidelines on diagnosis and treatment of PGP: "Pelvic girdle pain generally arises concerning pregnancy, trauma or reactive arthritis" [7]. Pain is experienced between the posterior iliac crest and the gluteal fold, particularly in the vicinity of the sacroiliac joints (SIJs). The pain may radiate from the posterior thigh. It can occur in conjunction with/or separately from the symphysis. For pregnancy-related PGP, the onset of symptoms occurs from approximately week 6 of the pregnancy and

reaches peak pain intensity between the 24th and 36th week of pregnancy [2, 8].

Research on pain in the lower back area during pregnancy, published in the past 20 years, reveals that a dissensus regarding nomenclature still prevails. Studies of the condition have not used the same definition of PGP [9]. Commonly, studies include participants with pain in the lumbopelvic region without distinguishing PGP from low back pain [9]. PGP has been called symptom-giving pelvic girdle relaxation [10], peripartum pelvic pain [11], pelvic joint instability [12], posterior pelvic pain [13], pelvic instability, pregnancy-related lumbopelvic pain [14], pregnancy-related low back pain [15], and pregnancy-related pelvic girdle pain [16].

PGP can be divided into five subgroups according to joint involvement: symphysiolysis (separation of the symphysis pubis), one-sided SIJ syndrome (pain at one SIJ), double-sided SIJ syndrome (pain at both SIJs), the pelvic girdle syndrome (PGS) (in which both the symphysis and SIJs are affected), and a miscellaneous group [3, 7, 16-18]. The miscellaneous group is defined as inconsistent objective findings of daily pain in ≤ 1 pelvic joint [17]. The pelvic girdle syndrome (PGS) group

has the worst prognosis: 21 % continue to have pain two years after delivery [17]. The symphysiolysis group have a 100% chance for a full recovery, not later than six months after delivery [17, 19]. The groups with one-sided and/or double-sided SIJ syndrome also have a chance for full recovery in no later than 12 and 18 months, respectively [17]. No figures for recovery exist for the various group.

In this thesis, the term pelvic girdle pain (PGP) will be used, following the definition from the European guidelines on diagnosis and treatment of PGP above.

1.3 Incidence and prevalence

The incidence and prevalence of PGP vary depending on the definition, the diagnostic means utilized, and the design of the study [20]. Most of the literature reporting a prevalence (= the number of existing cases in a certain time period) and describing the epidemiological characteristics of PGP have been conducted in Europe [9]. Studies have reported prevalence rates ranging from as low as 7 % to as high as 84 % [8, 21-36]. However, these studies have not used the same guidelines to classify women with PGP. Some studies are based on self-report measures alone,

such as pain location drawings and questionnaires [21, 25, 26, 30, 31, 37]. In contrast, others have used physical examination as well as self-reported measures to confirm the classification of PGP [8, 22, 24, 27, 29, 32]. Some studies are prospective and some retrospective, which makes comparison problematic. Retrospective designs are prone to recall bias, which may explain the large variability in the published data on prevalence rates [38]. The incidence (= number of new cases in a certain time period) of pelvic girdle pain in pregnancy is unknown. Estimates from low-level evidence are contradictory, ranging from approximately 4 % to 84 % [32, 39-41], is higher in late pregnancy [21, 24, 26, 32, 42] and among women with a higher BMI [26].

1.4 Aetiology

Several etiological factors have been suggested for pregnancy-related PGP; biomechanical [22, 43-45], hormonal [8, 10, 46-49], metabolic [50], genetic [27, 40, 51, 52], and biopsychosocial factors [19, 53]. Biomechanical factors in combination with hormonal factors are proposed as the most plausible hypothesis [3, 7, 52].

1.4.1 Stability of the pelvis

In the European guidelines on diagnosis and treatment of PGP, a definition of optimal stability of the pelvis is described as: "The effective accommodation of the joints to each specific load demand through an adequately tailored joint compression, as a function of gravity, coordinated muscle and ligament forces, to produce effective joint reaction forces under changing conditions" [7]. Optimal stability of the pelvis consists of form and force closure (Figure 1) [54]. Form closure is due to the fit of the irregular surfaces of the sacrum and iliac bones physically locking the sacrum into the pelvic ring (arrows) between the two iliac bones (Panel A) [54]. The dorsal interosseous ligament maintains the integrity of the joint. Force closure is the compressive

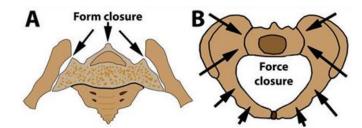


Figure 1. Form and force closure of the pelvic ring

effect exerted on the pelvic ring by the coordinated contraction of the abdominopelvic muscles, which shut the sacrum between the iliac bones and stops it from rotating outward (Panel B) [54].

1.4.2 Biomechanical factors in PGP

In general, bracing the SIJ by sufficient force closure of lumbopelvic muscles and the nutation of the ilium are thus essential for an efficient load transfer to the legs [55]. A systematic review found that patients with PGP have increased motion in their pelvic joints compared with healthy pregnant controls [56]. This increased motion in the pelvic joints diminishes load transmission efficiency and increases the shear forces across the joints [3]. These increased shear forces might be responsible for pain in pregnant women with PGP [57].

Significantly reduced strength of the transversus abdominis (TrA), lumbar multifidus, internal oblique muscles, the pelvic floor musculature, and insufficient coordination of all lumbopelvic muscles is often observed in patients with PGP [58-61]. This impairment in muscle strength and coordination is caused by abdominal stretching and a shift of body gravity centre when PGP commences in the second and third

trimester of pregnancy, resulting in reduced force closure [62, 63]. Which, in turn, can generate neuromuscular compensatory strategies [62, 63], namely the butt-gripping and the chest-gripping strategy. In the butt-gripping procedure, there is an overuse of the posterior buttock muscles. In the chest-gripping approach, the external oblique is in overuse and compensating for the underuse of the TrA [63]. These actions are hypothesized to increase sheared forces in the SIJ, thus being accountable for pain [3, 56].

Researchers investigated resultant pain regarding the SIJ as a pain generator when injecting the joint with an irritant solution in healthy subjects [64, 65]. Their finding agrees with referred pain patterns reported on direct SIJ capsular stimulation, with decreased pain when treated by injecting an anaesthetic into the SIJ. Such referred pain patterns are often observed in women with PGP [66], suggesting that the SIJ may be a source of pain in PGP.

Pregnant women with PGP have also shown hypersensitivity in superficial and deep tissue in the lumbopelvic region and distant to it, indicating widespread hypersensitivity to the SIJ [67]. However, the

stage of pregnancy does not correlate with a self-reported disability, pain, or hypersensitivity, indicating that these symptoms likely are related to several factors, including altered biomechanics (of somatic and visceral tissues). Emotional health, poorer sleep quality, and changes in hormonal status are factors [67].

1.4.3 Hormonal factors in PGP

Hormones may be involved in several different factors related to PGP, including modulation of pain and collagen synthesis, as well as inflammatory processes [19, 52]. During pregnancy, the gonadal hormones enhance pain sensitivity directly, potentially by modulating the responses of primary afferents on neurons of the dorsal horn and at supraspinal locations [63] and indirectly through their influence on emotional status [68].

At present, there is little evidence concerning the likely involvement of high levels of relaxin in the elevated laxity of pelvic joints. It is known that hormonal changes associated with pregnancy are compensated for by adequate changes in the force of sacroiliac closure [7, 58]. Furthermore, the widening of the symphysis in response to high levels

of relaxin can be physiological if it does not exceed 10 mm [69, 70]. A wider gap can be viewed as a pathological consequence of the inadequate force of the sacroiliac closure [56, 71].

The number of previous deliveries has also been found to be associated with a risk of PGP [25]. Pain associated with a previous pregnancy or delivery may increase sensitivity to pain in the pelvic girdle in a subsequent pregnancy [16], like increased sensitivity to pain resulting from previous pain is well known [72]. A link between early menarche and PGP has been suggested [47], believed to be due to the influence of pre-pregnancy hormonal factors rather than altered hormones during pregnancy.

1.4.4 Psychological factors in PGP

Research has shown that emotional states play a significant role in pregnancy [73]. Psychosocial factors have long been associated with chronic pain, and the bio-psycho-social model has become the leading theory of the development and management of chronic pain [74]. Psychosocial factors have also been demonstrated to play a crucial role in the transition from acute and sub-acute pain to chronicity [75-77]. In

patients with musculoskeletal pain, psychosocial factors appear to exacerbate the clinical component of pain [78, 79]. They have shown to influence future disability, pain, self-reported improvement after treatment in LBP patients [80-84].

Even though pregnancy itself negatively influences health-related quality of life, lumbopelvic pain increases this influence [85]. Pregnancy-related lumbopelvic pain has also been shown to have a great negative emotional and psychological impact on women [86]. This impact is often associated with dominant psychological factors (somatization, catastrophizing, pathological fear and/or elevated anxiety, depression), as well as social factors (such as a history of sexual abuse) [52]. Daily stress is a demonstrated risk factor for pregnancy-related lumbopelvic pain [87]. Women with postpartum depressive symptoms are three times more likely to report lumbopelvic pain than those without [88].

1.4.5 Miscellaneous factors

PGP association with metabolic comorbidities such as diabetes has been reported, but the underlying etiological mechanism has not been

identified [89]. Epidemiologic research elucidates that women with PGP are more likely to have a mother or sister with PGP [27, 40].

In summary, the stability of the pelvis during pregnancy is dependent on form closure and adequate motor control, being potentially compromised by the reduced or excessive force of closure as well as influenced by emotions.

1.5 Diagnosis

The diagnosis of PGP can only be reached after the exclusion of lumbar causes. The specific clinical tests must reproduce pelvic pain or functional disturbance. In the European Guidelines on diagnosis and treatment of PGP, the recommended evaluated tests for diagnosing PGP have a very high specificity indicating that the patient does not suffer from PGP if they are negative [7]. However, the sensitivity is low. Hence it is recommended to perform all the recommended tests and not rule out PGP if one test is negative. Tests recommended for PGP clinical examination are for pain in the SIJ: Posterior pelvic pain provocation test (P4), Patrick's Faber test, palpation of the long dorsal SIJ ligament, and Gaenslen's test. For pain at the symphysis pubis, palpation of the

symphysis and the modified Trendelenburg test are recommended. Together with pain and disability scales, these diagnostic tests are useful in recording PGP symptoms, severity, and subgroup classification [90].



In order to distinguish between reduced force closure and excessive force closure [54], the Active Straight Leg Raise test (ASLR), illustrated here, is considered one of the most appropriate tests available for evaluating the functional stability of the pelvis [52]. The test is in a review even

referred to as "the golden standard for testing the functional ability of the pelvis" [19]. In a pregnant sample, the specificity of this test was 88 %,



and the sensitivity was moderate (54 %) [91]. If combined with the P4 test (illustrated here), the sensitivity increases to 68 %.

Reduced force closure represents pain associated with excessive strain to the SIJ, surrounding connective tissues and

myofascial structures due to ligamentous laxity [92] coupled with motor control deficits of muscles that control force closure of the SIJs [45]. This form of PGP presents with a positive ASLR test [45], as a delayed onset activity of the obliquus internus abdominis, multifidus, and gluteus maximus muscles discloses this motor control deficit. Thus, an alteration in lumbopelvic stabilization disrupts load transference through the pelvis [92].

Excessive force closure occurs when the peripheral nociceptive drive is generated by excessive, abnormal and sustained loading of SIJs, surrounding connective tissue, and myofascial structures from the excessive activation of the motor system local to the pelvis in response to a transfer of the increased weight load in the woman [93]. This form of PGP generates localized pain to the SIJs, the surrounding connective tissue, and in myofascial structures such as the pelvic floor and piriformis muscles [93]. It returns a positive P4 test and a negative ASLR test.

1.6 Risk factors

Risk factors for developing PGP consistently found in research are previous pelvic or lower back pain [94, 95] and a history of trauma to the

back or pelvis [16, 31]. Multiparity [33, 94], increased body mass index, physically demanding work, emotional distress, and smoking also increase women's risk [7, 9, 25, 96].

Probable risk factors are increased workload, inactive lifestyle, higher age in pregnancy, generalized joint hypermobility [97, 98], and stress [2, 7, 25, 62, 99]. Research has shown that physically active women, regularly engaging in high-impact exercises before the first pregnancy, have a reduced risk of experiencing PGP in pregnancy [100].

The recommended diagnostic tests described above may also serve as prognostic tests and indicate the risk of disability and future pain in pregnant women [101]. Research shows that women with pain in all three pelvic joints (bilateral SIJs and symphysis) and with many positive tests have a markedly worse prognosis than women with other combinations of self-reported pain location [17]. These results suggest that a clinical examination, including a few tests performed in early pregnancy, may identify women at risk of a more severe PGP late in pregnancy [102].

1.7 Prognostic factors

High pain intensity [32], severe pelvic pain in three pelvic joints [102], the use of crutches [103], other pain conditions [104], menarche at a young age [47], previous low back pain [32], comorbidities [104], obesity [103], multiparity [105] and experience of emotional distress [104], are factors identified to influence the prognosis in pelvic girdle pain during pregnancy negatively.

Studies of testing have shown that pain in the pelvic joints, bilateral positive P4 tests, and certain positive pain provocation tests in the early stages of pregnancy are significantly associated with disability and pain intensity at gestation week 30 [102]. A poor ASLR performance and localized pressure pain hypersensitivity in the pelvis during pregnancy is correlated with low physical health-related quality of life postpartum and pain quality [106]. Distress in early pregnancy is also significantly associated with disability at gestation week 30, but not with pain intensity [32]. However, the ability to manage emotional distress during pregnancy, and a belief in improvement, may prevent the persistence of pelvic girdle pain postpartum [107].

1.8 Clinical course

Research on the clinical course of PGP in pregnancy is scarce. The data have usually been collected at the baseline and at one or more follow-ups [39, 108] which have shown that the onset of PGP varies significantly, from the end of the first trimester to a couple of months postdelivery, including the labour stage. A peak of symptoms seems to exist closer to the third trimester between the 24th and 36th weeks of pregnancy. With only a few measurement points in time, stability may be indicated, and a fluctuating PGP course may be undetected. Hence prospective frequent data collection is warranted to describe the clinical course accurately. It has been suggested that a clinical examination including a few tests performed in early pregnancy, with follow-ups over time, may identify women at risk of a more severe course of PGP late in pregnancy [102].

1.9 Consequences

PGP during pregnancy is associated with depressive symptoms and greatly affects the experience of being pregnant, roles in relationships and social context [86]. For women with young children, PGP negatively

affects the role of being a mother, a situation that further strains the pregnancy experience [109]. Women with PGP have less day to day mobility than women with back pain only and require crutches or wheelchairs more frequently [41, 110]. In addition, emotional distress is significantly associated with disability [32].

PGP is one of the major causes of sick leave in pregnancy [16, 27, 87, 111, 112]. Despite an increasing number of women working throughout their reproductive years, only a small number of studies regarding the frequency and duration of sick leave during pregnancy and prevalence estimates differ according to study methodology and populations examined [112-116].

PGP accounts for up to 72% of sick leave in pregnancy with an average length of 12-15 weeks [16, 27, 87, 111, 112]. Women with the involvement of several joints and a high pain level also have longer sick-leave duration than others [102], making PGP during pregnancy a major public health issue [27]. Different occupational groups may have various sick leave patterns, and occupational factors may contribute to sick leave

in 50% of pregnancies [112]. Sick leave patterns may be linked to sick leave benefits without fully explaining them [112].

1.10 Gaps in knowledge about PGP

Although PGP can significantly impact pregnant women's health and quality of life, varying definitions of approach diagnosis and study designs have estimated its incidence and prevalence problematic [9, 19]. One additional challenge in this respect is the wide range of outcomes measured, from self-report symptoms alone (e.g., location and severity of the pain as indicated from questionnaires [21, 25, 26, 30, 31, 37] or in combination with physical examinations [8, 22, 24, 27, 29, 102]. Moreover, most demographic and clinical characterizations of subgroups of patients demonstrating different PGP symptoms have not been longitudinal, with repeated data collection. More standardized research is required to identify women at risk of developing PGP during pregnancy and identify the predictors of deleterious clinical courses. In addition, even though PGP is a major cause of sick leave during pregnancy, relatively little is known about this connection. For instance, can differences in tolerance levels for pain and/or demographic and psychosocial characteristics and clinical symptoms

explain why some women take sick leave while others do not [20, 117]? Furthermore, in connection with prospective PGP studies are data usually collected at baseline with only one or few follow-ups. The PGP course during pregnancy thus remains to be examined in detail [15, 32, 118-120].

1.11 The three major objectives of the current project

- 1: To determine the prevalence of PGP during pregnancy in a random population of women and factors that may be associated with the development of this condition.
- 2: To explore factors that influence taking sick leave due to PGP during pregnancy, including pain-related activities of daily living and the nature of employment, including the physical workload involved.
- 3: To examine whether pregnant women with PGP who have been subgrouped based on two clinical tests differ regarding demographic characteristics and/or the clinical course of PGP (i.e., the number of days per week with bothersome symptoms) during the second half of their pregnancy.

2.1 Study aims of the retrospective study (Papers I and II)

The objectives of the study were to investigate the cumulative prevalence of LBP, PGP and combined lumbopelvic pain (LBPP) during pregnancy, including features possibly associated with the development of PGP in an unselected population of women. Further objectives were to explore the frequency of sick leave in pregnancy due to PGP, assess the relationship between different types of pain-related activities of daily living, examine physical workload, type of work concerning sick leave, and explore factors that make women less likely to take sick leave for PGP.

2.1.2 Setting and inclusion criteria

The data collected in the retrospective study were conducted at Stavanger University Hospital maternity ward in March – July 2009. The hospital has the only birth department in the southern part of the county of Rogaland, with a population of approximately 330 000 inhabitants. The annual number of deliveries at the hospital varies between 4 400 and 4800. Inclusion criteria were a singleton pregnancy of at least 36 weeks

and good competence in the Norwegian language. Within 24 hours after delivery, the women received verbal and written information about the study from a midwife. Participation was voluntary, but all eligible women were encouraged to participate in obtaining the inclusion of an unselected sample.

2.1.3 Instrument and variables

The women filled in a questionnaire specially designed by the research group, based on previous studies and the experience of the team. The questionnaire contained demographic information, questions regarding pain distribution in the pelvic girdle area, pain-related activities of daily living (ADL), sick leave in general and due to PGP, and frequency of exercising before and during pregnancy. The questionnaire (in Norwegian) can be found in Appendices.

2.1.4 Pain variables

The women marked the location of the pain on illustrations of the pelvic girdle and low back included in the questionnaire package. The pelvic girdle and the low back were labelled and separated according to boundaries described in the European guidelines for the diagnosis and treatment of PGP [7].

The question on pain intensity ("Rate in each square, representing each month in pregnancy, the average pelvic pain you have experienced") was rated retrospectively on a numerical rating scale (NRS) [121] from 0 to 100, to collect information on the presentation of symptoms and the peak intensity of pain during pregnancy. In this study, the score was anchored at 0, meaning "No pain", and 100 meaning "Unbearable pain". For the analyses, "average pain PGP intensity" was calculated as the mean of the values reported in all months.

Information on pain-related ADL was collected through the Oswestry Disability Index (ODI) [122], which, at the time, was one of the principal condition-specific outcome measures for defining disabling effects from spinal disorders and PGP. A patient-completed questionnaire gives a subjective percentage score of the level of function (disability) in 10 ADLs in patients with low back pain [122]. Every activity contains six statements on how well the activity is performed. The statements are scored from 0 to 5. The scores for all questions answered are summed, then multiplied by two to obtain the index (range 0–100). Zero is equated with no disability, and 100 is the maximum disability possible.

2.1.5 Work-related variables

The questionnaire also provided information on the total number of years of education (including elementary school), the level of physical workload (measured with five answer categories ranging from 'sedentary' to 'heavy', following a scale used in the Stockholm Public Health questionnaire [123]. The type of work (in free text) and work satisfaction (a five-level scale runs from very bad to very good) [124, 125].

2.1.6 Sick leave variables

Sick leave was estimated in two different ways. First, the women were asked about their total number of weeks of full-time sick leave during pregnancy and the total number of weeks with part-time sick leave and sick leave percentage. In the analyses, weeks of sick leave in total were calculated by adding the full-time sick leave weeks to the part-time weeks adjusted for sick leave percentage. After reporting the total amount of sick leave, the women were asked to specify the primary cause of their sick leave. Second, in the section concerning pain intensity during pregnancy, the women were asked whether they had been on sick

leave due to PGP in any month of the pregnancy and indicated when. It was, therefore, not possible to determine the number of consecutive weeks of 100 % sick leave due to any specific cause from the available information. For instance, several women only reported "pain" without any specific pain area details as the primary cause of sick leave in pregnancy. All the available information was combined in the analyses to establish if the women were on sick leave due to PGP. If the women reported any sick leave due to PGP in any month of the pregnancy, they were classified as having sick leave due to PGP. The women, who explicitly stated that PGP was the primary cause of their sick leave, but who did not indicate sick leave due to PGP in any specific month of pregnancy in the questionnaire's pain intensity-section were also classified as having sick leave due to PGP.

2.1.7 Assessment instruments, retrospective data collection 24 hours after giving birth

Instruments	and	Description	of	answer	Reference
variables of int	erest	options			
Questionnaire					

Education (comm)	N1	
Education (years)	Number of years (including	
	elementary school)	
Physical workload	Five levels; from very easy	[123]
	to very heavy. Sixth option:	
	not working	
Profession	Free text	
Job satisfaction	5 level Likert scale; from	[126]
	very bad to very good	
Weeks on sick leave	Number of weeks, or: not	
	been on sick leave	
Weeks on full time and	Number of weeks	
part-time, including		
percentage sick leave		
Cause for sick leave?	Free text	
Height	Centimetres	
Weight (most recently)	Kg	
Weight before	Kg	
pregnancy		
Weight before delivery	Kg	
Depressed until now in	Four categories: Never,	
this pregnancy	on/off, often, almost all the	
	time	
If you have been	Nine options: Month 1-9	
depressed: In which		
month/s?		
1		

Chronic disease?	Yes/No	
If yes, which disease?	Free text	
The number of previous	Number	
births?		
Pelvic pain in previous	Yes/No	
pregnancies?		
Hormonal treatment to	Yes/No	
achieve this pregnancy?		
Regular exercising (2-	Yes/No	
3x/week) before this		
pregnancy?		
Regular exercising (2-	Yes/No	
3x/week) in this		
pregnancy?		
Injuries to the pelvis?	Yes/No	
Type of injury	Free text	
Low back or pelvic pain	Yes/No	
before this pregnancy?		
Low back pain onset in	Nine options: Month 1-9	
which month?		
Low back pain location	Pain drawing	[127-129]
The low back pain level	NRS 1-100 for each month	[121]
in months 1-9		

Month/s with sick leave	Nine options: Month/s 1-9	
With sick leave	Nine options. Month's 1-9	
for low back pain in this		
pregnancy?		
Pelvic pain onset in	Nine options: Month 1-9	
which month		
Pelvic pain location	Pain drawing; pelvis	[40]
	back/front	
The pelvic pain level in	NRS 0-100 for each month	[121]
months 1-9?		
Month/s with sick leave	Nine options: Month/s 1-9	
for pelvic pain in this		
pregnancy?		
Oswestry Disability		
Index (ODI)		
An instrument	A subjective percentage	[122]
measuring level of	score of the level of function	
disability in activities of	in 10 ADLs. Every activity	
daily living (ADL)	contains six statements,	
	scored 0 to 5.	

