## Improving Postpartum Healthcare and Health Outcomes of Women with a History of Gestational Diabetes Mellitus

PhD Thesis of Dr Emer Van Ryswyk

Discipline of Obstetrics and Gynaecology School of Paediatrics and Reproductive Health The University of Adelaide, Australia

Submitted for the degree of Doctor of Philosophy in March 2015

## Table of Contents

Abstract	3
Thesis Declaration	6
Acknowledgements	7
<i>Chapter:</i> Literature Review	8
Published Paper: Clinician views and knowledge regarding healthcare provision in the	
postpartum period for women with recent gestational diabetes: A systematic review of	
qualitative/survey studies	30
<i>Chapter:</i> Women's views and knowledge regarding healthcare seeking for gestational diabetes in the postpartum period: a systematic review of qualitative/survey studies	42
<b>Published Protocol:</b> The DIAMIND study: postpartum SMS reminders to women who had gestational diabetes mellitus to test for type 2 diabetes: a randomised controlled trial	nave
study protocol	72
Accepted Paper: Postpartum SMS reminders to women who have experienced gestation	al
diabetes to test for type 2 diabetes: the DIAMIND randomised trial	79
<i>Chapter:</i> Predictors of OGTT completion in the DIAMIND Study	97
Submitted Paper: Women's views on postpartum testing for type 2 diabetes after gestat	ional
diabetes: six month follow-up for the DIAMIND RCT	106
Conclusions	122
Bibliography	129
Appendices	161

## Abstract

#### Background

Women who have had gestational diabetes mellitus (GDM) are at increased risk of type 2 diabetes (T2DM), and are recommended to have T2DM screening in the postpartum period, although this screening is often not undertaken. This thesis examines how postpartum care for women with GDM may be improved.

#### Methods

Two systematic reviews of qualitative/survey studies examine:

- (1) Clinicians' views and knowledge relating to provision of healthcare in the postpartum period for GDM.
- (2) Women's views and knowledge relating to healthcare seeking after GDM.

A randomised controlled trial and two nested studies assess:

- Postpartum SMS reminders to women who have experienced GDM to test for T2DM: The DIAMIND Trial
- (2) Predictors of postpartum diabetes screening in the DIAMIND Trial
- (3) Barriers and facilitators to postpartum diabetes testing.

#### Results

The systematic review on clinician's views included 13 studies (4435 clinicians). Key themes included adequacy of knowledge of risk of T2DM, and differing perceptions of the value of postpartum screening. Women faced obstacles to accessing healthcare, and a need for improved GDM education. Studies reported shortfalls in systems to ensure communication of the GDM diagnosis and postpartum screening.

The systematic review on women's views included 42 studies (7949 women). Nonjudgemental, well-coordinated care was preferable. Perception of T2DM risk increased with time from their GDM diagnosis, family history of T2DM and other risk factors for GDM. Children's needs took priority over their own healthcare. A need for a more pro-active approach to postpartum care was identified.

The DIAMIND Trial found that SMS reminders did not increase attendance for an oral glucose tolerance test (OGTT) within six months postpartum, with 104 (77.6% of 134) women attending in the six week group and 103 (76.8% of 134) women attending in the control group (RR 1.01, 95% CI 0.89-1.15).

Women were more likely to complete OGTTs if they were of Asian ethnicity (P =0.007), had a bachelor's degree (P = 0.036), and if they did not smoke prior to pregnancy (P = 0.045). Women were less likely to attend if they had gained excessive weight during their pregnancy (P = 0.004) or were Caucasian (P = 0.001).

208 women (75%) returned their questionnaires. Preferred reminder types were *SMS* (67%), *email* (17%), *postal* (12%) and *voice call* (1%). Common barriers to postpartum glucose testing included: *not having enough time* (73%), *inadequate or non-availability of childcare* (30%), and a *need to focus on the health of the baby* (30%). The most common facilitator for postpartum testing was *having a shorter test* (33%).

#### Conclusions

Postpartum care for women with GDM could be improved through systematic communication of the diagnosis, clear responsibilities for postpartum care, better GDM education and minimisation of healthcare cost barriers. Non-judgemental, holistic and pro-active care is preferable.

4

An OGTT postpartum attendance "ceiling effect" may explain the non-increase in attendance in the six week SMS reminder group. Lack of time and caring responsibilities were barriers to OGTT completion. Further research is needed on reasons for postpartum non-attendance to facilitate diabetes detection and prevention.

## Thesis Declaration

I certify that this work contains no material which has been accepted for the award of any other degree or diploma in my name, in any university or other tertiary institution and, to the best of my knowledge and belief, contains no material previously published or written by another person, except where due reference has been made in the text. In addition, I certify that no part of this work will, in the future, be used in a submission in my name, for any other degree or diploma in any university or other tertiary institution without the prior approval of the University of Adelaide and where applicable, any partner institution responsible for the joint-award of this degree. I give consent to this copy of my thesis when deposited in the University Library, being made available for loan and photocopying, subject to the provisions of the Copyright Act 1968.

The author acknowledges that copyright of published works contained within this thesis resides with the copyright holder(s) of those works. I also give permission for the digital version of my thesis to be made available on the web, via the University's digital research repository, the Library Search and also through web search engines, unless permission has been granted by the University to restrict access for a period of time.

Signature:\_\_\_\_\_

Date:\_\_\_\_

## Acknowledgements

There are many people who have supported and encouraged me throughout my candidature to whom I am very grateful. I would like to thank my supervisors, Professor Caroline Crowther, Ms Philippa Middleton and Professor William Hague for their excellent guidance and constructive feedback. I would like to thank the Robinson Research Institute, School of Paediatrics and Reproductive Health and Discipline of Obstetrics and Gynaecology for the opportunity to undertake this PhD.

To my family, particularly my amazing husband Ronald and my parents Ann and Edward, thank you for your love, support and unwavering belief in me.

I would like to thank all of the midwives of the postnatal ward, and the diabetes educators Maree Thus and Kerry Boylan for their recruitment advice, encouragement and assistance.

I would like to thank Michael Draper for his assistance in development of the search strategy of the two systematic reviews included in this thesis.

I am also very appreciative of the friendly encouragement and professional assistance of the staff and students working in the School of Paediatrics and Reproductive Health.

I am grateful for the financial support I received from the HCF Health and Medical Research Foundation for the DIAMIND Study, as well as the Faculty of Health Sciences for the PhD Scholarship.

Finally, I would like to thank all of the women who volunteered to take part in the DIAMIND trial who kindly gave their time and views, and made the study possible.

### Literature Review

Gestational diabetes mellitus (GDM) is a form of diabetes, or carbohydrate intolerance, which is first diagnosed during pregnancy (World Health Organization 2013). Women who have had GDM are at higher risk of type 2 diabetes mellitus (T2DM) in the future; they are also at risk of GDM in future pregnancies (Kim et al 2007;Kim et al 2002;Lee et al 2007). Due to this increased risk of T2DM, clinical practice guidelines recommend screening for T2DM in the postpartum period (American Diabetes Association 2014;Nankervis et al 2014;South Australian Perinatal Practice Guidelines 2012;The American College of Obstetricians and Gynecologists 2013;Thompson et al 2013;Walker 2008). However, screening rates are moderately low, and vary considerably between settings (Tovar et al 2011). It is therefore important to investigate both the causes of these low rates, as well as possible solutions.

#### GDM: A brief history

Prior to development of the term GDM, and diagnostic and screening recommendations, it was observed that women with poor obstetric histories (e.g. a large baby or fetal loss) were more likely to have high blood glucose during a later pregnancy and to go on to develop T2DM and that treatment of the high blood glucose improved obstetric outcomes; thus research was conducted into methods of detection of diabetes during pregnancy (Carrington et al 1957). The O'Sullivan and Mahan criteria was subsequently published in 1964; this criteria for diagnosing diabetes in pregnancy was based on results from 100 gram oral glucose tolerance tests (OGTTs) performed in 752 mainly second- and third-trimester pregnant women. Four venous glucose values were measured (fasting, one hour, two hours and three hours); the results were normally distributed, and the predictive value for future diabetes was validated by applying them to a second population of 1013 non-pregnant women who had been tested during a previous pregnancy and followed up for up to eight years. O'Sullivan and Mahan concluded from their results that the mean values plus two standard deviations

(rounded to the nearest 5mg/dL) were the most appropriate limits for diagnosing diabetes in pregnancy (O'Sullivan and Mahan 1964).

In 1979, the National Diabetes Data Group in the United States recommended that GDM be recognised as a condition with its own diagnostic criteria, and that high risk pregnant women be screened (many risk factors were specified by the group and included family history of diabetes in a first degree relative, history of stillbirth and maternal obesity) (National Diabetes Data Group 1979). Then in 1980, the American Diabetes Association recommended that all pregnant women be screened between the 24<sup>th</sup> and 28<sup>th</sup> weeks of pregnancy, using a 50 gram oral glucose challenge test (OGCT), followed by a 100 gram oral glucose tolerance test (OGTT). Women would be diagnosed with GDM if both tests had abnormal results (Freinkel and Josimovich 1980). Many countries developed their own diagnostic criteria, primarily based on expert opinion (Cundy et al 2014). In 2005 and 2009, two randomised controlled trials were published which demonstrated benefits of treatment of GDM (Crowther et al 2005;Landon et al 2009). Then, in 2008, the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) study was published, and the findings of this study led to new diagnostic recommendations (Hapo Study Cooperative Research Group et al 2008;International Association of Diabetes Pregnancy Study Groups Consensus Panel et al 2010). There is currently ongoing debate around the diagnostic criteria for GDM, as briefly described in the following paragraphs.

#### GDM diagnostic criteria: Arguments for changing the diagnostic threshold

The HAPO study, a prospective, multicentre cohort study, with 23,216 pregnant women, found that there was a continuous relationship between maternal glycaemia at 24 to 28 weeks, and adverse outcomes for mothers and babies. The results of the study remained significant after adjusting for possible confounders such as maternal obesity. The results of the HAPO study, and results from other studies on the same topic (Hillier et al 2007;Jensen et al 2001;Jensen et al 2008;Pettitt and Knowler 1998;Pettitt et al 1980;Sacks et al 1995;Sermer et

9

al 1995), including two randomised controlled trials (Crowther et al 2005;Landon et al 2009) were considered by the International Association of Diabetes and Pregnancy Study Groups (IADPSG) (International Association of Diabetes Pregnancy Study Groups Consensus Panel et al 2010), and led to recommendations of new diagnostic criteria. The key differences in the IADPSG recommendations compared with previous criteria were use of a one-step (OGTT only) rather than two-step approach to diagnosis, and adjustment of the glucose thresholds for diagnosis to levels associated with 1.75-fold increase in risk above the mean (from HAPO study results) for birth weight, cord C peptide concentration, and percentage body fat > 90<sup>th</sup> centile. Only one abnormal value would then be needed for diagnosis of GDM (Table 1).

Organisation	Target group	Method of screening	Diagnostic thresholds
IADPSG	All women	75 g OGTT (one-step)	Fasting $\geq$ 5.1 mmol/L
			1 hour $\geq$ 10.0mmol/L
			2 hours $\geq$ 8.5mmol/L

 Table 1: IADPSG Recommended GDM Diagnostic Criteria

The new diagnostic criteria were endorsed by the Australasian Diabetes in Pregnancy Society (ADIPS) and the American Diabetes Association (American Diabetes Association 2014), as well as associations in France and China (Cundy et al 2014;Nankervis et al 2013;Nankervis and Conn 2013). The World Health Organisation (WHO) also endorsed the recommendations, but rated the quality of the evidence as very low, and rated the strength of their recommendation as weak (World Health Organization 2013).

#### GDM diagnostic criteria: Arguments against decreasing diagnostic threshold

Not all clinicians, researchers and associations with an interest in GDM agree with the proposed changes to the GDM diagnostic criteria (Cundy et al 2014;Langer et al 2013). For example, the US College of Obstetricians and Gynaecologists has not endorsed the change (Cundy et al 2014;The American College of Obstetricians and Gynecologists 2013), and the 2013, Eunice Kennedy Shriver National Institute of Child Health and Human Development Consensus Development Conference on diagnosing GDM recommended that clinicians

continue to use a two-step approach to screen for and diagnose GDM because there would be a certain rise in health care costs despite insufficient evidence for clinically significant improvements in outcomes (Vandorsten et al 2013). The Canadian Diabetes Association (CDA) has also recommended retaining a two-step approach as their preferred option for GDM diagnosis (Thompson et al 2013).

Numerous arguments for and against adopting the new criteria have been made (Cundy et al 2014;d'Emden 2014;International Association of Diabetes Pregnancy Study Groups Consensus Panel et al 2010;Kevat et al 2014;Langer et al 2013). The new criteria, where adopted, may significantly increase the number of women being diagnosed with GDM. It is important that costs and benefits of the proposed criteria are further explored through well-designed research, particularly randomised controlled trials.

#### GDM Diagnosis and Prevalence

In 2010, the Australian Institute of Health and Welfare (AIHW) released the first national report on the impact of diabetes in pregnancy on Australian women and their babies (Australian Institute of Health and Welfare 2010). This report examined maternal characteristics in comparison with perinatal interventions and outcomes, using data from both the National Hospital Morbidity Database and the National Perinatal Data Collection database. The AIHW report found that the overall population prevalence of GDM in Australia (using ADIPS 1998 criteria (Hoffman et al 1998)) was 5%.

The prevalence of GDM is not only influenced by diagnostic criteria, but also by the presence of individual risk factors of the women becoming pregnant in a particular setting. The risk factors for GDM overlap with those for T2DM, and include overweight and obesity, use of medications such as corticosteroids, increased age, previous GDM, previous macrosomic babies, polycystic ovarian syndrome and certain ethnicities including Asian, Indian subcontinent, Aboriginal, Torres Strait Islander, Pacific Islander, Maori, Middle Eastern and non-white African (Nankervis et al 2013;Teh et al 2011). The prevalence of GDM in Australia is likely to grow with increases in maternal obesity and age, migration and use of altered GDM diagnostic criteria (Nankervis and Conn 2013;Scheil et al 2013).

#### Women who have had GDM are at greatly increased risk of T2DM

Women who have had GDM are at higher risk of development of T2DM in the future (Bellamy et al 2009). A systematic review and meta-analysis published in 2009 found that women who have had GDM (diagnosed prior to IADPSG criteria) are at least seven times as likely as those who had a normoglycaemic pregnancy to develop T2DM (RR 7.43, 95% CI 4.79-11.51) (Bellamy et al 2009). This systematic review included retrospective and prospective cohort studies from 1960 to 2009, with data from women who had GDM as well as from women who had normoglycaemic pregnancies. The length of follow-up in the included studies ranged from 16 weeks to 28 years (Bellamy et al 2009).

The methodological quality of the systematic review described above may be assessed using the AMSTAR tool, which is validated, has 11 criteria, and was developed based on previous research into assessment of the methodological quality of systematic reviews (Shea et al 2009). The review was generally well designed and reported, with a score of 9/11 (high quality), according to AMSTAR criteria; the review had an "a priori" design (established research question and inclusion criteria prior to conduct). Methods included duplicate study selection and data extraction, and a comprehensive literature search was performed. Other aspects, such as the likelihood of publication bias, were addressed. There was significant unexplained heterogeneity in the meta-analysis, and studies were included regardless of their rate of follow-up, which may have resulted in a biased effect estimate (Kristman et al 2004), but the increased risk of T2DM was unequivocally demonstrated.

#### Why postpartum follow-up of GDM is important

Postpartum follow-up of women who have experienced GDM allows early detection of T2DM, as well as an opportunity for prevention of T2DM in women who are known to be at higher risk.

#### The benefits of early detection of T2DM

Glucose testing in the postpartum period allows those who have developed T2DM to be identified and treated early, with a view to reducing the risk of complications of pregestational diabetes (diabetes present before pregnancy) (Thompson et al 2013) and long term complications. Identification of T2DM before future pregnancies allows for optimised blood glucose control in the early stage of pregnancy (Thompson et al 2013). Maternal hyperglycaemia during early pregnancy increases the risk of several complications for the mother and her baby such as congenital malformations (e.g. abnormalities of the heart and central nervous system, which are increased three-fold), miscarriage, stillbirth, fetal macrosomia, requirement for induction of labour or caesarean section and birth trauma for mother and baby (Bell et al 2008;Feig et al 2006;Macintosh et al 2006;Thompson et al 2013).

For women, in the long term, treating T2DM using exercise, dietary and pharmacological methods helps to reduce morbidity, including macrovascular disease (atherosclerosis) and microvascular disease (retinopathy, nephropathy, and neuropathy) and mortality (Bailey et al 2005;Hordern et al 2012;McCulloch 2012).

#### The benefits of being able to prevent T2DM in women who are at high risk

All women who have had GDM are at increased risk of T2DM, so for women who have not yet developed T2DM, postpartum appointments present an opportunity for T2DM prevention. This may lead to numerous health and cost benefits. The overall financial cost of T2DM in Australia was estimated to be \$12.4 billion in 2008, and has been rising (Access Economics 2008). Direct medical costs increase greatly with progression from impaired glucose tolerance to diabetes with complications (Herman 2011). Lifestyle and pharmacological interventions to prevent T2DM have been shown to be cost-effective (Herman 2011).

For example, both metformin and lifestyle modification were shown to reduce the development of T2DM in the Diabetes Prevention Program (DPP) trial. The DPP trial included 3234 people with increased risk of T2DM (i.e. elevated fasting and post-load plasma glucose concentrations), and was conducted in 27 centres in the United States (Knowler et al 2002); the primary goals of the DPP trial were to determine whether a lifestyle intervention or metformin would prevent or delay the onset of T2DM, whether the interventions differed in effectiveness, and also whether their efficacy differs according to age, sex, or ethnicity. Participants were followed-up for an average 2.8 years. The lifestyle intervention reduced the incidence of T2DM by 58 percent (95% CI 48 to 66) and metformin reduced it by 31 percent (95% CI 17 to 43); the incidence of diabetes was 11.0, 7.8, and 4.8 cases per 100 person-years in the placebo, metformin, and lifestyle groups, respectively (Table 2)

Participants	Overall Incidence		R	eduction in Inciden	ce	
	(cases/100 person-years)		Percent (95% CI)			
	Placebo	Metformin	Lifestyle vs	Lifestyle vs	Metformin vs	Lifestyle vs
			Placebo	Placebo	Placebo	Metformin
3234	11.0	7.8	4.8	58 (48 to 66)	31 (17 to 43)	39 (24 to 51)

 Table 2: Overall Incidence and Reduction of Incidence of T2DM in the DPP Trial (Knowler et al 2002)

The DPP lifestyle intervention goal was for participants to achieve and maintain a 7% initial body weight reduction through low-fat diet and physical activity. The participants were taught by case managers individually for the first 24 weeks after enrolment about diet, exercise and behaviour modification, with subsequent individual sessions (usually monthly) and group sessions with the case managers to reinforce the behavioural changes.

In 2008, a sub-group analysis of the DPP trial was published in which the results from women with a history of GDM (hGDM) were compared with results from women without hGDM (Ratner et al 2008). The findings were that women with hGDM were more likely to develop T2DM overall (71% higher incidence), and that both intensive lifestyle and metformin therapy reduced the incidence of T2DM in women with hGDM by approximately 50% compared with the placebo group.

Further research studies examining lifestyle interventions, specifically for women who have had GDM, are necessary (Bentley-Lewis et al 2008; Pan et al 1997; Tuomilehto et al 2001). One such study is the Mothers After Gestational Diabetes in Australia Diabetes Prevention Program (MAGDA-DPP; Shih et al 2013), which is a randomised controlled trial for assessment of a structured diabetes prevention program for post-GDM women (compared with usual care). The diabetes prevention program in the MAGDA study aims to encourage participants to achieve goals of: (1) reducing fat intake (no more than 30% of energy from fat); (2) decreasing saturated fat intake (no more than 10% of energy from saturated fat); (3) increasing fibre intake (at least 15 gram per 1000 kcal); (4) increasing physical activity (at least 30 minutes of moderate exercise per day); and (5) reducing body weight (at least 5% of body weight reduction within 12 months). Another Diabetes Prevention Program trial specifically for women who have experienced GDM is currently being conducted in Northern California, The GEM (Gestational Diabetes' Effects on Moms) study. The GEM study follows on from a feasibility study conducted by the same research group of a pregnancy and postpartum print and telephone lifestyle intervention based on the DPP curriculum (Ferrara et al 2011). The GEM study is a cluster randomised trial of a DPP-derived lifestyle intervention (delivered via telephone) versus usual care (Ferrara et al 2014). The results of these trials are not yet available.

Breastfeeding may also decrease the risk of development of T2DM in women who have had GDM (Feig 2012). Two large cohort studies have shown decreased rates of T2DM in women who breastfed their babies compared with those who did not (Stuebe et al 2005;Liu et al 2010). Another trial has found that longer duration of breastfeeding was associated with lower incidence of metabolic syndrome in women with a history of GDM (after controlling for BMI and sociodemographic and lifestyle traits) (Gunderson et al 2010).

### Guidelines relating to follow-up of GDM in the postpartum period

Due to the increased likelihood of T2DM, many guidelines, including those listed in the table below, recommend postpartum diabetes screening for women who have had GDM.

Guideline	Screening Recommendations	Comments
Australasian Diabetes in	Early screening:	This guideline is based on evidence
Pregnancy Society	75g 2-hr OGTT, preferably at 6-12	and consensus opinion of Council
(Nankervis et al 2014)	weeks post-partum, with classification	members of the ADIPS. These
	according to the WHO criteria.	guidelines have been endorsed by the
	Longer term:	Royal Australian and New Zealand
	If contemplating another pregnancy:	College of Obstetricians and
	OGTT annually. If not, then depending	Gynaecologists (RANZCOG).
	on the clinical circumstances either an	
	OGTT, HbA1c or FPG at 1-3 years.	
South Australian	Early screening:	These guidelines are based on
Perinatal Practice	75 gram OGTT at 6-12 weeks post-	systematic review of evidence, and
Guidelines 2012	partum.	have been approved by the South
	Longer term:	Australian Department of Health.
	1-2 yearly OGTT.	
American Diabetes	Early screening:	This guideline is evidence based. It is
Association 2014	Screen women with GDM for	annually revised by the ADA's
	persistent diabetes at 6-12 weeks	multidisciplinary Professional Practice
	postpartum, using a test other than	Committee. This guideline was
	A1C.	approved by the Executive Committee
	Longer term:	of ADA's Board of Directors, which
	3 yearly testing using OGTT.	includes health care professionals,
		scientists, and lay people. Feedback
		from the larger clinical community was
		also incorporated. No industry support
		was used to fund the development of
		the guideline.
The American College of	Early screening:	This guideline represents an
Obstetricians and	75 gram OGTT or FPG 6-12 weeks	assessment on the issue by the
Gynecologists 2013	postpartum.	American College of Obstetricians and
	Longer term:	Gynaecologists' Committee on
	3 yearly testing using OGTT	Obstetric Practice.
Canadian Diabetes	Early screening:	These guidelines are evidence based
Association (Thompson	75 gram OGTT at 6 weeks to 6 months	and developed by the Clinical &
et al 2013)	postpartum	Scientific Section of the Canadian

 Table 3: Postpartum glucose testing guideline recommendations

	Longer term:	Diabetes Association, a
	No recommendation	multidisciplinary team of experts
		working as volunteers to develop the
		guideline. Financial assistance for
		guideline was provided by industry
		sponsors in the form of unrestricted
		educational grants
The UK National	Early screening:	NICE guidelines aim to be evidence
Institute for Health and	FPG (not an OGTT) should be	based. They are developed by an
Clinical Excellence	performed at 6 weeks.	independent committee of experts
(NICE) guideline	Longer term:	including clinicians, women, carers and
(Walker 2008)	FPG annually.	health economists, with UK National
		Health Service funding. The committee
		also consult with an associated public
		citizen's council.
New Zealand Ministry of	Early screening:	The guideline has been commissioned
Health	Primary care provider should offer	and funded by the Ministry of Health.
2014	screening for T2DM at three months	It was developed by a multidisciplinary
	after birth using HbA1c.	Guideline development team, and was
	Longer term:	evidence based.
	Repeat the HbA1c test in one year.	

How often does postpartum follow-up of women with GDM occur?

There is evidence that rates of postpartum screening for T2DM are low-to-moderate (Tovar et al 2011). A systematic review by Tovar et al 2011 examining the rates of postpartum blood glucose screening reported in papers published between 2008 and 2010 (11 included studies, with 32,240 women) found that 34-73% of women with past GDM completed postpartum screening. The length of follow-up within studies varied, and all but one study had follow-up of greater than 12 weeks. The screening rate did not increase over time across studies. There were a few limitations of this review according to an AMSTAR assessment, although generally the review appeared methodologically sound. Limitations included: no list of excluded studies, unclear methods for assessment of quality in included studies, no exploration of publication bias and no exploration of conflict of interest. Most of the included studies were conducted in healthcare settings with interventions in place to try to increase the

rates of postpartum follow-up, so it is possible that other settings may have had lower rates (Tovar et al 2011).

Two Australian studies were included in the Tovar et al (2011) systematic review. These Australian studies reported relatively higher rates of postpartum screening compared with other included studies (Morrison et al 2009;Swan et al 2010). Morrison and colleagues (Morrison et al 2009) examined rates of postpartum glucose testing using a cross-sectional survey of 1372 women diagnosed with GDM between 2003 and 2005, sampled from the National Diabetes Services Scheme Database (NDSS) in Australia; they found that any form of postnatal glucose testing was completed at 6 to 8 weeks by 60.9% of respondents and within 6 months by 73.2% of respondents. However, only 27.3% had had an OGTT at 6-8 weeks. Furthermore, the overall survey response rate was 36%, and it is possible that there was response bias towards potentially more motivated women. In fact, the authors noted that, if none of the 14,521 women from the NDSS dataset who did not take part in the survey had returned for post-partum screening, the rate of any type of postnatal testing would actually be as low as 6.3%. This highlights the difficulty of accurately estimating the actual rate of uptake of testing, as well as the clear need to increase the rates of oral glucose tolerance testing in Australia. In the other Australian study, postal questionnaires were sent to 210 women in regional Victoria, and 61% of respondents had a postpartum OGTT (response rate was 40%) (Swan et al 2010). If all non-respondents did not have an OGTT, the actual rate may have been as low as 24%. Interestingly, this study found that a higher proportion of women living in smaller rural areas had a postpartum OGTT compared with women living in large rural areas (82.5% vs. 48.9%, p < 0.05). The authors speculated that this may have been due to the women in smaller rural areas seeing a family doctor who was more likely to be aware of the family diabetes risk.