2.2 Study aims of the prospective study (Paper III)

The objective of this study was to explore if pregnant women with probable PGP, sub-grouped according to the clinical tests recommended in the European guidelines, differed in demographic and clinical characteristics at mid-pregnancy and the weekly number of days with bothersome symptoms through the second half of pregnancy. The hypothesis was that sacroiliac dysfunction and failing force closure diagnosed at mid-pregnancy might predict a course of bothersome symptoms through the second half of pregnancy.

2.2.1 Setting and inclusion criteria

Data collection was conducted at the obstetric outpatient clinic, Stavanger University Hospital, Norway, from mid-March to mid-June 2010. Pregnant women who had their second-trimester routine ultrasound examination in pregnancy week 18 were asked by a midwife about their experience of pain in the lumbopelvic region and were informed about the study. The inclusion criteria were ongoing lumbopelvic pain or isolated pelvic pain, singleton pregnancy and good proficiency in the Norwegian language.

2.2.2 Instruments and variables

On acceptance to participate, the women were asked to sign a letter of consent. They were given an envelope with questionnaires on demographic and clinical features, used in a previous retrospective study on pelvic girdle pain [20, 28] to fill in at home. A chiropractor consultation for a physical examination was arranged, and the women were asked to bring the filled-in questionnaires with them to the examination. For comparative purposes, women without pain symptoms were informed about the study, given a letter of consent to fill in if they accepted to join the study, and a questionnaire on demographic features to fill in and hand to the receptionist on departure. All consenting women were followed from week 18 of their pregnancy to week six postpartum with weekly automated text messages (SMS).

Two licensed chiropractors performed a physical examination of the pelvic region, including diagnostic tests recommended in the European guidelines for diagnosing and treating pelvic girdle pain [7] and a neurologic examination of the lower extremities. The results of their examination were recorded as PGP or not PGP diagnosis.

2.2.3 Pain variables

The women marked the pain location on drawings with the pelvis and the low back separated. Pain intensity was rated on a numerical rating scale (NRS) from 0 to 100, anchored at 0, meaning "No pain" and meaning 100 "Unbearable pain" [121]. Information on pain-related ADL was collected through the Oswestry Disability Index (ODI) [122]. At the time of data collection, the ODI was one of the main outcome measures for defining the disabling effects of spinal disorders and PGP [7, 122].

2.2.4 Work-related variables

Answers to a question on job satisfaction were recorded on a 5-point Likert scale with increments in two opposite directions ('Very bad' and 'Very good') and a neutral point in the middle [123].

2.2.5 SMS-tracking

Every Sunday, the women received an automated SMS asking how many days the previous week they had experienced bothersome pelvic pain [130]. The question was repeated 24 hours later if there was no reply [130]. The question should be answered with one single number between

0 and 7 [130]. The response was automatically entered into a database, which contained continuous information updates from each participant throughout the study [130].

2.2.6 Assessment instruments and procedures at baseline; week 18

Instruments and	Description of answer	Reference
variables of	options	
interest		
Group-designed		[28]
questionnaire		
Education (years)	Number of years (including	
	elementary school)	
Physical workload	Five levels; from very easy to	[123]
	very heavy. Sixth option: not	
	working	
Profession	Free text	
Job satisfaction	5 level Likert scale; from	[124, 125]
	very bad to very good	
Weeks on sick leave/	Number of weeks	
not been on sick leave		
Weeks on full time	Number of weeks	
and part-time,		

Free text	
Centimetres	
Kg	
Kg	
Four categories: Never,	[131]
on/off, often, almost all the	
time	
Five categories: Weeks 1-4,	
5-8, 9-12, 13-16, 17-20.	
Yes/No	
Free text	
Number.	
Yes/No	
Yes/No	
	Centimetres Kg Kg Four categories: Never, on/off, often, almost all the time Five categories: Weeks 1-4, 5-8, 9-12, 13-16, 17-20. Yes/No Free text Number.

Regular exercising (2-	Yes/No	
3x/week) before this		
pregnancy?		
1 0 1	Yes/No	
Regular exercising (2-	Tes/NO	
3x/week) in this		
pregnancy?		
Injuries to the pelvis?	Yes/No	
Type of injury	Free text	
Low back pain until	Yes/No	
now in this		
pregnancy?		
Pelvic pain until now	Yes/No	
in this pregnancy?		
Low back pain onset	Five categories: Weeks 1-4,	
in which weeks?	5-8 ,9-12 ,13-16, 17-20	
Low back pain	Pain drawing	[127-129]
location		
Low back pain level in	NRS 0-100	[121]
weeks 1-4, 5-8, 9-12,		
13-16, 17-20?		
Sick leave for low	Five categories: Weeks 1-4,	
back pain in this	5-8, 9-12,13-16, 17-20	
pregnancy?		
Pelvic pain onset in	Five categories: Weeks 1-4,	
which weeks?	5-8, 9-12,13-16, 17-20	

Pelvic pain location	Pain drawing; pelvis	[40]
1	back/front	
Pelvic pain level in	NRS 0-100	[121]
	1113 0-100	[121]
weeks 1-4,5-8,9-		
12,13-16,17-20?		
Sick leave for pelvic	Five categories: Weeks 1-4,	
pain in this	5-8, 9-12, 13-16, 17-20	
pregnancy?		
Modified	Positive/Negative	[22]
Trendelenburg`s test		
Active straight leg	6-point scale; From "not	[132]
raise (ASLR)	difficult at all" to "unable to	
	do."	
Posterior pelvic pain	Positive/Negative	[133]
provocation test (P4)		
Gaenslen`s test	Positive/Negative	[95]
Patrick`s FABER test	Positive/Negative	[22]
Long dorsal sacroiliac	Positive/Negative	[22]
ligament test		
Symphysis palpation	Positive/Negative	[22]
test		
Oswestry Disability		
Index (ODI)		
An instrument	A subjective percentage score	[122]
measuring level of	of the level of function based	
		1

disability in activities	on 10 ADL's. Every activity	
of daily living (ADL)	contains six statements,	
	scored from 0 to 5.	
SMS-Track		
No. of days with	Answer with a single number	[130]
bothersome pelvic	0-7.	
pain in the previous		
week?		

2.3 Analysis of the retrospective data (Papers I and II)

2.3.1 Paper I

The objective of this paper was to examine the prevalence and incidence of LBP, PGP and LBPP during pregnancy, including features possibly associated with the development of PGP in an unselected population of pregnant women. In this analysis, we studied the influence of the following variables on LBP, PGP and LBPP during pregnancy:

2.3.2 Independent variables in Paper I				
Pre-pregnancy variables	Pregnancy variables			
-BMI before pregnancy [134]	-Age at delivery [41]			
-LBP in previous pregnancies [2]	-BMI at delivery [134]			
-PGP in previous pregnancies	-Exercised at least 2-3			
[135]	times/week during pregnancy			
-LBP in the year before pregnancy	[135]			
[87]	-Weeks of full_time sick leave			
-PGP in the year before pregnancy	during pregnancy [127]			
[2]	-Received treatment for LBP			
-Exercised at least 2-3 times/week	and/or PGP during pregnancy			
before pregnancy [135]	[40]			
-Number of years of education [40]	-ODI [7, 101]			
-Physically heavy work [2]	-Moderate and severe pain			
-Number of previous births [87]	distribution according to pain			
	drawings [30]			

Descriptive data on demographic and clinical features were presented by mean values and standard deviations for continuous variables and frequencies for categorical variables. The dependent variable: pain symptoms, was classified into three pain levels through analyses of variance. The cut-off point with the largest F ratio between mild and

moderate and severe pain was found at 35 in the NRS (0-100). Thus, patients were grouped into three pain categories:

- -No pain (NRS = 0)
- -Mild pain (NRS \leq 35)
- -Moderate and severe pain (NRS >35).

Kruskal-Wallis statistics were used to explore these pain groups for differences regarding pre-pregnant and pregnant variables. Multivariate hierarchical logistic regression analysis was used to calculate whether the pre-pregnancy and pregnancy variables could predict moderate to severe PGP (with no pain as the reference category). Mild pain was omitted from the analysis because of presumed low clinical interest for their ADL. In the first block of the analysis, age, educational level, and the number of previous births were entered, followed by a block containing the average LBP level throughout the pregnancy. The last block used backwards stepwise regression using the likelihood ratio removal criterion, including the variables workload, BMI before the pregnancy, BMI at birth, feelings of depression during pregnancy, physical activity before, and physical activity during pregnancy. Thus,

the first two blocks served mainly as statistical controls with the forced entry of all variables before exploring the final block variables.

All analyses were performed in SPSS 16 (IBM, New York, NY), and results were considered significant at $P \le .05$.

2.3.3 Paper II

The primary objective of this paper was to examine the frequency of sick leave in pregnancy due to PGP and to assess the relationship with different types of pain-related ADLs, physical workload, and type of work. Variables entered into the analysis were age [136], years of education [40], BMI before pregnancy [134], number of total sick leave weeks during pregnancy [127], physical workload [2], work satisfaction [21], average PGP [2], average LBP [137], depressed [131], no. of previous births [40], regular exercise 2-3 times per week before pregnancy [135], seated work [138], Oswestry disability index; 10 items [41].

A further objective was to explore factors associated with less sick leave due to PGP by contrasting two groups of women with PGP, differing by having been versus not having been on sick leave for their pain. The final objective was to explore the relative contribution of PGP to the total amount of sick leave in pregnancy.

In this paper, the women were classified into three groups (the dependent variable):

- -Women who did not report sick leave
- -Women who reported sick leave but without indicating PGP as the cause
- -Women who reported sick leave and stated PGP as a partial origin of their sick leave.

Descriptive data on demographic and clinical features were reported as mean values, standard deviations for continuous variables, and as frequencies for categorical variables.

For comparisons of the three groups with different sick leave patterns, the non-parametric Kruskal-Wallis statistics were used, applying Bonferroni correction to counteract multiple comparisons. The next step in the analysis was pairwise follow-ups with the group who had sick leave due to PGP as a reference whenever significant omnibus group differences were found in the previous Kruskal-Wallis test. For categorical data, chi-square statistics were calculated, and 2×2 table

follow-ups were used for pairwise comparisons between the group with sick-leave due to PGP vs the other groups.

Multinomial logistic regression analysis was then performed to investigate the independent input of variables hypothesized to affect sick leave due to PGP. Forced entry was implemented for the variables age, education, parity, and average PGP to adjust for them in the final model. In an exploratory approach, single items from ODI (except sex and pain intensity) were entered in a stepwise procedure together with the variable's workload, work satisfaction and seated work (= working in a sitting position), using a likelihood ratio-based criterion with p<.05 for entry and p<.10 for exclusion.

A sequential linear regression analysis was performed, using the total number of calculated weeks of sick leave (weeks of 100 % sick leave + weeks of part-time sick leave multiplied by sick leave percentage) for any reason as a dependent variable to explore the factors linked with the total amount of sick leave in pregnancy:

-In the first block, the grand mean of monthly reported PGP was entered to analyze the unadjusted effect of PGP on weeks of sick leave.

- -In the next block, all appropriate ODI items were entered using a stepwise procedure (p<.05 to enter, p<.01 to omit a variable).
- -In block 3, the variables years of education, pre-pregnant BMI, workload, age, standing work, and mobile work were entered, using the same stepwise procedure as in block 2.
- -Finally, the variables work satisfaction and depression in pregnancy were entered with a stepwise procedure.

Only block 1 contained a forced entry variable, average PGP, as the objective was to explore unadjusted and adjusted effects of PGP on weeks of sick leave. In order to explore factors that may diminish the influence of PGP on sick leave, all women with PGP who did not report sick leave in pregnancy were identified.

When calculating causal effects using observational data, it is desirable to replicate a randomized experiment as closely as possible by obtaining intervention, in this case, sick leave, and control groups with comparable covariate distributions. This goal can often be obtained by choosing well-matched samples of these groups, thereby reducing bias due to the covariates [139]. Thus, a macro was written in Microsoft Excel (Visual Basic) then selected a random woman having been on sick leave, who

matched the mean PGP score of a woman with no sick leave. If a perfect match was not found, a difference of +/-1 point on the PGP score was approved. If still no match was found, the subject was rejected. Hence, this procedure resulted in two equal groups regarding mean PGP, but with and without sick leave. The same variables were compared in these two equal groups for the sick leave due to PGP vs no sick leave and sick leave due to other reasons groups in a Mann-Whitney U test. Effect sizes (the strength of the relationship between two variables) were reported as standardized mean differences (Cohen's D), using Bonferroni correction, which can be interpreted as small (around 0.3), medium (around 0.5) and large (0.8 to infinity) [140].

The analyses were conducted using SPSS 21 (IBM, New York, NY), and results were considered significant at p<.05.

2.4 Analysis of the prospective data

2.4.1 Paper III

The objective of this paper was to examine if pregnant women with probable PGP, sub-grouped following the results from two valid and reliable clinical tests recommended in the European guidelines, differ in demographic and clinical characteristics at mid-pregnancy and the weekly number of days with bothersome symptoms through the second half of pregnancy. The hypothesis was that sacroiliac dysfunction and failing force closure diagnosed at mid-pregnancy might predict a course of bothersome symptoms through the second half of pregnancy.

Demographic descriptive data were shown as mean and median values with standard deviation for continuous variables and frequencies for categorical variables.

For univariate comparisons between symptomatic and asymptomatic subgroups, the non-parametric Kruskal-Wallis statistic was performed. Categorical predictors in our model were four groups, following the results from the ASLR and P4 tests:

- -P4 positive
- -ASLR positive
- -Both P4 and ASLR positive
- -ASLR and P4 negative

The time (pregnancy week) and the interaction term between time and test group explore whether the trajectory of SMS-reported number of bothersome days differed between the test groups. Age, parity, and BMI

before pregnancy have previously shown significant association with PGP and were added to our model [134].

The longitudinal trajectory of the SMS- responses were modelled using a generalized estimating equations (GEE) approach, extending the generalized linear model to correlated longitudinal data and clustered data within subjects. The within-subject dependencies resulting from repeated measurement were modelled, assuming an autoregressive relationship in the working correlation matrix. As the outcome variable was count data (weekly number of bothersome days with pain), the Poisson distribution was assumed with a log-link function.

A robust non-parametric Brown-Forsythe ANOVA then examined equality of variances and data distribution within and between test groups. Data were analyzed using SPSS software (version 22.0; SPSS Inc, Chicago, IL, USA). A p-value < 0.05 was considered statistically significant.

2.5 Ethics

Both projects (retrospective and prospective) were carried out following the Helsinki Declaration II (https://www.who.int/bulletin/archives/79 (4)

<u>373.pdf</u>). The Regional Ethics Committee approved the retrospective study of Western Norway (rek-vest, ref.no. 2009/356-CAG). The Regional Ethics Committee approved the prospective research project of Northern Norway (rek-nord, ref. no. 2010/174).

All participants received written and oral information about each study's aim and the test procedures in the prospective study. All individuals provided written informed consent before participation. They were informed that participation was voluntary and that they could discontinue participation in the prospective study at any point without explanation. No risks to the women in the studies were identified.

3 RESULTS

3.1 The retrospective data collection (Papers I and II)

At Stavanger University Hospital, 1204, women gave birth during the inclusion period. All women were invited to participate, and after exclusions, 994 women were eligible for the study. However, 336 women did not return a questionnaire, and 89 did not fully complete the questionnaire. Thus, the final study population consisted of 569 women, 58% of the total possible sample.

To examine if the study sample was representative of the source population, we compared to age and parity from the study sample with all women who gave birth during the study period and found an almost perfect match.

3.1.1 Demographic features

The women's mean age was 30 years, mean years of education was 14.7 years, and slightly more than one-third of the women were primiparous. The mean BMI was 23.8, and almost half of the women exercised 2 to 3 times per week

before pregnancy. The mean amount of sick leave during the pregnancy was 9.6 weeks.

3.1.2 Pain

Approximately a quarter of the participants did not report any LBPP during their pregnancy. A further 13% had only experienced mild pains, while the cumulative prevalence for moderate and severe pain during pregnancy was 57.4% (n = 327). Moderate and severe combined LBPP was experienced by 21.6% (n = 123), moderate and severe PGP was experienced by 26.0% (n = 148), and almost 10% of the women (n = 56) had moderate and severe pain in the lumbar area. Twenty-three per cent of all women, 40 % of those who developed moderate and severe pain, reported such pain already after five months of pregnancy.

3.1.3 Pain distribution

Table 1 below illustrates the distribution of pain in the pelvic area for all women reporting pain in this region, divided into women with moderate and severe LBPP and those with moderate and severe PGP alone. More than half of all women

(52%) experienced pain at the symphysis, and circa 20% experienced pain only in this area. Approximately 33% had pain at all three pelvic joints, and 24% had pain at one sacroiliac joint alone. Almost half the women with moderate and severe lumbopelvic pain experienced pain at all three pelvic joints.

Table 1: Prevalence of moderate and severe pain distribution according to the pain drawings among women reporting pain during pregnancy.

Pain location	All women with	Moderate and	Moderate and
	pelvic pain	severe	severe
	N=348	pelvic girdle pain	lumbopelvic pain
		N=148	N=123
At all pelvic joints, N	113 (32.5)	40 (27.0)	58 (47.2)
(%)			
At symphysis, N (%)	67 (19.3)	33 (22.3)	18 (14.6)
At one SI joint, N (%)	83 (23.9)	38 (25.7)	24 (19.5)
At both SI joints, N (%)	60 (17.2)	27 (16.2)	16 (13.0)
Other areas, N (%)	7 (2.0)	1 (0.7)	2 (1.6)
Missing data, N (%)	18 (5.2)	9 (6.1)	5 (4.1)

3.1.4 Potential risk factors

In the multivariate analysis, we examined the impact of clinical and demographic variables on moderate to severe PGP. The first block of the binary logistic regression analyses was not statistically significant (p = 0.379), indicating that age, education, and the number of previous births did not contribute to the prediction of moderate to severe PGP. The following block, however, was highly significant (p≤0.001), showing that a high level of LBP reduced the risk of PGP (p≤0.001; odds ratio: 0.845, [CI; 0.798-0.894]). The final block containing previously entered variables, and the variables remaining after the backward stepwise procedure (BMI before pregnancy and physical activity before pregnancy), was also highly significant (p≤0.001). The resulting omnibus logistic regression model was significant (p≤0.001, Nagelkerke R2 =0.319). Predictor variables in the full multivariate model were average LBP (p≤0.001, odds ratio; 0.837, [CI: 0.790-0.887]), BMI before pregnancy (p=0.011, odds ratio: 1.074, [CI: 1.016-1.134]), and physical activity before pregnancy (p=0.015, odds ratio; 1.826, [CI: 1.126-2.960]). So, both higher BMI before pregnancy and higher physical activity levels

before pregnancy were independent potential risk factors for PGP after controlling for age, education, number of previous pregnancies, and LBP. Average LBP during pregnancy reduced the risk of moderate to severe PGP.

3.1.5 Sick-leave and disability

Women with moderate to severe LBPP had a mean sick-leave period of 15.5 weeks, those with moderate to severe PGP 10.7 weeks, and women with moderate to severe LBP 9.1 weeks. Women with mild pain had a mean sick-leave period of 6.5 weeks, indicating that experiencing moderate to severe pain had different clinical consequences from experiencing less pain. The same pattern was found for disability, measured with the ODI. Women with PGP were seeking more care than women with lumbar pain, and those with moderate to severe pain received more treatment than those with mild pain.

In the analysis to explore the frequency of sick leave in pregnancy due to PGP and to assess the relationship of different types of pain-related ADLs, physical workload, and type of work with sick leave due to PGP, one woman was excluded from the analyses (she did not report having a

job, profession, nor workload). This analysis thus consisted of 568 women. Of these, 165 (29 %) reported that they had experienced isolated PGP during the pregnancy. The sample's demographic and descriptive statistics for the variables used in the multivariate analyses are shown in Table 2. Several significant differences were found between subjects who reported sick leave due to PGP vs those who did not.

Table 2: Demographic data and descriptive statistics of women with and without sick leave for variables used in the multivariate analyses.

	Sick leave for PGP		No sid	No sick leave		eave for reason	p****
			N = 13	39 (24%)	N = 23	36 <i>(4</i> 2%)	
	N = 19	3 (34%)					
Age in years,	29.7	(4.3)	30.5	(4.8)	29.8	(5.0)	=.254
mean (SD)							
Education in	14.5	(2.4)	15.7	(2.4) *	15.4	(2.6) *	<.001
years, mean (SD)							
BMI before	24.8	(4.6)	23.1	(3.6) *	23.4	(4.2) *	<.001
pregnancy, mean							
(SD)							
Total sick leave,	10.8	(9.4)	0.0	(0.0) *	8.4	(8.9) *	<.001
mean number of							
weeks (SD)							
Workload,	3.0	(1.1)	2.4	(1.1) *	2.6	(1.2) *	<.001
mean score (SD)							
Work satisfaction,	4.4	(8.0)	4.6	(0.7) *	4.3	(0.9)	<.001
mean score (SD)							
Mean pain PGP,	26.8	(15.1)	6.7	(10.4) *	6.1	(10.0) *	<.001
(SD)							

Results

Mean pain LBP (SD)	13.2	(16.9)	4.7	(9.1) *	6.6	(11.4) *	<.001
Pain-related ADL (ODI), mean score (SD)	1.9	(0.8)	0.9	(0.9) *	1.0	(0.8) *	<.001
Depressed, mean score (SD)	1.4	(0.5)	1.3	(0.5)	1.4	(0.6)	=.055
No. of previous births, mean (SD)	1.00	(0.06)	0.94	(0.09)	0.79	(0.05)	<.05
Regular exercise before pregnancy, N (%)	63 (33 %)		67 (49 %) **		94 (40 %)		=.013
Seated work, N (%)	51 (27 %)	68 (4	9 %) *	81 (34 %)	<.001

PGP Pelvic girdle pain, LBP Low back pain. Pairwise comparison with sick-leave for PGP: *p < .001, **p < .01, ***p < .05 ****Kruskal-Wallis omnibus test. Bonferroni-corrected alpha = 0.0038

In Table 3, a comparison was made for all single items from the ODI between the group with sick leave due to PGP, the group without sick leave, and sick leave due to other causes. Effect sizes were reported to enable a direct comparison using a standardized scale. All ODI-items were significantly higher in the group who had been on sick leave for PGP than in both the other groups. The effect sizes were all moderate to large (Cohen's d > 0.6).

Table 3: Comparison of disability in activities (ODI single items) between women with sick leave due to PGP, women with no sick leave, and women with sick leave due to other causes.

	Sick leave for	No sid	No sick leave		Sick leave for other	
	N=190	N:	=96	N=154		
ODI item	Mean (SD)	Mean (SD)	E.S.ª	Mean (SD)	E.S.ª	
Pain intensity	2.76 (0.86)	1.67 (1.13)	1.142	1.81 (0.99)	1.024	
Personal care	1.23 (1.40)	0.59 (0.97)	0.591	0.53 (0.98)	0.655	
Lifting	2.18 (1.19)	0.95 (1.12)	1.056	1.30 (1.16)	0.751	
Walking	1.63 (0.99)	0.65 (0.94)	1.007	0.85 (1.05)	0.762	
Sitting	1.68 (0.96)	0.80 (0.98)	0.905	1.07 (1.11)	0.589	
Standing	2.44 (1.24)	1.17 (1.28)	1.018	1.48 (1.27)	0.769	
Sleeping	1.67 (1.02)	0.82 (0.88)	0.872	1.04 (0.88)	0.657	
Sex	1.75 (1.50)	0.76 (1.19)	0.707	0.76 (1.30)	0.701	
Social function	1.89 (1.26)	0.83 (1.17)	0.862	0.79 (1.14)	0.911	
Travelling	1.63 (1.26)	0.61 (1.00)	0.860	0.82 (1.11)	0.680	

E.S.: Effect size (Cohen's d. >0.8 is considered large)

3.1.6 Factors associated with sick leave due to PGP

Individual risk factors with odds-ratios and confidence intervals resulting from the multinomial regression analysis are shown in Table 4

 $^{^{\}rm a}$ All differences of means were statistically significant, assuming a Bonferroni corrected alpha of p < .005

below. All results refer to the group with sick leave due to PGP as the reference category. The estimated pseudo-R2 was relatively high (Nagelkerke R2 = .40), and the total correct classification percentage was 62 %. Work satisfaction and lower scores for the ODI-items lifting, sleep, and average pain intensity significantly classified individuals to the no sick leave group.

Table 4: Risk factors for sick leave during pregnancy, with sick leave due to PGP as the reference category. For each factor, the odds ratio and p-value (in brackets) are displayed.

	Age	Education	Pelvic pain	No of previous births	ODI lifting	ODI sleep	ODI social life	Work satisfaction
Sick	1.051	1.054	0.951	0.760	0.708	0.916	0.785	0.960
leave	(0.129)	(0.262)	(0.000)	(0.128)	(0.020)	(0.622)	(0.105)	(0.814)
due to								
other								
reason								
No sick	1.056	1.113	0.955	0.915	0.622	0.521	1.206	1.607
leave	(0.157)	(0.074)	(0.001)	(0.667)	(0.011)	(800.0)	(0.294)	(0.049)

The matching procedure resulted in two groups, with 50 subjects in each group. The group with sick leave due to PGP and the group with no sick leave had similar PGP intensities (approximately 18/100). Univariate Mann–Whitney U test revealed that the group with no sick leave (the

coping group) on average had longer education (15.8 vs 14.8 years), p = 0.022 and higher work-satisfaction (4.66 vs 4.32), p = 0.014. Finally, the scores on several ODI items were lower in the coping group (with no sick leave), as seen in Table 5.

The methods section in Paper II describes an analysis using sequential linear regression with the total number of weeks as the dependent variable. This analysis is not included in the result section due to an unfortunate error, and the analysis was not performed as planned.

Table 5: Disability (ODI scores) in women with and without sick leave for PGP. For each item, the mean score and SD (in brackets) are displayed.

ODI item	No sick leave for PGP	Sick leave for PGP	E.S.	p
Pain intensity	2.30 (0.84)	2.33 (0.88)	0.039	=.954
Personal care	0.90 (1.11)	0.94 (1.14)	0.033	=.889
Lifting	1.40 (1.21)	1.90 (1.29)	0.395	=.044
Walking	0.96 (1.01)	1.44 (0.97)	0.483	=.011
Sitting	0.96 (1.00)	1.46 (0.87)	0.528	=.003*
Standing	1.50 (1.33)	2.00 (1.22)	0.392	=.031
Sleeping	1.12 (0.85)	1.48 (1.03)	0.379	=.113
Sex	1.18 (1.41)	1.23 (1.49)	0.037	=.969
Social function	1.26 (1.27)	1.33 (1.31)	0.056	=.677
Travelling	0.92 (1.12)	1.25 (1.23)	0.280	=.180

ES Effect size (Cohen's d); Bonferroni-corrected alpha = 0.005

In a quasi-experiment, a matching procedure (i.e., when the intervention is not randomly assigned) enables comparison of groups with similar characteristics to estimate the effect of an intervention, the outcome in this process differ from the ODI-results presented in Table 3 as the effect sizes between the groups are very different for the different items. If a strict Bonferroni-correction is applied, only the ODI score for sitting is significantly higher in the group with sick leave due to PGP (p = 0.003) compared to the group not on sick leave.

3.2 The prospective data collection (Paper III)

Five hundred and six women agreed to participate in this study, but three were excluded due to incomplete data. At the ultrasound examination in pregnancy week 18, 42% (212/503) of the women reported pain in the lumbopelvic region. A clinical examination revealed that 39% (196/503) of the women fulfilled the criteria for a probable PGP diagnosis, and 27% (137/503) showed a positive response to ASLR and P4 tests, which were the tests most frequently found positive, followed by the long dorsal ligament test and the symphysis provocation test.

A further 12 women reported pelvic pain but did not get any of the recommended clinical tests positive. Hence they were placed in the ASLR and P4 tests negative group.

3.2.1 Demographic and clinical features

In Paper III does Tables 1 and 2 illustrate significant differences in some demographic and clinical features at baseline and test outcomes between the women with and without pelvic pain. Columns a - e add up to 503. Column f contains all positive ASLR and P4 test, i.e., the sum of columns c - e, for comparison with columns a and b. Column b contains women with other tests positive, which are described in paper III on page 2 under the subheading "Sequence of stability and pain provocation tests for PGP". The women reported pelvic pain, but all tests, including ASLR and P4, were negative, hence they were placed in column b.

Women with positive P4 and ASLR tests experienced a heavier workload. They also presented with a higher BMI at week 18, exercised less both before and during pregnancy, and reported a higher rate of feeling depressed during the pregnancy.

Physical disability (ODI) and pain level (NRS) at week 18 were higher in women with positive tests than in women reporting pain but having negative P4 and ASLR tests. Among women with a positive ASLR but negative P4 test, the highest number of previous pregnancies was reported.

Most of the women were with a positive ASLR and negative P4 test (58%). Almost half of the women with both P4 and ASLR tests positive (47%) had been on sick leave during pregnancy. Among women with a positive P4 and negative ASLR, 38 % had been on sick leave, and among women with both tests negative, 30% had been on sick leave during pregnancy.

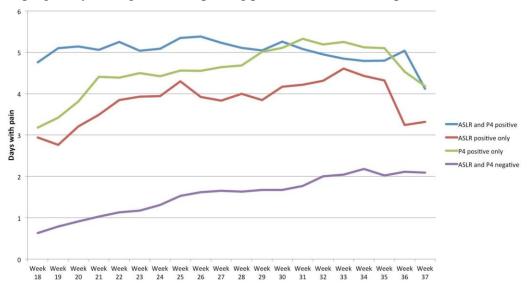
3.2.2 SMS-tracking

The SMS response rate was 75% (2148 responses to 2877 sent messages). Due to a declining response at the end of the pregnancy, we stopped our analysis at week 38. A GEE analysis revealed that all entered variables except age were significant predictors for the number of days with bothersome pelvic pain. Further, there was a significant interaction between the diagnostic group and time, implying that the number of days with bothersome pelvic pain developed differently for the different test groups.

3.2.3 PGP course

The estimated weekly mean number of days with bothersome pelvic pain for the different test groups is presented in Figure 1 on page 60. Women with both P4 and ASLR tests positive experienced a high weekly mean number of days (4.8 days) with bothersome pelvic pain from week 18 and throughout the pregnancy. Women with both tests negative showed a steadily increasing number of bothersome days throughout the pregnancy, from 0.5 days in week 18 to 2 days in week 37. The group with a P4 positive and an ASLR negative test had approximately three days of bothersome pelvic pain in week 18, which was considerably lower than the group with both tests positive. However, the number of days with pain rapidly increased. In week 29, the mean number of days with bothersome pelvic pain equalled the group with both tests positive (5.1 days) and matched this group.

Figure 1: Estimated mean number of bothersome days in the latter half of pregnancy among women reporting pain in the lumbar and pelvic area.



Women with a positive ASLR and a negative P4 test also showed three days of bothersome pelvic pain in week 18. However, they never reached the mean number of bothersome days reported by women with P4 test positive and ASLR negative and both P4 and ASLR tests positive.

3.2.4 Factors predicting the number of bothersome days per week

The estimated rate for experiencing bothersome days was 7.5 times higher in women with both ASLR and P4 tests positive than women with both tests negative (Table 6). Women with both tests positive had twice the number of bothersome days per week than the negative test group. For women with either P4 or ASLR test positive, the mean number of bothersome days was lower but still approximately 1.5 times higher than for women with both tests negative. For every pregnancy, the mean number of bothersome days increased by 13.5%. Even a slightly higher BMI had a significant impact on the mean number of bothersome days. Age had no impact on this outcome.

Table 6. Parameter estimates.

			95% Wald Confidence Interval		Hypothesis Test				95% Wald Confidence Interval for Exp(B)	
Paramet er	В	Std. Error	Lowe r	Uppe r	Wald Chi- square	d f	Sig.	Exp(B)	Lowe r	Upper
P4 and ASLR positive	2.02	.1581	1.71 2	2.33 1	163.48 1	1	0.00	7.549	5.53 7	10.29 1
ASLR positive	1.54 0	.229 7	1.09 0	1.99 1	44.992	1	0.00	4.667	2.97 5	7.319
P4 positive	1.61 7	.183 2	1.25 8	1.97 6	77.852	1	0.00	5.037	3.51 7	7.213
Negative tests	0 *							1		
Age	- 0.00 9	.006 7	- 0.02 2	0.00 4	1.807	1	0.17 9	0.991	0.97 8	1.004
No. of births	0.13 5	.041 5	0.05 3	0.21 6	10.551	1	0.00 1	1.144	1.05 5	1.241
BMI before pregnanc y	0.01 3	.006 5	0.00	0.02 6	3.961	1	0.04 7	1.013	1.00	1.026

4 DISCUSSION

Here, PGP incidence and prevalence were difficult to estimate because of the variability in terminology and diagnostic approaches in this field, which have changed during these studies. The project was initially based on the term "pelvic pain (PP) in pregnancy", illustrated in the first paper, with a plan to follow the diagnostic procedure introduced in the European guidelines and conduct thorough examinations for the correct diagnosis of PGP in the second data collection [7]. However, despite the new term pelvic girdle pain, PGP, and its recommended diagnostic testing procedure, a diverse nomenclature still prevails, with and without physical testing, including pelvic girdle syndrome [89, 141], pregnancyrelated lumbopelvic pain [14], pregnancy-related symphysis pubis dysfunction [142], pregnancy-related low back and pelvic pain [143], pregnancy-induced low back pain [144], posterior pelvic girdle pain (PPGP) [145], and pregnancy-related pelvic girdle pain [146]. Although the term PGP may be somewhat too narrow, focusing solely on the -pain associated with this condition without considering how it affects a woman's activity level and participation in daily activities, we chose to

employ this term in our remaining studies since it is the term recommended by the European guidelines [7].

4.1 Methodological considerations

4.1.1 Retrospective study

The first of this project's two data collections were retrospective. Although retrospective studies in this field are not uncommon, we could not identify any study comparing postpartum recall of pelvic pain in pregnancy with prospectively prepartum recorded pain data. Whereas some studies find assessing pain retrospectively unreliable [126, 147, 148], others report acceptable validity levels up to a 3-months recall period [149, 150]. It has been reported that pain is usually overestimated when pain intensity is high and underestimated when it is low [126, 148, 151]. Pain and disability recall were found to be influenced by current pain and disability levels. In contrast, the influence of pain relief and disability reporting at the initial consultation one year earlier were less influential [152]. However, recall of chronic pain in terms of its average intensity, interference with activities (disability due to pain), number of

days with pain and number of days with activity limitation in the previous six months show acceptable validity levels [150].

Pain recall seems to be good on a group level, but between and within individuals, the variability between monthly, quarterly, and yearly retrospective measurements are high [153]. Social support from, e.g., colleagues make you talk more about your pain and thereby increase the awareness of the pain and thus lead to an over-reporting of symptoms [153]. A meta-analysis aimed at determining whether pregnancy is associated with objective declines in cognitive functioning included 20 studies [154]. The conclusion was that performance related to memory, and executive functioning was significantly poorer in pregnant than in control women, particularly during the third trimester.

A literature review of studies using objective memory testing suggests that a mild antepartum decline in explicit verbal recall occurs in some women [155]. Diminished memory function may occur in a specific subset of pregnant women who display depressive symptoms associated with pregnancy [155].

Thus, the retrospectively collected data on pain during pregnancy may, in some circumstances, be biased. There was, unfortunately, no data in the project on the participants' social support or depression during pregnancy to allow controlling for these factors.

The retrospective data collection was completed for four months, in which all women (n=1204) giving birth at Stavanger University Hospital were asked to participate in the study. The hospital has the only birth department in the southern part of the county of Rogaland, Norway, with a population of about 330,000 inhabitants. The 994 women eligible for the study were found to match the general delivery database at the hospital regarding the age and parity of all women who gave birth during the study period. Thus, we are confident that the study sample represents women giving birth in the county of Rogaland.

Three-hundred and thirty-six (n=336) women did not return a questionnaire, and 89 did not complete a received questionnaire, resulting in a relatively low response rate (569/994, 57%), probably reflecting other priorities of the women in the stressful situation shortly after delivery. The response rate could also be caused by a particular

lower interest in the study by women not experiencing any pain during pregnancy, introducing a bias toward overestimating the frequency of moderate to severe pelvic and lumbar pain Norwegian population. Research has shown that studies with low response rates, even as low as 20%, may generate more accurate results than studies with higher response rates of 60% to 70% [156]. Reviews of response rates ranging from 5 % to 54 % have also reported that studies with a much lower response were often only marginally less accurate than those with higher response rates [157, 158]. A low response rate does not naturally mean low validity. It only illustrates a potentially greater risk for this. Response rates remain informative and are on their own not a good representative for study validity [159].

The impact of PGP on general functioning was measured through patients' self-reports. Most of the instruments frequently utilized in clinical studies in this field were developed and tested for psychometric properties in patients with LBP, including disability instruments such as the Oswestry Disability Index (ODI) [122] and the Disability Rating Index (DRI) [160]. A condition-specific instrument for PGP, the Pelvic Girdle Questionnaire (PGQ), was developed one year after this project's

data collection was completed. The PGQ assesses activity limitations and symptoms and is found to have good internal consistency, test-retest reliability, good construct validity, satisfactory discriminant validity, and reliability [161]. It is important to provide evidence for the comparative performance of instruments to inform future selection in research, and the PGQ has been used extensively in recent years [34, 67, 162-164].

An existing suitable questionnaire for the retrospective data collection objectives could not be found at the time of data collection. Instead, the research team, consisting of competent researchers in obstetrics, psychology, and neurology, constructed a questionnaire based on their expertise and experience, using validated scales (Appendix 1). Answers in the completed questionnaires did not produce any extreme outliers, so we are confident of their face and content validity.

In a Swedish study with a similar objective and design as in Paper I, 72% of pregnant women had LBP or PGP during pregnancy [40]. In our study, 71% had experienced pain in the lumbopelvic area. Identical results from two neighbouring countries suggest the validity of the findings in Paper I. However, it is important to note that studies using questionnaires that have not been validated in the population of interest may be subject to

measurement error, and any conclusions drawn should be interpreted in the light of this fact.

4.1.2 Prospective study

In striving for a homogenous study population, all women were consecutively recruited for data collection at the same stage in pregnancy, week 18, which is the baseline in this prospective data collection. On arrival for their 18th-week routine ultrasound examination, they were all screened by independent midwives for possible pelvic pain. Outcomes of the diagnostic tests recommended in the European guidelines for the diagnosis of PGP were systematically recorded. Two tests at baseline had by far the highest number positive reactions, the two with the highest specificity and highest sensitivity for sacroiliac joint pain and functional pelvic test: the posterior pelvic pain provocation test and the active straight leg raise test [7], i.e., diagnostic tests for excessive force closure and reduced force closure respectively.

Women with these two tests positive were followed with weekly SMS, introduced for the first time in this research field. The SMS instrument had previously been used in research on low back pain and had shown good recall and compliance and instant, easy data handling. This project

experienced good patient compliance and data handling, but with a lower response rate of 75% due to failing technology in some telecommunication providers. The SMS-responses provided information on the number of days each week the participants had experienced bothersome PGP.

The text messaging monitoring objective was to follow up the examination at baseline with a short question once per week, which could quickly be replied to with a one-digit answer. The initial intention was to send a question on the women's current level of pain. However, an earlier study on LBP, using the SMS technology, had successfully administered the term "bothersome" instead of "pain" [130]. The measure "bothersomeness" has shown associations with measures of pain, disability, psychological health, and work absence. It has predicted six months outcome in patients with severe LBP [165]. Thus, instead of a question using the measures "pain" or painful", the SMS question in this prospective study had the following wording: 'How many days during the previous week has your pelvic pain been bothersome?' [120]. The participants were informed at baseline about the SMS question and

how to reply. This technology seemed to be user-friendly as compliance was high.

4.1.3 Matching procedure

In order to investigate factors that may modify the effect of PGP on sick leave, a matching procedure of retrospective data (Paper II) was conducted. This procedure aimed to find pairs who differed with respect to sick leave but who matched on certain baseline characteristics to assess the effect of potential coping on sick leave. The matching procedure was carried out without obstacles. Challenges were to find random women been on sick leave, matching the average PGP score of women without sick leave. However, a macro found matching pairs, creating two groups with 50 subjects in each. The implication for previous results was that the scores on several ODI-items now differed from earlier analysis. Effect sizes between the groups now showed that women with longer education, higher work satisfaction and fewer problems with sitting, walking, and standing, were less likely to take sick leave in pregnancy, despite the same pain intensity as women being on sick leave.

4.2 Discussion of results

4.2.1 Paper I

The retrospective data collection results revealed that nearly half (47.6%) of the women in Rogaland, Norway, experienced moderate and severe pain in the lumbopelvic region during pregnancy. More specifically, moderate and severe combined LBPP was experienced by 21.6% (n = 123), and moderate and severe PGP was experienced by 26.0% (n = 148). Similarly, detailed pain information is sparse in the literature. However, this finding concurs with an earlier retrospective study with an identical design on LBPP in pregnancy, in which 23.2% of the women were rating 7 and above on VAS [40]. One other small retrospective study on PGP in athletes reported pain location but not pain severity [166]. They reported pain location mainly at the pubic symphysis (33,3%) and both sacroiliac joints (29,6%). Corresponding numbers in this thesis project was 19,3% and 17,2%, respectively. The primary pain location in this project (reported by 32,5%) was at all three pelvic joints simultaneously (a pelvic girdle syndrome), reported by 22,5 % of the athletes [166]. One other study has reported that disability in pelvic girdle pain is associated with pain location and that pain at the pubic symphysis combined with

bilateral pain at sacroiliac joints produced the largest impact on Disability Rating Index [102].

One reason for the difference in symptom distribution between the athletes mentioned above and the women in this thesis project may be that a high number of years of regular leisure physical activity before pregnancy decreases the risk of developing LBPP [135]. It has also been reported that women who exercise regularly and engage in high impact exercises up to five times weekly before the first pregnancy may have a reduced risk of pelvic girdle pain in pregnancy [100]. The mechanisms by which pre-pregnancy exercise influences the risk of pelvic girdle pain remains unknown. However, it is well known that both aerobic exercise and resistance training have a hypoalgesic effect on pain in healthy nonpregnant individuals and chronic pain patients [167]. Though exercise's long-term effect on pain remains unclear, women who exercise regularly pre-pregnancy are more likely to continue to exercise throughout pregnancy [168]. It is believed that the relationship between leisure-time exercise and pelvic girdle pain depends on both frequency and types of exercise [100].

The retrospective data also indicate that women with moderate and severe combined LBPP had approximately 50% more sick leave days than women with PGP or LBP at the same pain level, and 2 - 3 times more sick leave days than women without pain or with only mild pain. A result not surprising since the most frequently reported cause of sick leave among pregnant women in other studies are combined LBPP [112, 113]. However, previous studies have not categorized the level of pain for either LBPP or PGP with sick leave during pregnancy.

A relatively recent cross-sectional study comparing 12 European countries found that 50.6% of the women had been on sick leave at some point during pregnancy. Nonetheless, the rates varied greatly, ranging from 31.7% (Sweden) to 71.3% (Poland) [113]. Duration of sick leave also showed considerable variation, with one Norwegian study reporting 50% of pregnant women being off work between 4 and 16 weeks [112]. This retrospective data collection disclosed that 34% of women reported a mean sick leave duration of 9,6 weeks for pain in the lumbopelvic region, which concurs relatively well with other studies regarding prevalence and duration.