Another systematic review, published in 2013, aimed to identify approaches associated with higher postpartum glucose testing rates. The authors categorised studies into either (1) "Usual Care" (data obtained via retrospective chart or database review) or (2) "Active Care"

(reporting results of studies with a proactive plan enacted to improve testing rates, such as calling or posting reminders to women or their physicians). This systematic review, with 54 included studies, found that the use of proactive contact programs increased postpartum testing rates (Carson et al 2013). With regards to AMSTAR assessment, the research aims and inclusion criteria were clearly stated. There was duplicated study selection and data extraction, and a comprehensive literature search (four databases) was performed. However, the review did not systematically assess the quality of the included studies, or assess the likelihood and implications of positive publication bias. There was also a lack of information on the types of active care within the included studies.

# Patient/provider adherence to screening according to clinical practice guidelines: a theoretical framework

Examination of what is known about patient and clinician adherence to screening from previous research may form a useful theoretical framework upon which to base further exploration of the factors influencing screening for T2DM in the postpartum period.

One of the best studied areas of screening in relation to patient and physician adherence is in cancer detection (Subramanian et al 2004; Limmer et al 2014). Cancer screening adherence has the potential for reduction in morbidity and mortality with early detection, in conjunction with varied rates of uptake (Subramanian et al 2004; Limmer et al 2014).

Physician recommendation of the screening test has been frequently reported as an important predictor of patient adherence with breast and colorectal cancer screening recommendations, and less frequently with cervical cancer screening (Subramanian et al 2004; Limmer et al 2014). Physician recommendation of postpartum T2DM screening for women who have had GDM may be an important positive predictor of screening.

Other factors identified as positively impacting on patient adherence to cancer screening include secure financial status, having health insurance, being married, tertiary education attainment, family history of cancer or presence of known risk factors for cancer, good psychological health and positive perception of health care system interactions and treatment efficacy (Subramanian et al 2004; Limmer et al 2014).

According to research into cancer screening, several factors influence the likelihood that physicians will recommend screening tests, including their past experience of patient adherence, patient demographics, their perception of test efficacy, their training, knowledge and agreement with screening guidelines and barriers to provision of care such as lack of equipment or poor reimbursement (Subramanian et al 2004).

#### What is known to influence postpartum screening rates?

It is likely that there would be some overlap in the above factors influencing attendance for postpartum diabetes screening, particularly physician recommendation, financial and sociodemographic factors and the presence of known risk factors, such as family history.

#### The influence of sociodemographic factors on postpartum follow-up

There have been somewhat conflicting results as to if and how age, ethnicity, marital status, parity and education influence the likelihood of postpartum glucose testing testing (Keely 2012;Tovar et al 2011). However, there is some evidence that predictors of higher postpartum screening attendance may be older age, nulliparity, higher income, higher education, Asian ethnicity, being a recipient of prenatal care, and antenatal treatment with insulin (Tovar et al 2011).

#### Women's views relating to postpartum follow-up of GDM

Many studies have sought insight into barriers and enablers to postpartum follow-up from the perspective of women, through interviews, focus groups and surveys. Given the number of studies and the importance of the topic, synthesis of their results is likely to be very valuable for improving postpartum care; such a systematic review has been conducted as part of this

thesis. Some examples of the included studies follow. Keely and colleagues (2010) asked 36 women why they did not complete oral glucose tolerance testing in a follow-up survey relating to their randomised controlled trial of postnatal postal reminders (Clark et al 2009). The most common reason for non-completion was time pressure (20 women, 55%), followed by lost lab requisition forms (7 women, 19%). In another study (n = 88), the most commonly reported barrier was test inconvenience (36%), followed by not knowing a follow-up test was required (28%) (Sterne et al 2011). In the same study, women said that reminders (39%) and raised awareness (31%) were likely to act as facilitators to screening. Bennett and colleagues (n = 22) found recent birth experiences, baby's health issues, adjustment to a new baby, emotional stress, postpartum mood symptoms and lack of time for self-care all contributed to low attendance for postpartum care. Other barriers included negative experiences with medical services such as long waiting periods and receiving care from multiple providers. Facilitators included availability of child care at the time of the appointment, a positive connection with clinical and office staff as well as a desire to have an appointment for either a check-up, to discuss family planning and/or obtain clearance for return to work. At a local level, barriers to completion of postpartum glucose testing identified in previous studies, were assessed in a follow-up survey for the women who participated in the DIAMIND Study (Heatley et al 2013).

Perception of increased risk of T2DM may be an important factor influencing motivation to undertake postpartum glucose testing (Keely 2012). Research has shown that some women with GDM do not perceive themselves as being at high risk for T2DM (Malcolm et al 2009;Morrison et al 2010). Therefore, changing perception of risk and providing further education to women regarding postpartum follow-up may result in increased attendance. A recent retrospective study examined the efficacy of an educational intervention for women to improve postpartum follow-up (Stasenko et al 2011). This study provided education about the importance of postpartum follow-up, delivered by a trained diabetes educator (registered nurse). The women were informed about their increased risk of T2DM, and provided with a

21

handout containing follow-up information and instructions on how to obtain an OGTT prior to their postpartum visit. This education resulted in an increase in postpartum blood glucose testing (with either fasting blood glucose or an OGTT) from 33% in the 2002 to 2006 cohort before the educational intervention began, to 53% in 2007 to 2009 (p <0.001) when the intervention was in place.

#### Healthcare factors: Providers and the health care system

Numerous studies have examined the factors influencing provision of healthcare for women with GDM through surveying or interviewing clinicians (Baker et al 2009;Divakar and Manyonda 2011;Doran and Davis 2010;Gabbe et al 1998;Hunsberger et al 2012;Keely et al 2010;Ko et al 2013;Oza-Frank 2012;Persson et al 2011;Pierce et al 2011;Power et al 2013;Stuebe et al 2010;Weaver 2004). It is important that the results of these studies are applied to improving postpartum care, and given that the results have not been previously synthesised, a systematic review of these studies was conducted as part of this thesis (Van Ryswyk et al 2014; paper contained in this thesis).

Communication of the diagnosis of GDM to those responsible for postnatal follow-up care is likely to be an important factor influencing rates of follow-up testing. A study assessed documentation of GDM history in the electronic problem list of the Brigham Women's Hospital (BWH) in Boston Massachusetts (Stuebe et al 2010). In this study, primary care providers and obstetric care providers were asked if they used the electronic problem list for communication; 93% of primary care providers (127 surveyed) and 81% of obstetric care providers (80 surveyed) said they did. The study team found that 772 women with GDM gave birth at BWH. Of those, only 58% (450/772) had any entry on their problem list, with 18% (141/772) having the correct code of "diabetes of pregnancy" and 8% (65/772) having a free text entry indicating the diagnosis of GDM. This low rate of documentation of the diagnosis of GDM may be more widespread than just this individual hospital.

Another study examined rates of referrals for follow-up, rather than documentation of the diagnosis of GDM. This study was a retrospective chart review (n = 2617) at a major academic hospital in Philadelphia, USA. The study found that only 20% of women with GDM had documented orders from an obstetrician for glucose screening tests. When referrals to primary care physicians was included as a method of providing this follow-up, only 33% of women had either documented orders for postpartum screening from obstetricians or referral to a primary care provider for postpartum follow-up of GDM (Almario et al 2008). Thus, rates of postpartum follow-up testing are affected by referrals from care providers, not just by whether or not women choose to return for recommended testing.

#### Which postpartum glucose test should be used?

Current Australian guidelines recommend use of an OGTT at 6-8 weeks postpartum for T2DM screening (Nankervis et al 2013). However, OGTTs have several disadvantages in terms of ease of completion: the test itself takes at least two hours to complete, it requires overnight fasting, and the glucose drink is sometimes poorly tolerated and can lead to nausea and vomiting (d'Emden 2014). Other, shorter tests may be more acceptable to women and more likely to increase test completion rates. Recent research has focussed on the possibility of using HbA1c, which requires no fasting and no waiting time, specifically for postpartum T2DM screening (Benaiges et al 2013;Garcia de Guadiana Romualdo et al 2012;Gingras et al 2013;Katreddy et al 2013;Kim et al 2011;Megia et al 2012;Noctor et al 2013;Picon et al 2012).

Guidelines in New Zealand have been updated to recommend postpartum screening using HbA1c at three months after birth, with repeated annual testing if results are normal (Ministry of Health 2014). Australasian Diabetes in Pregnancy Society guidelines still recommend use of OGTTs for early postpartum screening (6-12 weeks) and for women contemplating another pregnancy, but recommend HbA1c as an option for subsequent screening (Nankervis et al 2014).

23

Since 2002, the South Australian GDM Recall Register sent registered South Australian women a reminder to undertake glucose testing at 15 months. This recall register has now been replaced by the National Gestational Diabetes Register, which sends registered women postal reminders at 12-16 weeks after their expected due date (The National Diabetes Services Scheme 2012), and the SA Perinatal Practice Guidelines recommend that all women with GDM (and Medicare-coverage) should be offered the opportunity to join this register (South Australian Perinatal Practice Guidelines 2012). Chittleborough et al (2009) studied the effectiveness of the South Australian GDM Recall Register and found that of the 429 women who had been sent their first reminder letter (at 15 months), 46.4% had returned the update form and 56.3% of those who returned the update form had undertaken a glucose test for diabetes. If none of the 429 women who did not return the update form had the test, the actual rate of testing may be as low as 26%. It is also worth noting the high variability of recruitment rates to the South Australian register over the years; a high of 71.6% of eligible women were recruited in 2003, and a low of 26.6% of those eligible were recruited in 2006 (Table 4). The authors of the Chittleborough (2009) paper speculated that reasons for this variability in recruitment rates may have been due to time constraints during appointments, change in staff, and differences in staff efforts to recruit.

Year	Total number of women	Number of women on	Register recruitment
	with GDM	register	
2002	38	26	68.4%
2003	74	53	71.6%
2004	62	31	50.0%
2005	123	48	39.9%
2006	268	72	26.9%
2007	309	199	64.4%

 Table 4: Recruitment to the GDM Recall Register at participating hospitals (Chittleborough et al 2010)

In 2014, a Cochrane systematic review was published on the topic of reminder systems for women with previous GDM to increase uptake of testing for T2DM or impaired glucose tolerance (Middleton and Crowther 2014). This Cochrane review, with last search in April 2013, found that there was only one trial available for assessment (Clark et al 2009), and that, whilst this trial showed increased OGTT completion in the reminder arms, the trial had unclear risk of bias in most assessment criteria (including allocation concealment, blinding of women, attrition bias, selective reporting, and baseline imbalance) and was therefore of low quality evidence for efficacy.

This trial (Clark et al 2009) included 256 women, regardless of age, who attended the High Risk Obstetrical Unit (Ottawa Hospital, Ontario, Canada) between 29 August 2002 and 31 March 2005, for treatment of GDM, who provided written informed consent. The trial had four arms: (1) reminder sent to primary care physician and woman, (2) reminder sent to primary care physician only, (3) reminder sent to woman only, and (4) usual care, no reminder. The postal reminders were sent at three months after birth. The primary outcome of the trial was the proportion of women who underwent OGTT within 1 year after birth, and the results from 223 women were analysed (33 were excluded due to loss to follow-up). Oral glucose tolerance test rates were significantly increased in all three reminder groups compared with the no reminder group (Table 5).

Reminder Group	Proportion screened with OGTT
Physician and woman	49 of 81 women (60.5%)
Physician only	16 of 31 women (51.6%)
Woman only	42 of 76 women (55.3%)
No reminder	5 of 35 women (14.3%)

Table 5: Primary outcome of RCT of postal reminders for postpartum OGTTs (Clark et al 2009)Reminder GroupProportion screened with OGTT

The same research team then implemented a women only reminder system into routine care (Shea et al 2011), and conducted a non-randomised study examining the effects of implementation of the reminder system at two different healthcare sites in their region compared with no reminder at a third healthcare site. The primary outcome was the proportion of women who were screened for T2DM with an OGTT within 6 months of

delivery. The study included all women who had antenatal GDM education classes at one of the three healthcare sites included in the study (the Ottawa Hospital General Campus, the Queensway Carleton Hospital and The Ottawa Hospital Civic Campus). The GDM education classes gave information on the development of T2DM after birth, as well as other aspects of GDM. Women were excluded from the study if they did not have Ontario Health Insurance Plan (OHIP) coverage (provincial health insurance that is universally available without copayment, covers all physician visits, medical care and diagnostic testing). Whilst the study had ethical approval, it was not clear whether informed consent was gained from the included women.

Three months after delivery, women in one study site were mailed a reminder that included information on the importance of diabetes screening and a laboratory requisition for an OGTT at a non-hospital based laboratory; women at a second study site were either sent a letter with a laboratory requisition (for OGTT) or phoned, or both. Women at a third site were not sent a postal reminder or given a reminder phone call. None of the sites provided routine postpartum GDM-specific follow-up visits.

The results of their implementation of a women only reminder system into routine care, with regards to postpartum oral glucose tolerance testing within 6 months after birth, were lower than those found in the previous randomised controlled trial of postal reminders (Clark et al 2009). 23.3% (21/90) at the first Shea study site completed OGTTs, compared with 55.3% in the women only arm of the Clark RCT, while 36.4% (20/55) at the second Shea site completed OGTTs. The rate of OGTT completion at this site was higher than at the non-reminder site (13.7%, 16/117, p = 0.03), as was the combined reminder site rate (p = 0.01) (Table 6).

Table 6: Results of implementation of postal women's reminder Shea et al 2011)

Site	Reminder Type	OGTT completion	
		n	%
The Ottawa Hospital- General campus	Mailed information and lab requisition for OGTT at 3 months postpartum	21/90	23.3
Queensway Carleton Hospital	Mailed information and lab requisition for OGTT/phoned/both at 3 months postpartum	20/55	36.4
The Ottawa Hospital - Civic campus	None	16/117	13.7

A small number of non-RCT studies have also examined reminder systems for postpartum blood glucose testing, with positive results. For example, a Finnish prospective observational study found that a special call or reminder from their central hospital increased the rates of postpartum OGTT uptake (OR 13.4 [4.6-38.1], P < 0.001) (Korpi-Hyovalti et al 2012). Also, a Canadian study (Toronto), examined the efficacy of a physician checklist for increasing postpartum screening in women who have had GDM. They found by retrospective chart review that the checklist was associated with a 3 fold increase in odds of being screened postpartum, and an almost 4 fold increase in postpartum follow-up visits (OR 2.99, 95% CI 1.84–4.85 and OR 3.71, 95% CI 2.26–6.11) (Lega et al 2012).

Given the apparent efficacy of postal based reminders, it follows that SMS-text reminders may be an effective way to increase postpartum oral glucose testing rates. In Australia, in 2011, there were at least 28 million mobile phone subscriptions (i.e. 6 million more subscriptions than people) (Budde and McNamara 2011). With the very high rate of mobile phone usage in, it is likely that most women of reproductive age have access to a mobile phone. In addition, text messages may be less easy to lose than mail reminders, more likely to be read than emails, and less invasive, time consuming and expensive than voice calls.

A Cochrane review found that there is low to moderate quality evidence that mobile phone text messaging reminders increase attendance at healthcare appointments compared with no reminders, or postal reminders (Gurol-Urganci et al 2013). This review included randomised controlled trials assessing mobile phone reminders for healthcare appointments, in which it was possible to assess effects of mobile phone messaging independent of other technologies or interventions. A total of eight randomised controlled trials, involving 6615 participants, were included. The authors concluded that there was moderate quality evidence (7 studies, n =5841) of efficacy of text message reminders, compared with no reminders, for increasing healthcare appointment attendance (RR 1.14, 95% CI 1.03 to 1.26), as well as moderate quality evidence that mobile text reminders had a similar impact to phone call reminders (three studies, 2509 participants, RR 0.99, 95% CI 0.95 to 1.02). There was low quality evidence from just one study, with n = 291 participants, that mobile text message reminders combined with postal reminders alone (RR 1.10, 95% CI 1.02 to 1.19). This potential to increase rates of attendance may translate into increased rates of women's attendance for oral glucose tolerance testing in the postpartum period for women who have had gestational diabetes. The authors recommended that further high quality randomised trials of mobile phone messaging reminders are needed

#### Summary

In summary, GDM is an increasingly common health condition that indicates a substantially increased lifetime risk for development of T2DM, and the postpartum period is an important time for early T2DM detection and intervention to prevent T2DM. Screening in the postpartum period for T2DM is recommended, although frequently women do not complete or are not offered screening. It is important to investigate both the causes of the low postpartum screening rate, as well as methods of improving follow-up. Mobile (SMS) reminder systems have been found to be effective for increasing attendance at various healthcare appointments in some studies, but the quality of the existing evidence is low-moderate, and there is only low quality evidence from one randomised controlled trial that postal reminders increase completion of oral glucose tolerance testing in women who have had GDM. A combination of decreased use of the postal system and very high use of mobile

28

phones indicates that mobile text reminders may be preferable and possibly more effective for increasing postpartum glucose test attendance. The research covered in the following chapters further explores reasons for sub-optimal postpartum follow-up, with systematic synthesis of the results of qualitative/survey studies on the topic from the perspective of both clinicians and women. The efficacy of an SMS text reminder system specifically for women who have experienced GDM to increase postpartum glucose test completion is also explored via a randomised controlled trial, as are the barriers and facilitators to postpartum testing from the perspective of the women in the study.

#### Thesis Purpose

The following chapters examine several related aims, all of which seek to answer the research questions: (1) why is postpartum follow-up of gestational diabetes, particularly screening for T2DM sub-optimal, and (2) what can be done to improve this situation?

Aim 1: To examine the factors that influence postpartum follow-up of women with gestational diabetes, from the perspective of clinicians.

Aim 2: To examine the factors that influence postpartum healthcare seeking for women who have had gestational diabetes, from the perspective of women.

*Aim 3: To test whether a SMS reminder system may help improve the rate of attendance for postpartum oral glucose tolerance testing – The DIAMIND Trial.* 

Aim 4: To examine other factors that may have influenced completion of glucose testing in the DIAMIND Trial.

Aim 5: To ascertain the views of women in the DIAMIND Trial relating to barriers and facilitators to postpartum OGTT completion

# Statement of Authorship

Title of Paper	Clinician views and knowledge regarding healthcare provision in the postpartum period
Publication Status	$\odot$ Published, O Accepted for Publication, O Submitted for Publication, O Publication style
Publication Details	Van Ryswyk E, Middleton P, Hague W, Crowther C. Clinician views and knowledge regarding healthcare provision in the postpartum period for women with recent gestational diabetes: a systematic review of qualitative/survey studies. Diabetes Research and Clinical Practice. 106(3): 401-11.DOI: 10.1016/j.diabres.2014.09.001

#### **Author Contributions**

By signing the Statement of Authorship, each author certifies that their stated contribution to the publication is accurate and that permission is granted for the publication to be included in the candidate's thesis.

Name of Principal Author (Candidate)	Emer Van Rysywk	
Contribution to the Paper	Led review conception and design, prepared and registered protocol, developed search strategy with research librarian and conducted the review. Wrote drafts of the manuscript for publication.	
Signature	Date 20/2/15	

Name of Co-Author	Philippa Middleton	
Contribution to the Paper	Contributed to review conception and design, supervised and participated in conduct of the review and edited drafts of the manuscript for publication.	
Signature	Date 19215	

Name of Co-Author	William Hague
Contribution to the Paper	Contributed to review design, and edited drafts of the manuscript for publication.
Signature	Date 26 2 15

Name of Co-Author	Caroline Crowther
Contribution to the Paper	Contributed to review design, supervised conduct of the review, and edited drafts of the manuscript for publication.
Signature	Date KIZIS

Van Ryswyk, E., Middleton, P., Hague, W. & Crowther, C. (2014) Clinician views and knowledge regarding healthcare provision in the postpartum period for women with recent gestational diabetes: A systematic review of qualitative/survey studies. *Diabetes Research and Clinical Practice, v. 106 (3), pp. 401-411* 

NOTE:

This publication is included on pages 31 - 41 in the print copy of the thesis held in the University of Adelaide Library.

It is also available online to authorised users at:

http://dx.doi.org/10.1016/j.diabres.2014.09.001

## Women's views and knowledge regarding healthcare seeking for gestational diabetes in the postpartum period: a systematic review of qualitative/survey studies

#### Abstract

Aim: To identify factors influencing postpartum healthcare seeking, from the perspective of women who have experienced gestational diabetes mellitus (GDM).

Methods: Systematic review that searched PubMed, Web of Science, EMBASE and CINAHL on 27<sup>th</sup> February 2013. Qualitative studies and surveys, with women as participants, which reported pre-specified outcomes, including barriers and facilitators to healthcare seeking for GDM after birth, were included. For each included study, two authors independently assessed quality and undertook thematic synthesis.

Results: 42 studies were included, with data from 7949 women. Studies were conducted in the United States, Australia, Europe, the United Kingdom, Canada, Brazil, Vietnam and Tonga. For some women, being diagnosed with GDM was a concerning or upsetting experience. Antenatal care for GDM was sometimes a very positive experience, whilst for others it was a more difficult or confusing experience. There was a need for more specific information about GDM to be available around the time of diagnosis. Holistic, non-judgemental and positive care was preferred.

While women were often knowledgeable about their type 2 diabetes risk and about ways of preventing it, they faced multiple barriers to undertaking such preventive behaviours. Lifestyle change support was needed. Women would like healthcare providers to take a more pro-active approach to postpartum care.

Conclusions: Improved GDM education and materials coupled with the provision of holistic, non-judgemental and pro-active care from diagnosis of the condition through all stages of

postpartum follow-up may increase healthcare seeking by women with GDM in the postpartum period, which may facilitate both prevention and early diagnosis of type 2 diabetes.

#### Introduction

Gestational diabetes mellitus (GDM) is strongly associated with future risk of type 2 diabetes mellitus (T2DM) (Bellamy et al 2009; Feig et al 2008). The risk of development of T2DM in the first 10 years for women after GDM has been reported to be between 20% and 50%, and the lifetime risk may be greater than 70% (Feig et al 2008;Kim et al 2002;Lee et al 2007). International and national organisations recommend postpartum screening following GDM (American Diabetes Association 2013; Canadian Diabetes Association Clinical Practice Guidelines Expert Committee 2013; IADPSG Consensus Panel 2010; Nankervis et al 2014; National Institute for Health and Care Excellence 2008). Previous research shows that many women who have experienced GDM do not complete, or are not offered, adequate follow up for their GDM, particularly postpartum diabetes screening (Dietz et al 2008;McGovern et al 2014;Tovar et al 2011). This deficiency in postpartum follow-up for women with GDM has been recognised by clinicians and researchers, with calls for improvements in the United States (Gabbe et al 2012) and Australia, with researchers highlighting fragmentation and inconsistencies in postpartum care and the need for a professional group to take responsibility for care of women who have had GDM (Wilkinson et al 2014).

Considerable research has been conducted into factors influencing healthcare seeking and preventive behaviour from the perspective of women with a history of GDM, although only a fraction of this research has been systematically synthesised. A previous meta-synthesis of 16 qualitative studies relating to women's experiences of GDM, their perception of their risk of T2DM and their views on T2DM prevention reported on several themes. These included an emotional response to their GDM diagnosis, loss of a "normal" pregnancy experience, the importance of "personal control" in relation to GDM management, a motivation to have the best possible glucose control or GDM management in order to do their best for the baby, as well as the importance of adequate information relating to GDM and healthcare support (Parsons et al 2014).

The objective of this systematic review is to identify factors that influence postpartum healthcare seeking for women who have experienced GDM through synthesis of results from qualitative and survey studies.

#### Methods

The protocol for this review is registered with the international systematic review register PROSPERO 2013:CRD42013003599.

#### Search strategy

PubMed, EMBASE, Web of Science, and CINAHL were searched from inception to 27<sup>th</sup> February 2013, with no date or language restrictions. The full search strategy is available in Appendix 1. Reference lists of included studies were searched for additional studies. Titles and abstracts were examined for eligibility by one author. The full text of studies that appeared to meet inclusion criteria were assessed independently by two authors, with a final list of included studies resulting from discussion and consensus between two authors.

#### Selection of studies

Studies were selected for inclusion in the review if they met the following criteria: (a) qualitative study, or survey; (b) participants were women who have experienced GDM; and (c) reported at least one of the following pre-specified outcomes:

#### Primary outcomes

- (1) Barriers to postpartum healthcare seeking, for women who have experienced GDM
- (2) Facilitators to postpartum healthcare seeking, for women who have experienced GDM

#### Secondary outcomes

- (1) Knowledge of risk of type 2 diabetes
- (2) Perception of risk of type 2 diabetes
- (3) Attitudes towards postpartum follow up of GDM
- (4) Attitudes towards postpartum blood glucose testing

- (5) Attitudes towards reminders for follow up or blood glucose testing
- (6) Knowledge of complications of type 2 diabetes (for mothers and/or babies)
- (7) Knowledge of how to prevent type 2 diabetes in the future

#### Data extraction

At least two authors (EVR, ES, PM) independently extracted data on the characteristics, prespecified outcomes, and funding sources of the included studies.

#### Quality assessment

Quality assessment was carried out using the Critical Appraisal Skills Programme (CASP) checklist for qualitative studies (Critical Appraisal Skills Programme). Any differences between assessments were resolved through discussion amongst authors.

#### Data synthesis

The qualitative synthesis method chosen was thematic synthesis, due to its successful application in previous systematic reviews of qualitative studies (Schumann et al 2012;Thomas and Harden 2008;Tong et al 2009). The authors of this paper also used thematic synthesis in a companion review of healthcare provision by clinicians (Van Ryswyk et al 2014). The independently extracted outcome data from each included study were compared and finalised by two authors. Potential themes and theme categories were then identified, discussed and modified until a final decision regarding each was reached.

#### Results

#### Study selection process

This systematic review is reported in accordance with the PRISMA statement (Moher et al 2009). 1249 abstracts and full-text articles were identified. After removal of duplicates and exclusion on the basis of title and abstract alone, two authors assessed the records of 68 studies (with 81 abstract/full-text records) for inclusion. Twenty seven studies were excluded; for the list of excluded studies, with reasons, refer to Appendix 2. The abstract for one
additional study was identified during assessment, and was also included (Wylie et al 2011). As a result of this process, 42 studies (with 55 records) were included (Figure 1). For the full list of included study references, including abstracts, please refer to Appendix 3.