The results showed that work satisfaction, problems with lifting, sleeping, and pain intensity predicted sick leave for women with PGP. This result concurs with previous research where factors related to sick leave were sleeping problems, hyperemesis, chronic pain before or during pregnancy, workplace conflicts, multiparity, previous depression, insomnia and lower education [112]. One other study reported that disability, pain intensity and occupation were associated with sick leave due to lumbopelvic pain [117]. Thus, pain intensity, sleeping problems, and type of work/work satisfaction seem to be common factors for sick leave in pregnancy.

For analysis of function, this project utilized ODI in which "moderate disability" is considered in the scoring window of 21 – 40 [122]. The analysis regarding the general level of function during pregnancy displayed moderate ODI values of 21.92% for LBP, 33.24% for PGP, and 37.66% for LBPP. This outcome of moderate disability concurs with previous research conducted in Australia in which the mean ODI scores for women with both LBP and PGP was 33.5%, thus a higher disability level than women with PGP (26%), or LBP alone (18%) [31].

In this retrospective part of the thesis, 71% of the women had experienced pain in the lumbopelvic region. A previous Swedish study with a similar design and identical result lends strength to the validity of these findings [40].

4.2.2 Paper II

Further analysis of the data collected retrospectively demonstrated that women with long education, high work satisfaction, and little problem with sitting, walking, and standing, were less likely to take sick leave during pregnancy compared to women with short education, low work satisfaction and problems with sitting, standing, and walking, despite having the same pain intensity. These findings are partly in agreement with a recent study reporting that an advanced degree education was protective against sick leave for pregnant women [115]. Patients with lower educational levels, suffering from pain in the lumbopelvic area, appear to have less knowledge about self-care compared to patients with higher education [169].

The retrospective data analysis revealed that ADL's were significantly more difficult to carry out for pregnant women on sick leave for PGP than for women on sick leave for other causes. Independent significant risk factors were lower education, heavy workload, and low work satisfaction. This find concurs with previous research in which PGP is reported to affect daily activities, work ability and quality of life [24, 41, 170].

The primary reason for long-term sick leave, more than 14 consecutive days, seem to be conflicts at work, high workload, reproductive occupational hazards, and different national sick leave policies across Europe during pregnancy [112, 113, 171, 172]. Recently published studies have displayed that nearly 10% of women with long-term sick leave during pregnancy reported such work-related reasons for sick leave [113, 115]. Furthermore, studies within the Danish national birth cohort have found that a psychosocially demanding work environment and some physically demanding work measures are associated with an increased risk of pelvic pain in pregnancy [173]. Also, working night

shifts during pregnancy significantly shifts longer than 12 hours and increases sick leave risk [174].

The matching procedure displayed that the group with no sick leave had, on average, longer education and higher work-satisfaction and the scores on several ODI items were lower in this group compared to the group of women who were on sick leave. This procedure also revealed unexpected differences in pain tolerance among those who did not take sick leave. What generates these differences is unclear, but may be due to education, work situation, and/or work posture.

Sitting was found to be the most challenging activity for women on sick leave. In a recent study, the prevalence of pain in the sitting position was in 73% for patients with a SIJ disorder, higher than for patients with lumbar disc herniation (49%) and lumbar spinal canal stenosis (20%) [175]. It is proposed that, in the sitting position, as the ischial tuberosity is fixed on the seat, distortion in the SIJ occurs, and pain develops [175]. The difficulty in performing a seated task during gestation may not be a unique effect of biomechanical adaptations due to pregnancy and result

from the environmental context [176, 177]. Patients with good social support may have access to good personal care, thus reducing their perceived pain and interfering with daily activities [178].

Our study found a higher prevalence of LBPP in women in the third trimester [20]. Such findings may be explained by the increase in muscle and ligament overload, especially during the third gestational trimester, when it is greater due to hormonal activity and the gravid uterus' growth [179].

An unfortunate unprecise use of PGP terms in Paper II may be misleading and needs clarification since they represent different constructs. "PGP score" (page 3) refers to pelvic pain score from the questionnaire, and "PGP intensity" (page 4) and "PGP intensity score" (page 6) refers to the ODI item "Pain intensity".

4.2.3 Paper III

This research appears to be the first, and to date, the only, prospective longitudinal study in which women with PGP in pregnancy have had a clinical examination mid-pregnancy and then have been followed with

regular short message service. A literature search has not presented any paper outside this thesis demonstrating future longitudinal PGP trajectories antepartum. Thus, it is the first attempt to follow the course of PGP subclassified groups through the second half of pregnancy.

Our analysis revealed that if both P4 and ASLR tests were positive in pregnancy week 18, a persistent PGP of more than five days/week throughout the remainder of the pregnancy could be predicted. If either test was positive in week 18, a similar course was shown. However, women with a positive P4 test only revealed a more uncomfortable course than women with a positive ASLR test. The latter never reached the bothersome levels of the P4-ASLR test group and the P4 only test group. Thus, women with the highest number of bothersome days per week with PGP were exposed to long dorsal sacroiliac ligament (LDL) strain (as indicated by the positive P4 test). Overloaded sacroiliac joints (SIJs) (as indicated by the positive P4 test), accompanied by reduced sacroiliac force closure (i.e., loss of functional control due to loss of co-contraction of the lumbopelvic muscles) (as the ASLR test was positive).

Although women who had a positive ASLR test and negative P4 test at baseline presented a comparatively low mean number of bothersome

days with pain, they also had the highest mean number of previous pregnancies and the highest mean rate of pelvic pain in previous pregnancies. Our data also revealed that they exercised more frequently both before and during the present pregnancy compared to women in the other positive test groups. Regarding the level of pain, a recent systematic review found that prenatal exercise is an effective treatment to decrease the severity of LBP, PGP and LBPP during pregnancy [180]. Although the prenatal exercise was not found to decrease the odds of LBP, PGP and LBPP during pregnancy results from a meta-analysis provided evidence that different types of exercise performed alone or in combination, such as yoga, general or specific strengthening exercise and aerobics performed anywhere from once per week to once per day, significantly reduced the severity of LBP, PGP or LBPP related symptoms during pregnancy[180].

Considering that LBPP affects more than half of pregnant women and is associated with disability, depression, reduced quality of life and higher prevalence of sick leave during pregnancy [181], exercise may offer a cost-effective self-management strategy option to expecting

mothers as part of a multimodal approach to decrease symptom severity [181].

Since sufficient force closure of the sacroiliac joints requires appropriate muscular, ligamentous, and fascial interaction, women with pelvic pain in previous pregnancies, may have experienced improved muscle activation, recovered function and decreased pain from exercise. Additionally, pain prevention and rehabilitation experience in previous pregnancies may work as an incitement to engage in physical activity and regular exercise, both before and during pregnancy.

Although the mechanisms through which exercise may reduce pain severity remains unclear, it is suggested that physical activity lessens the degree of biomechanical change occurring as pregnancy advances, such as decreasing the load on the spine, increasing joint stabilization and contributing to better spinal alignment and segmental motion [182]. From a more general standpoint, exercise may help reverse trunk muscle imbalance [60] or initiate a pain desensitization process leading to increased pain detection threshold [183].

Our analysis also revealed a significant difference in depression between women with different positive clinical tests. Distress has previously been identified as a factor associated with a higher likelihood of PGP in pregnancy, with a higher BMI and a higher gestational age [26]. One previous study found that distress contributes to disability but not to pain intensity [32]. Approximately 10% of women experience postpartum depression, with nearly 25% of them still in treatment after one year [184]. One review also provided strong evidence that physically active women experience significantly fewer symptoms of depression during the postpartum period compared with their inactive counterparts [185]. Nevertheless, some individuals seem to tolerate pain better, have less catastrophizing tendencies and show more positive social response to pain, regardless of exposure to stressful circumstances and/or internal distress [186].

The parameter estimates revealed that a slightly higher than average prepregnancy BMI significantly impacted the mean number of bothersome days. However, with a difference of only 1,3 %, the clinical relevance is questionable. Interestingly, in a recent published study on PGP in pregnancy does women with a combination of generalized joint hypermobility (GJH) and BMI ≥25 kg/m² report higher pain compared to women with normal joint mobility and BMI <25 kg/m² [187]. Nonetheless, the researchers state that the association between GJH and BMI should be interpreted with caution due to the small study sample.

Finally, women with the possibility to control their own work situation have better health during pregnancy than women without such options. As indicated in this study, and confirmed in other research, many pregnant women seem to benefit from exercise as it increases pain tolerance, improves or maintains physical fitness, helps with weight management, reduces the risk of gestational diabetes in obese women, and enhances psychological well-being [100, 188-190].

4.3 What does this thesis contribute to our knowledge about PGP?

Our present findings confirm and re-emphasize that PGP can cause major impairment of a pregnant woman's health and well-being, as well as constituting a burden to society as a frequent and major cause of sick leave. We retrospectively showed how PGP affects a pregnant woman's ADL, revealing unexpected differences in tolerance for pain among those who did not take sick leave. Those who took less sick leave demonstrated higher pain tolerance, which was associated with their level of education and their situation and/or posture at work.

In addition, we demonstrate here that if both ASLR and P4 tests are positive mid-pregnancy, persistent and bothersome PGP can be expected for more than five days each week throughout the remainder of the pregnancy. The number of days each week with bothersome pelvic pain increases for every additional pregnancy.

4.4 Clinical implications

Clinicians need to be aware of the potential risk factors for PGP identified here, i.e., less education, a heavy workload and dissatisfaction with one's work. Moreover, the application of ASLR and P4 tests to assess the likelihood of persistent PGP will help the caregiver, together with his/her patient, design a treatment plan that can alleviate the pain involving, e.g., exercise, management of stress and a reduced workload.

4.5 Implications for future research in this area

Usage of the SMS-Track system to collect research data, an apparent strength of the present work, is highly recommended. This approach results in a high response rate and provides information concerning the participant's situation instantly. In addition, the immediate recording of the responses minimizes further handling of the data and, thereby, the risk of error.

The potential risk factors for PGP identified here should now be examined in a prospective study that controls for confounders and includes the clinical examination of pregnant women's sub-groups offered different therapeutic interventions in a blinded design.

Some women with PGP appear to tolerate their symptoms more effectively than others, e.g., the women with higher education and more work satisfaction took less sick leave, even though their pain level was the same as in the sub-groups who take more sick leave. We also recommend the performance of a prospective longitudinal study designed to see if appropriate modification of workplaces can reduce the amount of sick leave taken by women who experience PGP.

Finally, a gold standard for diagnosing PGP, particularly for the number of clinical tests to be performed, is still lacking. Further efforts to identify predictive and preventive factors, as well as diagnostic tests for PGP during pregnancy, are warranted. For example, it would be interesting to determine whether women with a history of PGP have elevated pain-related anxiety that influences their experience of pain.

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6 Erratum

In Paper III, "Discussion" section, it is written "As indicated in this study and confirmed in previous studies, most pregnant women seem to benefit from exercise as it increases pain tolerance, improves or maintains physical fitness, helps with weight management, reduces the risk of gestational diabetes in obese women, and enhances psychological well-being". The paragraph should read "As indicated in this study, and confirmed in other research, many pregnant women seem to benefit from exercise as it increases pain tolerance, improves or maintains physical fitness, helps with weight management, reduces the risk of gestational diabetes in obese women, and enhances psychological well-being."

7 PAPERS

7.1 Paper I

PREVALENCE OF LOW BACK AND PELVIC PAIN DURING PREGNANCY IN A NORWEGIAN POPULATION

Stefan Malmqvist, DC, MSc, a Inger Kjaermann, MSChir, a Knut Andersen, DC, b Inger Okland, MD, c Kolbjøm Brønnick, PhD, d and Jan Petter Larsen, MD, PhD e

ABSTRACT

Objective: The purpose of this study was to investigate the cumulative prevalence of low back pain (LBP), pelvic pain (PP), and lumbopelvic pain during pregnancy, including features possibly associated with development

pregnancy-related PP, in an unselected population of women.

Methods: A retrospective cohort study was conducted in which all women giving birth at Stavanger University hospital in a 4-month period were asked to participate and to fill in a questionnaire on demographic features, pain, disability, and Oswestry Disability Index. Inclusion criteria were singleton pregnancy of at least 36 weeks and competence in the Norwegian language.

Results: Nearly 50% of the women experienced moderate and severe PP during pregnancy. Approximately 50%

results: Nearry 30% of the women experienced inductate and severe Pr during pregnancy. Approximately 30% of them had PP syndrome, whereas the other half experienced lumbopelvic pain. Ten percent of the women experienced moderate and severe LBP alone. These pain syndromes increased sick leave and impaired general level of function during pregnancy. Approximately 50% of women with PP had pain in the area of the symphysis. The analysis of risk factors did not present a unidirectional and clear picture.

Conclusions: Pelvic pain in pregnant women is a health care challenge in which moderate and severe pain develops rather early and has important implications for society. The observed associations between possible causative factors and readed according to the property of the p

and moderate and severe LBP and PP in this study may, together with results from other studies, bring some valuable insights into their multifactorial influences and provide background information for future studies. (J Manipulative

Key Indexing Terms: Pelvic Pain; Low Back Pain; Pregnancy; Risk Factors; Retrospective Studies

ow back pain (LBP) and pelvic pain (PP) are common conditions in many cultures during pregnancy. 1-8 Two Swedish studies presented a

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prevalence of 54% for PP, 17% for LBP, and 29% to 72% for combined symptoms. ^{1,2} One Dutch study showed a prevalence of 7% for self-reported PP during pregnancy, whereas an Iranian study revealed that 28% of pregnant women had PP, 13% had LBP, and 8% had combined symptoms. ^{3,4} An international study concluded that PP in pregnancy does not vary according to geography or socioeconomy and presented a PP prevalence of 49% in Sweden, 66% in Tanzania, 77% in Finland, and 81% in Zanzibar, with an overall similarity of symptom location and degree of pain. 5 Furthermore, an Australian retrospective study showed that 35% of women had experienced LBP during pregnancy⁶; in a Danish study, Albert et al found a 20% cumulative prevalence of isolated PP in pregnancy, and two-thirds of the pregnant women report LBP in a cross-sectional study in the United States by Skaggs et al. 8 Interestingly, most women report their first episode ever of LBP to occur during pregnancy. 9,10
Although these are common complaints in pregnancy,

the etiology is still unknown, and the pathophysiology is unclear. In addition, low back and pelvic disorders during

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Malmqvist et al Cumulative Pelvic Pain During Pregnancy

pregnancy are considered a major public health issue.11 Sick leave for LBP or PP during pregnancy has been shown to occur in 37% to 72%, and the period of sick leave is on average between 12 and 15 weeks. 9-11

The wide range of the reported frequency of these complaints during pregnancy may be caused by methodo-logical differences. Most of the previous studies are based on selected populations of pregnant women, and a clearly defined clinical diagnosis of PP is missing. In lack of a clear definition, a diverse nomenclature for these conditions has been used: pelvic insufficiency, LBP, lumbosacral pain, symphysiolysis, pelvic syndrome, posterior PP, and pregnancy-related PP. 11,12

Guidelines for identification and classification of pregnancy-related PP have been established in later years, based on physical examination and history taking. 13 However, studies of risk factors related to the development of PP during pregnancy have not yet been able to reveal one single dominant causative factor, but several different physical and psychosocial factors have been found to correlate with self-reported pain. 3,9,13-15 An increased abdominal diameter, higher body mass index (BMI), muscle dysfunction, and fetal weight are clearly associated with LBP and PP during pregnancy. ¹⁶⁻¹⁸ A general increase in mobility of joints during pregnancy has also been described, ¹⁹ and Sipko et al²⁰ found that the most frequently irritated ligaments during pregnancy are the interspinous, iliolumbar, and sacroiliac.

However, some suggest that pain during pregnancy is not only explained by biomechanical factors alone;

psychosocial factors too seem to be important. 21,22

The objectives of this study were to investigate the cumulative prevalence of LBP, PP, and combined lumbopelvic pain (LBPP) in an unselected population of women, giving birth during a 4-month period, and to study clinical and demographic features possibly associated with the development of pregnancy-related PP.

METHODS

This study is a retrospective longitudinal cohort study with data collection over the period of March to June 2009, at the maternity ward of Stavanger University Hospital, Norway. All women giving birth at the hospital during this period were asked to participate and to fill in the questionnaire. Inclusion criteria were a term singleton pregnancy of at least 36 weeks and good competence in the Norwegian language. The hospital has the only birth department in the southern part of the county of Rogaland, with a population of approximately 330000 inhabitants. The annual number of deliveries at the hospital varies between 4400 and 4800.

Within 24 hours after delivery, the women received both oral and written information from a midwife. Participation

was voluntary, but to obtain inclusion of an unselected sample, all women were encouraged to give their informed consent to participate. The study was carried out in accordance with the Helsinki Declaration II and was approved by the Regional Ethics Committee of Western Norway. All subjects consented to participate in this study.

To assess if the study population was a representative sample of the delivering women, we compared demograph and clinical characteristics of the study population with that of the general delivery database at the hospital.

The women completed a questionnaire on demographic features and pain, disability, and exercising before and during pregnancy. The questionnaire was produced and specially designed by the research group, based on previous studies and the experience of the team.

Information on presence of pain, pain distribution, and level of pain intensity was collected for both previous and present pregnancies. Pain intensity for both LBP and PP was rated retrospectively on a numerical rating scale (NRS) from 0 to 100 for each month of the pregnancy, to collect information on appearance of symptoms and peak intensity pain during pregnancy. There are 2 versions of the NRS: 0 to 10 and 0 to 100, and in this study, the latter one was used to record a detailed pain level. 23,24 Reports on pain distribution were obtained by asking for

drawings on figures of the lower back and pelvic areas. Studies evaluating pelvic girdle pain occasionally use pain drawings, in spite of clinical experience showing that some women have difficulties in anatomically locating the pain on a drawing. ^{2,3,17,25,26} In this study, there were 3 figures: 1 low back and 2 pelvic (front and back), all with explanation of the regions involved. The pain drawings were used to differentiate between low back and PP and to identify location of pain in either area. ²⁷

Information on disability was collected through the Oswestry Disability Index (ODI), which is one of the principal condition-specific outcome measures for defining disabling effects from spinal disorders. ^{28,29}

The questionnaires also provided information on number of years of education, level of physical work load (a 5-level scale running from "very light" to "very heavy"—a score of 3 or more was used to characterize women with a heavy work load), work satisfaction (a 5-level scale running from "very bad" to "very good"—a score of 3 or less was used to describe women with a low satisfaction at work), sick leave during pregnancy (time periods and percentages), height, and weight before pregnancy and at delivery. Body mass index was calculated and used in the data analyses.

Further variables included were number of previous

births, pain in previous pregnancies, pain during the last year before pregnancy, and exercising habits ("Did you exercise regularly, at least 2-3 times per week before pregnancy/during pregnancy?").

Demographic and clinical variables	No pain	Mild pain; all types	Moderate to severe LBP	Moderate to severe PP	Moderate to severe LBPP	All women
n (%) "	158 (27.8)	73 (12.8)	56 (9.8)	148 (26.0)	123 (21.6)	569
Education years, mean (SD)	14.9 (3.6)	15.6 (2.9)	14.6 (3.6)	15.0 (2.9)	14.1 (3.5)	14.7 (3.3)
BMI before pregnancy, mean (SD)	22.8 (3.6)	23.3 (3.4)	23.1 (4.1)	24.5 (4.8)*	24.9 (4.5) *	23.8 (4.2)
LBP in previous pregnancies, n (%)	11 (6.9)	18 (24.6)	20 (35.7)*	24 (16.2)*	56 (45.5)*	124 (21.8)
PP in previous pregnancies, n (%)	12 (7.6)	21 (28.7)	10 (17.8)	64 (43.4)*	54 (43.9)*	165 (29.0)
LBP in the year before pregnancy, n (%)	13 (8.2)	12 (16.4)	19 (33.9)	35 (23.6)	53 (43.0)*	135 (23.7)
PP in the year before pregnancy, n (%)	6 (3.8)	3 (4.1)	2 (3.6)	23 (15.5)*	15 (12.2)*	51 (9.0)
Exercised at least 2-3 times a week before pregnancy, n (%)	68 (43.0)	34 (46.6)	25 (44.6)	44 (29.7)*	47 (38.2)	224 (39.4)
Physically heavy work, n (%)	31 (19.6)	18 (24.6)	18 (32.1)	42 (28.4)	49 (39.8)*	162 (28.4)

^{*} Nonparametric test showing significance in difference between asymptomatic group and symptomatic groups at P ≤ .05 level.
a Eleven women did not fill in the scales for low back and PP.

Statistical Analyses

Descriptive data are presented by mean values and standard deviations for continuous variables and by frequencies for categorical variables. The data on pain symptoms were classified into 3 pain level groups through analyses of variance. Cutoff point with the largest F ratio between mild and moderate and severe pain was found at 35 in the NRS (0-100). Thus, for each type of pain variable, the patients were grouped into 3 categories according to their symptoms: One group with no pain (NRS = 0), one group with mild pain (NRS \leq 35), and one group with moderate and severe pain (NRS > 35). Kruskal-Wallis statistics were used to explore the group differences between pain distri-

bution and prepregnant and pregnant values.

Multivariate hierarchical binary logistic regression analysis was used to assess whether the prepregnancy and pregnancy variables could predict moderate to severe PP as contrasted with no pain. Mild pain was omitted from the analysis. In the first block of the analysis, age, education level, and number of previous births were entered, followed by a block including average level of LBP throughout the by a lock including actual even of 12th integritors in pregnancy, and finally, the last block used backward step-wise regression using the likelihood ratio removal criterion, with the variables workload, BMI before the pregnancy, BMI at birth, feelings of depression during pregnancy, physical activity before, and physical activity during pregnancy. Thus, the first 2 blocks served mainly as statistical controls with forced entry of all variables, before entering the variables to be explored in the final block. All analyses were performed in SPSS 16 (IBM, New York, NY), and results were considered significant at $P \leq .05$.

RESULTS

Study Population

Women who gave birth at Stavanger University Hospital during the inclusion period (n = 1204) were invited to participate in the study. After exclusions, there were 994 vomen eligible for the study. In addition, 336 women did

not return a questionnaire, and 89 did not complete a received questionnaire. The final study population thus consisted of 569 women, which was 58% of the total possible sample.

To examine if the study population was representative, we compared age and parity variables from the study population with that of all women who gave birth during the study period and found a near complete match (results

Tables 1 and 2 show demographic and clinical data for the women included in the study. The mean age of the population was 30 years, mean years of education was 14.7 years, and slightly more than one-third of the women were giving birth for the first time. The mean BMI was 23.8 kg/m², and close to half of the women exercised 2 to 3 times per week before pregnancy. The mean amount of sick leave during the pregnancy was 9.6 weeks.

Frequency of Moderate and Severe Pain in Lumbar and Pelvic Area During Pregnancy

Approximately a quarter of the women did not report any LBPP during the pregnancy. A further 13% had only experienced mild pains. For moderate and severe pain during pregnancy, the cumulative prevalence was 57.4% (n = 327) (Table 3). Moderate and severe combined LBPP was experienced by 21.6% (n = 123), and moderate and severe PP was experienced by 26.0% (n = 148). Almost 10% of the women (n = 56) had moderate and severe pain solely in the lumbar area. Twenty-three percent of all women, that is 40 % of those who developed moderate and severe pain, had such pain already after 5 months of pregnancy (Table 3).

Table 4 shows the distribution of pain in the pelvic area

for all women with PP, for women with moderate and severe combined LBPP, and for those with moderate and severe PP alone. Among all women with such pain, more than half of them (55%) experienced pain at the symphysis. Approximately 20% experienced pain only in this

Table 2. Demographic and clinical features of the participating women according to perceived types of pain in the lumbopelvic area during pregnancy

Demographic and clinical variables	No pain	Mild pain; all types	Moderate to severe LBP	Moderate to severe PP	Moderate to severe LBPP	All women
n (%)*	158 (27.8)	73 (12.8)	56 (9.8)	148 (26.0)	123 (21.6)	569
Age at delivery (y), mean (SD)	30.0 (4.8)	30.1 (4.8)	29.4 (6.2)*	30.5 (4.2)	29.6 (4.4)	30.0 (4.7)
BMI at delivery, mean (SD)	27.7 (3.7)	27.9 (3.2)	28.2 (4.4)	29.6 (5.1)	30.3 (4.8) *	28.8 (4.5)
Primiparous, n (%)	71 (45)	36 (49.3)	25 (44.6)	43 (29.0)	44 (35.8)	219 (38.5)
Exercised at least 2-3 times a week during pregnancy, n (%)	43 (27.2)	20 (27.4)	9 (16.0)	28 (18.9)	12 (9.7) *	117 (20.6)
Weeks of fulltime sick leave during pregnancy, mean (SD)	5.4 (8.82)	6.5 (8.62)	9.1 (10.13)	10.7 (12.26)	15.5 (15.25)	9.6 (12.20)
Received treatment for LBP and/or PP, n (%)	_	14 (19.2)	12 (21.4)	63 (42.6)	64 (52)	156 (27.4)
Mean ODI score (n)	4.68% (35)	14.11% (70)	21.92% (52)	33.24% (145)	37.66% (118)	27.57% (429)

Nonparametric test showing significance in difference between asymptomatic group and symptomatic groups at $P \le .05$ level.

Table 3. Prevalence of moderate and severe pain at different stages of pregnancy

Prevalence	Lumbar pain	95% CI lower	95% CI upper	PP	95% CI lower	95% CI upper	LBPP	95% CI lower	95% CI upper
At 5 mo pregnancy	18 (3.1%)	0.567	1.040	75 (13.2%)	1.098	1.375	39 (6.8%)	0.522	0.697
At 9 mo pregnancy	36 (6.3%)	1.189	1.632	107 (18.8%)	1.459	1.690	95 (16.7%)	0.839	0.949
Cumulative	56 (9.8%)	0.074	0.123	148 (26.0%)	1.098	1.375	123 (21.6%)	0.182	0.250

Approximately 40% had pain at all 3 pelvic joints, and 20% had pain at 1 sacroiliac joint alone. For those with LBP in addition to PP, a pain distribution to all 3 pelvic joints was more common.

Table 2 also shows that women with moderate and severe LBPP have an average sick-leave period of 15.5 weeks, moderate to severe PP 10.7 weeks, and women with moderate to severe LBP 9.1 weeks. Women with mild pain revealed a mean sick-leave period of 6.5 weeks, indicating that moderate to severe pain was clinically different from those with less pain. The same pattern was also found for disability as measured with the ODI. For seeking treatment for LBP and PP during the pregnancy, PP was a stronger driver for this than lumbar pain, and again, moderate to severe pain needed more treatment than mild pain (Table 2).

Potential Risk Factors for Lumbar and PP During Pregnancy

Nonparametric test for exploring group differences between demographic and clinical features, before and during pregnancy for the participating women, are shown according to pain categories in Tables 1 and 2. In the multivariate analysis, we examined the impact of different clinical and demographic variables on all women with moderate and severe PP vs those without any pain. The first block of the binary logistic regression analyses was not statistically significant (P=.379), indicating that age, education, and number of previous births did not contribute to the prediction of moderate to severe PP. Block 2,

however, was highly significant ($P \le .001$), as a high level of LBP reduced the risk of PP (P < .001; odds ratio, 0.845; 95% confidence interval [CI], 0.798-0.894). The final step of the final block consisted of the previously entered variables and the variables remaining (BMI before pregnancy and physical activity before the pregnancy) after the backward stepwise procedure and was also highly significant (P=.001). The resulting omnibus logistic regression model was significant (P<.001, Nagelkerke $R^2=.319$). Average LBP (P<.001; odds ratio, .837; 95% CI, 0.790-0.887), BMI before pregnancy (P = .011; odds ratio, 1.074; 95% CI, 1.016-1.134), and physical activity before pregnancy (P = .015; odds ratio, 1.826; 95% CI, 1.126-2.960) were significant predictor variables in the full multivariate model. Thus, both higher BMI before pregnancy and more physical activity before pregnancy were independent potential risk factors for PP after controlling for age, education, number of previous pregnancies, and LBP. Low back pain reduced the risk of moderate to severe PP.

DISCUSSION

This study shows that nearly 50% of Norwegian women experience moderate and severe PP during pregnancy. Approximately half of them had only PP syndrome, whereas the other half experienced PP combined with LBP. In addition, 10% of the women experienced moderate

Eleven women did not fill in the scales for LBP and PP.

Table 4. Prevalence of moderate and severe pain distribution according to pain drawings

Pain prevalence	All women with PP (348)	95% CI lower	95% CI upper	Moderate and severe PP (148)	95% CI lower	95% CI lower	Moderate and severe LBPP (123)	95% CI lower	95% CI upper
At all pelvic joints	113 (32.5%)	0.804	0.931	40 (27.0%)	0.264	0.44	58 (47.2%)	0.420	0.607
At symphysis	67 (19.3%)	0.656	0.866	33 (22.3%)	0.370	0.615	18 (14.6%)	0.160	0.378
At 1 SI joint	83 (23.9%)	0.651	0.842	38 (25.7%)	0.348	0.567	24 (19.5%)	0.190	0.389
At both SI joints	60 (17.2%)	0.599	0.834	27 (16.2%)	0.320	0.580	16 (13.0%)	0.151	0.382
Other areas	7 (2.0%)	-0.066	0.923	1 (0.7%)	-0.207	0.492	2 (1.6%)	-0.166	0.737
Missing data	18 (5.2%)	0.734	0.824	9 (6.1%)	0.368	0.475	5 (4.1%)	0.306	0.410

SI, sacroiliac

and severe LBP alone. These pain syndromes increased the number of weeks of sick leave and impaired the general level of function during pregnancy. Approximately 50% of women with PP had pain in the area of the symphysis. Analysis of risk factors did not present a unidirectional and clear picture. The features that showed a significant association with moderate and severe PP had rather low odds ratios, but the observed influence of a higher BMI before pregnancy may have implications for preventive measures for this important complaint of pregnancy.

for this important complaint of pregnancy.

Several methodological issues must be considered when measuring the frequency of a disease or complaint in a population. A high diagnostic accuracy of the studied disorder is needed, and the result also depends on the use of an unselected patient population. Unfortunately, in both this and in previous studies, ^{2-6,17} pregnancy-related LBP and PP have not been examined using standardized measurements that have received an international acceptance. This is partly because of the self-perceived nature of the complaints themselves and, as in this study, the use of retrospectively collected data on pain during pregnancy. In addition, the European guidelines for pelvic girdle pain require clinical examinations. ¹³ To perform examinations several times throughout pregnancy could have improved the diagnostic accuracy but was not part of the study design and would have been a major undertaking. Nonetheless, we believe that our approach by using moderate and severe pain as criteria for the diagnosis of PP and LBPP is an improvement in study design as compared with previous works. ^{2,3,5,6,17} This strategy will reduce the methodological problems related to recall bias and the impact by minor pains experienced only for shorter periods. ³⁰⁻³²

Our aim was to enroll all women who fulfilled the inclusion criteria during a given period. Unfortunately, our response rate was rather low, probably reflecting other priorities of the women in the stressful situation shortly after delivery. We did, however, introduce several measures to obtain a high awareness of the importance of the study both among the women and midwives. The rather low response rate could be caused by a selective lower interest in the study by women who did not have any pain during pregnancy and therefore introducing a bias toward overestimating the frequency of moderate to severe pelvic and lumbar pain in this Norwegian population. Although this

cannot be ruled out completely, we found that the age and parity of the studied population were identical to the general population of women who give birth at the hospital. We therefore believe the results from this study to be representative for pregnant women in the study area and most probably also in Norway.

In this study, we examined the cumulative prevalence of moderate to severe pain in the lumbopelvic area during pregnancy. A very high proportion (57%) of pregnant women had such pain. In a Swedish study, 72% of pregnant women had LBP or PP during pregnancy. In our study, 71% had experienced any pain. The identical results from 2 different Scandinavian countries lend strength to the validity of our findings. Furthermore, these complaints did have consequences for both sick leave and daily functioning. Those with moderate and severe pain had 2 to 3 times more sick leave days than women without or with only mild pain. In addition, the ODI showed that these complaints have a major impact on the functioning of pregnant women. These findings underline the importance of PP in pregnancy for women and society.

Limitations

Although the retrospective data in this study is hampered by some uncertainty, it revealed that most women who developed moderate and severe PP also had this at the termination of pregnancy and that already 40% had such pain after 5 months of pregnancy. This shows that these pains develop rather early and persist throughout pregnancy.

In this study, we used pain drawings to reveal the pain distribution. The lack of clinical examinations limits the value of these observations, but interestingly, more than 50% experienced pain at the symphysis and nearly 25% had predominating unilateral pain at 1 sacroiliac joint.

The observed associations between possible causative factors and moderate and severe PP did not provide clear answers. The demographic and clinical features from before and during pregnancy that were evaluated in this study could only account for a minor influence. This indicates that pregnancy-related mechanisms not examined in this study may have an important impact on the development of PP. Nevertheless, our observations together with results from

other studies may bring some valuable insights into the multifactorial influences on PP in pregnancy.

Our study has shown that PP is a major health care challenge in pregnant women with implications for society. Our results may provide important background information on potential risk factors for PP, which in future studies should be investigated in a prospective study, controlling for confounders, and include clinical examinations of sub groups of pregnant women who are offered therapeutic interventions in a blinded design.

Practical Applications

- Nearly 50% of the women in this study experienced moderate and severe PP during pregnancy.
- · Pelvic pain and LBPP increased the number of weeks of sick leave and impaired the general level
- of function during pregnancy.

 Analysis of risk factors did not present a unidirectional and clear picture.

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Funding Sources and Potential Conflicts of Interest

No funding sources or conflicts of interest were reported for this study

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7.2 Paper II

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RESEARCH ARTICLE

Open Access

The association between pelvic girdle pain and sick leave during pregnancy; a retrospective study of a Norwegian population



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Background: The incidence of pelvic girdle pain (PGP) in pregnancy is wide ranged depending on definition, the utilised diagnostic means, and the design of the studies. PGP during pregnancy has negative effects on activities of daily living and causes long sick leave, which makes it a major public health issue. Our objectives were to explore the frequency of sick leave in pregnancy due to PGP, assess the relationship between different types of pain-related activities of daily living, examine physical workload, type of work in relation to sick leave, and to explore factors that make women less likely to take sick leave for PGP.

Methods: All women giving birth at the maternity ward of Stavanger University Hospital, Norway, were asked to participate and complete a questionnaire on demographic features, PGP, pain-related activities of daily living, sick leave in general and for PGP, frequency of exercising before and during pregnancy. Drawings of pelvic girdle and low back area were used for the localization of pain. PGP intensity was then rated retrospectively on a numerical rating scale. Non-parametric tests, multinomial logistic regression and sequential linear regression analysis were used in the statistical analysis.

Results: PGP is a frequent and major cause of sick leave during pregnancy among Norwegian women, which is also reflected in activities of daily living as measured with scores on all Oswestry disability index items. In the and relected in activities or long in ling as measured with expression of account of a multivariate analysis of factors related to sick leave and PGP we found that work satisfaction, problems with lifting and sleeping, and pain intensity were risk factors for sick leave. In addition, women with longer education, higher work satisfaction and fewer problems with sitting, walking and standing, were less likely to take sick leave in pregnancy, despite the same pain intensity as women being on sick leave.

Conclusions: A coping factor in pregnant women with PGP was discovered, most likely dependant on education, associated with work situation and/or work posture, which decreases sick leave. We recommend these issues to be further examined in a prospective longitudinal study since it may have important implications for sick leave frequency during pregnancy.

Keywords: Pelvic girdle pain, Sick leave, Pregnancy, Coping

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Background

Pelvic girdle pain (PGP) in pregnancy is an unclear and poorly understood condition. There is no official acknowledged nomenclature, hence an abundance of different terminologies have been used in studies to describe it [1, 2]. The European guidelines for the diagnosis and treatment of PGP describe the localization opain as**... experienced between the posterior iliac crest and the gluteal fold, particularly in the vicinity of the sacroiliac joints (SIJ). The pain may radiate in the posterior thigh and can also occur in conjunction with/or separately in the symphysis" [3].

The incidence of pain in the pelvic girdle region in pregnancy ranges from 4 to 76 % depending on the definition, the utilised diagnostic means, and the design of the studies [4]. One review study concludes that approximately 45 % of all pregnant women suffer from lumbopelvic pain in pregnancy, and approximately 20 % of these have isolated [PG 11].

of these have isolated PGP [1].
PGP accounts for 37–72 % of sick-leave in pregnancy and length of the sick leave had been found to be 12–15 weeks on average [4–8]. Other pregnancy complications may also occur in women with PGP, leading to sick leave [7]. However, women with a high degree of self-reported PGP have longer sick-leave duration than others, and these pain symptoms were in one study reported to bring about 80 % of sick leaves during pregnancy. The authors argued that this makes PGP during pregnancy a major public health issue [7].

Several risk factors for developing PGP during pregnancy have been identified, such as work load, previous PGP, and previous trauma to the pelvis [1, 3, 7–10]. PGP during pregnancy also has negative effects on activities of daily living (ADL) [7–11]. On average, these women do not experience continuous PGP, but activities of daily living such as walking, standing, sitting, lying down, or changing positions may become painful after some time. There is commonly a difficulty in walking fast or over long distances [1]. Disabling problems among women with lumbopelvic pain during pregnancy range from 21 to 81 % (median 28 %) [1]. A moderate to high Oswestry Disability Index (ODI) in women with pelvic pain during pregnancy has been identified as a risk factor for persistent PGP 3–6 months after birth [12].

The objectives of this study were to explore the frequency of sick leave in pregnancy due to PGP and to assess the relationship of different types of pain-related ADLs, physical workload, and type of work with sick leave due to PGP. Further, we wanted to explore factors that induce less sick leave due to PGP, by contrasting women who were matched for PGP, but differed by having been versus not having been on sick leave for PGP. The final objective was to explore the relative contribution of PGP to total amount of sick leave in pregnancy.

Methods

The data used in this study were collected in the period March – June 2009 at the maternity ward of Stavanger University Hospital, Norway [13]. All women giving birth at the hospital during this period were asked to participate and to fill in a study questionnaire. Inclusion criteria were a term singleton pregnancy of at least 36 weeks and good competence in the Norwegian language. The hospital has the only birth department in the southern part of the county of Rogaland, with a population of about 330,000 inhabitants.

tion of about 330,000 inhabitants.

Within 24 h after the delivery, the women received both oral and written information about the study from a midwife. To obtain inclusion of an unselected sample, all women were encouraged to give their informed consent to participate. The study was carried out in accordance with the Helsinki Declaration II and was approved by the Regional Ethics Committee of Western Norway.

To assess if the study population was a representative sample of the delivering women, we compared demographic and clinical characteristics of the study population with that of the general delivery database at the hospital and found a nearly complete match [13].

hospital and found a nearly complete match [13]. The women completed a questionnaire on demographic features, pain in pelvic girdle area, pain-related ADL, sick leave in general and due to PGP, and frequency of exercising before and during pregnancy. The questionnaire was produced and specially designed by the research group, based on previous studies and the experience of the team [13].

The women marked location of pain on drawings of the pelvic girdle and low back included in the question-naire package. The pelvic girdle and the low back were labelled, and separated according to boundaries described in the European guidelines for the diagnosis and treatment of PGP [3]. Pain intensity for PGP was then rated retrospectively on a numerical rating scale (NRS) from 0 to 100, for each month of the pregnancy, in order to collect information on appearance of symptoms and peak intensity pain during pregnancy. In this study the score 0 meant "No pain" and 100 "Unbearable pain", and average pain intensity for PGP was calculated from the values reported in all months.

Information on pain-related ADL was collected through the Oswestry Disability Index (ODI), which is one of the principal condition-specific outcome measures for defining disabling effects from spinal disorders and pelvic girdle pain [3, 14, 15]

The questionnaires also provided information on number of years of education, level of physical work load (a five level scale running from very light to very heavy), type of work (in free text but coded into mainly seated, mainly standing, mainly mobile), work satisfaction (a five level scale running from very bad to very good), and height and

weight before pregnancy and weight at delivery. Further variables included number of previous births, exercising habits before pregnancy, defined as regular exercise at least twice weekly (yes or no). Body mass index (BMI) was calculated as weight/length².

In Norway the employer is obliged to pay sick pay for the first 16 calendar days (employer's period) [16]. After that, the Norwegian Labour and Welfare Service (NAV) will take over the responsibility. The employee is required to notify the employer as soon as possible of any absence due to illness. The duty to report includes only information regarding the absence. The obligation to pay sick pay commences from the day the employer is notified about the absence, unless the employee has been prevented from notifying the employer immediately.

In the present study sick leave was assessed in two dif-

ferent ways. First, the women were asked about the total number of weeks of full-time sick leave in the pregnancy, as well as the total number of weeks with part-time sick leave and sick leave percentage. Weeks of sick leave in total were calculated by adding the full-time sick leave weeks to the part time weeks adjusted for sick leave percentage. After reporting the total amount of sick leave, the women were asked to indicate a primary cause of their sick leave. Second, in association with the section of the questionnaire concerning NRS of pain during the pregnancy, the women were also asked whether they had been on sick leave due to PGP in any month of the pregnancy and to indicate when. It was not possible to determine the number of consecutive weeks of 100 % sick leave due to any specific cause from the available information. For instance, several women only reported "pain" without any specific details as the main cause of sick-leave in pregnancy. To determine whether the women had sick leave due to PGP, we combined the available information. If the women reported any sick leave due to PGP in any month of the pregnancy, they were classified as having sick leave due PGP. Women, who explicitly stated that PGP was the primary cause of their sick leave, but who did not indicate sick leave due to PGP in any specific month of pregnancy in the NRS-section of the questionnaire, were also classified as having sick leave due to PGP. This procedure resulted in three groups: women with no re-ported sick leave, women with sick leave but without any indication of PGP being the cause, and finally, women who had sick leave and reported PGP as the cause of, at least, parts of their sick leave.

Descriptive data are presented as mean values and standard deviations for continuous variables and as frequencies for categorical variables. For comparisons between groups the non-parametric Kruskal-Wallis statistics were used, applying Bonferroni correction. Pairwise follow-ups were performed with the group who had sick leave due to PGP as reference, whenever significant omnibus group differences were found. For categorical data, chi-square statistics were computed, and 2×2 table follow-ups were used for pairwise comparisons between the group with sick-leave due to PGP vs the other groups.

Multinomial logistic regression analysis was used to investigate the independent contribution of variables hypothesized to affect whether the women had been on sick leave due to PGP. Forced entry was used with age, education, parity, and average PGP in order to adjust for these variables in the final model. As an exploratory approach, single items from ODI (except sex and pain intensity) were entered in a stepwise procedure together with workload, work satisfaction and seated work, using a likelihood ratio based criterion with p < .05 for entry and p < .10 for removal.

In order to explore factors associated with the total amount of sick leave in pregnancy, a sequential linear regression analysis was conducted, using total number of calculated weeks of sick leave (weeks of 100 % sick-leave + weeks of part-time sick leave multiplied by sick leave percentage) for any reason as dependent variable. In the first block, the grand average of monthly reported PGP was entered, in order to analyse the unadjusted effect of PGP on weeks of sick leave. In block 2, all relevant ODI items were entered using a stepwise procedure (p < .05 to enter, p < .01 to exclude a variable). In block 3, years of education, pre-pregnant BMI, workload, age, standing work and mobile work were entered, using the same type of stepwise procedure as in block 2. Finally, in block 4, work satisfaction and depression in pregnancy were entered, also with a stepwise procedure. Thus, only block 1 included a forced entry variable, average PGP, as we wanted to explore unadjusted and adjusted effects of PGP on weeks of sick leave.

In order to investigate factors that may decrease the effect of PGP on sick leave, we identified all women with PGP who did not have any sick leave in pregnancy. A macro written in Microsoft Excel (Visual Basic) then chose a random woman having been on sick leave, and who matched the average PGP score of a woman without sick leave. If a perfect match was not found, a difference of +/-1 point on the PGP score was accepted. If still no match was found, the subject was discarded. Hence, this procedure chose two equivalent groups with regard to average PGP, but with and without sick leave. We then compared these groups on the same variables as for the sick leave due to PGP, vs no sick leave, vs sick leave due to the research groups.

leave due to other reasons groups.

Effect sizes were reported as standardized mean differences (Cohen's D), using Bonferroni correction, which can be interpreted as small (around 0.3) medium (around 0.5) and large (0.8 to infinity) [17].

All analyses were performed using SPSS 21 (IBM, New York, NY), and results were considered significant at p < .05.

Results

Study population

In all, 1204 women who gave birth at Stavanger University Hospital during the inclusion period were invited to participate in the study. After exclusions, there were 994 women eligible for the study. In addition, 336 women did not return a questionnaire and 89 did not complete provided questionnaire. One present did not the provided programment of the provided programment of the provided programment. having a job or profession, nor any workload, and was excluded from the analyses. The final study population thus consisted of 568 women. Of these, 165 (29 %) reported that they had experienced isolated PGP during the pregnancy.

the pregnancy.