## Study characteristics

A summary of the characteristics of the included studies is provided in Table 1. The 42 included studies had data from 7949 women, who spoke languages including English, Hindi, Bengali, Vietnamese, Cantonese, Mandarin, Filipino, Spanish, Danish, Arabic, Swedish, French, German and Portuguese. The included studies were conducted via interviews (18 studies), a combination of interviews and surveys (four studies), interviews and focus groups (three studies), survey-only (15 studies), or focus groups (two studies). Twelve of the studies were conducted in the United States, ten in Australia, nine in Europe/UK, seven in Canada, two in Brazil, one in Vietnam and one in Tonga. The results from one group of six studies by the same author (Hjelm) were grouped together as one study, as the studies appeared to have included results from the same women across the six studies, albeit with different comparison groups; this was done to reduce inappropriate amplification of views from the same women.

#### Quality assessment

All included studies were appraised using the 10-item CASP checklist for qualitative studies (Critical Appraisal Skills Programme) (Table 2). Two studies met all of the criteria for quality assessment (Bieda 2009;Doran and Davis 2010). In all other studies, it was unclear whether one or more quality assessment criteria were met, and in 11 studies less than half of the quality assessment criteria were clearly met. Seven studies were not in full text format, limiting the information available for quality assessment; six studies were abstracts only (Bell et al 2011;Hoy-Rosas and Lancaster 2011;Remsberg 2012;Segall-Gutierrez et al 2011;Wylie et al 2012), whilst one study was published in commentary format (Keely et al 2012).

Most included studies had clearly stated aims (n = 36/42), and warranted a qualitative approach (40/42). Whilst most studies had well described and justified research design (n =

47

26/42), several did not meet this criteria, due to being in abstract or commentary format, with insufficient details of the research design reported, or through having the design reported but not adequately justified by the authors. Just over half of the included studies had adequately described recruitment strategies, but nineteen studies had insufficient details of their strategies reported to make a clear assessment of their method. Fourteen study records did not include details on their data collection methods; six were in abstract format. The other eight studies either did not discuss data saturation, did not provide the actual questions asked in their data collection process or did not explain why their data collection methods were the most appropriate for their research question.

It was evident in just nine of the included studies that the authors had critically examined their own potential bias and role (relationship criteria) in formulation of the research questions, methods and their response to events in the study. Just over half of the included studies (n = 22) had clearly taken ethical issues into consideration, although in the remaining studies there were either insufficient details of how the research was explained to participants for the reader to assess whether ethical standards were maintained, or the researchers did not discuss ethical issues such as informed consent.

Data analysis was sufficiently rigorous in most of the included studies (n = 26). Sixteen studies had either insufficient information reported to allow assessment of the data analysis process and results, or it was unclear whether the authors had critically examined their own potential bias in the selection of data for presentation.

Most studies (n = 26) had explicit findings, with adequate discussion of the credibility of results, and how the results related to the original research question. Studies mainly had well described value and research implications (n = 29).

## Data synthesis

Seven theme categories were identified: experiences relating to antenatal GDM diagnosis and management; maternal role; perception and knowledge of risk of T2DM; how healthcare is provided; barriers to healthcare access; T2DM detection postpartum; and T2DM prevention.

# Experiences of GDM diagnosis and management

Four studies reported that for some women, being diagnosed with GDM was a concerning or upsetting experience, particularly where the women were concerned about the possibility of future complications of type 2 diabetes:

"When I heard the diagnosis, it was scary. I panicked (Lawson and Rajaram 1994)."

"I was really depressed for weeks. I cried and cried. It was like: Oh no, I don't have

diabetes! I mean, the whole thing was very scary (Lawson and Rajaram 1994)."

Women who had previously experienced medically complicated pregnancies had more subtle reactions to the diagnosis (Lawson and Rajaram 1994).

A diagnosis of GDM sometimes resulted in a change in or loss of identity as a healthy person with a "perfect" or "normal" pregnancy (Lawson and Rajaram 1994), and due to a pressure to conform to this ideal, women sometimes felt the need to hide their diagnosis:

"I took my blood at LaMaze classes (pregnancy and birth classes), and it was interesting, because I wanted to hide it, even though other gestational diabetics could have been in the class (Lawson and Rajaram 1994)."

Women had varied experiences of their antenatal GDM care. Some women had positive experiences of their care, with increased confidence relating to the lifestyle changes they had been able to make, and their now good understanding of GDM and its management. Others experienced antenatal management of GDM as confusing or difficult:

"They told me I'm supposed to be on this crazy diet where there's only lettuce and vegetables and they put me on this diet that made my blood go up to like 20. And I told them, *I'm not going to go on that diet because the diet you guys put me on—the way you're talking, you guys are killing me quicker than [when] I was eating my fries!* (Neufeld 2011)"

Some women would have preferred for pregnancy and GDM be care to be better integrated, or more holistic, rather than segmented:

"...both the pregnancy and the diabetes need to be focused, so that it is not just the diabetes when you come here [diabetes clinic] and that it is not just the pregnancy when you come to the midwife...that it feels like co-operation...and...there are a lot of specialists involved (Hjelm et al 2007)."

"...not that they [clinicians] mean to be, but they are very clinical and removed and don't seem to understand, you are attempting to deliver a healthy baby and manage and plan the rest of your life; whereas they are with you for 20 minutes and are attempting to determine why you decided to eat your toast with jam! (Nicklas et al 2011)"

A small number of women expressed a preference for more positive, constructive care:

"I found myself very annoyed at the clinicians because I always felt they were a tinge judgmental about the GDM and had a lot of assumptions. Any meeting with them started with, 'Now you have to change your lifestyle,' and I thought: *You don't know what my lifestyle is, so how do you know what is bad or what needs to change?* I may already know and be changing what I need to in order to be healthy. I am not a child (Nicklas et al 2011)."

"GDM comes with a whole team of professionals, but what is missing is a place to bounce off how to move forward [after delivery] with life ideas in a positive surrounding, as opposed to looking back at mistakes (Nicklas et al 2011)."

The information available for GDM was an aspect of care that women viewed as being deficient. Women sought information on their GDM from multiple non-medical sources, and were often able to find much information on T2DM, but little information specific for GDM.

Furthermore, the education and information was sometimes in a different language or needed to be adapted to the woman's own culture.

# <u>Maternal role</u>

The maternal role sometimes presented a barrier to care seeking, with children's needs taking priority over care seeking:

"I don't really spend too much time thinking about [my risk for diabetes]. Because I've got two kids under four and I am too busy to spend my day worrying about [my health] (Razee et al 2010)."

Similarly, time constraints, lack of sleep or fatigue, maternal attachment and adjustment to a new baby all played a role. Some women found it difficult to attend postpartum care due to breastfeeding, and may have been more receptive to health messages relating to diabetes after their babies have been weaned.

# Perception and knowledge of risk of T2DM

More often than not, women had good or frequent knowledge of their risk of T2DM:

"I hope it will be over after the delivery, but it is latent, there is an increased risk of diabetes (Hjelm et al 2005)."

In eight studies, women had poorer knowledge of their risk:

"The doctor did discuss some of the risks for the baby, but they didn't discuss what my risks would be of getting diabetes later (Collier et al 2011)."

Women reportedly used differing primary sources of information relating to T2DM risk, with some women getting most of their information from health professionals, and others from their families. A small number of studies reported a need to increase public awareness of the link between GDM and T2DM.

# Perceptions and emotional associations

There was variation in perception of future T2DM risk, with women's personal risk perceptions being increased with longer time since diagnosis, family history of T2DM, and other known risk factors for T2DM. For some, the development of chronic T2DM was "dreaded" (Lawson and Rajaram 1994), especially the prospect of needing lifelong insulin, or of having complications that may have been witnessed in others:

"I don't know if I could handle giving myself shots forever. I could not handle it. Someone would have to give me shots all my life (Lawson and Rajaram 1994)." "I guess one thing . . . is fear. People coming back at 6 weeks, they know they have to do that blood sugar check-up, and they probably fear that . . . they're gonna have to continue doing blood sugars, and, continue with their diets like they were during the pregnancy (Bennett et al 2011)."

The hope that diabetes may no longer be present after birth made it easier to tolerate the condition and its strict management. Some women saw GDM as an indicator of a need to take positive steps to try to prevent T2DM, whilst others felt that they lacked control over the development of T2DM, or perceived that it was inevitable.

# How healthcare is provided

Some women felt a sense of postpartum abandonment, after intensive antenatal management of their GDM:

"You're kind of left on your own. I don't know, it's kind of hard to elaborate on something like that, like I feel I was abandoned. Like, OK, what am I supposed to do? (Evans et al 2010)"

Similarly, some women expressed that there was a need for clinicians to take a more proactive approach to postpartum care, including recommending the diabetes screening test, providing advice on self-blood glucose monitoring, making follow-up appointments for monitoring and support of T2DM prevention. Reminders to facilitate attendance for care were considered to be helpful. Positive connections with clinical staff were important facilitators to care attendance, and women valued trustworthiness, professional competence and cultural sensitivity in their interactions with healthcare providers.

# Barriers to healthcare access

Socioeconomically disadvantaged women often faced barriers to accessing care, such as lack of health insurance, out of pocket costs. Women also experienced long waiting times for appointments, and a clash between their own working hours and the opening hours of healthcare clinics. For some, there was not enough time with providers, and women sometimes did not have their healthcare questions answered:

"A lot of times those doctors will see so many Medicaid patients. They don't want to answer your questions (Collier et al 2011)."

"A doctor only has you for so many minutes. They don't explain in detail how you should take care of the gestational diabetes (Collier et al 2011)."

Not having a specific primary care provider was also a barrier for some.

# T2DM detection postpartum

Postpartum testing for T2DM was an emotional experience, with feelings ranging from fear and sadness at the prospect of a diagnosis of T2DM to great happiness at receiving a normal result. Numerous reasons for non-completion of T2DM screening in the postpartum period were apparent. Some, but not all women understood the need for follow-up T2DM screening. Other women had continued to perform self-blood glucose monitoring after birth and therefore did not see a need to attend for glucose screening; others forget the test, became pregnant again or lost their laboratory request form. Some women experienced postpartum mood symptoms or had no desire to take the test. The oral glucose tolerance test was a barrier for some women, with a more convenient, pleasant test being desired. Women were more likely to attend for T2DM screening if they had a desire to know their blood glucose status, and also when their test was able to be coupled with other reasons for postpartum clinic attendance, such as clearance for their return to work, birth-related health checks and family planning.

### T2DM prevention

Education about T2DM prevention was evenly distributed between the family, health care providers and the media. Women often had a positive attitude towards T2DM prevention, along with knowledge of how to prevent T2DM, with many having knowledge of the role of diet, exercise and weight control in diabetes prevention:

"It's good to learn about it, otherwise the way I was going, definitely I would have diabetes 2 (sic). I didn't know about it but now I can control myself and . . .Yeah, because the way I'm not having anything, I was having heaps of sugars every day (Carolan et al 2012)."

Motivators for lifestyle change included high risk perception and fear of future GDM and T2DM. Many had a high awareness of the need to take steps to prevent T2DM. However, numerous barriers existed to achieving and maintaining these preventive changes. Barriers to healthy eating included cost, lack of time, being unsure which foods to eat, a focus on food in social situations, lack of motivation and personal or cultural food preferences. For some women, there was a lack of knowledge of how to prevent T2DM. Given the numerous barriers to lifestyle change, lifestyle change support both in the short and long term was important, although the types of support required varied between women e.g. social support was needed by some, whilst financial or professional support was required by others.

# Discussion

# Summary of the main results

The diagnosis of GDM was for some women, concerning or upsetting experience. Following the diagnosis, women sought information from multiple sources and found that there was a

lack of specific information on GDM compared with other forms of diabetes. Some women had difficult or confusing experiences relating to antenatal management of their GDM, whilst other women had more positive experiences of antenatal care.

Knowledge of the risk of T2DM was common, although in some studies women had poorer knowledge and could benefit from better education. There was wide variation in perception of future T2DM risk, with some women believing development of T2DM to be inevitable. Women had increased perception of risk of T2DM with increased time from their GDM diagnosis, family history of T2DM and other known risk factors for GDM. Women worried about receiving a diagnosis of chronic diabetes, especially in relation to the prospect of lifelong insulin or the possibility of complications.

The maternal role played an important part in determining attendance for postpartum care, with children's needs often taking priority over care seeking. A need for clinicians to take a more pro-active approach to postpartum care was identified. This included recommending the diabetes screening test to women, providing advice on blood glucose self-monitoring after birth, and making follow-up appointments relating to T2DM prevention. Women also expressed positive views relating to reminders to facilitate care.

Knowledge of how to prevent T2DM, including the role of diet, exercise and weight control was common amongst women in a third of studies. In a smaller number of studies, women lacked knowledge relating to T2DM prevention, and could have benefited from better education. Women's views on T2DM prevention likely also played an important role in postpartum healthcare seeking, with many women having a positive attitude towards T2DM prevention. Motivators for lifestyle changes included high risk perception and fear of future GDM and T2DM. Although, the lifestyle changes required, particularly healthy eating, were often difficult to achieve and maintain on a long term basis, and women often described a need for lifestyle change support.

55

# Agreements and disagreements with other studies or reviews

Similar themes were found relating to barriers to postpartum diabetes screening as those described in another recent systematic review, which investigated "determinants" and barriers to GDM care from initial screening to postpartum follow-up, and which included studies from searching only one database, PubMed. Their barriers to attendance for postpartum care included time constraints and a focus on the baby's health (Nielsen et al 2014), although only two qualitative studies specific to postpartum diabetes screening were included in their systematic review (Bennett et al 2011;Keely et al 2010). The same systematic review (Nielsen et al 2014) included several studies relating to barriers to having a healthy lifestyle in the postpartum period, and there were similarities in the themes identified, particularly in relation to a need for lifestyle change support.

Our systematic review had some similar findings to the meta-synthesis of 16 qualitative studies by Parsons et al 2014 (Parsons et al 2014) examining women's experiences of GDM. One similar finding included a significant emotional response to the GDM diagnosis; the emotions in response to the GDM diagnosis described in the qualitative meta-synthesis were shock, denial, sadness, fear and difficulty coming to terms with a condition for which there were no symptoms (Parsons et al 2014). Other similar findings were a preference for holistic and non-judgemental care; a lack of specific information on GDM; a wide variety of perception of type 2 diabetes risk; and several similar barriers to adopting a healthier lifestyle.

## Strengths and limitations

The search strategy for this systematic review did not use date or language restrictions, encompassed four databases, and there were well defined pre-specified outcomes. This systematic review included several additional studies relevant to postpartum care for women with GDM than a comparable synthesis of qualitative studies (Parsons et al 2014). While there was variation in locations and settings of the included studies, there were common themes across studies. For each included study, two authors independently extracted data, and appraised study quality using a CASP checklist, which is a commonly used tool for

56

quality assessment (Critical Appraisal Skills Programme). More than half of the included studies were of good to excellent quality, meeting seven or more of the ten quality assessment criteria. Nine studies (>20%) were average to moderate quality, meeting five to six criteria. Just ten studies met less than half of the criteria (six of which were in abstract or commentary format and contributed little to the overall findings of the review, and two of which were in languages other than English).

# Conclusion

### Implications for practice

Care provision from diagnosis of GDM through to postpartum follow-up may influence the likelihood of women's seeking and participation in care after they give birth. It is important to recognise the potential emotional impact of the GDM diagnosis, with associated loss of the "normal" or "perfect" pregnancy experience, and for adequate support such as counselling to be available. Having culturally and language-appropriate written information on GDM is similarly important. Following the diagnosis, some women indicated a preference for provision of non-judgemental, holistic and positively focussed care.

In the postpartum period, care could be improved by healthcare providers taking a more proactive approach, including recommending the diabetes screening test, providing advice on blood glucose self-monitoring, making follow-up appointments for monitoring and support of T2DM prevention, and providing reminders to facilitate attendance. Other critical factors are improving the education given to women regarding their risk of type 2 diabetes and how to prevent the condition.

# Implications for research

Further research is required on how to best manage the emotional impacts of GDM such that women find being diagnosed with GDM to be a more positive experience. Production and assessment of educational materials is important. Methods for provision of more holistic care require further exploration. Systematic methods of improving follow-up care and support for women who have experienced GDM into diabetes prevention is necessary.

# Acknowledgements

We are grateful to Michael Draper, who provided guidance and assistance with development of the search strategy for this review, to Dr Caroline Schneeberger, who kindly translated an included study, and also to Dr Mojgan Vatani, who assisted with assessment of a study in Swedish.

### Figure 1: A flow chart of study selection, with excluded studies reasons included in the flow chart



#	Study ID	Location	Type of study	Participants (women with GDM)	Data collection period (year)
	Australia				
-	Bandyopadhyay 2011	Melbourne	Interviews	17 South Asian (Hindi, Bengali or English speaking)	Antenatal, Postpartum (2009)
5	Carolan 2013	Melbourne	Interviews/focus group	15 English speaking	Antenatal (NS)
3	Graco 2009	Victoria	Interviews	10 English speaking women	Postpartum (2004)
4	Doran 2008	New South Wales	Survey and interviews	38 Survey (subset of 8 Interviewed) English speaking	Antenatal, Postpartum (unclear)
5	Morrison 2010	National	Survey	1176 English speaking	Postpartum (2003-5)
9	Carolan 2010	Melbourne	Survey	143 Vietnamese, Indian, Chinese, Filipino or Caucasian	Antenatal (2007)
				(interpreters utilised)	
7	Razee 2010	Sydney	Interviews	57 (20 Arabic, 20 Cantonese/Mandarin & 17 English speaking)	Postpartum (2006)
8	Smith 2005/Zehl 2008	Sydney	Survey	244 English speaking	Postpartum (unclear)
6	Sterne 2011	Queensland	Survey	88 English speaking	Postpartum (2006-7)
10	Swan 2007	Victoria	Survey	53 women (either in English, or with interpreters)	Postpartum (2004-5)
	United States				
11	Jones 2012	"A South-western	Interviews and survey	22 self-identified American Indian	Postpartum (unclear)
		State"			
12	Hoy-Rosas 2011	New York	Interviews	3 (1 Latina, 1 Asian, 1 African-American)	Postpartum (unclear)
13	Bennett 2011	Baltimore	Interviews	22 English speaking	Postpartum (unclear 2008-9)
14	Collier 2011	Atlanta	Focus group	54 English or Spanish speaking	Postpartum (2004-5)
15	Downs 2006	Pennsylvania	Survey	28 English speaking	Postpartum (2004)
16	Bieda 2009	Michigan	Interviews	25 African-American (English speaking)	Antenatal (unclear)
17	Kim 2007 Risk	Michigan	Survey	217 English speaking	Postpartum (unclear)

18	Kim 2007 Racial	National	Survey	4718 English speaking	Antenatal, Postpartum (2001-3)
19	Lawson 1994	Kentucky	Interviews	17 English speaking	Antenatal (unclear)
20	Nicklas 2011	Boston	Interviews/focus group	25 English speaking	Postpartum (2009)
21	Remsberg 2012	Ohio	Survey	73 (language not specified - abstract only)	Postpartum (unclear)
22	Segall-Gutierrez 2011	California	Survey	75 English or Spanish speaking	Postpartum (unclear)
	Canada				
23	Evans 2005	Ontario	Interviews	12 English speaking	Antenatal, Postpartum (unclear)
24	Evans 2010	Ontario	Interviews and survey	13 English speaking	Postpartum (unclear)
25	Feig 1998	Toronto	Survey	65 English speaking	Postpartum (1996)
26	Gaudreau 2012	Quebec	Interviews	7 Algonquin (Native American)	Postpartum (2006)
27	Keely 2010	Ottawa	Survey	140 English speaking	Postpartum (2002-5)
28	Keely 2012	Ottawa	Survey	51 English speaking	Antenatal (2010-11)
29	Neufeld 2011	Winnipeg	Interview	29 Aboriginal (Canadian)	Antenatal, Postpartum (2006-7)
	Europe/UK				
30	Stage 2004	Denmark	Survey	121 Danish speaking (Caucasian)	Postpartum (unclear)
31	Hjelm 2005-12	Lund, Sweden	Interview	14 born in the Middle East	Antenatal, Postpartum (2000-1)
	(all Hjelm studies i.e.			10 African born	
	2005, 2007, 2008,			• 13 Swedish born (Clinic A – Hospital Endocrinology)	
	2009, 2012, 2011/12b)			10 Swedish born (Clinic B – Specialist Maternity Clinic)	
32	Lindmark 2010	Sweden	Interview	10 Swedish speaking	Postpartum
33	Persson 2010	Sweden	Interview	10 Swedish speaking	Antenatal (1998, 2000, 2006)
34	Clarke 2012	Paris, France	Survey	124 French speaking	Postpartum (2008-9)
35	Trutnovsky 2012	Austria (not AUS)	Interview and survey	45 German speaking	Antenatal (unclear)
36	Bell 2011	UK (NS)	Interviews	31 (abstract only)	NS (unclear)

37	Wylie 2011	UK (Plymouth)	Interviews	12 (abstract only)	Antenatal (unclear)	
38	Wylie 2012	UK (Plymouth)	Interviews	12 (abstract only)	Unclear	
	South America					
39	Saloman 2004	Minas Gerais Brazil	Interviews	9 Portuguese speaking	Antenatal (2003)	
40	Soares 2006	Minas Gerais Brazil	Interviews/focus group	56 interviewed and 7 in focus group (Portuguese speaking)	Unclear (translated from Dorthomese)	
					1 oluguese)	
	Asia					
41	Doran 2010	Nuku'alofa, Tonga	Interviews	11 English speaking	Postpartum (2006)	
42	Hirst 2012	Ho Chi Minh,	Focus group	34 Vietnamese women	Antenatal (2010-11)	
		Vietnam				
14	busiding and mot spacified	USIV P				

Abbreviations are: not specified (NS).

Assessmen	
Onality	( , , , , , , , , , , , , , , , , , , ,
ASP	
C	Ì

	Tonga Vietnam	Doran Hirst	2010 2012	•	•	•	•	•	•	•	•	•	•
	Austria	Trutnovsky	2012	•	÷	÷	<del></del>	<del></del>	<del></del>	•	<del></del>	<mark></mark>	•
		Hjelm	2005-12	•	•	•	<mark>:</mark>	•	•	•	•	•	•
rveys)	Sweden	Lindmark	2010	•	•	٠	•	•	<mark>∾</mark>	•	•	•	÷
groups (±su		Persson	2010	•	•	•	~	•	•	2	•	•	•
Studies containing either interviews or focus	Kingdom	Wylie	2012	•	•	~	<mark>~</mark>	~	<mark>~</mark>	<mark>~</mark>	<mark>~</mark>	<mark>~</mark>	<mark>~</mark>
	nited Kingdo	Wylie	2011	•	•	~	<u>~</u>	~	<del>~</del>	~	<mark>~</mark>	~	~
	Unit	Bell	2011	•	•	~	~	~	~	~	~	~	~
		Razee	2010	•	•	•	~	~	~	~	•	•	•
	ralia	Doran	2008	•	•	•	•	•	<mark>~</mark>	•	•	<mark>~</mark>	•
		Graco	2009	•	•	•	<u>~</u>	•	~	~	•	•	•
	Austr	Carolan	2013	•	•	•	•	•	~	•	•	•	•
		Bandyopadhyay	2011	+	÷	÷	+	÷	<b>~</b> •	•	+	+	÷
	Location	CASP		Clear aims	Qualitative	Design	Recruitment	Collection	Relationship	Ethics	Analysis	Findings	Value

Location				USA						Canada		Bra	zil
CASP	Bennett	Collier	Bieda	Lawson	Nicklas	Hoy-Rosas	Jones	Evans	Evans	Gaudrea	Neufeld	Saloman	Soares
	2011	2011	2009	1994	2011	2011	2012	2005	2010	2012	2011	2004	2006
Clear aims	•	•	•	••	•	~	•	••	•	•	•	•	<mark>.</mark>
Qualitative	•	•	•	~	•	•	•	•	•	•	•	•	<mark>.</mark>
Design	•	•	•	••	•	~	•	•	•	•	•	•	<mark>.</mark>
Recruitment	•	•	•	•	~	~	•	•	<mark>~·</mark>	~	•	<u>~</u>	<mark>.</mark>
Collection	•	•	•	•	•	~	•	•	<mark>~·</mark>	•	•	•	<mark>.</mark>
Relationship	•	~	•	••	~	~	<mark>~</mark>	•	<mark>~·</mark>	~	~	<u>~</u>	<mark>.</mark>
Ethics	è	+	•	<mark>.</mark>	•	~	•	<mark>~</mark>	<mark>~</mark>	•	•	•	Ł
Analysis	<mark>.</mark>	•	•	•	•	~	•	•	<mark>~·</mark>	•	~	<u>~</u>	<mark>.</mark>
Findings	è	+	•	•	•	~	•	Ł	<mark>~</mark>	•	~	<mark>2</mark>	Ł
Value	+	+	•	•	•	~	•	2	•	•	•	<mark>2</mark>	2

	Denmark	Stage	2004	<u>~</u>	•	<u>~</u>	·	•	••	••	•	•	•	
	France	Clarke	2012	~	•	•	~	•	•	•	~		•	
		Keely	2012	•	•	~	•	~	<mark>~</mark>	~	•	•	<mark>.</mark>	
	Canada	Keely	2010	•	•	•	<mark>.</mark>	<mark>~</mark>	<mark>.</mark>	<mark>~</mark>	•	+	2	
		Feig	1998	•	•	•	•	•	<mark>.</mark>	<mark>~</mark>	<mark>.</mark>	+	•	ez.
		Segall-G.	2011	•	•	•	•	•	<mark>.</mark>	•	<mark>.</mark>	•	<mark>?</mark>	= Segall-Gutieri
urvey type	United States	Remsberg	2012	•	•	~	~	~	<mark>~</mark>	~	<del>。</del>	Ł	•	. Segall- $G = S\epsilon$
Studies which are exclusively		Kim	2007b	•	•	~	~	~	~	~	•	•	•	ria not met
		Downs	2006	•	•	<mark>∼•</mark>	~	•	~	•	•	+	•	= Crite
		Kim	2007a	•	•	<mark>~</mark>	<mark>~</mark>	~	<mark>.</mark>	~	•	•	•	as met.
		Swan	2007	•	•	•	•	•	•	•	•	+	ŧ	criterion was
		Carolan	2010	•	•	•	•	•	•	••		2	•	r whether ci
	ustralia	Smith	2005	•	•	•	÷	•	<mark>~</mark>	•	•	÷	•	= Unclea
	Α	Sterne	2011	•	•	~	•	•	~	•	•	÷	•	rion. 🔁
		Morrison	2010	•	•	•	•	•	~	•	•	•	•	ssessment crite.
	Location	CASP		Clear aims	Qualitative	Design	Recruitment	Collection	Relationship	Ethics	Analysis	Findings	Value	= Met quality as

themes
ъ
an
\$
ie.
1
5
3
Ē
3
e
8
Ð
д
C.
le
þ
đ
F

eriences relating to antenatal GDM diagnosis management	Australia	United States	Canada	South America	Europe/UK	Asia
t or confusing experience relating to antenatal ment of GDM		Bennett 2011 Lawson 1994	Neufeld 2011		Hjelm 2005-12 Trutnovsky 2012 Persson 2010	
nformation specific for GDM wanted		Collier 2011		Saloman 2004	Hjelm 2005-12 Lindmark 2010	Hirst 2012
n or sadness about the GDM diagnosis and its titions (may be worse in women with greater edge of T2DM complications, and lessened in with previously complicated pregnancies)		Bieda 2009 Lawson 1994		Saloman 2004	Lindmark 2010	
e experiences of antenatal GDM care sed confidence relating to lifestyle changes, nderstanding of GDM and its management)					Hjelm 2005-12 Lindmark 2010 Trustnovsky 2012 Persson 2010	
education must be adapted to culture and ge		Collier 2011	Gaudreau 2012		Hjelm 2005-12	
n obtained information on GDM from multiple edical sources		Collier 2011 Lawson 1994				Hirst 2012
e in identity from being a healthy person and normal pregnancy experience		Lawson 1994			Persson 2010	
nce for care to be holistic, non-judgemental cussed on positive change		Nicklas 2011			Hjelm 2005-12	
notivation for GDM treatment to do the best for vy					Trutnovsky 2012	

Maternal role	Australia	United States	Canada	South America	Europe/UK	Asia
Women still undertaking most house work, leaving less time for self-preventive care	Razee 2010					
Time constraints		Bieda 2009 Nicklas 2011	Keely 2010			
Child's needs taking priority over care seeking	Razee 2010	Nicklas 2011 Bennett 2011		Soares 2006	Hjelm 2005-12 Bell 2011	
Breast-feeding, with more receptivity to health messages after weaning		Bennett 2011	Keely 2010		Bell 2011	
Lack of sleep or fatigue		Bennett 2011 Nicklas 2011			Bell 2011	
Maternal attachment and adjustment to the new baby	Bell 2011 Bennett 2011					
Perception and knowledge of risk of T2DM	Australia	United States	Canada	South America	Europe/UK	Asia
Good or frequent knowledge of risk of T2DM	Bandy. 2011 Graco 2009 Morrison 2010 Razee 2010	Bieda 2009 Jones 2012 Kim 2007a Remsberg 2012 Segall-Guteriezz 2011	Feig 1998	Saloman 2004	Hjelm 2005-12 Lindmark 2010 Wylie 2012 Persson 2010	
Poor knowledge of risk of T2DM	Carolan 2013	Bieda 2009 Collier 2011	Feig 1998		Hjelm 2005-12 Lindmark 2010 Clarke 2012 Wylie 2012 Trutnovsky 2012	
Need to increase public awareness of link between GDM and T2DM	Doran 2008					Doran 2010

							Asia					
			Hjelm 2005-12	Persson 2010	Persson 2010 Hjelm 2005-12	Stage 2004	Europe/UK	Wylie 2011	Hjelm 2005-12			Hjelm 2005-12
	Saloman 2004,	Saloman 2004		Saloman 2004	Saloman 2004		South America					
		Neufeld 2011					Canada	Keely 2010	Evans 2010	Keely 2010 Keely 2012		
Bieda 2009	Bieda 2009 Kim 2007a	Bennett 2011 Bieda 2009 Lawson 1994	Kim 2007a	Bieda 2009	Lawson 1994	Kim 2007a Nicklas 2011	United States	Nicklas 2011				Bennett 2011
	Morrison 2010 Sterne 2011			Doran 2008		Morrison 2010	Australia	Doran 2008 Sterne 2011	Graco 2009	Doran 2008 Razee 20104 Sterne 2011	о Wall 2007	
Differing primary sources of information on future risk - Health professionals for some, family for others	Women had increased perception of risk of T2DM with increased time from diagnosis, family history of T2DM and other known risk factors for GDM	Dread of chronicity of diabetes, including the prospect of lifelong insulin, and distress relating to witnessing complications of T2DM in others	GDM perceived as an indicator of the need to take steps to prevent T2DM	Perception of inevitability or no control over development of T2DM (sometimes due to religious beliefs)	The hope that diabetes may no longer be present after birth made it easier to tolerate the condition and its strict management	Much variation in perception of future diabetes risk	How healthcare is provided	Need for clinicians to take a more pro-active approach to postpartum care (recommending the	diabetes screening test, providing advice on blood glucose self-monitoring, making follow-up appointments for T2DM prevention)	Positive views on reminders to facilitate attendance for care		Positive connection with clinical staff important

. competence and arded			Gaudreau 2012		Hjelm 2005-12	
	Graco 2009		Evans 2010		Lindmark 2010	
	Australia	United States	Canada	South America	Europe/UK	Asia
		Kim 2007b Segall-Gutierezz 2011 Collier 2011				
		Bennett 2011 Collier 2011			Hjelm 2005-12	
	Doran 2008	Collier 2011				Hirst 2012
					Bell 2011 Hjelm 2005-12	
		Kim 2007b	Keely 2012			
A	ustralia	United States	Canada	South America	Europe/UK	Asia
		Bennett 2011	Keely 2010		Hjelm 2005-12	
•1	Sterne 2011		Keely 2010		Hjelm 2005-12	
<b>H 3</b> 1	3andy. 2011 Sterne 2011				Hjelm 2005-12	

				Asia									Doran 2010				
				Europe/UK	Hjelm 2005-12	Lindmark 2010	Hielm 2005-12	Lindmark 2010				Hjelm 2005-12		Bell 2011	Hjelm 2005-12 Lindmark 2010	Bell 2011	Lindmark 2010
Soares 2006				South America								Saloman 2004				Soares 2006	
Keely 2010	Keely 2010			Canada	Evans 2005	Evans 2010 Neufeld 2011	Gandrean 2012	Neufeld 2011	Evans 2005	Evans 2010			Evans 2010	Evans 2010			
		Bennett 2011		United States	Lawson 1994	Jones 2012	Iones 2012	Kim 2007a	Lawson 1994	Hoy-Rosas 2011	, Remsberg 2012	Bieda 2009 Kim 2007a	Hoy-Rosas 2011 Nicklas 2010	Jones 2012	Nicklas 2011	Nicklas 2010	Bieda 2009
Sterne 2011	Sterne 2011		Carolan 2010	Australia	Carolan 2013	Doran 2008 Swan 2007	Carolan 2013	Doran 2008	Swan 2007			Swan 2007		Swan 2007		Smith 2005	
Other reasons for non-attendance at T2DM screening (forgetting the test, new pregnancy, mood symptoms, lost laboratory test form, no desire to take the test)	OGTT difficult to complete (more convenient, pleasant test as a facilitator)	Reasons for postpartum clinic attendance other than T2DM risk (clearance for work, birth-related issues, family planning)	Follow up care well understood by some	T2DM prevention	Positive attitude towards T2DM prevention		Common knowledge of how to nrevent T2DM	including the role of diet, exercise, weight control				High awareness of need to take steps to prevent T2DM	Motivators for lifestyle change included high risk perception and fear of future GDM and T2DM	Difficult to achieve or maintain healthy lifestyle (esp	pregnancy motivated benaviour change)	Lifestyle change support (long-term) is critical but	types of desired support varies (e.g. social or financial or professional support)

t, focus on food in Jones 2012 Lindmark 2010 ation, personal food
tices) Nicklas 2011

Bandy. = Bandyopadhyay.

# Statement of Authorship

Title of Paper	The DIAMIND study: postpartum SMS reminders to women who have had gestational diabetes to test for type 2 diabetes: a randomised controlled trial - study protocol
Publication Status	Published, O Accepted for Publication, O Submitted for Publication, O Publication style
Publication Details	Heatley E, Middleton P, Hague W and Crowther C, The DIAMIND study: postpartum SMS reminders to women who have had gestational diabetes mellitus to test for type 2 diabetes: a randomised controlled trial – study protocol, BMC Pregnancy and Childbirth 2013, 13:92.

# **Author Contributions**

By signing the Statement of Authorship, each author certifies that their stated contribution to the publication is accurate and that permission is granted for the publication to be included in the candidate's thesis.

Name of Principal Author (Candidate)	Emer Van Rysywk (maiden surname = Heatley)				
Contribution to the Paper	Contributed to study design. Wrote initial and subsequent drafts of the manuscript for publication.				
Signature	Date 20/2/15				

Name of Co-Author	Philippa Middleton				
Contribution to the Paper	Conceived study idea, wrote and led initial study design and protocol for ethics submission and edited the manuscript for publication.				
Signature	Date 1925				

Name of Co-Author	William Hague
Contribution to the Paper	Participated in study design and edited the manuscript for publication.
Signature	Date 26 falir

Name of Co-Author	Caroline Crowther					
Contribution to the Paper	Contributed to study design and trial protocol, participated in ethics submission and edited the manuscript for publication.					
Signature	Date 15/2/15					

# **STUDY PROTOCOL**



**Open Access** 

# The DIAMIND study: postpartum SMS reminders to women who have had gestational diabetes mellitus to test for type 2 diabetes: a randomised controlled trial – study protocol

Emer Heatley<sup>1\*</sup>, Philippa Middleton<sup>1</sup>, William Hague<sup>1</sup> and Caroline Crowther<sup>1,2</sup>

# Abstract

**Background:** Postpartum follow up of women who have been found to have gestational diabetes during pregnancy is essential because of the strong association of gestational diabetes with subsequent type 2 diabetes. Postal reminders have been shown to increase significantly attendance for oral glucose tolerance testing postpartum. It is possible that a short message service (text) reminder system may also be effective. This trial aims to assess whether a text message reminder system for women who have experienced gestational diabetes in their index pregnancy will increase attendance for oral glucose tolerance testing within six months after birth.

**Methods/Design:** *Design*: Single centre (Women's and Children's Hospital, South Australia), parallel group randomised controlled trial.

Inclusion criteria: Women diagnosed with gestational diabetes in their index pregnancy (oral glucose tolerance test with fasting glucose  $\geq$  5.5 mmol/L and/or two hour glucose  $\geq$  7.8 mmol/L), with access to a mobile phone, whose capillary blood glucose profile measurements prior to postnatal discharge are all normal (fasting glucose < 6.0 mmol/L, postprandial glucoses < 8.0 mmol/L).

*Exclusion criteria*: Pregestational diabetes mellitus, triplet/higher order multiple birth or stillbirth in the index pregnancy, requirement for interpreter.

*Trial entry and randomisation*: Allocation to intervention will be undertaken using a telephone randomisation service (computer-generated random number sequence generation, with balanced variable blocks, and stratification by insulin requirement).

*Study groups*: Women in the intervention group will receive a text reminder to attend for an oral glucose tolerance test at 6 weeks postpartum, with further reminders at 3 months and 6 months if they do not respond to indicate test completion. Women in the control group will receive a single text message reminder at 6 months postpartum. *Blinding*: Baseline data collection will be undertaken blinded. Blinding of participants and blinded collection of primary outcome data will not be possible for this study.

*Primary study outcome*: Attendance for the oral glucose tolerance test within 6 months postpartum.

Sample size: 276 subjects will be required to show an 18% absolute increase in the rate of attendance ( $\alpha$ =0.05 two tailed,  $\beta$ =80%, 5% loss to follow up) from 37% to 55% in the intervention group.

(Continued on next page)

Institute, The University of Adelaide, 72 King William Road North, Adelaide, SA 5006, Australia

Full list of author information is available at the end of the article



© 2013 Heatley et al.; licensee BioMed Central Ltd. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

<sup>\*</sup> Correspondence: emer.heatley@adelaide.edu.au

<sup>&</sup>lt;sup>1</sup>Australian Research Centre for Health of Women and Babies, Robinson

#### (Continued from previous page)

**Discussion:** Given the heightened risk of impaired glucose tolerance and type 2 diabetes in women who have had gestational diabetes, ensuring the highest possible rate of attendance for postpartum glucose tolerance testing, so that early diagnosis and intervention can occur, is important. A text message reminder system may prove to be an effective method for achieving improved attendance for such testing. This randomised controlled trial will assess whether such a system will increase rates of attendance for postpartum oral glucose tolerance testing in women who have experienced gestational diabetes.

Trial Registration: Australian New Zealand Clinical Trials Registry - ACTRN12612000621819

**Keywords:** Gestational diabetes mellitus, Reminder system, SMS text reminder, Randomised controlled trial, Postpartum care, Oral glucose tolerance test, Type 2 diabetes mellitus

#### Background

#### Introduction

Gestational diabetes mellitus (GDM) has been defined as carbohydrate intolerance of variable severity with onset or first recognition during pregnancy [1]. GDM affects 5% of pregnancies in Australia, and the prevalence of GDM is likely to rise with increases in maternal age and obesity [2]. Women who have had GDM are at higher risk for development of type 2 diabetes mellitus (DM) in the future compared with women who have had normal blood glucose values during pregnancy, as well as being at increased risk of GDM in future pregnancies [3-7]. Therefore, many clinical practice guidelines recommend screening for type 2 DM and impaired glucose tolerance in the postpartum period [8-13]. Such follow up is important, given the welldocumented risks to women and their babies resulting from type 2 DM and GDM [14], the availability of interventions for prevention of type 2 DM and recurrent GDM such as lifestyle changes, use of metformin, and encouragement of breastfeeding [15], and because direct medical costs increase greatly with progression from impaired glucose tolerance to type 2 DM with complications [16]. However, follow up rates for GDM tend to be sub-optimal and need to be improved [17]. Increasing the rate of follow up with an oral glucose tolerance test (OGTT) is important, given that it is the most sensitive test for detection of impaired glucose tolerance and type 2 DM within 6 months postpartum [18].

# Reminder systems for increasing postpartum follow up of GDM

Reminder systems have been used to improve healthcare with positive results, and some research studies have assessed the effect of reminders on increasing attendance for follow up of GDM. For example, implementation of a postpartum patient reminder system for women who had experienced GDM into routine care at two hospital sites in Canada (The Ottawa Hospital, General Campus and the Queensway Carleton Hospital) resulted in higher rates of completion of oral glucose tolerance testing by 6 months postpartum, with a rate of 28% (41/145) compared with 14% (16/117 women) in the site where reminders were not used (The Ottawa Hospital, Civic Campus) (p = 0.01) [19].

However, the rate of completion in both sites where reminders were used was considerably lower than the rates of OGTT completion observed in the randomised controlled trial (RCT) of postal reminders previously conducted by the same team at The Ottawa Hospital in Canada [20]. In the RCT, OGTT completion rates were 60% (49/81 women) in the physician and patient reminder group, 55% (42/76 women) in the patient-only reminder group, 52% (16/31) in the physician-only reminder group and 14% (5/35 women) in the group with no reminders sent (p < 0.05). It should be noted that OGTT completion was measured up until six months postpartum in the implementation study, compared with up to one year postpartum in the RCT.

A small number of non-randomised studies have examined reminder systems for postpartum blood glucose testing in women who have experienced GDM. A Finnish prospective observational study found that a phone call reminder by a nurse increased rates of postpartum oral glucose tolerance testing at one year after birth (odds ratio 13.4, 95% confidence interval 4.6–38.1) [21]. Another study examined the efficacy of a checklist as a physician reminder for increasing postpartum screening for type 2 DM in women who had GDM at the Endocrine Obstetrics Clinic of the Women's College Hospital in Toronto, Canada. In this study, retrospective chart review revealed that use of the reminder checklist for physicians was associated with a 3 fold increase in odds of a woman being screened with an OGTT, as measured at  $\geq$  6 months postpartum (odds ratio 2.99, 95% confidence interval 1.84-4.85) [22].

Following on from a previous study investigating trends in postpartum glucose test ordering by clinicians and completion by women (with recent GDM) that found increasing but suboptimal rates at Kaiser Permanente Northwest (KPNW, a large non-profit health organisation in western Oregon and Washington state) [23], another study at KPNW examined the efficacy of several interventions aimed at increasing the proportion of postpartum glucose test ordering by clinicians and test completion by women (including revising the nursing protocol for pregnant women with GDM, improving the electronic medical record system, educating clinical staff and providing additional reminders to women who did not complete the test within 3 months of delivery) [24]. Orders for postpartum glucose screening increased from 77% of (155/200) women in the pre-implementation period, to 89% (159/ 179) in the post-implementation period (p = 0.004), and completion of postpartum glucose screening increased from 60% (pre-implementation) to 72% (post-implementation) (hazard ratio, 1.37; 95% confidence interval, 1.07–1.75).

#### Postpartum reminders for GDM follow up in Australia

The South Australian Gestational Diabetes Recall Register was established in July 2002 [25]. This register sent a reminder letter at 15 months after birth to women who had experienced GDM to encourage oral glucose tolerance testing. A study examining the efficacy of the register found that, of the 429 women who had been sent their first reminder letter (at 15 months), 56% had undertaken a glucose test for diabetes (response rate 46%) [25]. There was considerable variation in the rate of recruitment of eligible women to the register, with a nadir of 27% in 2006, and a peak of 72% in 2003. The authors of this study speculated that reasons for this variability in recruitment rates may have included time constraints during appointments, change in staff, and differences in staff efforts to recruit to the register. The South Australian register has now been replaced by a national register, the National Gestational Diabetes Register [26]. Women resident in Australia and eligible for a Medicare card are recruited to this register at the time of registering for the National Diabetes Services Scheme (NDSS) following a diagnosis of GDM, and the register sends a follow-up reminder letter to such women to visit a general practitioner to arrange an OGTT at 12-16 weeks after the expected due date (provided to the register at the time of registration) and an information booklet called Life after Gestational Diabetes.

# Short message service (SMS) reminders – the reminders of the future?

In Australia in 2011, there were at least 28 million mobile phone subscriptions (i.e. 6 million more subscriptions than people) [27]. With the very high rate of mobile phone usage, it is likely that almost all women of reproductive age have access to mobile phones. Given the high rate of mobile phone use in Australia, and the low cost of SMS messages, a reminder system that utilises SMS technology might prove to be a cost-effective way of increasing the number of women in Australia with a recent history of GDM, who could then be prompted to undertake oral glucose tolerance testing in the postpartum period. Several studies in other areas of health care have indicated that SMS reminders can increase appointment attendance rates [28-31]. This potential to increase rates of attendance may translate into increased rates of oral glucose tolerance testing in the postpartum period for women who have had GDM.

#### Aims and objectives

The primary aim of this RCT is to determine whether an SMS reminder system will significantly increase attendance for oral glucose tolerance testing by 6 months postpartum in women who have recently experienced GDM.

#### Hypotheses

The primary hypothesis is that a SMS reminder system for women who have recently had GDM will increase the number of women who complete oral glucose tolerance testing by 6 months postpartum.

#### **Methods/Design**

#### Ethics statement

Ethics approval was obtained from the Women's and Children's Health Network Human Research Ethics Committee (REC2200/8/2015).

#### Study design

Single centre (Women's and Children's Hospital, South Australia), parallel group randomised controlled trial.

#### Inclusion criteria

Women diagnosed with GDM in their index pregnancy (positive 75 g OGTT with fasting glucose  $\geq$  5.5 mmol/L and/or two hour glucose  $\geq$  7.8 mmol/L), with access to a personal mobile phone, whose capillary blood glucose profile measurements prior to hospital discharge after giving birth are normal (fasting blood glucose < 6.0 mmol/L), who provide written, informed consent, will be included in the trial.

#### **Exclusion criteria**

Pregestational diabetes mellitus, triplet/higher order multiple birth or stillbirth in the index pregnancy or requirement for interpreter.

#### Trial entry

Women who are potentially eligible for the study will be approached in the postnatal ward, counselled and given the study information sheet. They will be entered into the trial if they give written consent and have normal blood glucose profile results prior to discharge from hospital.

#### Study groups and management

Eligible women will be randomised into one of two study groups: either the '6 week (intervention) group' or the '6 month (control) group'.

#### Randomisation

Randomisation will be carried out using a telephone randomisation service. The randomisation schedule has balanced variable blocks and has been prepared by an investigator not involved in recruitment or clinical care. Randomisation will be stratified by antenatal requirement for drug therapy to treat GDM.

#### **Treatment schedules**

#### Intervention (6 week reminder) group

Women in the intervention group will be sent a SMS reminder at six weeks after the birth of their baby: "*Hi* (*Participant Name*), *This is a reminder from the DIAMIND* study team for you to have your oral glucose tolerance test for type 2 diabetes. Please let us know when you have done the test, and what the results were by texting us on (study number) or emailing us at (study email address) - Thanks for participating and best wishes". If the participant responds to say she has completed the test, no further text reminders will be sent. If not, a further text reminder will be sent at three and six months (same message).

#### Control (6 month reminder) group

Women in the control group will receive no text reminders for the first 6 months of the study period. A single text message reminder (same text as for intervention group) will be sent to these women at 6 months postpartum (measured from date of birth of baby).

#### Primary study outcomes

Oral glucose tolerance test undertaken by 6 months postpartum

#### Secondary study outcomes

Fasting blood glucose test undertaken by 6 months postpartum

Glycated haemoglobin (HbA1c) test undertaken by 6 months postpartum

#### Data collection

#### Baseline data collection

Baseline data will be collected to assess the similarity between the two groups in terms of factors that may influence attendance for postpartum oral glucose tolerance testing. At trial entry, information will be collected on demographic characteristics of the participants as well as smoking history, current BMI at booking and previous pregnancy outcomes. With regards to GDM, women will be asked whether or not they were given the opportunity to join the National Diabetes Services Scheme (NDSS) and therefore the National Gestational Diabetes Register, whether they joined, and where they intend to have their postpartum OGTT completed. Data regarding control of GDM (dietary control only or requirement for metformin or insulin), diagnostic OGTT date and results, and complications at birth relating to GDM (requirement for induction, caesarean section, perineal injury, blood loss) will also be collected.

Baseline data relating to the health of the newborn(s) will be collected: singleton or twin, birth order (if twin), gestational age at birth, birth weight (grams), time of birth, gender, Apgar scores, mode of birth (normal vaginal birth, operative vaginal birth, caesarean section), nerve palsy, bone fracture, newborn hypoglycaemia (plasma glucose  $\leq 2.0 \text{ mmol/L}$ ), neonatal intensive care admission, respiratory distress syndrome, neonatal jaundice requiring phototherapy, and death prior to first discharge.

The following information will be collected at hospital discharge: breastfeeding status (given the link between breastfeeding and reduced risk of type 2 DM [32,33], as well as the possible influence that breastfeeding may have on the mother's ability to attend for oral glucose tolerance testing [34]), mention of GDM in the problem list of the discharge summary and whether or not follow up oral glucose tolerance testing was recommended in the discharge summary.

#### Outcome data collection: 6 months postpartum

All women in the study will be asked to complete a questionnaire at 6 months after the birth of the baby either by post or by email (using Survey Monkey), depending on their expressed preference at trial entry, to ascertain whether an OGTT was undertaken within the first 6 months, or whether a fasting blood glucose or glycated haemoglobin (HbA1c) test was used instead. The questionnaire also asks for the date and results of these tests, where known. The date and results of the OGTT will be confirmed using the hospital clinical information system, or by contact with the participant's general practitioner, where necessary. The questionnaire also examines women's attitudes towards the OGTT, as well as reasons for not being able to undertake the test, if applicable. Where questionnaires are not returned within 2 weeks, participants will be contacted by telephone, and then again in another 2 weeks if the questionnaires are still not received by the study team. During this telephone contact, participants will be asked whether or not they have completed an OGTT within the six months since they gave birth.

#### Sample size

The baseline rate of OGTT uptake used in the sample size calculation for the proposed trial is 37%; this rate is at the lower end of the range in the recent review by Tovar and colleagues (2011) [17]. Using a figure from the lower end of this range is a sound estimate, given that the health centres in the study had reminder systems in place, and that some results came from surveys with moderately low

response rates [17]. Data from the South Australian Gestational Diabetes Mellitus Recall Register indicates that the actual rate may be somewhere between 26 and 56 percent [25].

The Stata version 10.0 sample size calculator has been used to calculate the target sample size.

The figures entered were:

- baseline uptake of postnatal OGTT 37%
- projected 18% absolute improvement to 55% (48% relative increase)
- power 80%
- significance (two-tailed) 5%

This resulted in a calculation that the study will require 262 women (131 in each arm). With a predicted up to 5% loss to follow up, 276 women will be required.

#### Analyses and reporting of results

Baseline characteristics of all randomised women will be compared descriptively between the study groups. Outcome comparisons will be made according to the treatment allocation at randomisation on an 'intention to treat' basis. Categorical variables will be reported as risk ratios with corresponding 95% confidence intervals. Continuous outcomes will be reported as mean (and standard deviation) for normally distributed results, or median (interquartile range) for results which are not normally distributed. All model assumptions will be assessed. Statistical significance will be defined at the 0.05 level using a two-sided comparative test.

#### Discussion

This randomised controlled trial of a SMS reminder system to improve the rate of attendance for follow up oral glucose tolerance testing in women who have experienced GDM is important given the high, and rising, rate of GDM, and the strong association between GDM and subsequent type 2 DM worldwide. In Australia, this is a particularly timely trial, given the recent establishment of the National Gestational Diabetes Register, the increasingly ubiquitous use of mobile phones, and the decreasing use of postal services ("snail mail") for communication. Such a reminder system may prove to be a cost-effective measure to reduce future rates of type 2 DM in Australian women, and is therefore of public health as well as obstetric importance. Such a system has the potential to be implemented locally or on a wider scale to improve the health of all women who have experienced GDM.

#### Abbreviations

DM: Diabetes mellitus; GDM: Gestational diabetes mellitus; HbA1c: Haemoglobin A1c; OGTT: Oral glucose tolerance test;

RCT: Randomised controlled trial; SMS: Short message service.

#### **Competing interests**

The author(s) declare that they have no competing interests.

#### Authors' contributions

EH, PM, WH and CAC are all members of the DIAMIND Study Group. The primary investigator of the DIAMIND Study (EH) prepared the initial draft of the DIAMIND protocol. All members of the DIAMIND study team participated in the design of the study. The DIAMIND Study Group participated in the protocol development, commented on drafts of the protocol, and have read and approved the final draft of the protocol. All authors read and approved the final manuscript.

#### Acknowledgements

This study is supported by a 2 year health services research grant from the HCF Health and Medical Research Foundation.