The sample's demographic data and descriptive statistics for the variables used in the multivariate analyses are shown in Table 1. Several significant differences are shown between subjects who reported sick leave due to PGP vs. those who did not [Table 1]. In Table 2 we compare the group with sick leave due to PGP with the no sick leave group and the group with sick leave due to other causes on single items from the ODI. We report effect sizes to enable a direct comparison using a standardized scale [Table 2].

All ODI-items were significantly higher in the group who had been on sick leave for PGP than in both the other groups. The effect sizes were all moderate to large (Cohen's d > 0.6).

Factors associated with sick leave due to PGP

Individual risk factors with odds-ratios and confidence limits resulting from the multinomial regression analysis are shown in Table 3.

are snown in Table 3. All results in Table 3 refer to the group with sick leave due to PGP as reference category. The estimated pseudo R^2 was quite high (Nagelkerke $R^2 = 40$) and the total correct classification percentage was 62 %. We see that work satisfaction, as well as lower scores for ODI-lifting, sleep and average pain intensity, significantly predicted efficiency as the product of the product o affiliation to the no sick leave group. The group with sick leave due to other reasons had lower score on average pain intensity and ODI-lifting [Table 3].

Coping with the effects of pelvic girdle pain

The matching procedure resulted in two groups with 50 subjects in each group. The group with sick leave due to PGP and the group with no sick leave (coping group) had similar PGP intensities of approximately 18. Univariate Mann–Whitney U test revealed that the coping group on average had longer education (15.8 vs 14.8 years), p = .022 and higher work-satisfaction (4.66 vs 4.32), p = .014. Finally, the scores on several ODI-

items were lower in the coping group [Table 4].

These results differ from the ODI-results in Table 2, as the effect sizes between the groups are very different for the different items. If a strict Bonferroni-correction is applied, only the ODI score for sitting is higher in the group with sick leave due to PGP (p<.003). However, also walking and standing differ if uncorrected p-values are applied.

Table 1 Descriptive statistics

	Sick leave for PGP	No sick leave	Sick leave for other reason	p****	Total
N (96)	193 (34 %)	139 (24 %)	236 (42 %)		568 (100 %)
Age	29.7 (4.3)	30.5 (4.8)	29.8 (5.0)	=.254	30.0 (4.7)
Education	14.5 (2.4)	15.7 (2.4)*	15.4 (2.6)*	<.001	15.1 (2.6)
BMI before pregnancy	24.8 (4.6)	23.1 (3.6)*	23.4 (4.2)*	<.001	23.8 (4.3)
Total sick leave weeks	10.8 (9.4)	0.0 (0.0)*	8.4 (8.9)*	<.001	7.2 (9.0)
Workload	3.0 (1.1)	2.4 (1.1)*	2.6 (1.2)*	<.001	2.6 (1.1)
Work satisfaction	4.4 (0.8)	4.6 (0.7)*	4.3 (0.9)	<.001	4.4 (0.8)
Average PGP	26.8 (15.1)	6.7 (10.4)*	6.1 (10.0)*	<.001	13.3 (15.5)
Average LBP	13.2 (16.9)	4.7 (9.1)*	6.6 (11.4)*	<.001	8.4 (13.4)
Pain-related ADL (ODI)	1.9 (0.8)	0.9 (0.9)*	1.0 (0.8)*	<.001	1.4 (0.9)
Depressed	1.4 (0.5)	1.3 (0.5)	1.4 (0.6)	=.055	1.4 (0.5)
Mean no. of previous births	1.00 (0.06)	0.94 (0.09)	0.79 (0.05)***	<.05	0.90 (1.00)
Regular exercise before pregnancy	63 (33 %)	67 (49 %)**	94 (40 %)	=.013	224 (39 %)
Seated work	51 (27 %)	68 (49 %)*	81 (34 %)	<.001	200 (35 %)

PGP Pelvic girdle pain, LBP Low back pain. Pairwise comparison with sick-leave for PGP: *p < .001, **p < .01, ***p < .05 ****Kruskal-Wallis omnibus test. Bonferroni-corrected alpha = 0.0038

Table 2 Oswestry disability index items

	Sick leave for PGP	No sick leave		Sick leave for other	er reason	Total
	N = 190	N = 96		N = 154		N = 440
ODI item	Mean (SD)	Mean (SD)	E.S.ª	Mean (SD)	E.S. ^a	Mean (SD)
Pain intensity	2.76 (0.86)	1.67 (1.13)	1.142	1.81 (0.99)	1.024	2.19 (1.09)
Personal care	1.23 (1.40)	0.59 (0.97)	0.591	0.53 (0.98)	0.655	0.85 (1.10)
Lifting	2.18 (1.19)	0.95 (1.12)	1.056	1.30 (1.16)	0.751	1.60 (1.27)
Walking	1.63 (0.99)	0.65 (0.94)	1.007	0.85 (1.05)	0.762	1.15 (1.09)
Sitting	1.68 (0.96)	0.80 (0.98)	0.905	1.07 (1.11)	0.589	1.28 (1.08)
Standing	2.44 (1.24)	1.17 (1.28)	1.018	1.48 (1.27)	0.769	1.83 (1.37)
Sleeping	1.67 (1.02)	0.82 (0.88)	0.872	1.04 (0.88)	0.657	1.26 (1.01)
Sex	1.75 (1.50)	0.76 (1.19)	0.707	0.76 (1.30)	0.701	1.18 (1.45)
Social function	1.89 (1.26)	0.83 (1.17)	0.862	0.79 (1.14)	0.911	1.27 (1.31)
Travelling	1.63 (1.26)	0.61 (1.00)	0.860	0.82 (1.11)	0.680	1.12 (1.24)

This study shows that PGP is frequent and a major cause of sick leave during pregnancy among Norwegian women, which is also reflected in problems with ADL as measured with scores on all ODI items. In the multivariate analysis of factors related to sick leave and PGP we found that work satisfaction, problems with lifting and sleeping, and pain intensity predicted sick leave. In addition, we found that women with longer education,

Table 3 Multinomial regression with sick leave due to PGP as

	P	Odds ratio	C.L. Low	C.L. High
No sick-leave				
Age	.157	1.056	.979	1.138
Education	.074	1.113	.990	1.252
Pelvic pain	.001	.955	.930	.981
No. of previous births	.667	.915	.612	1.369
ODI: Lifting	.011	.622	.432	.895
ODI: Sleep	.008	.521	.321	.846
ODI: Social life	.294	1.206	.850	1.713
Work Satisfaction	.049	1.607	1.001	2.580
Sick-leave due to other re	ason			
Age	.129	1.051	.985	1.122
Education	.262	1.054	.961	1.157
Pelvic pain	.000	.951	.932	.971
No. of previous births	.128	.760	.533	1.083
ODI: Lifting	.020	.708	.530	.946
ODI: Sleep	.622	.916	.646	1.299
ODI: Social life	.105	.785	.586	1.052
Work Satisfaction	.814	.960	.681	1.352

C.L.: 95 % confidence limits

higher work satisfaction and less problems with sitting, walking and standing, were less likely to take sick leave during pregnancy, despite having the same pain intensity as women being on sick leave. These findings may have implications for planning of measures that could reduce

mightautists for painting of ineastics untar could reduce sick leave among pregnant women.

Most studies describe percentages of pregnant populations on sick leave, and mainly use lumbopelvic pain as a general term, describing the location of pain. Few specify length of sick leave or differentiate PGP from low back pain for a more specific location of pain. Also, the back pain for a more specific location of pain. Also, the majority of studies are prospective or cross-sectional. Our study was retrospective and shows a sick leave percentage and span similar to other studies with the same methodology. In our study, 34 % of the women had been on a median of eleven weeks sick leave for PGP during the pregnancy. In a Swedish retrospective study, sick

Table 4 ODI in women who had PGP, with and without sick

icave for f or				
ODI item	No sick-leave for PGP	Sick-leave for PGP	E.S.	p
Pain intensity	2.30 (0.84)	2.33 (0.88)	0.039	=.954
Personal care	0.90 (1.11)	0.94 (1.14)	0.033	=.889
Lifting	1.40 (1.21)	1.90 (1.29)	0.395	=,044
Walking	0.96 (1.01)	1.44 (0.97)	0.483	=,011
Sitting	0.96 (1.00)	1.46 (0.87)	0.528	=.003
Standing	1,50 (1.33)	2.00 (1.22)	0.392	=.031
Sleeping	1.12 (0.85)	1.48 (1.03)	0.379	=.113
Sex	1.18 (1.41)	1.23 (1.49)	0.037	=.969
Social function	1.26 (1.27)	1.33 (1.31)	0.056	=,677
Travelling	0.92 (1.12)	1.25 (1.23)	0.280	=.180

ES Effect size (Cohen's d); Bonferroni-corrected alpha = 0.005

E.S.: Effect size (Cohen's d, >0.8 is considered large) ^aAll differences of means statistically significant assuming a Bonferroni corrected alpha of p < .005

leave for pain in the pelvic girdle area was reported among 48 % of the pregnant women, with a mean span of sick leave of 12 weeks [11]. In another retrospective study, 41 % had been on sick leave for PGP, but duration was not described [18]. Taken together, our and previous studies underline the importance of PGP as a major cause of sick leave during pregnancy.

A previous Norwegian study showed that sick leave increases for each trimester in pregnancy. It revealed that fatigue and sleep problems seem to be the main risk factor for sick leave, followed by nausea, vomiting, exercising less than weekly, chronic PGP before or during pregnancy, conflicts in the workplace, and lower education [18]. In contrast, Mogren in a retrospective study found lumbopelvic pain to be the main cause of sick leave during pregnancy [11]. This finding is supported by our study, and Robinson and co-workers who showed that almost a third of all delivering women were sick listed due to PGP during pregnancy [18]. A result confirming that PGP accounts for a great part of sick leave during pregnancy.

during pregnancy.

Our study shows that ADL were significantly more difficult to carry out for pregnant women on sick leave for PGP than for women on sick leave for other causes. We found independent significant risk factors to be lower education, heavy workload and low work satisfaction. This association has previously been shown in three other studies in which pregnant women with demanding working conditions presented increased incidence of PGP in pregnancy, while those with opportunity to adjust their work pace reported a better health status than women without this possibility [19–21]. Risk indicators for long-term sick leave during pregnancy have been shown to be less qualified work and heavy work load [22]. Lower education level has also been found to associate with higher pain intensity during pregnancy [23].

The ODI score on sitting scored the highest significant effect size in the group with sick-leave due to PGP (p < .003). It is known that prolonged sitting may alter the passive stiffness of the lumbar spine. Erector spinae muscles fatigue induces a shift in load-sharing towards passive stabilizing structures. Loss of muscle contribution together with or without laxity in the viscoelastic tissues may have a substantial impact on post fatigue stability [24].

In our study, a matching procedure revealed that the group with sick leave due to PGP had a lower PGP intensity score than the group with no sick leave. The reason for this unexpected find may be that pain is a complex construct that contributes to profound physical and psychological dysfunction, and the experience of it is modulated by physical and psychological factors [25]. Following the bio-psychosocial model, emotional

distress may predispose people to experience pain or may be a moderator that amplifies or inhibits the severity of pain [26]. Traumatic experiences related to pregnancy seem to be associated with lumbopelvic pain and physical ability up to 6 months after delivery [27].

Certain individuals can stand the pain, have less catastrophizing tendencies, show more positive social responses to pain, and more organized health care and medication patterns. These coping skills are displayed as effective functioning despite exposure to stressful circumstances and/or internal distress [28]. Most important psychological contributors to individual well-being and coping stress responses are positive emotions, which appear to buffer individual reactivity to pain [25].

In a retrospective study, with a similar design to ours, the authors found that pregnancy seems to be a period during which a sick-listed is prone to be influenced by attitudes and coping strategies in order to achieve needed rest prior to delivery [29]. They concluded that certain social conditions and attitudes are likely to explain why pregnant women are on sick leave [29]. Chang and co-workers suggest that education and interventions targeting passive coping and stimulating resilience may be useful to prevent PGP during pregnancy turning chronic [23]. In Korea, a back-pain-reducing program was effective in reducing the intensity of back pain experienced by pregnant women [30].

rienced by pregnant women [30].

Our findings show that women may benefit from a pre-pregnancy and pregnancy strategy program decreasing the risk for pregnant women to end up on sick leave for PGP. According to our results the strategy should contain information about the value of pre-pregnancy regular physical exercise in order to withstand physical workload during pregnancy. In order to boost coping towards PGP and sick leave, information about physical, physiological and psychological changes and challenges in pregnancy, should be included. Employers should through an incentive be encouraged to ergonomically adapt the pregnant employees' workplace in order to decrease work load and thus maintain, or even improve, work satisfaction.

There are limitations in the present study. We introduced several tactics to obtain a high awareness of the importance of the study, both among the women and assisting midwives, with the objective to enrol all women who fulfilled the inclusion criteria during a given time period. Nevertheless, our response rate was rather low, in spite of aforementioned initiatives, but we did find that age and parity of the studied population was identical to the general population of women that gives birth at the hospital. We therefore believe the results from this study to be representative for pregnant women in the study area, and most probably also in the rest of Norway.

Ideally, to use the term PGP a physical examination is required, but it could not be done in this study. Another limitation is that all our data are based on questionnaires with retrospective data collection. We are aware that recalling pain and disability experienced long time ago is considered an unreliable way to collect information. Regarding the occurrence and duration of sickness absence during pregnancy has the agreement between self-report and a public registry been investigated [31]. Mainly because of low precision the agreement on the duration of sickness absence was poor, but the agreement regarding the occurrence of sickness absence was good.

Conclusions

PGP is a frequent and major cause of sick leave in pregnancy. We have retrospectively identified how it affects pregnant women's ADL, and found unexpected differences in pain appreciation between women on sick leave and not on sick leave during pregnancy. A coping factor seem to be present, most likely dependent on education, associated with work situation and/or work posture, which decreases sick leave. We recommend that these issues should be further examined in a prospective longitudinal study since it may have important implications for sick leave frequency during pregnancy.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions
SM, IK, JPL, KA and IID planned and designed the study, SM, IK and KA clistributed and collected the questionnaires. KB performed the statistic analyses and together with SM, JPL and I/O drafted the manuscript. All authors read and approved the final manuscript.

Authors' information

Availability of data and materials

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7.3 Paper III

Open access Research

BMJ Open Can a bothersome course of pelvic pain from mid-pregnancy to birth be predicted? A Norwegian prospective longitudinal SMS-Track study

Stefan Malmqvist, ^{1,2} Inger Kjaermann, ^{1,3} Knut Andersen, ^{1,3} Anne Marie Gausel, ^{1,3} Inger Økland, ⁴ Jan Petter Larsen, ^{1,5} Kolbjorn S Bronnick⁶

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ABSTRACT

Objective To explore if pregnant women with pelvic girdle pain (PGP), subgrouped following the results from two clinical tests with high validity and reliability, differ in demographic characteristics and weekly amount of days with bothersome symptoms through the second half of

Design A prospective longitudinal cohort study Participants Pregnant women with pelvic and lumbopelvic pain due for their second-trimester routine ultrasound examination.

Setting Obstetric outpatient clinic at Stavanger University

Hospital, Norway,

Methods Women reporting pelvic and lumbopelvic pain completed a questionnaire on demographic and clinical features. They were clinically examined following a test procedure recommended in the European guidelines for the diagnosis and treatment of PGP. Women without pain symptoms completed a questionnaire on demographic data. All women were followed weekly through an SMS-Track survey until delivery.

Primary and secondary outcome measures The outcome measures were the results from clinical diagnostic tests for PGP and the number of days per week with bothersome pelvic pain.

Results 503 women participated. 42% (212/503) reported pain in the lumbopelvic region and 39% (196/503) fulfilled the criteria for a probable PGP diagnosis. 27% (137/503) reported both the posterior pelvic pain provocation (P4) and the active straight leg raise (ASLR) tests positive at baseline in week 18, revealing 7.55 (95% Cl 5.54 to 10.29) times higher mean number of days with bothersome pelvic pain compared with women with both tests negative. They presented the highest scores for workload, depressed mood, pain level, body mass index. Oswestry Disability Index and the number of previous pregnancies. Exercising regularly before and during pregnancy was more common in women with negative tests

Conclusion If both P4 and ASLR tests were positive mid-pregnancy, a persistent bothersome pelvic pain of more than 5 days per week throughout the remainder of pregnancy could be predicted. Increased individual control over work situation and an active lifestyle, including regular exercise before and during pregnancy, may serve as a PGP prophylactic.

Strengths and limitations of this study

- We used a prospective design with SMS-Track system in data collection, providing instant data on participants' situation, with automatically recorded responses in a database, which minimises further
- data handling and risk of error.

 We applied clinical diagnostic tests with high validity and reliability, recommended in the international guidelines for the diagnosis of pelvic musculoskele-
- tal affliction in pregnancy.

 There were frequent problems in reaching the participants through some of the phone providers, which led to SMS-Track data missing at random, but
- which led to SMS-Track data missing at Tranoom, but a generalised estimating equation analytic approach compensates for missing data in these instances. The retrospectively collected information on pelvic pain in previous pregnancies and pelvic pain before pregnancy may produce bias.

INTRODUCTION

Pelvic girdle pain (PGP) during pregnancy affects approximately half of all pregnant women, and for 25%–30% the condition becomes severe.^{1 2} The aetiology of PGP is still unknown, and the underlying mechanisms have not been fully investigated.1 2 Researchers have explored the physical, psychological and socioeconomic implications of PGP during pregnancy.³ Pain-related restrictions on physical activity have been described, both during pregnancy and after childbirth, and the psychological impact on perceived health, sexual life and quality of life has been explored, as well as the prevalence of sick leave due to PGP.³⁻⁶

PGP is classified into specific (caused by trauma) or non-specific (multifactorial).³ Several clinical tests are needed to diagnose the latter, including pain provocation and functional ability tests. However, there is still no 'gold standard' for diagnosing PGP. The European guidelines present evidence-based



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recommendations for the diagnosis and treatment of PGP, but inconsistencies on the definition as well as treatment still prevail. $^{3.7}$

Classification of PGP can, according to guidelines, only be reached after lumbar causes have been excluded through a clinical examination. All tests recommended in the European guidelines have a very high specificity, but generally a low sensitivity. Hence, it is recommended to perform all the tests, as one negative test is not sufficient to rule out PGP. The posterior pelvic pain provocation test (P4), for diagnosing sacroiliac joint dysfunction, and the active straight leg raise test (ASLR), for detecting failing force closure, have shown high validity and reliability. 8-10 In a Swedish study, substantial agreement between examiners using ASLR and P4 tests was found in discriminating non-specific lumbopelvic pain into lumbar pain and PGP in pregnant women. Il Together with a description of pain location, these tests are considered relevant when evaluating affliction in pregnant women likely to have PGP. 12

So far, the longitudinal course of PGP in pregnancy is incompletely examined. In prospective studies data are usually collected at baseline and at one or a few follow-ups. Measuring only at a few points in time may indicate stability in the examined condition, and a fluctuating course may be missed. A difference could reflect only a temporary fluctuation in an otherwise stable condition. Accordingly, a more frequent data collection is warranted to accurately describe the clinical course. Mobile phones and text messages have previously been found feasible when collecting frequent longitudinal data in clinical settings. ^{13–13} Phones are usually at hand in daily life; hence, this method yields a high response rate for weekly measures.

The objective of this study was to explore if pregnant women with probable PGP, subgrouped following the results from two valid and reliable clinical tests recommended in the European guidelines, differ in demographic and clinical characteristics at mid-pregnancy and in weekly amount of days with bothersome symptoms through the second half of pregnancy. The hypothesis was that sacroiliac dysfunction and failing force closure diagnosed at mid-pregnancy may predict a course of bothersome symptoms through the second half of pregnancy.

METHODS

This is a prospective longitudinal cohort study of pregnant women who had their second-trimester routine ultrasound examination in pregnancy week 18 at an obstetric outpatient clinic at Stavanger University Hospital, Norway, from mid-March to mid-June 2010. At the hospital, all the women were asked by a midwife about their experience of pain in the lumbopelvic region. The inclusion criteria were current lumbopelvic pain or isolated pelvic pain, singleton pregnancy and good proficiency in the Norwegian language. Women who met the criteria were informed about the study, handed a letter of consent to fill in if they agreed to participate, and an

envelope with questionnaires on demographic and clinical data to complete at home. An appointment with a chiropractor for a physical examination was arranged, and the women were asked to bring the completed questionnaires with them to the consultation. Women without pain symptoms were informed about the study, handed a letter of consent to fill in if they agreed to participate, and a questionnaire on demographic data to complete and leave at the reception on departure. All consenting women were followed from week 18 with weekly, automated text messages (SMS-Track).

Two chiropractors (SM and IK) performed a physical examination of the pelvic region, including diagnostic tests recommended in the European guidelines for the diagnosis and treatment of PGP, and a neurological examination of the lower extremities.

Sequence of stability and pain provocation tests for PGP Active straight leg raise

The test is performed with the patient in a supine position with a straight leg and the feet 20 cm apart. The test is performed after the instruction 'try to raise your legs, one after the other, above the couch for 20 cm without bending the knee'. The patient is asked to score impairment on a 6-point scale: not difficult at all=0; minimally difficult=1; somewhat difficult; difficult=2; fairly difficult=3; very difficult=4; unable to do=5. The scores on both sides are added, so that the total score range from 0x 10.09.