#### Author details

<sup>1</sup>Australian Research Centre for Health of Women and Babies, Robinson Institute, The University of Adelaide, 72 King William Road North, Adelaide, SA 5006, Australia. <sup>2</sup>The Liggins Institute The University of Auckland, Private Bag 92019, Victoria Street West, Auckland 1142, New Zealand.

#### Received: 28 March 2013 Accepted: 8 April 2013 Published: 12 April 2013

#### References

- Hoffman L, Nolan C, Wilson JD, Oats JJ, Simmons D: Gestational diabetes mellitus-management guidelines. The Australasian diabetes in pregnancy society. *Med J Aust* 1998, 169(2):93–97.
- 2. AIHW: Diabetes in pregnancy: its impact on Australian women and their babies. Australian Institute of Health and Welfare: Canberra; 2010.
- Kim C, Newton KM, Knopp RH: Gestational diabetes and the incidence of type 2 diabetes: a systematic review. Diabetes Care 2002, 25(10):1862–1868.
- Kim C, Berger DK, Chamany S: Recurrence of gestational diabetes mellitus: a systematic review. Diabetes Care 2007, 30(5):1314–1319.
- Lee AJ, Hiscock RJ, Wein P, Walker SP, Permezel M: Gestational diabetes mellitus: clinical predictors and long-term risk of developing type 2 diabetes: a retrospective cohort study using survival analysis. *Diabetes Care* 2007, 30(4):878–883.
- Bottalico JN: Recurrent gestational diabetes: risk factors, diagnosis, management, and implications. Semin Perinatol 2007, 31(3):176–184.
- Bellamy L, Casas JP, Hingorani AD, Williams D: Type 2 diabetes mellitus after gestational diabetes: a systematic review and meta-analysis. *Lancet* 2009, 373(9677):1773–1779.
- Meltzer S, Leiter L, Daneman D, Gerstein HC, Lau D, Ludwig S, Yale JF, Zinman B, Lillie D: Clinical practice guidelines for the management of diabetes in Canada. Canadian Diabetes Association. CMAJ 1998, 159 (Suppl 8):1–29.
- Simmons DS, Walters BN, Wein P, Cheung NW: Guidelines for the management of gestational diabetes mellitus revisited. *Med J Aust* 2002, 176(7):352.
- South Australian Perinatal Practice Guidelines Chapter 65 Diabetes mellitus and abnormal glucose tolerance: [http://www.health.sa.gov.au/ppg/Default. aspx?PageContentID=2116&tabid=100]
- ACOG: ACOG Committee Opinion No. 435: postpartum screening for abnormal glucose tolerance in women who had gestational diabetes mellitus. Obstet Gynecol 2009, 113(6):1419–1421.
- Metzger BE, Buchanan TA, Coustan DR, de Leiva A, Dunger DB, Hadden DR, Hod M, Kitzmiller JL, Kjos SL, Oats JN, et al: Summary and recommendations of the Fifth International Workshop-Conference on Gestational Diabetes Mellitus. Diabetes Care 2007, 30(Suppl 2):S251–260.
- 13. Diabetes CareStandards of medical care in diabetes--2013. 2013, 36(1):11-66.
- 14. Ali S, Dornhorst A: Diabetes in pregnancy: health risks and management. *Postgrad Med J* 2011, **87**(1028):417–427.
- 15. Feig DS: Avoiding the slippery slope: preventing the development of diabetes in women with a history of gestational diabetes. *Diabetes Metab Res Rev* 2012, **28**(4):317–320. doi:10.1002/dmrr.2276.
- Herman WH: The economics of diabetes prevention. Med Clin North Am 2011, 95(2):373–384. viii.
- Tovar A, Chasan-Taber L, Eggleston E, Oken E: Postpartum screening for diabetes among women with a history of gestational diabetes mellitus. *Preventing chronic disease* 2011, 8(6):A124.

- Reinblatt SL, Morin L, Meltzer SJ: The importance of a postpartum 75 g oral glucose tolerance test in women with gestational diabetes. J Obstet Gynaecol Can 2006, 28(8):690–694.
- Shea AK, Shah BR, Clark HD, Malcolm J, Walker M, Karovitch A, Keely EJ: The effectiveness of implementing a reminder system into routine clinical practice: does it increase postpartum screening in women with gestational diabetes? *Chronic diseases in Canada* 2011, 31(2):58–64.
- Clark HD, Graham ID, Karovitch A, Keely EJ: Do postal reminders increase postpartum screening of diabetes mellitus in women with gestational diabetes mellitus? A randomized controlled trial. Am J Obstet Gynecol 2009, 200(6):631–637.
- Korpi-Hyovalti E, Laaksonen DE, Schwab U, Heinonen S, Niskanen L: How can we increase postpartum glucose screening in women at high risk for gestational diabetes mellitus? *Int J Endocrinol* 2012, 2012:519267.
- Lega IC, McLaughlin H, Coroneos M, Handley-Derry F, Donovan N, Lipscombe LL: A physician reminder to improve postpartum diabetes screening in women with gestational diabetes mellitus. *Diabetes Res Clin Pract* 2012, 95(3):352–357.
- Dietz PM, Vesco KK, Callaghan WM, Bachman DJ, Bruce FC, Berg CJ, England LJ, Hornbrook MC: Postpartum screening for diabetes after a gestational diabetes mellitus-affected pregnancy. *Obstet Gynecol* 2008, 112(4):868–874.
- Vesco KK, Dietz PM, Bulkley J, Bruce FC, Callaghan WM, England L, Kimes T, Bachman DJ, Hartinger KJ, Hornbrook MC: A system-based intervention to improve postpartum diabetes screening among women with gestational diabetes. Am J Obstet Gynecol 2012, 207(4):281–286.
- Chittleborough CR, Baldock KL, Taylor AW, Hague WM, Willson T, Martin W, Wood J, Phillips PJ: Long-term follow-up of women with gestational diabetes mellitus: the South Australian Gestational Diabetes Mellitus Recall Register. Aust N Z J Obstet Gynaecol 2010, 50(2):127–131.
- National Gestational Diabetes Register: [http://www.ndss.com.au/en/GD/ Diabetes-Register/].
- Australia Mobile Communications Statistics and Forecasts: [http://www. budde.com.au/Research/Australia-Mobile-Communications-Statistics-and-Forecasts.html].
- Car J, Gurol-Urganci I, de Jongh T, Vodopivec-Jamsek V, Atun R: Mobile phone messaging reminders for attendance at healthcare appointments. *Cochrane Database Syst Rev* 2012, 7, CD007458.
- Chen ZW, Fang LZ, Chen LY, Dai HL: Comparison of an SMS text messaging and phone reminder to improve attendance at a health promotion center: a randomized controlled trial. J Zhejiang Univ Sci B 2008, 9(1):34–38.
- 30. Downer SR, Meara JG, Da Costa AC: Use of SMS text messaging to improve outpatient attendance. *Med J Aust* 2005, **183**(7):366–368.
- Downer SR, Meara JG, Da Costa AC, Sethuraman K: SMS text messaging improves outpatient attendance. *Aust Health Rev* 2006, 30(3):389–396.
  Stuebe AM, Rich-Edwards JW, Willett WC, Manson JE, Michels KB: Duration of
- 32. Stuebe AM, Rich-Edwards JW, Willett WC, Manson JE, Michels KB: Duration of lactation and incidence of type 2 diabetes. *JAMA* 2005, **294**(20):2601–2610.
- Liu B, Jorm L, Banks E: Parity, breastfeeding, and the subsequent risk of maternal type 2 diabetes. *Diabetes Care* 2010, 33(6):1239–1241.
  Keely E, Clark H, Karovitch A, Graham I: Screening for type 2 diabetes
- 54. Reely E, Clark H, Raforicci A, Granam E Screening for type 2 diabetes following gestational diabetes: family physician and patient perspectives. Can Fam Physician 2010, 56(6):558–563.

#### doi:10.1186/1471-2393-13-92

**Cite this article as:** Heatley *et al.*: The DIAMIND study: postpartum SMS reminders to women who have had gestational diabetes mellitus to test for type 2 diabetes: a randomised controlled trial – study protocol. *BMC Pregnancy and Childbirth* 2013 13:92.

# Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at www.biomedcentral.com/submit

() BioMed Central

# Statement of Authorship

Title of Paper	Postpartum SMS reminders to women who have experienced gestational diabetes mellitus to test for type 2 diabetes: The DIAMIND randomised trial
Publication Status	O Published,
Publication Details	Van Ryswyk E, Middleton P, Hague W, Crowther C. Postpartum SMS reminders to women who have experienced gestational diabetes mellitus to test for type 2 diabetes: The DIAMIND randomised trial. Diabetic Medicine. Accepted Feb 2015.

# **Author Contributions**

By signing the Statement of Authorship, each author certifies that their stated contribution to the publication is accurate and that permission is granted for the publication to be included in the candidate's thesis.

Name of Principal Author (Candidate)	Emer Van Rysywk				
Contribution to the Paper	Contributed to study design. Recruited participants, collected and analysed data. Wrote manuscript drafts for publication.				
Signature	Date 20/2/15				

Name of Co-Author	Philippa Middleton					
Contribution to the Paper	Conceived study idea, wrote and led initial study design and protocol for ethics submission, participated in conduct of the study and interpretation of the data and edited the manuscript for publication.					
Signature	Date 19/2 15					

Name of Co-Author	William Hague
Contribution to the Paper	Contributed to study design, conduct of the study, interpretation of the data and editing of the manuscript for publication.
Signature	Date 26 2/17

Name of Co-Author	Caroline Crowther
Contribution to the Paper	Contributed to study design and trial protocol, participated in ethics submission and conduct of the study, interpretation of the data and edited the manuscript for publication.
Signature	Date 1572/15.

# Postpartum SMS reminders to women who have experienced gestational diabetes to test for type 2 diabetes: the DIAMIND randomised trial

# Abstract

Aims: This parallel group randomised controlled trial assessed whether an SMS reminder system for women, after gestational diabetes (GDM), would increase their attendance for an oral glucose tolerance test (OGTT) by six months postpartum.

Methods: Women were eligible for inclusion if they were diagnosed with GDM in their recent pregnancy, had a mobile phone and normal blood glucose profile prior to postnatal discharge from the Women's and Children's Hospital, Adelaide. A computer-generated random number sequence and telephone randomisation were used. 276 women were randomised. Women in the six week group (n = 140) were sent a text reminder to attend for an OGTT at six weeks postpartum, with further reminders at three and six months if required. Women in the control group (n = 136) received one text reminder at six months postpartum. Blinding was not feasible. The primary outcome was OGTT attendance within six months postpartum.

Results: Women in the six week group did not increase their attendance for an OGTT within six months postpartum compared with women in the control group, 104 (77.6% of 134) versus 103 (76.8% of 134), RR 1.01, 95% CI 0.89-1.15.

Conclusions: The SMS reminder system did not increase postpartum OGTT, fasting plasma glucose or HbA1c completion, although high rates of test completion were measured in both groups. Further research is required into factors influencing attendance for postpartum testing

from the perspective of women, and into optimal counselling relating to type 2 diabetes risk in the postpartum period for increasing postpartum glucose testing rates.

# Introduction

Women who have had GDM are at much higher risk of type 2 diabetes in the future; they are also at risk of recurrent GDM in future pregnancies (Kim et al 2007;Kim et al 2002). Due to this increased risk of type 2 diabetes, clinical practice guidelines recommend screening for prediabetes and type 2 diabetes in the postpartum period (American Diabetes Association 2014;Nankervis et al 2014;Thompson et al 2013). More specifically, the Australasian Diabetes in Pregnancy Society (ADIPS) guidelines recommend that women diagnosed with GDM should have a 75 gram oral glucose tolerance test (OGTT), preferably at six to twelve weeks postpartum, unless clinically contraindicated (Nankervis et al 2014). Women identified with prediabetes can then be counselled regarding type 2 diabetes prevention options (Diabetes Prevention Program Research Group 2009;Herman 2011;Knowler et al 2002;Ratner et al 2008). Identification of previously undiagnosed type 2 diabetes prior to a subsequent pregnancy allows treatment to prevent early pregnancy hyperglycaemia. This can reduce the risk of several complications for the mother and her baby (Bell et al 2008;Feig et al 2006;Macintosh et al 2006;Thompson et al 2013).

Postpartum screening rates are often reported to be low, and vary considerably between settings (Carson et al 2013;Tovar et al 2011). In Australia, women who are eligible for Medicare now have the option of joining the National Gestational Diabetes Register, which sends women postal reminders at 12-16 weeks after their expected due date. A previous Canadian trial of postal reminders for a postpartum oral glucose tolerance test (OGTT) in women who had GDM, found that postal reminders increased OGTT completion (Clark et al 2009). Short message service (SMS) reminders have been found to increase attendance for healthcare appointments in general (Gurol-Urganci et al 2013).

The DIAMIND study aimed to determine whether an SMS reminder system would increase attendance for an OGTT by six months postpartum in women who have had GDM.

82
#### Methods

#### Study design and population

The protocol for this single centre parallel group randomised controlled trial was published in 2013 (Heatley et al 2013), and the methods used followed this protocol. Ethical approval was obtained from the Women's and Children's Health Network Human Research Ethics Committee (REC2200/8/2015).

Women were eligible for inclusion in the trial if they were diagnosed with GDM in their most recent pregnancy (positive 75 gram OGTT with fasting glucose  $\geq 5.5$  mmol/L and/or two hour glucose  $\geq 7.8$  mmol/L), had access to a personal mobile phone, and had normal capillary blood glucose profile measurements prior to postnatal discharge from hospital (fasting plasma glucose < 6.0 mmol/L and 2 hour postprandial blood glucoses < 8.0 mmol/L). Women were excluded if they had pre-existing diabetes mellitus (type 1 or type 2 diabetes), a triplet/higher order multiple birth, requirement for an interpreter (due to text reminders being written in English), or if they had experienced a perinatal death in their most recent pregnancy.

The daily postnatal midwifery coordinator was consulted about women's eligibility. Women who were eligible were then approached and provided with verbal and written information about the study. Women were enrolled if they gave written informed consent and had a normal blood glucose profile prior to discharge from hospital. Recruitment for the DIAMIND Study took place from June 2012 until January 2014, when the pre-specified sample size was reached, in the postnatal ward of the Women's and Children's Hospital, South Australia, with follow-up of study outcomes completed by September 2014.

#### Randomisation procedures

Women were randomised into one of two study groups: either the 'six week group' or the control group. Allocation to study groups was carried out using a telephone randomisation service. The randomisation schedule was prepared by an investigator not involved in

recruitment or clinical care and used balanced variable blocks, with stratification by antenatal requirement for insulin therapy to treat GDM.

#### Treatment schedules

Women in the six week group were sent a SMS reminder at six weeks after the birth of their baby (Heatley et al 2013). Messages were sent automatically based on the date of birth of the baby using Clickatell bulk SMS gateway. Participants who responded to say they had completed the test were not sent further text reminders. All other women in the six week group were sent a further identical reminder at three and six months postpartum. A single text message reminder, using the same text as for the six week group, was sent to women in the control group at six months postpartum.

#### Baseline variables

Data were collected to assess the similarity between the two study groups in terms of factors that may have influenced attendance for a postpartum OGTT. At trial entry, information was collected on demographic characteristics of the participants, as well as smoking history, body mass index (BMI) and previous pregnancy outcomes. Women were asked whether or not they were offered the opportunity to join the National Diabetes Services Scheme (NDSS) and therefore the National Gestational Diabetes Register, whether they joined, and where they intended to have their postpartum OGTT. Data were collected regarding the date and the results of the antenatal diagnostic OGTT, control of GDM (dietary control only or requirement for metformin or insulin), and maternal complications at birth relating to GDM (requirement for induction, caesarean section, perineal injury, blood loss).

Health outcomes of the newborn(s) were collected that included: birth order (if twin), gestational age at birth, birth weight, time of birth, gender, Apgar scores, mode of birth, nerve palsy, bone fracture, newborn hypoglycaemia (plasma glucose  $\leq 2.0$  mmol/L), neonatal

intensive care admission, respiratory distress syndrome, neonatal jaundice requiring phototherapy, and neonatal death prior to first discharge.

Breastfeeding status at hospital discharge was collected given the link between breastfeeding and reduced risk of type 2 diabetes (Liu et al 2010;Stuebe et al 2005), as well as the influence that breastfeeding may have on the mother's ability to attend for an OGTT (Keely et al 2010). Inclusion of GDM in the problem list of the discharge summary and whether or not a follow up OGTT was recommended were also recorded.

#### Assessment of outcomes

The primary outcome was attendance for an OGTT by six months postpartum. Secondary outcomes were attendance for a fasting plasma glucose (FPG) test, or glycated haemoglobin (HbA1c) test by six months postpartum (if no attendance for OGTT recorded).

All women in the study were asked to complete a questionnaire at six months after the birth of their baby either by post or by email (using Survey Monkey), to ascertain whether an OGTT was undertaken within the first six months, or whether a FPG orHbA1c may have been undertaken instead. The questionnaire also asked for the date and results of these tests, where known. These results were confirmed by checking the participant's medical records. Women were contacted by telephone two weeks after the questionnaire over the phone, or to have the questionnaire sent to them again. Women were asked during this telephone contact if and where they undertook their postpartum OGTT. Non-responding participants were contacted again two weeks later, and offered the same options. A final reminder and copy of the questionnaire was mailed to the remaining non-responders after a further four weeks.

#### Statistical analysis

The sample size calculation used an estimated baseline rate of attendance for OGTT of 37%, at the lower end of the range in the review by Tovar and colleagues (Tovar et al 2011). This

was chosen because the health centres in the Tovar study often had reminder systems in place. The Stata version 10.0 sample size calculator was used to estimate the target sample size needed. To detect an 18% absolute improvement in attendance for OGTT from 37% to 55%, with 80% power, two-tailed significance level of 5%, and estimated 5% loss to follow up, it was estimated that 276 women would be required.

Baseline characteristics of all randomised women were compared descriptively between the study groups. Outcome comparisons were made according to the treatment group allocation at randomisation on an intention to treat basis. The primary and secondary outcomes were reported as risk ratios, with corresponding 95% confidence intervals, calculated using Epi Info 7 Software. Differences between categorical postpartum factors that may have influenced OGTT attendance were assessed using  $\chi_2$  test

#### Results

#### Recruitment and participant flow

A total of 554 women were assessed for inclusion in the trial. Of those women, 179 did not meet the inclusion criteria, 54 eligible women declined to participate, and 45 potentially eligible women were not counselled for other reasons (see Figure 1). Women were randomised into either the six week group (n = 140) or the control group (n = 136). 137 women in the six week group received their allocated reminders; two did not, due to mobile phone repairs in the early postpartum period, and one woman's mobile phone was unable to receive the text messages. All three were included in the analysis according to intention to treat. A total of 268 women (97%) were followed up to six months. Results for eight women (3%) were not available to be included in the analysis, due to being unable to be contacted after six months postpartum (n = 5), being no longer interested in the trial (n = 1), moving overseas (n = 1) or moving interstate (n = 1). Two participants had a 50 gram oral glucose challenge test (OGCT) rather than an OGTT; one was pregnant at the time, and the other

participant had an OGCT due to pathology testing centre error (both counted as OGTT noncompletions).

#### Sociodemographic characteristics of included women

There were no notable sociodemographic differences between the allocated study groups at trial entry (Table 1). The majority of women in each study group were between 30-39 years of age (61%), and were being treated within the public health system (95-6%). At trial entry, most of the women were either overweight (BMI 25.0-29.9 kg/m<sub>2</sub>) (~30%), or obese (BMI  $\geq$  30.0 kg/m<sub>2</sub>) (~40%), with only a fifth of the women being normal weight.

Ethnicities were similar between study groups as was socioeconomic status. Just under half (49%) of the women in each study group were Caucasian (of European descent), and most other women (47%) were of Asian descent (from all Asian countries, including the Indian subcontinent). There were five Indigenous Australian women, with three in the six week group (2%) and two in the control group (1.5%). Similarly, three women in the six week group and two women in the control group were from African countries (including Ethiopia, Liberia and Egypt).

There were high levels of socioeconomic disadvantage amongst participants, as judged according to postcode, with about half classified as disadvantaged or extremely disadvantaged. All participants had at least some secondary education, and half had a bachelor degree or higher.

Perinatal factors that may have influenced postpartum healthcare seeking: A comparison of study groups

There were no differences between the study groups with regards to perinatal factors (Table 2). Although, fewer women in the six week group had experienced a previous preterm birth (3%) than women in the control group (11%); the national rate of preterm birth in women

with GDM in Australia in 2005-7 was reported as 10% (Australian Institute of Health and Welfare 2010).

#### Outcomes and estimation

Primary and secondary outcome data were available for 268 participants (97%). Women in the six week group did not increase their attendance for an OGTT within six months after birth, with 104 (77.6%) women attending in the six week group and 103 (76.8%) women attending in the control group (RR 1.01, 95% CI 0.89-1.15) (Table 3).

Six women (4.5%) in the six week group, and five women (3.7%) in the control group attended for FPG tests; thus the intervention had no effect on the secondary outcome of FPG attendance within six months postpartum (RR 1.20, 95% CI 0.37 –3.84) (Table 3). Only one participant had an HbA1c test as their primary screening test (Table 3).

Finally, the SMS reminder for the six week group had no effect on the rate of completion of any of the tests combined (either OGTT or FPG or HbA1c) within six months postpartum, with 83% (n = 111) of the women in the six week group and 81% (n = 108) of the women in the control group having either test (RR 1.03, 95% CI 0.92 – 1.15).

#### Postpartum follow-up results: prediabetes and type 2 diabetes frequency

Overall, 11% of women were diagnosed with prediabetes and 2.3% diagnosed with type 2 diabetes by six months postpartum (Table 3).

#### Additional postpartum factors that may have influenced OGTT completion

Most women in both study groups ( $\geq$ 87%) self-reported that they had been offered an opportunity to join the Australian National Gestational Diabetes Register and had joined ( $\geq$ 83%) (National Diabetes Services Scheme 2014), and therefore would have received postal reminders at 12-16 weeks after their expected due date from this register (Table 3). Over 98% of women in each group receiving public medical care had a discharge summary forwarded to their postpartum care provider, with no difference seen between groups. The majority of these discharge summaries listed GDM in their problem list, although approximately 20% did not include recommendations of an OGTT in the follow-up plan.

#### Harms

No significant harms resulted from the study.

#### Discussion

SMS reminders at six weeks and three months postpartum were not found to affect the rate of attendance for postpartum screening for type 2 diabetes by six months after birth, with either OGTT, FPG or HbA1c tests. This is in contrast to the results of a previous Canadian randomised trial of postal reminders for women who had GDM (Clark et al 2009). In their study, completion of OGTTs was higher in those women who were sent a reminder (42 of 76 (55%), compared with five of 35 women (14%) in the control arm.

Within our trial, overall attendance for an OGTT within both study arms was more than 20% higher than previously reported rates of postpartum glucose testing in South Australia (Chittleborough et al 2010), and much higher than the vast majority of studies conducted worldwide (Carson et al 2013). Only a small number of studies, focussed on assessing rates of postpartum glucose intolerance or type 2 diabetes in women with recent GDM, have reported higher rates (Carson et al 2013). High rates of postpartum testing shows the positive influence of raised awareness of the need for postpartum screening amongst health professionals and women alike. The increase observed in our study may partially reflect the transition from the South Australian GDM Recall Register (established in July 2002) to the Australian National GDM recall register, which occurred just prior to the beginning of recruitment for the DIAMIND Study. A key difference between the function of these two registers was much earlier postal reminders from the National GDM Register at 3-4 months after birth, compared

with the South Australian GDM Recall Register that had provided reminders 15 months after birth (Chittleborough et al 2010).

The low use of HbA1c for type 2 diabetes screening is likely to reflect that, during the period of DIAMIND Study data collection, relevant Australian guidelines recommended use of an OGTT for postpartum type 2 diabetes screening (Nankervis et al 2014), and that Medicare reimbursement for HbA1c was only possible in people with established diabetes (d'Emden 2014).

It is noteworthy that a high proportion of women in the study received postal reminders from the national reminder scheme ( $\geq$ 83%). Furthermore, discharge summaries were completed and sent to the relevant clinicians in a very high proportion of cases (98% in each study group). Most summaries not only provided the diagnosis of GDM in the problem list (92%), but also recommended an OGTT in the follow-up treatment plan section (81%). This communication of the diagnosis of GDM is likely to have positively influenced rates of OGTT completion, as previous studies with clinicians' views have indicated that lack of communication of the diagnosis was a key factor preventing adequate postpartum healthcare provision for women with GDM (Van Ryswyk et al 2014). Most women in the study planned to attend for postpartum care with their general practitioners in the community (64%) rather than at the hospital, highlighting the importance of communication of the diagnosis to the relevant postpartum care providers.

The postpartum glucose test results from our study indicate the importance of screening relatively soon after birth; although, the rates of prediabetes (11%) and diabetes (2.3%) are at the lower end of the range of those found in previous studies of testing up to six months postpartum (prediabetes was reported in 13-32% of participants, and type 2 diabetes detected in 1-25% (Carson et al 2013). These lower rates were expected given that the women in the DIAMIND Study were only eligible if they had had a normal blood glucose profile before discharge after giving birth.

#### Limitations and generalisability

Rates of perinatal adverse outcomes known to be associated with GDM generally did not differ between study groups, indicating that the randomisation process was effective. As expected in GDM, women were slightly older (Kirke et al 2014), with higher BMIs (Torloni et al 2009) and more likely to be Asian (Teh et al 2011), compared with the overall population of women giving birth in South Australia (Scheil et al 2013) or Australia (Australian Institute of Health and Welfare 2010).

Most women in the study had been educated beyond secondary school. Almost half of the women in the study lived in either extremely disadvantaged (n = 64, 23%) or disadvantaged areas (n = 74, 27%), and a higher proportion of women in the study had received public rather (n = 264, 96%) than private care compared with other women giving birth in South Australia (71% public in 2011 (Scheil et al 2013)). This concurs with previous studies showing that socioeconomic disadvantage is linked with increased risk for GDM (Anna et al 2008). Women recruited to the DIAMIND trial were largely representative of women with GDM in South Australia.

Women were not eligible for recruitment to the study if they had an abnormal blood glucose profile after giving birth, if they had experienced a perinatal death, a triplet or higher order multiple pregnancy or if they required an interpreter. These women are likely to be at increased risk for development of type 2 diabetes compared with the eligible participants, and any reminder system implemented into care should include these women.

In any trial with behavioural outcomes, such as attendance for glucose testing, there is the potential for aspects of research participation to contribute to the observed frequency of the behavioural outcome (McCambridge et al 2014). Thus, it is possible that raised awareness of the risk of type 2 diabetes and the benefits of postpartum screening resulted from participation in the study, and that this contributed to an increase in attendance for postpartum glucose testing in both study groups.

Further research is required into the efficacy of a more convenient glucose test for postpartum screening, counselling relating to type 2 diabetes risk in the postpartum period, and better communication of the diagnosis of GDM, to optimise the rate of postpartum glucose testing.

#### Funding

This trial was registered with the ANZCTR - ACTRN12612000621819, and supported by the HCF Health and Medical Research Foundation.

#### Competing interests

None declared.

#### Acknowledgements

We are grateful to the diabetes educators, midwives and medical staff at the WCH for their advice on and assistance with recruitment.



Characteristic	6 Week Group	Control Group
	(n = 140)	(n = 136)
Age (years) <sup>a</sup>	32.1 (5.3)	32.8 (5.0)
≤ 19	1 (0.7%)	0 (0%)
20-29	50 (36%)	41 (30%)
30-39	80 (57%)	84 (62%)
≥ 40	9 (6.4%)	11 (8%)
Public care	133 (95%)	131 (96%)
BMI at trial entry (kg/m <sup>2</sup> ) <sup>b</sup>	29.2 (25.7 - 33.2)	29.0 (25.1 - 33.3)
< 18.5 (underweight)	1 (0.7%)	1 (0.7%)
18.5 – 24.9 (normal)	26 (19%)	28 (21%)
25.0 - 29.9 (overweight)	42 (30%)	39 (29%)
≥30.0 (obese)	52 (37%)	52 (38%)
Unknown	19 (14%)	16 (12%)
Ethnicity		
Caucasian	69 (49%)	67 (49%)
Asian	65 (46%)	65 (48%)
Indigenous Australian	3 (2%)	2 (1.5%)
Other	3 (2%)	2 (1.5%)
Socioeconomic Index <sup>c</sup>		
Extremely disadvantaged	31 (22%)	33 (24%)
Disadvantaged	38 (27%)	36 (26%)
Average	24 (17%)	20 (15%)
Advantaged	32 (23%)	30 (22%)
Most advantaged	15 (11%)	17 (13%)
Highest level of education <sup>d</sup>		
Postgraduate degree	24 (17%)	22 (16%)
Graduate diploma/certificate	2 (1.4%)	3 (2%)
Bachelor degree	42 (30%)	48 (35%)
Advanced diploma/diploma	16 (11%)	12 (9%)
Certificate level	20 (14%)	14 (10%)
Secondary education	36 (26%)	37 (27%)
(junior ± senior)		
Primary/other/unknown	0 (0%)	0 (0%)
Pre-pregnancy smoker	18 (13%)	20 (15%)

 Table 1: Comparison by study group of the sociodemographic characteristics of included women at trial entry

Values are number (%) unless otherwise indicated.

<sup>*a*</sup> Value is mean (standard deviation).

<sup>b</sup> Median (interquartile range).

<sup>c</sup> Socioeconomic index for area (SEIFA), where higher index scores indicate decreasing levels of social disadvantage. The index of relative socio-economic advantage and disadvantage was used. <sup>d</sup> Broad level of education from the Australian Standard Classification of Education 2001.

Maternal Data	6 Week Group	<b>Control Group</b>
	(n = 140)	(n = 136)
Treatment for GDM		
Diet only	105 (75%)	98 (72%)
Metformin only	26 (19%)	32 (24%)
Insulin only	5 (4%)	1 (<1%)
Metformin and insulin	3 (2%)	5 (4%)
None	1 (<1%)	0 (0%)
Diagnosis of GDM in index pregnancy		
GA at diagnosis (weeks +days) <sup>a</sup>	$28^{+3} (27^{+3} - 29^{+5})$	$28^{+4} (28^{+0} - 29^{+5})$
Fasting OGTT result (mmol/L) <sup>b</sup>	5.0 (0.9)	5.0 (0.7)
2 hour OGTT result (mmol/L)	8.8 (1.4)	8.8 (1.3)
Past obstetric history		
Previous pregnancy $\geq 20$ weeks GA	66 (47%)	67 (49%)
Preterm birth	4 (3%)	15 (11%)
Stillbirth	1 (<1%)	0 (0%)
Neonatal death	0 (0%)	0 (0%)
Index pregnancy		
Induction of labour	66 (47%)	56 (41%)
Perineal injury requiring suturing	50/85 (59%)	50/78 (64%)
(%vaginal births)		
Postpartum haemorrhage 600-999ml	26 (19%)	26 (19%)
Postpartum haemorrhage (>1000ml)	6 (4%)	8 (6%)
Mode of birth		
Any vaginal birth	85 (61%)	78 (57%)
Normal vaginal birth	69 (49%)	67 (49%)
Assisted vaginal birth <sup>c</sup>	16 (11%)	11 (8%)
Caesarean section	55 (39%)	58 (43%)
Caesarean section elective	27 (19%)	19 (14%)
Caesarean section emergency	28 (20%)	39 (29%)
Neonatal data (babies)	(n = 147)	(n = 140)
Birthweight (kg) <sup>b</sup>	3.2 (2.8-3.6)	3.2 (2.8-3.6)
Sets of twins <sup>d</sup>	7 (5%)	4 (3%)
GA at birth < 37 weeks	27 (18%)	26 (19%)
$GA$ at birth $34^{+0}$ - $36^{+6}$ weeks	19 (13%)	19 (14%)
GA at birth < 34 weeks	8 (5%)	7 (5%)
Macrosomia (>4000grams)	13 (9%)	13 (9%)
Birth injurv <sup>e</sup>	1 (<1%)	0(0%)
Respiratory distress syndrome	8 (5%)	5 (4%)
Angar $< 7$ at 5 minutes		5 (4%)
Neonatal hypoglycaemia <sup>f</sup>	25 (17%)	24 (17%)
Neonatal jaundiceg	14(100/2)	<u> </u>
NICLI admission	6 (10/0)	(70)

### Table 2: Comparison by study group of perinatal factors that may have influenced postpartum healthcare seeking

Unless otherwise specified, figures are number (%).

<sup>a</sup> Median (interquartile range).

<sup>b</sup> Mean (standard deviation).

<sup>c</sup> Assisted vaginal birth includes vaginal breech, forceps and ventouse.

<sup>d</sup> Number of sets of twins (i.e. number of twin babies is double this figure).

<sup>e</sup> Birth injury includes musculoskeletal or neurologic injury.

<sup>*f*</sup> Neonatal hypoglycaemia defined as  $\leq 2.0$  mmol/L at 1 or 4 hours after birth.

<sup>g</sup> Neonatal jaundice requiring phototherapy

Abbreviations are: gestational age (GA), oral glucose tolerance test (OGTT), neonatal intensive care unit (NICU).

	Overall	6 Week	Control	<b>Treatment Effect</b>
	(n = 268)	Group	Group	(95%CI)
		(n = 134)	(n = 134)	
Oral glucose tolerance test	207 (77%)	104 (78%)	103 (77%)	1.01 (0.89-1.15)
Fasting plasma glucose test	11 (4.1%)	6 (4%)	5 (4%)	1.20 (0.37 – 3.84)
HbA1c	1 (0.4%)	1 (0.7%)	0 (0%)	N/A
Overall attendance for testing	219 (82%)	111 (83%)	108 (81%)	1.03 (0.92-1.15)
Postpartum glucose test results	1	1		
<b>Overall results (either test)</b>	(n = 219)	(n = 111)	(n = 108)	N/A
Normal	184 (84%)	96 (86%)	88 (81%)	N/A
Prediabetes <sup>a</sup>	24 (11%)	11 (10%)	13 (12%)	N/A
Type 2 diabetes <sup>b</sup>	5 (2.3%)	0 (0%)	5 (4.6%)	N/A
Unknown	6 (2.7%)	4 (3.6%)	2 (1.8%)	N/A
Postpartum Care Practices	Overall	Intervention	Control	P-value
	(n = 276)	(n = 140)	(n = 136)	
Opportunity to join NDSS given	246 (89%)	122 (87%)	124 (91%)	0.28
Reminder from NDSS	235 (85%)	116 (83%)	119 (88%)	0.28
OGTT location at WCH <sup>c</sup>	99 (36%)	45 (32%)	54 (40%)	0.19
Breast-feeding at discharge	253 (92%)	126 (90%)	127 (93%)	0.31
Discharge summary <sup>d</sup>	(n = 264)	(n = 133)	(n = 131)	N/A
Available	259 (98%)	131 (98%)	128 (98%)	0.64
GDM in problem list	242 (92%)	125 (94%)	117 (89%)	0.17
OGTT recommended	212 (80%)	109 (82%)	103 (79%)	0.50

Table 3: Postpartum glucose testing and postpartum care practices

*Figures are number (%). Treatment effect is relative risk. P-value calculated using*  $X^2$  *test.* 

<sup>a</sup> Prediabetes defined as either impaired glucose tolerance (7.8-11.0 mmol/L.), impaired fasting

glucose (6.1 -6.9 mmol/L), or  $HbA1c \ge 5.7\%$ , equivalent to  $HbA1c \ge 38.7$  mmol/mol

<sup>b</sup>Type 2 diabetes was defined as fasting plasma glucose  $\geq$  7.0 mmol/L or 2 hour glucose tolerance

 $\geq$  11.1 mmol/L or HbA1c  $\geq$  6.5%, equivalent to HbA1c  $\geq$  47.5 mmol/mol.

<sup>c</sup> *Refers to the planned OGTT location specified by the participant at trial entry.* 

<sup>d</sup> Figures are for women who received public (not private) care.

*Abbreviations are National Diabetes Services Scheme (NDSS), Women's and Children's Hospital (WCH), gestational diabetes (GDM) oral glucose tolerance test (OGTT).* 

# Predictors of OGTT completion in the DIAMIND Study: Associations with sociodemographic, perinatal and postpartum factors

#### Introduction

Given that women with gestational diabetes (GDM) are at increased risk for prediabetes and type 2 diabetes (T2DM) (Bellamy et al 2009;Kim et al 2002), and that rates of postpartum screening for diabetes after GDM are reportedly low or moderate (Carson et al 2013;Tovar et al 2011), it is important to study the factors that may positively or negatively influence postpartum diabetes test completion. In particular, determining which groups of women are less likely to complete postpartum diabetes screening may allow improved care through pro-active strategies such as better tailored counselling, education and written information.

Predictors of postpartum diabetes screening have been examined in previous studies. The most commonly reported predictors were: insulin treatment during pregnancy (Almario et al 2008;Kerimoglu et al 2010;Kwong et al 2009;Lawrence et al 2010;Ogonowski and Miazgowski 2009;Stasenko et al 2010); being of older age (Ferrara et al 2009;Lawrence et al 2010;Stasenko et al 2010; Kwong et al 2009;Ogonowski and Miazgowski 2009): nulliparity or lower parity (Ferrara et al 2009;Kwong et al 2009;Lawrence et al 2010;Stasenko et al 2010), and higher education (Ferrara et al 2009;Kerimoglu et al 2010;Lawrence et al 2010). Less commonly reported predictors include GDM diagnosed earlier in pregnancy (Almario et al 2008;Ferrara et al 2009); more healthcare provider contacts after birth (Ferrara et al 2009;Lawrence et al 2010); living in small rural areas (Swan et al 2010); non-smoking status (Peticca et al 2014); sites of care with postal reminder systems (Peticca et al 2014); GDM diagnosis code in women's charts at discharge (Lawrence et al 2010); private health insurance

(Amorosa et al 2014); non-indigenous ethnicity (Chamberlain et al 2014); higher socioeconomic status or higher income (Chamberlain et al 2014;Lawrence et al 2010); being foreign born (Lawrence et al 2010); and family history of diabetes (Almario et al 2008).

The DIAMIND study measured the effects of a postpartum SMS reminder system on attendance by women for follow-up oral glucose tolerance test (OGTT) completion for T2DM, and also examined the views of the women in the study relating to postpartum diabetes screening. The protocol for DIAMIND is available in chapter 4 (Heatley et al 2013). The objective of this nested study is to examine the associations between OGTT completion and maternal sociodemographic characteristics, obstetric and perinatal outcomes, and postpartum healthcare factors experienced by women in the DIAMIND randomised controlled trial.

#### Methods

The protocol and results for the DIAMIND Study (randomised controlled trial and follow-up questionnaire) have been previously reported in full, including recruitment, data collection and outcome details (chapters 4-6). The association between sociodemographic, perinatal and postpartum factors and attendance for OGTTs of women in the DIAMIND Study was examined in this sub-analysis study using t-tests for continuous variables and  $X^2$  test for categorical variables. As previously described, the outcome data for the DIAMIND study was entered into a Microsoft Access database. The statistical software program Epi Info 7.4 was used for analysis.

#### Results

A total of 207 women completed postpartum OGTTs, with 61 not attending during the six month follow-up period; completion status for eight women was unknown.

In this study, women were more likely to complete OGTTs if they were of Asian ethnicity (P =0.007), had a bachelor's degree (P = 0.036), and if they did not smoke prior to pregnancy (P = 0.045) (Table 1). There was also a non-significant trend towards increased OGTT completion with postgraduate education (P = 0.095).

Women were less likely to attend if they had gained excessive weight during their pregnancy (P = 0.004) or were Caucasian (P = 0.001). No association was apparent between women's ages, public/private care status, body mass index (BMI) or level of socioeconomic disadvantage as indicated by postcode.

#### Perinatal factors

The frequency of perinatal factors, which had the potential to influence OGTT attendance, did not differ the OGTT completer and non-completers groups (Table 2); methods of antenatal control of GDM did not vary between these groups, nor did timing of the diagnosis of GDM, nor GDM diagnostic results. There were also no significant differences between past obstetric histories, labour complications, mode of birth or neonatal health outcomes. There was a trend towards increased attendance for postpartum diabetes testing when the mode of birth was emergency caesarean section (P = 0.093).

#### Postpartum factors

No differences were apparent between the OGTT completer and non-completer groups with regards to postpartum factors that may have influenced OGTT attendance (Table 3), with  $\geq$  85% of women receiving reminders from the NDSS in both groups, most women planning to have their OGTTs in the community rather than the WCH ( $\geq$  63%), and most women breast-feeding at discharge ( $\geq$  85%).

For women who received public medical care, there were also no differences between attending and non-attending groups with regards to completion of discharge summaries for clinicians providing postpartum care with GDM in the problem list and OGTTs recommended in most cases.

#### Discussion

#### Principal findings and comparison with other relevant studies

The predictors of postpartum diabetes screening completion in the DIAMIND Study (having a bachelor's degree, being non-smokers and being of Asian ethnicity) have all previously been found to be positively associated with postpartum glucose testing (Ferrara et al 2009;Kerimoglu et al 2010;Lawrence et al 2010; Peticca et al 2014).

The reasons for the higher rate of attendance for Asian women and lower rates of attendance amongst Caucasian women were unclear. At least two previous studies have found that Asian women were more likely to attend for postpartum testing (Lawrence et al 2010;Ferrara 2009;Tovar 2011). A study using data from the 2001-2003 Behavioural Risk Factor Surveillance System (BRFSS) survey in the United States, which assessed access to healthcare among women aged 18-44 years with a history of GDM, found that "Asian/Pacific Islander" women were the most advantaged in terms of health care access (Kim et al 2007). This was in comparison with "non-Hispanic white," "non-Hispanic African American," "Hispanic or Latina" and "Native American or Native Alaskan" women. In their study, healthcare access was measured in the survey by questions inquiring about lack of health insurance, the presence of cost barriers to physician visits in the past year, lack of a primary care provider, location of primary health care facility and lack of a physical examination within the past year. Additionally, Asian/Pacific Islander women were older and wealthier and had a lower body mass index than women who were "non-Hispanic whites," and were less often smokers. Both higher income (Chamberlain et al 2014;Lawrence et al 2010) and non-smoking status (Peticca et al 2014) have been found to be associated with increased attendance for postpartum diabetes testing.

Aside from healthcare access differences and other factors, it is possible that differences in risk perception led to an increased attendance amongst women of Asian descent and lower attendance amongst women of Caucasian descent. Women with a history of GDM who also have other known risk factors for T2DM, such as obesity and family history of diabetes, have been found to have higher personal risk perception than women without additional risk factors (Kim et al 2007;Morrison et al 2010; Salomon and Soares 2004; Sterne et al 2011). There is evidence that risk perception influences health behaviour (Brewer et al 2007), and more specifically that women with a history of GDM who perceive themselves to be at higher risk of T2DM have a greater intention to improve their own health behaviour (Kim et al 2007). This may translate into increased diabetes screening attendance for women with a history of GDM who have greater risk perception.

It has previously been pointed out that studies examining predictors of completion of postpartum diabetes screening have had conflicting results (Keely 2012); in particular, that some studies have identified that women with more severe GDM (i.e. needing insulin, higher diagnostic glucose levels during pregnancy) are more likely to undergo postpartum diabetes screening, whereas others have found that women with less serious hyperglycaemia and lower BMIs are more likely to have testing. In our study, there was no correlation between women's BMI at trial entry nor method of GDM control with their attendance for OGTTs. However, there was an association with excessive weight gain during pregnancy, although this outcome is subject to recall bias.

#### Limitations and generalisability

Whilst this sample of women is likely to be representative of women with GDM attending for care at the Women's and Children's hospital, as described in detail in chapter 5, this study had a relatively small sample size in comparison with other studies that have examined predictors

of postpartum diabetes screening, with some having sample sizes exceeding 10,000 women (Ferrara et al 2009;Lawrence et al 2010). The small sample may have therefore limited the ability of the study to identify predictors of screening. For example, the trend towards increased attendance with postgraduate degrees and emergency caesarean section may have become significant with a greater sample size, and other less common predictors may have been undetectable with this sample size.

Our study agreed with other studies on the small number of predictors of postpartum screening that were detected (higher education and non-smoking status); these factors may be common predictors of screening in other regions. However, the generalisability of this study to other populations is somewhat limited given the small sample size, and the fact that the study was carried out at a single centre. The results are therefore more useful for application at a local level.

#### Implications

The reasons for higher rates of postpartum OGTT completion by women of Asian descent and lower rates by women of Caucasian descent need to be further explored. Research should be conducted into interventions that may specifically improve postpartum screening amongst women known to be less likely to attend (e.g. lower socioeconomic status/income, smokers, and lower levels of education). Raising clinician awareness of predictors of diabetes screening and non-attendance may assist them to increase attendance through more targeted postpartum care and information provision for those women at greater risk of non-attendance.

Table 1: Sociodemographics of women wh	Table 1: Sociodemographics of women who attended for an OGTT compared with women who did no				
Characteristic	OGTT Attenders	OGTT	P-value		
	(n = 207)	Non-attenders			
		(n = 61)			
Age <sup>a</sup> (years)	32.6 (4.7)	32.3 (6.5)	0.662		
≤ 19	1 (0.5%)	0 (0.0%)	0.590		
20-29	68 (32.7%)	19 (31.6%)	0.881		
30-39	122 (58.9%)	36 (59.0%)	0.991		
$\geq$ 40	16 (7.7%)	4 (3.0%)	0.177		
Public patient	197 (95.2%)	59 (96.7%)	0.606		
Medicare eligible	196 (94.7%)	57 (93.3%)	0.710		
Body Mass Index <sup>a</sup> (kg/m <sup>2</sup> )	29.9 (6.7)	30.9 (6.3)	0.338		
< 18.5 (underweight)	1 (0.5%)	1 (1.7%)	0.347		
18.5 – 24.9 (normal)	44 (21.1%)	8 (13.3%)	0.177		
25.0 – 29.9 (overweight)	61 (29.3%)	19 (31.7%)	0.727		
≥30.0 (obese)	74 (35.6%)	27(45.0%)	0.184		
Unknown	27 (13.0%)	6 (9.8%)	0.287		
Excessive pregnancy weight gain <sup>b</sup>	49 (23.7%)	26 (42.6%)	0.004		
>18.0kg (underweight)	0 (0.0%)	0 (0.0%)	N/A		
>16.0kg (normal weight)	1 (0.5%)	2 (3.3%)	0.068		
>11.5kg (overweight)	18 (8.7%)	10 (13.4%)	0.084		
>9.0kg (obese)	30 (14.5%)	14 (23.0%)	0.117		
Unknown	47 (22.7%)	12 (19.7%)	0.615		
Ethnicity <sup>c</sup>					
Caucasian	91 (44.0%)	42 (68.8%)	0.001		
Asian	110 (52.8%)	20 (33.3%)	0.007		
Indigenous Australian	4 (1.9%)	1 (1.7%)	0.897		
African origin	2 (1.0%)	1 (1.7%)	0.647		
Socioeconomic Index for Area <sup>d</sup>		· ·			
Extremely disadvantaged	43 (20.7%)	18 (29.5%)	0.129		
Disadvantaged	53 (25.5%)	18 (29.5%)	0.623		
Average	35 (16.8%)	7 (11.7%)	0.332		
Advantaged	51 (24.5%)	11 (18.3%)	0.316		
Most advantaged	26 (12.5%)	6 (10.0%)	0.355		
Level of education <sup>e</sup>					
Postgraduate degree	40 (19.2%)	6 (10.0%)	0.095		
Graduate diploma/certificate	4 (1.9%)	1 (1.6%)	0.897		
Bachelor degree level	75 (36.0%)	13 (21.7%)	0.036*		
Advanced diploma/diploma	21 (10.0%)	4 (6.7%)	0.420		
Certificate level	23 (11.0%)	11 (18.3%)	0.136		
Secondary education	44 (21.6%)	26 (42.6%)	0.001*		
Primary education	0 (0.0%)	0 (0.0%)	N/A		
Pre-pregnancy smoker	24 (11.5%)	13 (21.7%)	0.045*		
107	(	· · · · · · · · · · · · · · · · · · ·	-		

Values are number (%) unless otherwise indicated.

<sup>*a*</sup> = Mean (standard deviation).

 $^{b}$  = Based on the recommendations for weight gain during pregnancy from the Institute of Medicine and National Research Council (US), and used recall pre-pregnancy weight and recall weight prior to birth according to study participants (Rasmussen KM et al 2009)

 $^{c}$  = Racial classifications from the Pregnancy Outcome in South Australia 2010 Report (Scheil et al 2013)  $^{d}$  = The index of relative socio-economic advantage and disadvantage was used (SEIFA 2011) (Australian Bureau of Statistics 2011)

<sup>e</sup> = Broad level of education from the Australian Standard Classification of Education 2001 (Trewin 2001)

Table 2	Perinatal	factors that	mav l	have i	influenced	OGTT	attendance
I abic 2	i ci matai	factors that	mayi	nave	mnuchecu	OULL	attenuance

	OGTT Attenders	Non-attenders	P-value
	(n = 207)	(n = 61)	
Method of control of GDM			
Diet only	151 (72.9%)	45 (73.8%)	0.899
Metformin only	45 (21.7%)	12 (19.7%)	0.729
Insulin only	5 (2.4%)	1 (1.6%)	0.719
Metformin and insulin	5 (2.4%)	3 (4.9%)	0.313
Diagnosis of GDM			
GA at diagnosis <sup>a</sup> (weeks <sup>+days</sup> )	$28^{+4} (27^{+4} - 29^{+4})$	$28^{+5} \left(27^{+0} - 30^{+4}\right)$	0.208
Fasting OGTT result <sup>b</sup> (mmol/L)	5.0 (0.9)	4.9 (0.6)	0.515
2 hour OGTT result <sup>b</sup> (mmol/L)	8.9 (1.4)	8.6 (1.4)	0.112
Past obstetric history			
Previous pregnancy $> 20$ weeks GA	100 (48.3%)	30 (49.2%)	0.905
Preterm birth	17 (8.2%)	2 (3.3%)	0.187
Neonatal death	0 (0.0%)	0 (0.0%)	N/A
Stillbirth	1 (0.5%)	0 (0.0%)	0.586
Index pregnancy			
Induction of labour	87 (42.0%)	30 (49.2%)	0.322
Perineal injury requiring suturing	74 (35.7%)	22 (36.1%)	0.964
Postpartum haemorrhage	54 (26.0%)	10 (16.4%)	0.119
600-999ml	42 (20.3%)	8 (13.1%)	0.206
≥1000ml	12 (5.8%)	2 (3.3%)	0.437
Neonatal data (babies)	OGTT Attenders	Non-attenders	P-value
	(n = 214)	(n = 64)	
Twins	7 sets (14 babies)	3 sets (6 babies)	0.441
GA at birth $< 37$ weeks	38 (17.7%)	14 (21.9%)	0.458
Birthweight <sup>a</sup>	3260 (2790-3625)	3150 (2765-3605)	0.576
Macrosomia (≥4000grams)	19 (8.9%)	7 (10.9%)	0.620
Birth injury <sup>c</sup>	1 (0.5%)	0 (0.0%)	0.583
Respiratory distress syndrome	11 (5.1%)	2 (3.1%)	0.503
Apgar < 7 at 1 minute	30 (14.0%)	6 (9.4%)	0.331
Apgar < 7 at 5 minutes	8 (3.7%)	1 (1.6%)	0.338
Neonatal hypoglycaemia	38 (17.7%)	10 (15.6%)	0.692
$(\leq 2.0 \text{ mmol/L at } 1 \text{ or } 4 \text{ hours})$			
Jaundice requiring phototherapy	21 (9.8%)	6 (9.4%)	0.917
NICU admission	5 (2.3%)	4 (6.3%)	0.120
Death prior to discharge	0 (0.0%)	0 (0.0%)	N/A
Mode of birth (babies)	(n = 214)	(n = 64)	
Any vaginal delivery	117 (54.7%)	42 (65.6%)	0.120
Normal vaginal	100 (46.7%)	33 (51.6%)	0.497
Assisted vaginal <sup>d</sup>	17 (7.9%)	9 (14.1%)	0.140
Vaginal breech	0 (0.0%)	0 (0.0%)	N/A
Forceps	11 (5.1%)	9 (14.1%)	0.015
Ventouse	6 (2.8%)	0 (0.0%)	0.175
Caesarean section	97 (45.3%)	22 (34.4%)	0.120
Elective	38 (17.7%)	11 (17.2%)	0.916
Emergency	59 (27.6%)	11 (17.2%)	0.093

Unless otherwise specified, figures are number (%).

a = Value is median (interquartile range)

<sup>b</sup> = Value is mean  $\pm$  standard deviation.

<sup>C</sup> = Birth injury includes musculoskeletal or neurological injury.

 $^{d}$  = Assisted vaginal birth includes vaginal breech, forceps, ventouse.

Abbreviations are: gestational age (GA), oral glucose tolerance test (OGTT).