Gaenslen's test

The patient, lying supine, flexes the knee and hip of the same side, the thigh being crowded against the abdomen with the aid of both the patient's hands clasped about the flexed knee. The patient is then brought well to the side of the table, and the opposite thigh is slowly hyperextended by the examiner with gradually increasing force by pressure of the examiner's hand on top of the knee. With the opposite hand, the examiner assists the patient in fixing the lumbar spine and pelvis by applying pressure over the patient's clasped hands. The test is positive if the patient experiences pain, either local or referred on the provoked side. ¹⁶

Long dorsal sacroiliac ligament test

The subject lies on her side with slight flexion in both the hip and knee joints. If the palpation causes pain that persists more than 5s after removal of the examiner's hand, it is recorded as pain. If the pain disappears within 5s, it is recorded as tenderness. ¹⁷

Modified Trendelenburg's test

The patient stands on one leg, and flexes the other at 90° in the hip and knee. If pain is experienced in the symphysis, the test is positive. ¹⁷

Patrick's FABER test

The subject lies supine. One leg is flexed, abducted and externally rotated (FABER, abbreviation of flexion

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abduction and external rotation) so that the heel rests on the opposite knee. If pain is felt in the sacroiliac joints or in the symphysis, the test is considered positive. ¹⁷

Posterior pelvic pain provocation test

The test is performed with the woman supine and the hip flexed to an angle of 90° on the side to be examined: a light manual pressure is applied to the patient's flexed knee along the longitudinal axis of the femur while the pelvis is stabilised by the examiner's other hand resting on the patient's contralateral superior anterior iliac spine. The test is positive when the patient feels a familiar well-localised pain deep in the gluteal area on the provoked side. A similar test is described as posterior shear or 'thigh trust'. ^{17 18}

Symphysis palpation test

The subject lies supine. The entire front side of the pubic symphysis is palpated gently. If the palpation causes pain that persists more than 5s after removal of the examiner's hand, it is recorded as pain. If the pain disappears within 5s, it is recorded as tenderness. ¹⁷

A demographic questionnaire used in an earlier study on pelvic pain in pregnancy was filled in at baseline. ¹⁹ The women marked the pain location on drawings with the pelvis and the low back demarcated. Pain intensity was rated on a Numerical Rating Scale (NRS) from 0 to 100, where 0 meant 'No pain' and 100 'Unbearable pain'. Information on pain-related activities of daily living (ADL) was collected through the Oswestry Disability Index (ODI), which at the time of the data collection was one of the principal outcome measures for defining disabling effects from spinal disorders and PGP.^{7 20} It is a patient-completed questionnaire which gives a subjective percentage score of the level of function (disability) in 10 ADLs in patients with low back pain. Every activity contains six statements on how it is performed. The statements are scored from 0 to 5, with the first statement scoring 0 through to the last at 5. The scores for all questions answered are summed, then multiplied by 2 to obtain the index (range 0–100). Zero is equated with no disability and 100 is the maximum disability possible.

Physical workload was measured through five answer categories ranging from 'sedentary' to ' heavy', following a scale used in Stockholm Public Health questionnaire. ²¹ The question on job satisfaction was a bipolar 5-point Likert scale with increments in two opposite directions ('Very bad' and 'Very good') and a neutral point in the middle ²²

Every Sunday the women were asked through a short message service (SMS) how many days the previous week they had experienced bothersome pelvic pain: 'How many days during the previous week has your pelvic pain been bothersome, (ie, affected your daily activities or routines)?' If there was no reply, the question was repeated 24 hours later. The question should be answered with one single number between 0 and 7. The response was automatically entered into a database, which collected the

continuous information from each participant over the duration of the study.

Demographic descriptive data are presented as median values with IQRs for continuous variables, and as frequencies for categorical variables. For univariate comparisons between symptomatic and asymptomatic subgroups, the non-parametric Kruskal-Wallis statistics were used. Categorical predictors in our model were four groups following the outcome from the ASLR and P4 tests (1: P4 positive, 2: ASLR positive, 3: both P4 and ASLR positive, 4: ASLR and P4 negative), time (pregnancy week), and the interaction term between time and test group for investigating whether the trajectory of SMS-reported number of bothersome days differed between the test groups. Other predictors in the model were age, number of previous deliveries and body mass index (BMI) before pregnancy. The longitudinal trajectory of the SMS-Track response

The longitudinal trajectory of the SMS-Track response was modelled using a generalised estimating equations (CEE) approach, extending the generalised linear model to correlated longitudinal data and clustered data within subjects. The within-subject dependencies resulting from repeated measurement were modelled assuming an autoregressive relationship in the working correlation matrix. As the outcome variable was count data (weekly number of bothersome days with pain), the Poisson distribution was assumed with a log-link function.

bution was assumed with a log-link function.

Data were analysed using SPSS V.22.0 software. The Strengthening the Reporting of Observational Studies in Epidemiology reporting guideline was used during the writing of this article.

Patient and public involvement

Patients and the public were not involved in developing the research questions, outcome measures, as well as in the design and conduct of the study, or in the recruitment of patients.

RESULTS

Overall, 506 women agreed to participate in this study. Three were excluded due to incomplete data. On ultrasound examination in pregnancy week 18 did 42% (212/503) of the women report pain in the lumbopelvic region. A clinical examination revealed that 39% (196/503) of the women fulfilled the criteria for a probable PCIP diagnosis, and 27% (137/503) showed positive response to ASLR and P4 tests. A further 12 women reported pelvic pain but did not respond to recommended clinical tests, and were therefore placed in the 'ASLR and P4 tests negative group'.

There were significant differences in some demographic and clinical features at baseline between the women with and without pelvic pain and with different test outcomes (tables 1–2).

Women with positive P4 and ASLR tests experienced heavier workload. They also presented higher BMI at week 18, exercised less both before and during pregnancy, and slightly more than one-third reported feeling depressed

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Table 1 De	mographic f	eatures of	Demographic features of women with and without positive ASLR and P4 tests at 18 weeks of pregnancy	and with	out positi	ve ASLR and	d P4 tests	s at 18 we	eks of pregr	ancy					
				ASLR and	ASLR and P4 tests positive	positive									
Variables	No pelvic pain a	P values a-b	P4 and ASLR tests negative b	P values	P values b-c	P4 only c	P values a-d	P values b−d	P values ASLR only b-d d	P values a-e	P values b-e	Both ASLR and P4 e	P values a-f	P values b-f	All ASLR and P4 f
(u) %	58 (291)		15 (75)			36 (50)			18 (24)			46 (63)			27 (137)
Age, median (IQR)	29.0 (26.0/34.0)	0.28	30.0 (28.0/34.0)	0.44	09.0	30.5 (26.7/33.2)	0.92	0.21	30.0 (26.0/33.0)	0.30	0.10	28.0 (24.0/34.0)	0.44	0.11	29.0 (25.7/33.0)
Education years, median (IQR)	15.0 (13.0/17.0)	0.64	15.0 (13.0/17.0)	0.55	0.89	15.0 (12.0/17.0)	0.55	0.49	15.0 (12.0/17.0)	0.41	0.33	15.0 (12.0/16.2)	0.46	0.36	15.0 (12.0/17.0)
Physical workload, median (IQR)	3.0 (1.0/4.0)	0.01	2.0 (1.0/4.0)	00:00	0.24	3.0 (2.0/4.0)	0.96	0.00	3.0 (2.0/3.0)	0.00	0.00	(3.0/4.0)	0.01	0.01	3.0 (2.0/4.0)
Work satisfaction, median (IQR)	5.0 (4.0/5.0)	0.32	4.0 (4.0/5.0)	0.88	0.66	4.5 (4.0/5.0)	0.96	0.80	5.0 (4.0/5.0)	0.91	0.74	4.0 (4.0/5.0)	0.52	0.38	4.5 (4.0/5.0)
BMI, median (IQR)	23.9 (22.3/27.0)	0.03	24.9 (22.7/27.9)	0.03	0.33	25.6 (23.7/29.9)	0.02	0.44	26.0 (24.0/30.4)	0.08	0.52	25.0 (22.0/28.0)	0.01	0.47	25.6 (22.8/28.5)
Number of births, median (IQR)	(0.0/1.0)	0.15	1.0 (0.0/2.0)	0.10	0.65	1.0 (0.0/1.5)	0.05	0.39	1.0 (0.0/2.0)	0.15	0.43	1.0 (0.0/1.0)	0.07	0.98	1.0 (0.0/1.0)
Exercising before pregnancy, % (n)	37 (108)	0.19	36 (27)	0.21	0.86	34 (17)	0.69	0.60	37.5 (9)	0.11	0.42	(17)	0.07	0.45	31 (43)
Exercising during pregnancy, % (n)	(60)	0.01	(14)	0.03	0.16	8 (4)	0.47	0.37	94 (4)	0.07	0.22	10	00:00	0.12	11 (15)

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lable 2 Offical	earnes		III and with	ASI R a	nd P4 tes	ASLR and P4 tests positive	200 000	Wood	Clinical features of women with and without positive ASLR and P4 tests at 18 weeks of pregnancy ASLR and P4 tests positive.						
Jarishlac	No pelvic	No pelvic P values	P4 and P ASLR tests values	P values	P values	P4	P values		P values ASLR only	P values	P values P values	Both ASLR and P4	P values	P values	All ASLR and P4
(u) %	58 (291)		15 (75)			36 (50)			18 (24)			46 (63)			27 (137)
Sick leave, % (n)	25 (74)	0.81	30 (23)	66.0	0.82	38 (19)	0.99	69.0	56 (14)	0.79	0.51	(30)	0.84	0.59	46 (63)
Feeling depressed, % (n)	20 (60)	0.04	21 (16)	0.03	0.36	30 (15)	0.08	0.02	38	0.01	0.01	43 (27)	0.01	0.01	37 (51)
PP previous pregnancies, % (n)	13 (37)	0.00	45 (34)	0.00	0.62	42 (21)	0.00	0.51	54 (13)	0.00	0.53	38 (24)	0.00	0.50	42 (58)
PP before pregnancy, % (n)	4 (12)	0.00	24 (18)	0.00	0.93	24 (12)	0.00	0.99	25 (6)	0.00	0.97	21 (13)	0.00	0.87	23 (31)
ODI week 18, median (IQR)	NA	0.02	16.0 (9.0/26.0)	1	0.51	16.0 (12.0/26.0)	ŋ.	0.00	22.0 (16.0/32.0)	1	0.00	28.0 (18.0/34.0)	1	0.00	24.0 (14.0/34.0)
PP week 18, median (IQR)	T.	NA A	20.0 (0.0/40.0)	10	0.00	35.0 (20.0/62.5)	T.	0.00	40.0 (27.7/57.5)	1	0.00	50.0 (25.0/75.0)	Í.	0.00	50.0 (20.0/66.2)
ASLR score, median (IQR)	î.	r	1	1	î.	NA	1.	0.00	2.0 (2.0/4.0)	1	0.00	3.0 (2.0/5.0)	T.	0.00	2.0 (0.0/4.0)
LDL test positive, % (n)	ī	1	33.3 (25)	1	0.01	76 (38)	1	0.04	75 (18)	1	0.01	(53)	1	0.00	(109)
Gaenslen's test positive, % (n)	1	ī	5.3 (4)	1	0.01	28 (14)	1	0.00	21 (5)	1	0.00	52 (33)	1	0.00	38 (52)
Symphysis test positive, % (n)	1	1	29.3 (22)	1	0.54	54 (27)	1	0.41	63 (15)	1	0.31	(38)	1	0.16	58 (80)
Modified Trendelenburg's test positive, % (n)	1	1	6.7	1	0.83	12 (6)	1	0.40	7+ (4)	1	0.23	21 (13)	r	0.29	17 (23)
FABER test positive, % (n)	1	1	4.0 (3)	1	0.01	26(13)	T.	0.00	33(8)	T,	0.00	41(26)	1	0.00	34(47)
SMS-Track, median (IQR)	0.0 (0.0/1.0)	0.00	3.0 (2.0/6.0)	0.00	0.01	5.0 (3.0/7.0)	0.00	0.00	4.5 (2.0/6.0)	0.00	0.00	6.0 (4.0/7.0)	0.00	0.00	(3.0/7.0)

IOR (25/75). Pvalue from non-parametric Kruskal-Wallis statistics. Subgroups marked with lower case letters from a to f for identification of p value from non-parametric test. ASI, active statistic lettere, flexion adduction and external rotation; LDL, long dorsal ligament test; NA, not applicable; ODl, Oswestry Disability Index; PA, posterior petvic pain prorocation test-Ppetvic pain.

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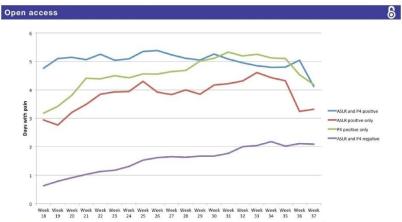


Figure 1 Estimated mean number of bothersome days in the latter half of pregnancy. If both P4 and ASLR tests were positive at week 18 (blue), the highest mean weekly number of days with bothersome pelvic pain throughout the second half could be expected. If both tests were negative, the mean number of bothersome days would never reach the amount reported by women with positive tests (purelp). Women with a positive P4 test and a negative ASLR test (green) presented approximately 3 days of bothersome pelvic pain in week 18, with the mean number of days increasing rapidly until week 29, equalling the group with both tests positive (blue). Interestingly, women with a positive ASLR and a negative P4 test (red) also showed 3 days of bothersome pelvic pain at week 18, but never reached the mean number of bothersome days reported by women with P4 and both tests positive. ASLR, active straight leg raise test; P4, posterior pelvic pain provocation test.

during the pregnancy. Physical disability (ODI) and pain level (NRS) at week 18 were considerably higher in women with positive tests than in women reporting pain but having negative P4 and ASLR tests (table 2).

Women with a positive ASLR, but negative P4 test, had

Women with a positive ASLR, but negative P4 test, had the highest number of previous pregnancies. Almost half of the women with both P4 and ASLR tests positive had been on sick leave during their pregnancy. Apart from the P4 and ASLR, the long dorsal ligament test showed the highest positive response rate, followed by the symphysis provocation test (table 2).

provocation test (table 2).

The SMS-Track response rate was 75% (2148 responses to 2877 sent messages). Due to a declining SMS-Track response at the end of the pregnancy, we stopped our SMS-Track analysis at week 38. A GEE analysis revealed that all entered variables, except 'age', were significant predictors for the number of days with bothersome pelvic pain, and there was a significant interaction between diagnostic group and time, implying that the time course of days with bothersome pelvic pain was different for the different test grouper.

different test groups.

The estimated weekly mean number of days with bothersome pelvic pain for the different test groups is presented in figure 1. Women with both P4 and ASLR tests positive experienced from week 18 a high weekly mean number of days (=5) with bothersome pelvic pain throughout the pregnancy. Women with both tests negative showed a steadily rising number of bothersome days

throughout the pregnancy, from 0.5 day in week 18, to 2 days in week 37. The group with a P4 positive and an ASLR negative test had approximately 3 days of bothersome pelvic pain in week 18, which was considerably lower than the group with both tests positive, but showed rapidly increasing number of days with pain. In week 29, the number of days with bothersome pelvic pain equalled that of the group with both tests positive, and thereafter matched this group. Women with a positive ASLR and a negative P4 test also showed 3 days of bothersome pelvic pain in week 18, but never reached the mean number of bothersome days reported by women with P4 and both tests positive.

The parameter estimates output showed the estimated rate for experiencing bothersome days to be 7.5 times higher in women with both ASLR and P4 tests positive, compared with the estimate for women with both tests negative. Women with both tests positive were estimated with twice the amount of bothersome days per week (table 3). For women with either P4 or ASLR test positive and a lower incidence rate, the mean amount of bothersome days was lower, but still estimated as approximately 1.5 times higher than for women with both tests negative. For every pregnancy, the mean number of bothersome days increased by 13.5%. Even a slightly higher BMI had a significant impact on the mean amount of bothersome days. Age had no impact.

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Table 3 Parameter estim	nates				
			95% Wald CI		
Parameter	В	SE	Lower	Upper	Significance
P4 and ASLR positive	2.021	0.1581	1.712	2.331	0.001
ASLR positive	1.540	0.2297	1.090	1.991	0.001
P4 positive	1.617	0.1832	1.258	1.976	0.001
Negative tests	0*	-	-	-	-
Age	-0.009	0.0067	-0.022	0.004	0.179
Number of births	0.135	0.0415	0.053	0.216	0.001
BMI before pregnancy	0.013	0.0065	0.000	0.026	0.047

*Set to 0 because this parameter is redundant.
ASLR, active straight leg raise test; BMI, body mass index; P4, posterior pelvic pain provocation test.

To our knowledge, this is the only study in which women with pelvic pain in pregnancy have been followed with SMS-Track. The main result of this study was that if both P4 and ASLR tests were positive in pregnancy week 18, a persistent pelvic pain of more than 5 days/ week throughout the remainder of pregnancy could be predicted. If either test was positive in week 18, a similar course was shown, but women with a positive P4 test revealed a more uncomfortable course than women with a positive ASLR test, who never reached the bothersome levels of the other groups. Robinson and coworkers²³ reported a similar outcome for the P4 test in a prospective cohort study on the association between sociodemographics, psychological and clinical factors measured at mid-pregnancy, and disability and pain intensity at week 30. However, their data showed no significant association between the ASLR test result and the disability and pain intensity in pregnancy week 30.

Although women who had a positive ASLR test and negative P4 test at baseline presented a comparatively low mean number of bothersome days with pain, they also had the highest mean number of previous pregnancies and the highest mean rate of pelvic pain in previous preg-nancies. Interestingly, our data also revealed that they exercised more frequently in comparison with women in the other positive test groups, both before and during the present pregnancy.

Interpretation

Since sufficient force closure of the sacroiliac joints requires appropriate muscular, ligamentous and fascial interac-tion, may women with pelvic pain in previous pregnancies have experienced that exercising improves muscle activa-tion, recovers function and decreases pain. 24-26 Additionally, experiences of pain prevention and rehabilitation in previous pregnancies may work as an incitement to engage in physical activity and regular exercise, both before and during pregnancy.

Our analysis also revealed a significant difference between the test groups in women described feeling depressed, and that a prepregnancy BMI slightly higher than average had a significant impact on the mean

number of bothersome days. Distress has previously been identified as a factor associated with a higher likelihood of PGP in pregnancy, as have a higher BMI and a higher gestational age.²⁷ One previous study found distress contributing to disability, but not to pain intensity.5

Nevertheless, some individuals seem to tolerate pain better, have less catastrophising tendencies and show more positive social response to pain, regardless of exposure to stressful circumstances and/or internal distress. ²⁸ Finally, women with a possibility to control their own work situation have better health during pregnancy than women without such chances. As indicated in this study and confirmed in previous studies, most pregnant women benefit from exercise since it increases pain tolerance, improves or maintains physical fitness, helps with weight management, reduces the risk of gestational diabetes in obesee women, and enhances psychological well-being. 29-32

A limitation of this study is the retrospectively collected information on pain in previous pregnancies and pain before pregnancy, which may produce biased results. Another limitation was found in the data collection via the SMS-Track system. In Norway, at the time of the study, there were more mobile phone service providers than in neighbouring countries, where SMS-Track studies previously had been successfully performed. Unfortunately, we had frequent problems reaching women through some of the providers. These problems led to data missing at random, but the GEE analytic approach may in these instances compensate for missing data. However, using the SMS-Track system in data collection is also a strength in our study, since it provides instant data on participants' situation, and responses are immediately recorded in a datasheet, which minimises further data handling and risk of error.

CONCLUSION

If both ASLR and P4 tests are positive at a clinical examination in mid-pregnancy, a course of persistent

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bothersome pelvic pain for more than 5 days per week throughout pregnancy may be predicted. The number of days per week with bothersome pelvic pain increases for every added pregnancy, but individual control over work situation and regular exercise may work as a PGP prophylactic since it invigorates a positive impact on optimal force closure of the pelvis, reduces risk of instability in the pelvic joints and enhances overall well-being. Since there is still no gold standard for diagnosing

PGP, particularly regarding the number of tests at the clinical examination, we recommend further research in this area, aiming at predictive, preventive and diagnostic measures for identifying women at risk of developing PGP in pregnancy. It would, for example, be interesting to see if women with a history of PGP have a higher pain-related anxiety and if it influences pain.

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Competing interests None declared.

Patient consent Obtained.

Ethics approval The study was carried out in accordance with the Helsinki Declaration II and was approved by the Regional Committee for Medical and Health Research Ethics, University of Bergen, Medical Faculty, Bergen, Norway (Ref Nr: 2010/174).