Table 3: Postpartum factors that may have influenced OGTT attendance

	OGTT Attenders	Non-attenders	P-value
	(n = 207)	(n = 61)	
Opportunity to join national GDM register	185 (89.4%)	54 (88.5%)	0.851
Reminder from national GDM register	175 (84.5%)	53 (86.9%)	0.651
OGTT planned at WCH	74 (35.7%)	23 (37.7%)	0.780
Breast-feeding at discharge	193 (93.2%)	53 (86.9%)	0.112
Public medical care	(n = 197)	(n = 59)	N/A
Discharge summary available	194 (98.5%)	57 (96.6%)	0.496
GDM in problem list	182 (92.4%)	53 (89.8%)	0.818
OGTT recommended	162 (82.2%)	43 (72.9%)	0.208

Unless otherwise specified, figures are number (%). Abbreviations are: Women's and Children's Hospital (WCH), gestational diabetes mellitus (GDM)

## Statement of Authorship

Title of Paper	Women's views on postpartum testing for type 2 diabetes after gestational diabetes: six month follow-up for the DIAMIND randomised controlled trial
Publication Status	O Published, O Accepted for Publication,
Publication Details	Van Ryswyk E, Middleton P, Hague W, Crowther C. Women's views on postpartum testing for type 2 diabetes after gestational diabetes: six month follow-up for the DIAMIND randomised controlled trial. Submitted to Primary Care Diabetes Dec 2014; currently under review.

#### **Author Contributions**

By signing the Statement of Authorship, each author certifies that their stated contribution to the publication is accurate and that permission is granted for the publication to be included in the candidate's thesis.

Name of Principal Author (Candidate)	Emer Van Rysywk
Contribution to the Paper	Contributed to study design. Prepared and sent questionnaires, made follow-up phone calls to participants, analysed results. Wrote manuscript drafts for publication.
Signature	Date 20/2/15

Name of Co-Author	Philippa Middleton
Contribution to the Paper	Conceived study idea, wrote and led initial study design and protocol for ethics submission, participated in conduct of the study and interpretation of the data and edited the manuscript for publication.
Signature	Date 19215

Name of Co-Author	William Hague
Contribution to the Paper	Contributed to study design, conduct of the study, interpretation of the data and editing of the manuscript for publication.
Signature	Date 262/15

Name of Co-Author	Caroline Crowther
Contribution to the Paper	Contributed to study design and trial protocol, participated in ethics submission and conduct of the study, interpretation of the data and edited the manuscript for publication.
Signature	Date 15/2/15-

# Women's views on postpartum testing for type 2 diabetes after gestational diabetes: six month follow-up for the DIAMIND randomised controlled trial

#### Abstract

**Background:** This study assessed the views of the women who participated in the DIAMIND randomised trial (of postpartum SMS reminders to test for type 2 diabetes after gestational diabetes) on their preferred type of postpartum reminder system and barriers and facilitators to completion of postpartum diabetes testing.

**Method**: A written questionnaire was sent to women who participated in the DIAMIND trial (n = 276) via post or email at six months after the birth of their baby.

**Results**: 208 women (75%) returned the study questionnaires. Preferred postpartum reminder types were: SMS (67%), email (17%), postal (12%) and voice call (1%). Women who had not yet completed an OGTT indicated that they planned to undertake one in the future (61%). Common barriers to postpartum OGTT completion included: not having enough time (73%), inadequate childcare (30%), and a need to focus on the health of the baby (30%). The most common facilitator was having a shorter test (33%).

**Conclusions**: Most women preferred postpartum SMS reminders, followed by email, postal and voice call reminders. Time constraints and test inconvenience were the most common barriers to postpartum test completion, with the option of a shorter postpartum test being the foremost facilitator.

#### Introduction

Women who have had gestational diabetes mellitus (GDM) are at significantly higher risk for the development of type 2 diabetes (T2DM) (Bellamy et al 2009;Kim et al 2002). Therefore, postpartum blood glucose screening is important to detect prediabetes and T2DM, allowing timely treatment (Nankervis et al 2014;South Australian Perinatal Practice Guidelines 2012;American Diabetes Association 2014;The American College of Obstetricians and Gynecologists 2013;Thompson et al 2013;Walker 2008).

However, often women are not offered or do not attend for postpartum screening for T2DM (Carson et al 2013;Tovar et al 2011). For example, in England, a study which examined postpartum glucose screening rates in women with a history of GDM, using a nationally representative sample from 127 urban and suburban primary care practices, found that just 18.5% of women had glucose screening within six months of birth (McGovern et al 2014). Similarly, a study from Boston, in the United States, found that just 23.4% of GDM affected women received any glucose test by six months postpartum (McCloskey et al 2014). Furthermore, a recent Australian study found rates of postpartum oral glucose tolerance testing of 25% in Indigenous women and 34% in non-Indigenous women at three years after birth (Chamberlain et al 2014).

Previous qualitative studies have identified reasons for non-attendance or non-completion from the perspective of women (Bennett et al 2011;Keely et al 2010;Sterne et al 2011). These reasons included: time pressures, lost laboratory forms, not knowing a test was necessary (Keely et al 2010;Sterne et al 2011), feelings of emotional stress whilst adjusting to a new baby and fear of a T2DM diagnosis (Bennett et al 2011;Sterne et al 2011).

In Australia, there is now a national scheme that sends postal reminders to women who have had GDM, and their general practitioners, at 12-16 weeks after the expected birth date (National Diabetes Services Scheme 2014). It is possible that a short message service (SMS) reminder system may be preferable to women, given the greatly decreasing use of the postal system in Australia and other developed countries (Australia Post 2013).

This paper reports the findings of the six month postpartum follow-up questionnaire for the DIAMIND randomised controlled trial (RCT) (n = 276), conducted at the Women's and Children's Hospital, Adelaide, Australia. This RCT examined the efficacy of an SMS reminder system, with reminders being sent to women who had recently experienced GDM, for increasing completion of an oral glucose tolerance test (OGTT) by six months postpartum (Heatley et al 2013).

The objectives of this follow-up study were to obtain the views of the women who participated in the DIAMIND RCT regarding: (1) their preferred type of postpartum reminder system (2) ease of completion and acceptability of their postpartum blood glucose test, (3) their intention to undertake postpartum blood glucose testing where not yet completed, and (4) their perceived barriers and facilitators to postpartum glucose testing.

#### Materials and Methods

#### Overall study design and participants

A questionnaire was designed to ascertain the views of participants of the DIAMIND RCT (Heatley et al 2013). This method of study was thought to be more acceptable to the women than focus groups or interviews, given their time constraints. The questionnaire was distributed to participants six months after the birth of their baby by post or email (using Survey Monkey), depending on the expressed preference at the time of recruitment. The DIAMIND RCT included women diagnosed with GDM in their most recent pregnancy (OGTT with fasting plasma glucose  $\geq 5.5$  mmol/L and/or two hour glucose  $\geq 7.8$  mmol/L), with access to a mobile phone, whose capillary blood glucose profile measurements prior to postnatal discharge were all normal (fasting glucose < 6.0 mmol/L, postprandial glucoses < 8.0 mmol/L) (Heatley et al 2013). Women were not eligible for the study if they had

pregestational diabetes mellitus, triplet/higher order multiple birth or stillbirth in the index pregnancy, or if they required an interpreter (SMS reminders were sent in English).

Recruitment for the DIAMIND RCT took place in the postnatal ward of the Women's and Children's Hospital, Adelaide, Australia. Women were counselled regarding the nature of the study and provided with a written information sheet.

Ethical approval for the DIAMIND study was obtained from the Women's and Children's Health Research Network Human Research Ethics committee.

#### Questionnaire content

The questionnaire was based on a review of the relevant literature, and the final questions were chosen by consensus amongst the investigators (for full questionnaire, please refer to the Appendix 4). Face validity of the questionnaire was checked with three non-investigators. The questionnaire was designed for ease of completion by women from both English and non-English speaking backgrounds.

Women were asked to choose (by ticking a box) their first preference for the type of postpartum reminder system. The options listed were: *Postal, SMS, Email, Voice Call,* or *Don't know.* Women who had completed postpartum glucose testing were asked for their opinion on the tests. Those who completed the OGTT were asked if: (1) it was easy to fast for the test, (2) it was easy to find time to take the test, and (3) overall whether they were happy with the test experience.

The same questions were asked of women who had completed alternative tests, such as the fasting plasma glucose (FPG) test and the HbA1c test (except for the question relating to fasting, which was not applicable to the HbA1c test).

A 5-point Likert scale was used, with the options being either *strongly disagree*, *disagree*, *neither agree nor disagree*, *agree* or *strongly agree*. These questions were included due to the

OGTT being previously reported to be a barrier in itself, with women saying the test is too long, inconvenient, and unpleasant (Sterne et al 2011). Also, it is possible that screening recommendations with regards to the type of glucose test, may change in the future for postpartum screening for T2DM. Recent research has examined use of HbA1c and/or fasting plasma glucose for this purpose, although timing, cut-off points and the long-term outcomes of changing the test needs to be investigated and balanced against the likelihood of increasing postpartum screening rates by using more convenient tests (Benaiges et al 2013;d'Emden 2014;Katreddy et al 2013;Kim et al 2011;Megia et al 2012;Nankervis and Conn 2013;Noctor et al 2013;Wilkinson et al 2014).

Women who had not completed a postpartum OGTT were asked whether they planned to have an OGTT in the near future (options were *Yes*, *No*, *Don't Know*). Women were asked to tick all of the boxes which applied to them regarding several previously reported barriers that may have prevented them for attending for an OGTT in the six months since the birth of their baby; those were concern or anxiety about being diagnosed with T2DM (Bennett et al 2011;Sterne et al 2011), baby's health (Bennett et al 2011), not having enough time (Keely et al 2010), feeling down or low (Bennett et al 2011), childcare not available/inadequate (Bennett et al 2011), transport not available/inadequate (Sterne et al 2011), perception of low risk of T2DM (Bennett et al 2011;Kim et al 2007;Morrison et al 2010), the test being too unpleasant (Sterne et al 2011) or not knowing where to go for the test.

Women who had not completed a postpartum OGTT were then asked two open ended questions:

(1) Are there any other reasons that you did not have an oral glucose tolerance test?(2) Is there anything that would have helped you to have had an oral glucose tolerance test in the six months after the birth of your baby?

#### Questionnaire follow-up

Women who did not return their questionnaire within two weeks were followed up with a telephone reminder call, and then again two weeks later where necessary. During these follow-up conversations, women were offered the option of completing the questionnaire over the phone, or having the questionnaire sent to them again via post or email. A final reminder was sent to non-completers of the questionnaire via post one month after the second telephone call, along with a paper copy of the questionnaire.

#### Analysis

Paper and Survey Monkey results were entered in a Microsoft Access database using an electronic data entry form created using Epi Info 7.4 (Centers for Disease Control and Prevention 2013). Statistical analysis was undertaken using Epi Info 7.4, to assess differences between questionnaire completers and non-completers.

#### Results

#### Questionnaire responders

275 women from the DIAMIND Study were sent the follow-up questionnaire (one participant requested that no questionnaire be sent due to time constraints), and 208 (75%) completed questionnaires either by email (n = 100) or post/telephone (n = 108). One participant returned their questionnaire with blank responses and was unable to be contacted for further clarification. For three women, responses to the questionnaire conflicted with the known test results, and so their barriers and/or facilitators were not included in the analysis.

Compared with women who returned their questionnaires, women who did not return the study questionnaire were more likely to have not attended for postpartum glucose testing within the six month study period, to have not undertaken education further than secondary schooling, and younger and/or of Indigenous Australian descent (Table 1). Neither maternal

body mass index (BMI) nor allocation to either study group (six week SMS reminders versus no reminder until six months) was associated with questionnaire completion.

#### Questionnaire results

207 women indicated their preferred postpartum reminder system, and there was little variation between the allocated treatment groups' responses. Most women (nearly 70%) selected SMS reminders as their preferred postpartum reminder type (Table 2). Email was preferred by about 17% of women, postal by 12% and voice call reminders by less than 1%.

A total of 168 women gave their views on their postpartum glucose test, of which 165 were able to be included (Table 3). Most women who completed a postpartum OGTT agreed or strongly agreed that it was easy to fast for (n = 137, 86%), easy to find time for (n = 100, 63%) and that they were happy with the test experience (n = 124, 78%). Women who had completed a FPG test (n = 5) rather than an OGTT were generally happy with their experience (n = 3, 60%): two of these women thought it was easy to fast for, and all thought it was easy to find time for. No views were available from women who completed an HbA1c test.

Some women (n = 33/69, 48% of those who did not complete OGTTs) gave their views relating to barriers to postpartum glucose testing (Table 4). The most frequently indicated barrier was *not having enough time* (n = 24/33, 73%), followed by *inadequate or non-availability of childcare* (n = 10/33, 30%), and a *need to focus on the health of the baby* (n = 10/33, 30%). Some women believed the test was *too long* (n = 6/33, 18%), that they were *at low risk of T2DM* (n = 5/33, 15%), and some women did not seek testing because of their *concern or anxiety relating to the possibility of being diagnosed with T2DM* (n = 5/33, 15%).

A small subset of the women who did not complete a postpartum OGTT (n = 15/69, 22%), provided information on what may have facilitated their attendance for postpartum OGTT completion (Table 4). A third of those women said that a shorter test would have made it easier for them to attend. Others suggested that having a health professional arrange the test, doing the test before discharge from hospital, or having a reminder for attendance would facilitate test completion. Most women who had not had an OGTT reported that they planned to complete one in the future (61%), with more women in the control group of the study indicating their intention to complete an OGTT than women in the six week reminder group.

#### Discussion

#### Principal findings and comparison with other relevant studies

Our study found that most women preferred SMS reminders over other forms of reminder systems, followed by email, then post, and lastly voice call reminders. Women's views on reminder systems have previously been elicited in a small sample of women attending either of two tertiary care sites for antenatal GDM education (n = 51) in Ottawa, Canada, with data collected from November 2010 until February 2011. They found that women's first preferences for postpartum reminder types were home phone/landline, followed by email, postcard, SMS message and voice message (Keely et al 2012); although, their study sample may not have been representative due to its small size, and the study did not collect the sociodemographic characteristics of the participants, so it was not possible to assess this aspect of their study. Furthermore, the differences in women's views between our findings and the Canadian study may be partially explained by antenatal versus postpartum data collection, with postpartum experiences and caring responsibilities possibly influencing preferences for reminder system types. It is also possible that preferences towards use of mobile phone SMS technology may have also increased with time from 2010 to 2012-14, and may differ between women in Canada and Australia.

Just over half of the women in the DIAMIND Study were still breast-feeding at six months postpartum and several others had breast-fed for six months and only recently stopped. Breast-feeding as a barrier to undertaking postpartum diabetes screening was only reported by one woman in both the DIAMIND Study and in the Canadian study of postal reminders for postpartum OGTTs (Keely et al 2012), and is therefore unlikely to be a barrier for most women.

Women who completed postpartum oral glucose tolerance testing agreed or strongly agreed that it was easy to fast for, easy to find time for, and that overall they were happy with the test experience. Many of the barriers to OGTT completion were related to lack of time and to the test itself. Taking time out from caring for their new baby and other children, in the absence of readily available and acceptable childcare, was also difficult for many women, so a shorter test would be preferable for this reason as well.

Suggested facilitators were often related to OGTT convenience, with women suggesting that having a shorter test, not having to arrange a separate appointment for the test, and being able to do the test in a more convenient location (such as home), would facilitate their glucose test completion. This suggests that for many women, the OGTT itself does not pose a barrier to T2DM screening, but for a minority it is a significant barrier. Changing from a two hour OGTT to an HbA1c test for T2DM screening would therefore probably increase completion of a postpartum test. With regards to convenience for women, the HbA1c test has several advantages when compared with a two hour OGTT, including not requiring the women to fast, consume a glucose drink, wait for two hours for final blood sampling and having more than one blood sample to be taken. Further research is required into the sensitivity, specificity and concordance of HbA1c compared with an OGTT for T2DM screening in the postpartum period (Duke et al 2015).

Postpartum counselling and education regarding the risk of T2DM is important, given the range of perception of risk from anxiety and concern about the possibility of being diagnosed with T2DM to perception of low risk of future T2DM development, both of which posed barriers for some women in the study. It is important that T2DM screening is linked with T2DM prevention interventions for the majority of women, who would not yet have developed the condition. There is ongoing research into interventions specifically for women

who have experienced GDM (Ferrara et al 2014;Shih et al 2013), and positive results have been found for sub-groups of women with a history of GDM in previous diabetes prevention studies (Knowler et al 2002;Ratner et al 2008;Rautio et al 2014).

Only five women gave their views on their FPG test, so only limited conclusions can be drawn from this aspect of the data. No women gave their views on HbA1c. The low use of HbA1c for T2DM screening, was largely because during the period of DIAMIND RCT data collection, relevant Australian guidelines recommended use of an OGTT for postpartum T2DM screening (Nankervis et al 2014;South Australian Perinatal Practice Guidelines 2012), and Medicare reimbursement for HbA1c was only possible in people with established diabetes (d'Emden 2014).

Over half of the women who had not had an OGTT by six months after birth reported they planned to undertake a test in the future, indicating that for some women, testing during the first six months postpartum may be difficult, but they do intend to have future diabetes screening. It is important that the opportunity to provide these women with screening is not missed, and the yearly reminders being provided by the Australian national GDM register may assist with this (National Diabetes Screening Scheme 2014).

#### Strengths and limitations of this study

The study questionnaire was designed to be user-friendly and visually appealing, and it contained mainly short, easy to answer questions with tick box option answers to cater for women with low health literacy and for whom English was not their first language. The moderate response rate for the questionnaire indicates that the views of most women in the DIAMIND RCT on the questionnaire topics are likely to be accurately represented (Draugalis and Plaza 2009). Our response rate was higher than the women's response rate in the follow-up survey of the Canadian RCT of postal reminders (63%, 140 of 223), the most comparable study to this one (Keely et al 2010).

Questions eliciting views on the postpartum blood glucose tests themselves were timely, given the possible move towards use of HbA1c for T2DM diagnosis in Australia (d'Emden 2014), as was the question relating to preferred reminder system type, given the fairly recent inception of the national Postal GDM reminder system (National Diabetes Services Scheme 2014). Women were given the opportunity, if they wished, to list further barriers and facilitators in addition to the suggested barriers. These are important to consider when planning strategies to increase uptake of postnatal T2DM testing.

#### Conclusion

Most women expressed a preference for SMS reminders over other methods of postpartum reminders, followed by email, postal and voice call reminders. Generally, women who completed oral glucose tolerance testing found the test easy to find time for, easy to fast for and were happy with their overall test experience. However, for women unable to attend, time constraints and test inconvenience were the most commonly cited barriers, and doing a shorter test was stated to be the main facilitator. This suggests that changing from an OGTT to an HbA1c test might facilitate an increase in the rate of postpartum glucose testing that would be clinically important, although further research is required into the accuracy of HbA1c for postpartum T2DM diagnosis. Screening for T2DM needs to be coupled with provision of effective counselling on T2DM risk, and risk reduction. Further research into T2DM prevention programs specific for women who have experienced GDM is important.

Maternal characteristics	Total	Responders	Non-	P-value
	(n = 276)	(n = 208)	responders	
			(n = 68)	
Age (years) <sup>a</sup>	32.5 (5.1)	32.9 (5.0)	31.2 (5.6)	0.020
BMI at trial entry (kg/m <sup>2</sup> ) <sup>a</sup>	30.2 (6.7)	30.2 (6.7)	30.3 (6.6)	0.910
<18.5	2 (0.7)	1 (0.5)	1 (1.5%)	0.413
18.5 – 24.9	55 (20)	42 (20)	13 (19%)	0.794
≥25.0	185 (67)	139 (67)	46 (68%)	0.901
Unknown	34 (12)	26 (13)	8 (12%)	0.873
Highest education level				
Post-secondary	203 (74)	164 (78)	39 (59)	0.001
Secondary	73 (26)	44 (21)	29 (43)	0.001
Ethnicity				0.003 <sup>b</sup>
Caucasian	136 (49)	109 (52)	27 (41)	0.069
Asian	130 (47)	96 (46)	34 (50)	0.581
Indigenous Australian	5 (2)	1 (0.5)	4 (6)	0.004
Other ethnicity	5 (2)	2 (1)	3 (4)	0.064
Diet control	203 (74)	157 (75)	46 (68)	0.203
Allocated treatment group				
6 week reminder	140 (51)	109 (52)	31 (46)	0.329
Control	136 (49)	99 (48)	37 (54)	0.329
OGTT completed	207 (75)	170 (82)	37 (54)	< 0.001
OGTT/FPG/HbA1c done	219 (79)	181 (87)	38 (56)	< 0.001
<b>Postpartum BMI</b> <sup>c</sup> (n = 194)	26.8 (23.1–32.2)		N/A	N/A
<b>Breastfeeding duration</b> $(n = 206)$				
Did not breast-feed	9 (5)	N/A	N/A	N/A
One week	10 (5)	N/A	N/A	N/A
One month	23 (11)	N/A	N/A	N/A
Three months	27 (13)	N/A	N/A	N/A
Six months	28 (14)	N/A	N/A	N/A
>Six months	106 (51)	N/A	N/A	N/A
Unsure/Don't know	3 (1.4)	N/A	N/A	N/A

*Figures are number (%).* a = Value is mean (SD). b = Calculated using  $X^2$  test for trend. c = Value is median (interquartile range). BMI = body mass index. OGTT = oral glucose tolerance test, FPG = fasting plasma glucose. N/A = not applicable.
Preferred reminder type	Overall	6 week group	Control group	P-value	
	(n = 207)	(n = 109)	n = 98		
Postal	25 (12)	13 (12)	12 (12)	0.944	
SMS	139 (67)	76 (70)	63 (65)	0.405	
Email	35 (17)	18 (16)	17 (17)	0.940	
Voice call	2 (1)	1 (<1)	1 (1)	0.587	
Unsure	6 (3)	1 (<1)	5 (5)	0.073	

Figures are number (%).

### Table 3: Women's views on their postpartum glucose test

Test	Strongly	Agree	Neither agree	Disagree	Strongly
	agree		nor disagree		Disagree
OGTT (n = 160)					
Easy to fast for	72 (45)	65 (41)	9 (6)	11 (7)	3 (2)
Easy to find time for	26 (16)	74 (46)	20 (13)	35 (22)	5 (3)
Happy with experience	37 (23)	87 (55)	27 (17)	5 (3)	4 (2.5)
HbA1c (n = 0)					
Easy to find time for	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)
Happy with experience	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)
Fasting plasma glucose (n	= 5)				
Easy to fast for	2 (40)	1 (20)	1 (20)	1 (20)	0 (0)
Easy to find time for	3 (60)	0 (0)	2 (40)	0 (0)	0 (0)
Happy with experience	3 (60)	1 (20)	1 (20)	0 (0)	0 (0)

Figures are number (%).

	Overall	6 Week	Control
		Group	Group
Barriers as listed on questionnaire	(n = 33)	(n =17)	(n = 15)
Concern or anxiety about being diagnosed with T2DM	5 (15)	4 (23)	1 (7)
Baby's health	10 (30)	4 (23)	6 (40)
Not enough time	24 (73)	14 (82)	10 (67)
Feeling down or low	4 (12)	2 (12)	2 (13)
Childcare not available or inadequate	10 (30)	3 (18)	7 (47)
Transport not available or inadequate	4 (12)	1 (6)	3 (20)
Did not know needed OGTT	1 (3)	0 (0)	1 (7)
Low perceived risk of type 2 diabetes	5 (15)	4 (23)	1 (7)
Test too unpleasant	2 (6)	1 (6)	1 (7)
Did not know where to go	0 (0)	0 (0)	0 (0)
Barriers suggested by women			
Test too long	6 (18)	4 (23)	2 (13)
Doctor wanted me to do a different	1 (3)	1 (6)	0 (0)
glucose test (not OGTT)			
Doctor access difficult (rural/remote)	1 (3)	0 (0)	1 (7)
Too difficult to attend while breast-feeding	1 (3)	0 (0)	1 (7)
6 weeks too soon for attendance	2 (6)	0 (0)	2 (13)
Overseas trip	1 (3)	0 (0)	1 (7)
Attended for OGTT but was given OGCT	1 (3)	0 (0)	1 (7)
No lab form	2 (6)	2 (12)	0 (0)
Facilitators suggested by women	(n = 15)	(n =8)	(n = 7)
Shorter test	5 (33)	3 (38)	2 (28)
Reminders	3 (20)	1 (13)	2 (28)
Do OGTT pre-discharge (after birth)	3 (20)	3 (38)	0 (0)
Arrange OGTT appointment for me	3 (20)	3 (38)	0 (0)
Do OGTT in own home	1 (7)	0 (0)	1 (14)
Partner available for assistance	1 (7)	0 (0)	1 (14)
Provide more information on how to get postpartum OGTT	1 (7)	1 (13)	0 (0)
done			
Plans for future OGTT completion	(n = 31)	(n =15)	(n = 16)
Planning to have OGTT	19 (61)	7 (47)	12 (75)
Not planning to have OGTT	4 (13)	2 (13)	2 (13)
Unsure	8 (26)	6 (40)	2 (13)

### Table 4: Barriers and facilitators to postpartum glucose testing

Figures are number (%). Women were able to choose multiple barriers. Some women suggested more than one facilitator.

## Final Chapter: Conclusions

Postpartum follow-up of women with GDM, to test for T2DM and to offer treatment or preventive options for diabetes, is an important area where further improvements can be made. There are well-documented low rates of postpartum diabetes test completion, and increasing evidence that follow-up is essential to ensure the best possible future health.

The results of the studies throughout this thesis give insight into the factors influencing postpartum follow-up of women with GDM from the perspective of both clinicians and women, as well as specifically examining whether an SMS reminder system may be a helpful intervention for improving postpartum diabetes screening. In concluding this thesis, the findings of these studies are summarised and final conclusions are made.

# Factors influencing postpartum follow-up of women with GDM, from the perspective of clinicians.

This systematic review collated clinician views and knowledge regarding postpartum healthcare provision for women who have experienced GDM, and incorporated results from qualitative and survey studies. The review found that most clinicians knew of the increased risk of T2DM in women with a history of GDM, but that there was a gap between this knowledge, and their actual practice of postpartum screening. Several barriers prevented optimal care provision including non-communication of the diagnosis of GDM, deficiencies in knowledge regarding relevant follow-up, being unsure who was responsible for postpartum care, and difficulty with collaboration. Often clinicians observed that healthcare opportunities were not taken up by women, but recognised deficiencies in the GDM education and support available for women, and that women faced significant cost and other barriers to attendance. Some clinicians thought that facilitators of provision of healthcare to women who had had GDM included creating an empowering relationship, providing advice about future risk of T2DM, and public health promotion relating to T2DM prevention.