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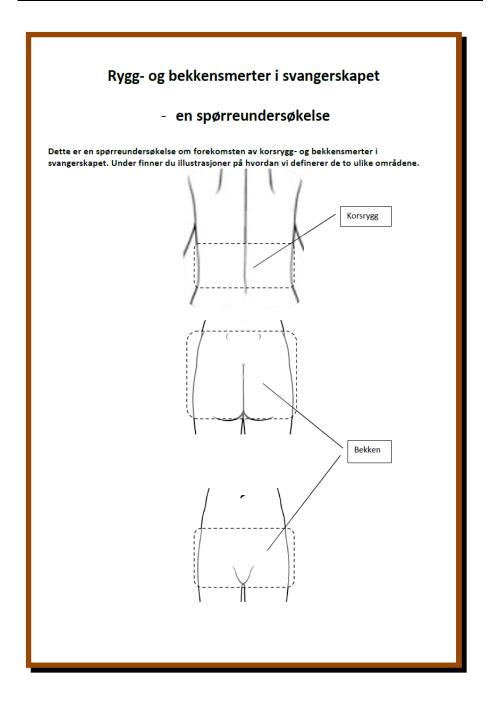
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8 APPENDICES

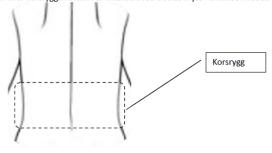
8.1 Questionnaire for the retrospective data collection



KYKE	g- og bekkensmerte	r i svangerskape	et - en spørreund	dersøkelse
Initialer	: Fødse	sdato:	Dagens dato:	
Hvor mange år Hvor fysisk tun Sett ett kryss.	s utdanning har du (inklu gt jobber du?	dert folkeskole/grun	nskole)?	<u>å</u> r
Veldig lett arbeid	Ganske lett arbeid	Verken lett eller tungt arbeid	Ganske tungt arbeid	Veldig tungt arbeid
3. Yrket ditt:	·			
 Hvor bra trives Sett ett kryss. 	du på din jobb eller der (du jobbet sist?		
Veldig dårlig	Ikke så bra	Verken bra eller dårlig	Ganske bra	Veldig bra
	%: antall u viktigste årsakene til sykr			
6. Oppgi den/de v	viktigste årsakene til sykr			
6. Oppgi den/de v	viktigste årsakene til sykr	neldingen(e):		
6. Oppgi den/de v 7. Din høyde: 8. Din vekt før sva	viktigste årsakene til sykr	neldingen(e):		
6. Oppgi den/de v 7. Din høyde: 8. Din vekt før sva 9. Din vekt like fø	viktigste årsakene til sykr cm angerskapet:kg ır fødselen:kg	neldingen(e):		
6. Oppgi den/de v 7. Din høyde: 8. Din vekt før sva 9. Din vekt like fø 10. Har du vært d	viktigste årsakene til sykr cm angerskapet:kg ır fødselen:kg	neldingen(e):	Nesten hele tider	
6. Oppgi den/de v 7. Din høyde: 8. Din vekt før sva 9. Din vekt like fø 10. Har du vært d Sett ett kryss.	viktigste årsakene til sykrcm angerskapet:kg or fødselen:kg leprimert i løpet av svang Av og til	gerskapet?	Nesten hele tider	

12. Har									
	du en kroni	sk sykdom:	Ja	N	ei				
13. Hvi	s ja; hvilken	sykdom:				_			
14. Ant	all tidligere	fødsler:	_						
a) K	orsryggsmei	erskap har d ter?Ja er?Ja	Nei						
16. Fikk	k du hormon	behandling f	or å bli grav	id før dette	svangerskape	et?Ja	Nei		
17. Tre	nte du regel	messig (mins	st 2-3 gange	r i uka) før sv	vangerskapet	?Ja	Nei		
18. Har	du trent reg	gelmessig (m	inst 2-3 gan	ger i uka) i s	vangerskapet	t?Ja	Nei		
19. Har	du hatt <u>von</u>	dt i korsrygg	<u>en siste år f</u>	<u>ør</u> svangersk	capet?	Ja	Nei		
20. Har	du hatt <u>von</u>	dt i bekkene	<u>t</u> siste år <u>før</u>	svangerska	pet?	Ja	Nei		
	du noen ga aNei	ng skadet ko	rsryggen elle	er bekkenet	slik at du må	tte oppsøk	e lege/sykehu	ıs?	
22. Hvi	s ja, hva slag	s type skade	hadde du?						
	du hatt von JaNe	,	en eller bek	kenet i løpet	t av dette sva	ingerskape	t?		
Hvis sv					er. Var svaret irsmålene un		spørsmål 23 tr	renger vi lit	t
Hvis sv mer inf Derso	ormasjon. \	/ennligst for	tsett med å :	svare på spø <u>// ERTER</u>			spørsmål 23 tr	renger vi liti	t
Hvis sva mer inf Dersc (har du	om du har ikke hatt ko	hatt KOR	SRYGGSN e, gå til spør	svare på spø MERTER rsmål 28)	irsmålene un	der.	spørsmål 23 tr yss i riktig rute		t
Hvis sv mer inf Derso (har du 24 . I hv	om du har ikke hatt ko	hatt KOR	SRYGGSN e, gå til spør	svare på spø MERTER rsmål 28)	irsmålene un	der.			t
Hvis sv mer inf Derso (har du 24 . I hv	ormasjon. N om du har ikke hatt ko rilken måned	hatt KOR rsryggsmert	SRYGGSN e, gå til spør apet begynt	NERTER Ssmål 28) e korsryggsr	orsmålene und merten? Mar	der. kér med kr	yss i riktig rute	е.	t
Hvis sv mer inf Derso (har du 24 . I hv	ormasjon. N om du har ikke hatt ko rilken måned	hatt KOR rsryggsmert	SRYGGSN e, gå til spør apet begynt	NERTER Ssmål 28) e korsryggsr	orsmålene und merten? Mar	der. kér med kr	yss i riktig rute	е.	t
Hvis sv mer inf Derso (har du 24 . I hv	ormasjon. \ om du har ikke hatt ko	hatt KOR rsryggsmert	SRYGGSN e, gå til spør apet begynt	NERTER Ssmål 28) e korsryggsr	orsmålene und merten? Mar	der. kér med kr	yss i riktig rute	е.	t
Hvis sva mer inf Derso (har du	ormasjon. \ om du har ikke hatt ko	hatt KOR rsryggsmert	SRYGGSN e, gå til spør apet begynt	NERTER Ssmål 28) e korsryggsr	orsmålene und merten? Mar	der. kér med kr	yss i riktig rute	е.	t

25. Vis på figuren nedenfor hvor korsryggsmerten var lokalisert ved å sette kryss i smerteområdet.



 $\textbf{26.} \ Mark\'{e}r\ i\ hver\ rute, som\ representerer\ hver\ måned\ i\ svangerskapet,\ i\ gjennomsnitt\ hvordan\ du\ har\ opplevd\ korsryggsmerten.$

Bruk tall fra 0-100 (0 = ingen smerte; 100 = uutholdelig smerte).

1	2	3	4	5	6	7	8	9

27. Dersom du har vært sykmeldt pga. korsryggsmerte: Markér nedenfor i hvilke måneder i ditt svangerskap du har vært sykmeldt. (Markér med kryss i riktig rute eller ruter.)

1	2	3	4	5	6	7	8	9

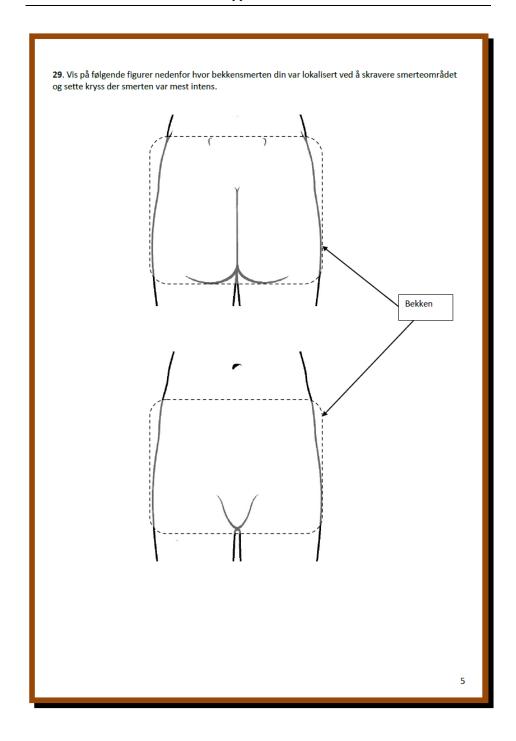
Dersom du har hatt BEKKENSMERTER

(Dersom du ikke har hatt bekkensmerte, kan du fortsette til spørsmål 32)

28. I hvilken måned i svangerskapet begynte <u>bekkensmertene</u>? Markér med kryss i riktig rute.

Į	1	2	3	4	5	6	7	8	9

4



Į.	2	3	4	5	6	7	8	9
31. Derson nar vært sy		t sykmeldt p	ga. bekkens	l merte: Mark	ér nedenfor	i hvilke mån	eder i svang	erskapet du
<u> </u>	2	3	4	5	6	7	8	9
vangerska					livet med di klarte meg p			
l	2	3	4	5	6	7	8	9
64. Har du Ja (Hvis Ne 65. Hvem d Le _j Kir	te/korsett	ferdig med	dette spørre		gg- eller bekl	sensmerter?		
	anuell terape dre behandl		l hvilke?					

Hvor god Verre I	d effekt, Ingen effekt mer enn e det bruk Ingen effekt lingen effekt du med lingen effekt	/utbytte Litt effekt en?) ga h ect varme Litt effekt et akupu Litt effekt	e hadde of God effekt vilke råd God effekt nktur? God effekt god effekt	du av rådene? Symptomfri Nei Symptomfri Ja Nei Symptomfri Symptomfri Symptomfri	JaNei (Hvis ja, hvilken terapeut?) (Hvis ja, hvilken terapeut?) Nei (Hvis ja, hvilken terapeut?)
Nerre I	lngen effekt mer enn e det bruk lngen effekt det bruk lngen effekt du med lngen effekt	Litt effekt en?) ga h et varme Litt effekt et akupu Litt effekt på bass Litt	God effekt vilke råd ??Ja God effekt nktur? God effekt engtreni God	Nei Symptomfri JaNei Symptomfri JaNei Symptomfri	(Hvis ja, hvilken terapeut?)
37. Ble di Verre Ii 839. Var di Verre Iii 640. Fikk ci Verre Iii	det bruk lingen effekt lingen effekt du med lingen effekt	effekt en?) ga h et varme Litt effekt et akupu Litt effekt på bass Litt	effekt vilke råd effekt God effekt God effekt engtreni God	Nei Symptomfri _JaNei Symptomfri	(Hvis ja, hvilken terapeut?)
37. Ble di Verre II 839. Var di Verre II 640 390. Var di Verre II 640 400. Fikk ci	det bruk Ingen effekt det bruk Ingen effekt du med Ingen effekt	en?) ga h et varme Litt effekt et akupu Litt effekt på bass Litt	vilke råd ??Ja God effekt nktur? _ God effekt engtreni God	Nei Symptomfri JaNei Symptomfri	(Hvis ja, hvilken terapeut?)
Verre II	Ingen effekt det bruk Ingen effekt du med Ingen effekt	Litt effekt et akupu Litt effekt på bass Litt	God effekt nktur? God effekt engtreni God	Symptomfri Ja Nei Symptomfri ing? Ja	(Hvis ja, hvilken terapeut?)
38. Ble de Verre III 1 1 1 1 1 1 1 1	effekt det bruk Ingen effekt du med Ingen effekt	effekt kt akupu Litt effekt på bass Litt	nktur?_ God effekt engtreni	Ja <u>Nei</u> Symptomfri ing? <u>Ja</u>	(Hvis ja, hvilken terapeut?)
38. Ble de Verre III 1 1 1 1 1 1 1 1	det bruk Ingen effekt du med Ingen effekt	t akupu Litt effekt på bass Litt	nktur? _ God effekt engtreni God	Symptomfri	
39. Var d Verre II 40. Fikk c Verre II	Ingen effekt du med Ingen effekt	Litt effekt på bass Litt	God effekt engtreni God	Symptomfri	
39. Var d Verre III	Ingen effekt du med Ingen effekt	Litt effekt på bass Litt	God effekt engtreni God	Symptomfri	
39. Var d Verre li e 40. Fikk d Verre li	du med Ingen effekt	på bass Litt	engtreni God	ng?Ja	Nei (Hvis ja, hvilken terapeut?)
Verre li 40. Fikk d Verre li	Ingen effekt	Litt	God		Nei (Hvis ja, hvilken terapeut?)
Verre li 40. Fikk d Verre li	Ingen effekt	Litt	God		Nei (Hvis ja, nviiken terapeutr)
40. Fikk o	effekt		ı	Symptomin	
40. Fikk o				1	
Verre I	december -				
				Nei	(Hvis ja, hvilken terapeut?)
€		Litt	God	Symptomfri	
	effekt	effekt	effekt		
41. Fikk o	du hjen	nmeøve	lser?J	aNei	(Hvis ja, hvilken terapeut?)
	Ingen	Litt	God	Symptomfri	, , ,
e	effekt	effekt	effekt		
42 Troni	ing mor	d voiladı	ning 2	la Noi	(Hvis ja, hvilken terapeut?)
Verre I		Litt	God	JaNei Symptomfri	(HVIS Ja, HVIIKEH LELAPEULT
		effekt	ı	Jypco	
			-	•	
43. Fikk o					(Hvis ja, hvilken terapeut?)
	Ingen effekt	Litt effekt	God effekt	Symptomfri	
	ellekt	EHERL	ellekt		
44. Ble d	let bruk	t TNS m	askin/st	røm?Ja _	_Nei (Hvis ja, hvilken terapeut?)
	_	Litt	God	Symptomfri	
e	effekt	effekt	effekt		
				ndling ?Ja r, Manuell ter	Nei peut. Andre utøvere ? Oppgi hvilke/n
	Ingen effekt	Litt effekt	God effekt	Symptomfri	

46. Har du hatt noen annen form for behandling?Ja	Nei
Hyis Ja, hya slags type behandling?	

Verre	Ingen	Litt	God	Symptomfri
	effekt	effekt	effekt	

- 47. Antall ganger du var til behandling?
- 48. I ca hvor mange uker fikk du behandling?
- 49. Har du en kommentar til behandlingen(e)?

50. Hvor godt fornøyd er du med behandlingen du fikk i svangerskapet? Fra null til ti (0=ikke fornøyd i det hele tatt, 10= veldig fornøyd). Sett ett kryss.

$$0-1-2-3-4-5-6-7-8-9-10$$

 $\textbf{51.} \ \ \text{Generelt hvor godt fornøyd er du med behandlingstilbudet for korsrygg- og bekkensmerter under svangerskapet? Fra null til ti (0=ikke fornøyd i det hele tatt, 10= veldig fornøyd). Sett ett kryss.}$

$$0-1-2-3-4-5-6-7-8-9-10$$

Takk for hjelpen så langt!

Husk å fylle ut de 2 andre skjemaene og!

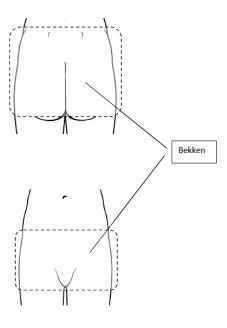
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8.2 Questionnaire for the prospective data collection

Rygg - og bekkensmerter i svangerskapet

Spørreskjema ved rutineundersøkelsen

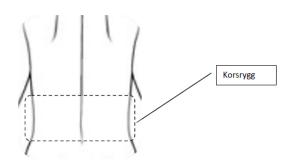
Vi ønsker å kartlegge forekomsten av bekkensmerter i svangerskapet. For å kunne tolke resultatene må vi gi en definisjon av hvor bekkenet er lokalisert. Nedenfor finner du en illustrasjon av hvor vi definerer bekkenområdet.



				
		ekkensmerter i s kjema ved rutineund		
Initialer	: Fødse	elsdato:	Dato i dag:	
	· In a land of talls	to a College leader (service	1-1-12	o_
	s utdanning har du (inklu	idert folkeskole/gruf	nnskole j r	ar
2. Hvor fysisk tung Sett ett kryss.	gt jobber du?			
Veldig lett arbeid	Ganske lett arbeid	Verken lett eller tungt arbeid	Ganske tungt arbeid	Veldig tungt arbeid
Jobber ikke				
4. Hvor bra trives Sett ett kryss.	du på din jobb eller der (du jobbet sist?		
Veldig dårlig	Ikke så bra	Verken bra eller dårlig	Ganske bra	Veldig bra
5. Hvor mange uk	er har du vært sykmeldt	til nå i svangerskapet	1?	
	er har du vært sykmeldt	til nå i svangerskapet	1?	
Har ikke vært s 100%:	ykmeldt antall uker.		t?	
Har ikke vært s 100%:	ykmeldt		1?	
Har ikke vært s 100%: Delvis	ykmeldt antall uker.	uker.		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v	ykmeldt antall uker. %: antall u iktigste årsakene til sykn	uker.		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde:	ykmeldt antall uker antall u %: antall u iktigste årsakene til sykn cm	uker.		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 	ykmeldt antall uker antall u %: antall u iktigste årsakene til sykn cm kg	uker. neldingen(e):		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde: 8. Din vekt:	ykmeldt antall uker antall uker antall ukiktigste årsakene til sykncmkg	uker. meldingen(e):		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde: 8. Din vekt:	ykmeldt antall uker antall u %: antall u iktigste årsakene til sykn cm kg	uker. meldingen(e):		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde: 8. Din vekt: 9. Din vekt før sva	ykmeldt antall uker antall uker antall ukiktigste årsakene til sykncmkg	uker. meldingen(e):		
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde: 8. Din vekt: 9. Din vekt før sva 10. Har du vært de Sett ett kryss.	ykmeldt antall uker kg kg kg kg kg eprimert til nå i svangers	uker. meldingen(e):	Nesten hele tiden	
Har ikke vært s 100%: Delvis 6. Oppgi den/de v 7. Din høyde: 8. Din vekt: 9. Din vekt før sva 10. Har du vært de Sett ett kryss.	ykmeldt antall uker antall uker antall ukiktigste årsakene til sykn cm kg kg kg eprimert til nå i svangers kg kg kg erimert til nå i svangers kg	uker. meldingen(e):	Nesten hele tiden	

12. Har du en kronisk sykdom:JaNei
13. Hvis ja; hvilken sykdom:
14. Antall tidligere fødsler:
15. Har du hatt bekkensmerter i tidligere svangerskap?JaNei
16. Fikk du hormonbehandling for å bli gravid før dette svangerskapet?JaNei
17. Trente du regelmessig (minst 2-3 ganger i uka) før svangerskapet?JaNei
18. Har du trent regelmessig (minst 2-3 ganger i uka) til nå i dette svangerskapet?JaNei
20. Har du hatt <u>vondt i bekkenet</u> siste år <u>før</u> svangerskapet?JaNei
21. Har du noen gang skadet bekkenet slik at du måtte oppsøke lege/sykehus?JaNei
22. Hvis ja, hva slags type skade hadde du?
23. Har du hatt vondt i korsryggen til nå i dette svangerskapet?JaNei
24. Har du hatt vondt i bekkenet til nå i dette svangerskapet?JaNei
(Hvis Ne i på spørsmålene 23 og 24 er du nå ferdig med dette spørreskjemaet)
Dersom du har hatt KORSRYGGSMERTER
(har du ikke hatt korsryggsmerte, gå til spørsmål 29)
25. I hvilke uker i svangerskapet begynte korsryggsmerten? Markér med kryss i riktig rute.
1-4 5-8 9-12 13-16 17-20

26. Vis på figuren nedenfor hvor korsryggsmerten var lokalisert ved å sette kryss i smerteområdet.



27. Markér i hver rute i gjennomsnitt hvordan du har opplevd korsryggsmerten. Bruk tall fra 0-100 (0 = ingen smerte; 100 = uutholdelig smerte).

1-4	5-8	9-12	13-16	17-20

28. Dersom du har vært sykmeldt pga. korsryggsmerte: Markér nedenfor i hvilke uker i ditt svangerskap du har vært sykmeldt. (Markér med kryss i riktig rute)

1-4	5-8	9-12	13-16	17-20

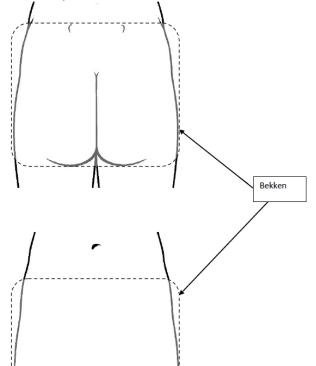
Dersom du har hatt BEKKENSMERTER

(Dersom du ikke har hatt bekkenplager, kan du gå til spørsmål 33)

29. I hvilke uker i svangerskapet begynte <u>bekkenplagene</u>? Markér med kryss i riktig rute.

1 /	E O	0.12	12 16	17 20

30. Vis på følgende figurer nedenfor hvor bekkensmerten din var/er lokalisert ved å skravere smerteområdet og sette kryss der smerten var/er mest intens.



31. Markér i hver rute, som representerer uker i svangerskapet, hvordan du i gjennomsnitt har opplevd bekkensmerten. Bruk tall fra $\,0-100\,$ (0 = ingen smerte; $\,100$ = uutholdelig smerte)

1-4	5-8	9-12	13-16	17-20	
-----	-----	------	-------	-------	--

32. Dersom du har vært sykmeldt pga. bekkensmerte: Markér nedenfor i hvilke uker i svangerskapet du har vært sykmeldt.

4.4	r 0	0.13	12.10	17.20
1-4	5-8	9-12	13-16	17-20

ingen prob	olem, jeg klai	rte meg pa e	Berniana, 10		, jeg matte na i	y	,
1-4	į	5-8	9-1	2	13-16		17-20
34. Har sm	nerten vært s	så sterk at di	u har trengt h	njelpemidler?			
Krykker		Ja	Nei				
Rullestol		Ja	Nei				
"Glidelake	n"	Ja	Nei				
Bekkenbe	lte/korsett	Ja	Nei				
35. Har du Ja	ı fått behand Nei		kensmerter i	i dette svangers	kapet?		
(Hvis N€	i er du nå fe	rdig med de	tte spørreskj	emaet)			
	oppsøkte du erapeut, Nar			ege, Kiroprakto	r, Fysioterapeu	t , Akupunl	ktør, Osteopat,
Manuell To Vi er intere eventuelle parentes h	erapeut, Nap essert i å vite e behandling nvem som gjo	e hva slags b smetoder so orde hva.	et)? ehandling du om er blitt bru I smerter?	ı har gjennomga ukt. Hvis du har	ått. Nedenfor fi	nner du sp	ørsmål om
Manuell To Vi er intere eventuelle parentes h	erapeut, Nap essert i å vite e behandling nvem som gjo m å takle hve	e hva slags b smetoder so orde hva.	et)? ehandling du om er blitt bru I smerter?	ı har gjennomga ukt. Hvis du har	ått. Nedenfor fi vært hos flere	nner du sp	ørsmål om
Vi er intereventuelle parentes h 37. Råd on Hvis ja,	erapeut, Nap essert i å vite e behandling nvem som gjo m å takle hve	orapat, anne e hva slags b smetoder so orde hva. erdagen med ype råd?	ehandling du om er blitt bru d smerter?	ı har gjennomga ukt. Hvis du har	ått. Nedenfor fi vært hos flere	nner du sp	ørsmål om
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Manuell To Vi er interneventuelle parentes h 37. Råd on Hvis ja, 38. Hvor g Verre 39. Ble det 40. Ble det	essert i å vite behandling nvem som gje m å takle hve , hva slags ty od effekt had lngen effekt t brukt varm	e hva slags b smetoder so orde hva. erdagen med ype råd? dde du av rå Litt effekt	ehandling du om er blitt bru d smerter? idene? God effekt Nei aNei ?JaN	u har gjennomga ukt. Hvis du har _Ja Symptomfri (Hvis j (Hvis j	ått. Nedenfor fi vært hos flere Nei ia, hva slags tera ia, hva slags tera	nner du sp terapeuter apeut? apeut?	ørsmål om r sett gjerne i

44. Trening med veiledning?	Ja	Nei	(Hvis ja, hva slags terapeut?	

45. Fikk du medikamenter? __Ja Nei (Hvis ja, hva slags terapeut?____

46. Ble det brukt TENS-maskin/strøm? __Ja __Nei (Hvis ja, hva slags terapeut?____

47. Fikk du manipulasjonsbehandling ? __Ja __Nei Hvis ja, av hvem: Kiropraktor, Manuell Terapeut, Osteopat, Naprapat eller annet?

48. Har du hatt noen annen form for behandling? __Ja __Nei Hvis Ja, hva slags type behandling?

49. Hvor mange ganger har du har vært til behandling til nå i svangerskapet?

50. I ca hvor mange uker har du fått behandling? _____

51. Hvor godt fornøyd er du med behandlingene du har fått i svangerskapet? Fra null til ti (0=ikke fornøyd i det hele tatt, 10= veldig fornøyd). Sett ett kryss.

$$0-1-2-3-4-5-6-7-8-9-10$$

52. Generelt hvor godt fornøyd er du med **behandlingstilbudet** for korsrygg- og bekkensmerter i svangerskapet?

Fra null til ti (0=ikke fornøyd i det hele tatt, 10= veldig fornøyd). Sett ett kryss.

$$0-1-2-3-4-5-6-7-8-9-10$$

Takk for all informasjon!