## Implications for practice and research: healthcare provision systematic review

Lack of communication of the diagnosis of GDM between care providers often contributes to lower rates of postpartum follow-up. Communication could be improved using systematic methods, such as documentation of GDM in diagnostic lists, having patient intake forms that ask about GDM history, or by utilising reminder systems for clinicians and/or women.

There is a need to clarify responsibility for follow up of women with GDM, to improve referral pathways between GDM-related care providers, and to ensure that clinician training covers all relevant aspects of postpartum screening.

For women who are diagnosed with GDM, it is important that they are provided with appropriate and timely verbal and written education on their condition and associated short-term and long-term risks. It is equally important to ensure that obstacles to women accessing healthcare, such as cost, are minimised.

More research is required into improvement of communication between clinicians regarding GDM diagnosis and care. It is important to investigate methods of education provision for women who have experienced GDM so that they can be optimally informed about their ongoing risk of T2DM. There is also a need to raise awareness of the risks of GDM and subsequent T2DM for women, using public health promotion methods.

## Factors influencing postpartum healthcare seeking for women who have had gestational diabetes, from the perspective of women

This systematic review synthesised views of women who have experienced GDM, relating to barriers and facilitators to postpartum healthcare seeking. Numerous contributory factors were found. The diagnosis of GDM was sometimes a worrisome or upsetting experience. Following the diagnosis, women sought information from multiple sources and found that there was a lack of specific information on GDM compared with other forms of diabetes. Some women had difficult or confusing experiences relating to antenatal management of their GDM, whilst other women had more positive experiences of antenatal care.

The maternal role played an important part in determining attendance for postpartum care, with children's needs often taking priority over their own care. A need for clinicians to take a more pro-active approach to postpartum care was identified.

Knowledge of the risk of T2DM was common, although in some studies women had poorer knowledge. There was much variation in perception of future T2DM risk. Women had increased perception of risk of T2DM with increased time from their GDM diagnosis, family history of T2DM and other known risk factors for GDM.

Women worried about the possibility of receiving a diagnosis of chronic diabetes. Knowledge of how to prevent T2DM, including the role of diet, exercise and weight control was common amongst women in a third of studies. In a smaller number of studies, women lacked knowledge relating to T2DM prevention, and could have benefited from better education. Many women had a positive attitude towards T2DM prevention. Motivators for lifestyle changes included high risk

124

perception and fear of future GDM and T2DM. Although the lifestyle changes required, particularly healthy eating, were difficult to achieve and maintain on a long term basis. Women often described a need for lifestyle change support.

Implications for practice and research: healthcare seeking systematic review

More research is required into how to best provide support relating to the emotional impacts of a GDM diagnosis. Production and assessment of educational materials is important. Methods for provision of more holistic care require further exploration. Further research into systematic methods of improving follow-up care for diabetes screening and prevention is necessary.

Findings from the DIAMIND Study: The DIAMIND SMS Reminders for T2DM Testing Trial, Predictors of Postpartum OGTT Completion, and Views of Women on Postpartum T2DM Screening

The DIAMIND Trial: Effects on an SMS reminder system on the rate of attendance for postpartum oral glucose tolerance testing.

The DIAMIND randomised controlled trial assessed the efficacy of an SMS reminder system for improving attendance for postpartum oral glucose tolerance tests (OGTTs), within six months after birth, for detection of T2DM and prediabetes in women who have recently had GDM. The trial found that SMS reminders at six weeks and three months postpartum did not affect the rate of attendance for postpartum screening with either OGTTs, fasting plasma glucose (FPG) tests or HbA1c by six months after birth. Although, overall attendance for an OGTT within both study arms was more than 20% higher than previously

reported rates of postpartum glucose testing (any test) in South Australia, and much higher than the vast majority of studies conducted worldwide.

Several factors are likely to have contributed to the relatively high rates of postpartum attendance in both groups, including raised awareness of participants and clinicians as a result of the trial, postal reminders received by most participants from the national GDM reminder register and high rates of communication of the diagnosis of GDM to postpartum care providers via discharge summaries.

### Predictors of completion of glucose testing (OGTTs) in the DIAMIND Trial.

This study assessed the predictors of postpartum completion of OGTTs by women in the DIAMIND study. The study found that having a bachelor's degree, being non-smokers and being of Asian ethnicity was associated with increased attendance, whilst Caucasian ethnicity and excessive weight gain during pregnancy were predictive of non-attendance. Both higher education and nonsmoking status had previously been found to be predictors of postpartum diabetes screening. The reasons for the higher rate of attendance for Asian women and lower rates of attendance amongst Caucasian women were unclear, and require further exploration. Previous studies have found that foreign born women were more like to attend for postpartum testing and that Asian/Pacific Islander women were more advantaged than women from other ethnic groups with regards to healthcare access factors and income, although our study did not explore whether these factors played a role in influencing attendance.

126

## Views of women in the DIAMIND Trial relating to barriers and facilitators to postpartum OGTT completion

Our six-month follow-up qualitative study for participants of the DIAMIND trial found that most women preferred SMS over other forms of reminders, followed by email, then post, and lastly voice call reminders. Many of the barriers to OGTT completion were related to lack of time and to the test itself. Taking time out from caring for their new baby and other children, in the absence of readily available and acceptable childcare, was also difficult for many women.

Suggested facilitators were often related to OGTT convenience, with women suggesting that having a shorter test, not having to arrange a separate appointment for the test, and being able to do the test in a more convenient location, such as their own home, would facilitate completion of their diabetes test. Over half of the women who had not had an OGTT by six months after birth reported they planned to undertake one in the future.

Women who completed an OGTT were generally happy with the test experience and found it easy to find time for and fast for. Only five women gave their views on their FPG test, and no women gave views on HbA1c, so firm conclusions could not be drawn on this aspect of the data.

### Implications for practice and research from the DIAMIND Study

SMS reminders cannot be recommended over postal reminders on the basis efficacy alone, although women's preferences for electronic forms of reminder systems should be taken into consideration. In order to maintain and further increase postpartum glucose screening, it is important to ensure that the diagnosis of GDM is communicated well between care providers, that postpartum reminders are given to women and that women and clinicians are aware of the benefits of postpartum diabetes screening.

For women who did not attend for postpartum diabetes screening, raising clinician awareness of predictors of diabetes screening, such as higher education and Asian ethnicity, and non-attendance, such as Caucasian ethnicity, smoking and excessive weight gain, may assist them to increase attendance through more targeted postpartum care and information provision for those women at greater risk of nonattendance. Further research is required into reasons for non-attendance by specific groups of women.

Given that the barriers to oral glucose tolerance test completion were sometimes related to lack of time and test inconvenience, it is important to further research the efficacy of a more convenient test, such as HbA1c, for this purpose.

### **Overall Conclusion**

In conclusion, postpartum care for women who have experienced GDM may be improved by provision of more holistic, pro-active and supportive care from diagnosis right through to postpartum follow-up. Communication of the diagnosis of GDM between healthcare providers involved in the care of individual women is essential, and women should continue to be provided with reminders for care. Further research is required on the advantages and disadvantages of using a more convenient glucose test for postpartum screening. Education for women with GDM relating to type 2 diabetes risk, screening and prevention requires improvement. Research should be conducted into interventions that may specifically improve postpartum screening amongst women known to be less likely to attend. Finally, further research is needed on how best to support diabetes prevention in women who have experienced GDM.

## Thesis Bibliography

Access Economics. The growing cost of obesity in 2008: three years on. Canberra: Report for Diabetes Australia, 2008.

ACOG: ACOG Committee Opinion No. 435: postpartum screening for abnormal glucose tolerance in women who had gestational diabetes mellitus. Obstet Gynecol 2009; 113(6):1419–1421.

Ali HI, Baynouna LM, Bernsen RM. Barriers and facilitators of weight management: perspectives of Arab women at risk for type 2 diabetes. Health Soc Care Comm 2010;18(2):219-28.

Ali S, Dornhorst A. Diabetes in pregnancy: health risks and management. Postgrad Med J 2011;87(1028):417–427.

Almario CV, Ecker T, Moroz LA, Bucovetsky L, Berghella V, Baxter JK. Obstetricians seldom provide postpartum diabetes screening for women with gestational diabetes. AJOG 2008;198(5):528.

American Diabetes Association. Standards of medical care in diabetes--2014. Diabetes Care 2014;37:S14-80.

American Diabetes Association. Standards of medical care in diabetes—2013. Diabetes Care 2013;36:S11–66.

Amorosa JM, Do S, Son M, Gilroy L, Gyamfi Bannerman C. Risk factors for poor compliance with postpartum oral glucose tolerance testing in women with gestational diabetes mellitus. Obstet Gynecol 2014;123,Suppl 1:135S. Anna V, van der Ploeg HP, Cheung NW, Huxley RR, Bauman AE. Sociodemographic correlates of the increasing trend in prevalence of gestational diabetes mellitus in a large population of women between 1995 and 2005. Diabetes Care 2008; 31:2288-2293.

Australia Post Annual Report 2013: Future Ready, in, 2013. Last accessed 10 Dec 2014. Available online: http://auspost.com.au/media/documents/annualreport-2012-2013.pdf

Australian Bureau of Statistics. 2033.0.55.001 - Census of Population and Housing: Socio-Economic Indexes for Areas (SEIFA), Australia, 2011. 2011.

Australian Institute of Health and Welfare. Diabetes in pregnancy: its impact on Australian women and their babies (Report). Diabetes series no. 14. Cat. no. CVD 52. Canberra: AIHW. Last Accessed 5 March 2015. Available online: http://www.aihw.gov.au/publication-detail/?id=6442472448.

Bailey CJ, Del Prato S, Eddy D, Zinman B, Global Partnership for Effective Diabetes M. Earlier intervention in type 2 diabetes: the case for achieving early and sustained glycaemic control. Int J Clin Pract 2005;59(11):1309-16.

Baker AM, Brody SC, Salisbury K, Schectman R, Hartmann KE. Postpartum glucose tolerance screening in women with gestational diabetes in the state of North Carolina. N C Med J 2009;70(1):14-9.

Bandyopadhyay M, Small R, Davey MA, Oats JJN, Forster DA, Aylward A. Lived experience of gestational diabetes mellitus among immigrant South Asian women in Australia. Aust N Z J Obstet Gynaecol 2011;51:360-4. Bell R, Bailey K, Cresswell T, Hawthorne G, Critchley J, Lewis-Barned N, Northern Diabetic Pregnancy Survey Steering G. Trends in prevalence and outcomes of pregnancy in women with pre-existing type I and type II diabetes. BJOG 2008;115(4):445-52.

Bell R, Lie MLS, Hayes L, Lewis-Barned N, May CR, White M. Preventing Type 2 diabetes after gestational diabetes: Women's experiences and implications for diabetes prevention interventions. Diabetic Medicine 2011;28:10.

Bellamy L, Casas JP, Hingorani AD, Williams D. Type 2 diabetes mellitus after gestational diabetes: a systematic review and meta-analysis. Lancet 2009;373(9677):1773-9.

Benaiges D, Chillaron JJ, Pedro-Botet J, Mas A, Puig de Dou J, Sagarra E, Carrera MJ, Goday A, Flores-Le Roux JA. Role of A1c in the postpartum screening of women with gestational diabetes. Gynecol Endocrinol 2013;29(7):687-90.

Bennett W, Levine D, Ennen C, Hill-Briggs F, Nicholson W, Carrese J, et al. Barriers to followup care in women with recent gestational diabetes mellitus: A qualitative study. Journal of General Internal Medicine 2010;25:S231.

Bennett WL, Ennen C, Carrese J, Hill-Briggs F, Levine D, Nicholson W, et al. Barriers to followup care in women with recent gestational diabetes mellitus: A qualitative study. Clinical and Translational Science 2010;3:S23.

Bennett WL, Ennen CS, Carrese JA, Hill-Briggs F, Levine DM, Nicholson WK, et al. Barriers to and facilitators of postpartum follow-up care in women with recent gestational diabetes mellitus: a qualitative study. J Women's Health 2011;20:239-45. Bentley-Lewis R, Levkoff S, Stuebe A, Seely EW. Gestational diabetes mellitus: postpartum opportunities for the diagnosis and prevention of type 2 diabetes mellitus. Nat Clin Pract Endocrinol Metab 2008;4(10):552-8.

Bieda J. Perceptions of Risk for the Development of Type 2 Diabetes in African-American Women with Gestational Diabetes. PhD thesis (electronic). University of Michigan, United States;2009.

Bottalico JN. Recurrent gestational diabetes: risk factors, diagnosis, management, and implications. Semin Perinatol 2007;31(3):176–184.

Brewer NT, Chapman GB, Gibbons FX, Gerrard M, McCaul KD, Weinstein ND. Metaanalysis of the relationship between risk perception and health behavior: the example of vaccination. Health Psychol 2007;26:136-45.

Budde P and McNamara S. Australia - Mobile Communications - Statistics and Forecasts. Secondary Australia - Mobile Communications - Statistics and Forecasts 2011. Available from: http://www.budde.com.au/Research/Australia-Mobile-Communications-Statistics-and-Forecasts.html

Canadian Diabetes Association Clinical Practice Guidelines Expert Committee. Canadian Diabetes Association 2013 Clinical Practice Guidelines for the Prevention and Management of Diabetes in Canada 2013;37:S1-S212.

Car J, Gurol-Urganci I, de Jongh T, Vodopivec-Jamsek V, Atun R. Mobile phone messaging reminders for attendance at healthcare appointments. Cochrane Database Syst Rev 2012;7: CD007458.

Carolan M, Steele C, Margetts H. Attitudes towards gestational diabetes among a multiethnic cohort in Australia. Journal of Clinical Nursing 2010;19:2446-53.

Carolan M, Steele C, Margetts H. Knowledge of gestational diabetes among a multi-ethnic cohort in Australia. Midwifery 2010;26:579-88.

Carolan M. Women's experiences of gestational diabetes self-management: a qualitative study. Midwifery 2013;29:637-45.

Carolan MC, Gill G, Steele C. Women's experiences of factors that facilitate or inhibit gestational diabetes self-management. BMC Pregnancy Childbirth 2012;12:99.

Carrington ER, Shuman CR, Reardon HS. Evaluation of the prediabetic state during pregnancy. Obstetr Gynecol 1957;9(6):664-9.

Carson MP, Frank MI, Keely E. Original research: postpartum testing rates among women with a history of gestational diabetes--systematic review. Primary Care Diabetes 2013;7(3):177-86.

Centers for Disease Control and Prevention, Epi Info 7.4, in, Available from: https://wwwn.cdc.gov/epiinfo/7/index.htm, 2013.

Chamberlain A, McLean A, Oats J, et al. Low Rates of Postpartum Glucose Screening Among Indigenous and non-Indigenous Women in Australia with Gestational Diabetes, Matern Child Health J 2014; [Epub ahead of print].

Chamberlain C, Fredericks B, McLean A, Oldenburg B, Mein J, Wolfe R. Associations with low rates of postpartum glucose screening after gestational diabetes among Indigenous and non-Indigenous Australian women. Aust N Z J Public Health 2014; doi: 10.1111/1753-6405.12285. [Epub ahead of print] Chen ZW, Fang LZ, Chen LY, Dai HL. Comparison of an SMS text messaging and phone reminder to improve attendance at a health promotion center: a randomized controlled trial. J Zhejiang Univ Sci B 2008;9(1):34–38.

Chittleborough CR, Baldock KL, Taylor AW, Hague WM, Willson T, Martin W, Wood J, Phillips PJ. Long-term follow-up of women with gestational diabetes mellitus: the South Australian Gestational Diabetes Mellitus Recall Register. Aust N Z J Obstet Gynaecol 2010;50(2):127-31.

Clark HD, Graham ID, Karovitch A, Keely EJ. Do postal reminders increase postpartum screening of diabetes mellitus in women with gestational diabetes mellitus? A randomized controlled trial. AJOG 2009;200(6):634 e1-7.

Clarke C, Girard G, Legardeur H, Mandelbrot L. Postpartum diabetes screening following gestational diabetes mellitus: Practices in a university hospital and focus on the role of the general practitioner. J Gynecol Obstet Biol Reprod (Paris) 2012;41:476-84.

Coffman S, Ray MA. African American women describe support processes during high-risk pregnancy and postpartum. J Obstet Gynecol Neonatal Nurs 2002;31(5):536-44.

Collier SA, Mulholland C, Williams J, Mersereau P, Turay K, Prue C. A qualitative study of perceived barriers to management of diabetes among women with a history of diabetes during pregnancy. J Women's Health 2011;20:1333-9.

Cosson E, Vittaz L, Bihan H, et al. A third of women who had gestational diabetes in Seine Saint Denis declare to realize a screening of dysglycemia in the postpartum period. First results of the IMPACT survey. Diabetes & Metabolism 2012;38:A44-A45.

Critical Appraisal Skills Programme. Qualitative Research Checklist. Available from http://www.casp-uk.net/wp-content/uploads/2011/11/CASP-Qualitative-Research-Checklist-31.05.13.pdf.

Crowther CA, Hiller JE, Moss JR, McPhee AJ, Jeffries WS, Robinson JS. Australian Carbohydrate Intolerance Study in Pregnant Women Trial G. Effect of treatment of gestational diabetes mellitus on pregnancy outcomes. NEJM 2005;352(24):2477-86.

Cundy T, Ackermann E, Ryan EA. Gestational diabetes: new criteria may triple the prevalence but effect on outcomes is unclear. BMJ 2014;348:1567.

Dahlberg A, Persson R, Wingardh K. Diabetes--pregnancy. An interview study of womens' experience of the problems it causes. Jordemodern 1981;94(10):339-60.

Daniells S, Grenyer BFS, Davis WS, Coleman KJ, Burgess JAP, Moses RG. Gestational diabetes mellitus - Is a diagnosis associated with an increase in maternal anxiety and stress in the short and intermediate term? Diabetes Care 2003;26(2):385-89.

d'Emden M. Glycated haemoglobin for the diagnosis of diabetes. Aust Prescr 2014;37(3):98-100.

d'Emden MC. Reassessment of the new diagnostic thresholds for gestational diabetes mellitus: an opportunity for improvement. Med J Aust 2014;201(4):209-11.

Diabetes Prevention Program Research Group, Knowler WC, Fowler SE, Hamman RF, Christophi CA, Hoffman HJ, et al. 10-year follow-up of diabetes incidence and weight loss in the Diabetes Prevention Program Outcomes Study. Lancet 2009; 374:1677-1686. Dietz PM, Vesco KK, Callaghan WM, Bachman DJ, Bruce FC, Berg CJ, et al. Postpartum screening for diabetes after a gestational diabetes mellitus-affected pregnancy. Obstet Gynecol 2008;112(4):868-74.

Divakar H and Manyonda I. Battling with rising prevalence of gestational diabetes mellitus: Screening and diagnosis. International Journal of Infertility and Fetal Medicine 2011;2(3):96-100.

Divakar H, Manyonda IT. Battling the rising prevalence of gestational diabetes in India: are clinicians on the right track. J Neonatal Perinat Med 2012;5:261–7.

Divakar H, Manyonda IT. Battling with rising prevalence of gestational diabetes mellitus are clinicians in India on the right track. Perinatology 2012;12:133–9.

Doran F and Davis K. Gestational diabetes mellitus in Tonga: Insights from healthcare professionals and women who experienced gestational diabetes mellitus. New Zealand Medical Journal 2010;123(1326):59-67.

Doran F, Davis K. Factors that influence physical activity for pregnant and postpartum women and implications for primary care. Australian Journal of Primary Health 2011;17(1):79-85.

Doran F. Gestational diabetes mellitus: Perspectives on lifestyle changes during pregnancy and post-partum, physical activity and the prevention of future type 2 diabetes. Aust J Prim Health 2008;14:85-92.

Downer SR, Meara JG, Da Costa AC, Sethuraman K. SMS text messaging improves outpatient attendance. Aust Health Rev 2006;30(3):389–396.

Downer SR, Meara JG, Da Costa AC. Use of SMS text messaging to improve outpatient attendance. Med J Aust 2005;183(7):366–368.

Downs DS, Ulbrecht JS. Understanding exercise beliefs and behaviors in women with gestational diabetes mellitus. Diabetes Care 2006;29:236-40.

Draugalis JR, Plaza CM. Best practices for survey research reports revisited: implications of target population, probability sampling, and response rate, Am J Pharm Educ 2009; 73:142.

Duke A, Yap C, Bradbury R, Hng T, Kim C, Wansbrough A, Cheung N. The discordance between HbA1c and glucose tolerance testing for the postpartum exclusion of diabetes following gestational diabetes. Diabetes Res Clin Pract 2015 Jan 20 [Epub ahead of print]; doi: 10.1016/j.diabres.2015.01.006.

Evans MK, O'Brien B. Gestational diabetes: The meaning of an at-risk pregnancy. Qual Health Res 2005;15:66-81.

Evans MK, Patrick LJ, Wellington CM. Health behaviours of postpartum women with a history of gestational diabetes. Can J Diabetes 2010;34:227-32.

Feig DS, Chen E, Naylor CD. Self-perceived health status of women three to five years after the diagnosis of gestational diabetes: A survey of cases and matched controls. Am J Obstet Gynecol 1998;178:386-93.

Feig DS, Razzaq A, Sykora K, Hux JE, Anderson GM. Trends in deliveries, prenatal care, and obstetrical complications in women with pregestational diabetes: a population-based study in Ontario, Canada, 1996-2001. Diabetes Care 2006;29(2):232-5.

Feig DS, Zinman B, Wang X, Hux JE. Risk of development of diabetes mellitus after diagnosis of gestational diabetes. CMAJ 2008;179:229-34.

Feig DS. Avoiding the slippery slope: preventing the development of diabetes in women with a history of gestational diabetes. Diabetes Metab Res Rev 2012; 28(4):317-20.

Ferrara A, Hedderson MM, Albright CL, Brown SD, Ehrlich SF, Caan BJ, Sternfeld B, Gordon NP, Schmittdiel JA, Gunderson EP, Mevi AA, Tsai AL, Ching J, Crites Y, Quesenberry CP, Jr. A pragmatic cluster randomized clinical trial of diabetes prevention strategies for women with gestational diabetes: design and rationale of the Gestational Diabetes' Effects on Moms (GEM) study. BMC Pregnancy Childbirth 2014;14:21.

Ferrara A, Hedderson MM, Albright CL, Ehrlich SF, Quesenberry CP, Jr., Peng T, Feng J, Ching J, Crites Y. A pregnancy and postpartum lifestyle intervention in women with gestational diabetes mellitus reduces diabetes risk factors: a feasibility randomized control trial. Diabetes Care 2011;34(7):1519-25.

Ferrara A, Peng T, Kim C. Trends in postpartum diabetes screening and subsequent diabetes and impaired fasting glucose among women with histories of gestational diabetes mellitus: A report from the Translating Research Into Action for Diabetes (TRIAD) Study. Diabetes Care 2009;32:269-74.

Freinkel N and Josimovich J. Summary and recommendations. Diabetes Care 1980;3:499-501.

Gabbe S, Hill L, Schmidt L, Schulkin J. Management of diabetes by obstetriciangynecologists. Obstet Gynecol 1998;91(5 Pt 1):643-7.

Gabbe SG, Landon MB, Warren-Boulton E, Fradkin J. Promoting health after gestational diabetes: a National Diabetes Education Program call to action. Obstet Gynecol 2012;119:171-6.

138

Garcia de Guadiana Romualdo, L., M. Gonzalez Morales, et al. The value of hemoglobin A1c for diagnosis of diabetes mellitus and other changes in carbohydrate metabolism in women with recent gestational diabetes mellitus. Endocrinol Nutr 2012;59(6): 362-366.

Gaudreau S, Michaud C. Cultural factors related to the maintenance of health behaviours in Algonquin women with a history of gestational diabetes. Chronic Dis Inj Can 2012;32:140-8.

Gingras V, Tchernof A, Weisnagel SJ, Robitaille J. Use of glycated hemoglobin and waist circumference for diabetic screening in women with a history of gestational diabetes. J Obstet Gynaecol Can 2013;35(9):810-5.

Graco M, Garrard J, Jasper AE. Participation in physical activity: perceptions of women with a previous history of gestational diabetes mellitus. Health Promot J Austr 2009;20:20-5.

Gunderson EP, Jacobs DR, Jr., Chiang V, Lewis CE, Feng J, Quesenberry CP, Jr., Sidney S. Duration of lactation and incidence of the metabolic syndrome in women of reproductive age according to gestational diabetes mellitus status: a 20-Year prospective study in CARDIA (Coronary Artery Risk Development in Young Adults). Diabetes 2010;59(2):495-504.

Gurol-Urganci I, de Jongh T, Vodopivec-Jamsek V, Atun R, Car J. Mobile phone messaging reminders for attendance at healthcare appointments. Cochrane Database Syst Rev 2013; 12:CD007458.

Hanna FWF, Peters JR, Harlow J, Jones PW. Discrepancy between postnatal and antenatal management of gestational diabetes in the U.K. Diabetes Care 2007;30:e64–70.

Hanna FWF, Peters JR, Harlow J, Jones PW. Gestational diabetes screening and glycaemic management; national survey on behalf of the Association of British Clinical Diabetologists. QJM 2008;101:777–84.

HAPO Study Cooperative Research Group, Metzger BE, Lowe LP, Dyer AR, Trimble ER, Chaovarindr U, Coustan DR, Hadden DR, McCance DR, Hod M, McIntyre HD, Oats JJ, Persson B, Rogers MS, Sacks DA. Hyperglycemia and adverse pregnancy outcomes. N Engl J Med 2008;358(19):1991-2002.

Heatley E, Middleton P, Hague W, Crowther C. The DIAMIND study: postpartum SMS reminders to women who have had gestational diabetes mellitus to test for type 2 diabetes: a randomised controlled trial - study protocol. BMC Pregnancy Childbirth 2013;13:92.

Herman WH. The economics of diabetes prevention. Med Clin North Am 2011;95(2):373-84, viii.

Hillier TA, Pedula KL, Schmidt MM, Mullen JA, Charles MA, Pettitt DJ. Childhood obesity and metabolic imprinting: the ongoing effects of maternal hyperglycemia. Diabetes Care 2007;30(9):2287-92.

Hirst JE, Tran TS, Do MAT, Morris JM, Jeffery HE. Gestational diabetes in vietnam: Little knowledge may be a dangerous thing. Int J Gynecol Obstet 2012;119:S369.

Hirst JE, Tran TS, Do MAT, Rowena F, Morris JM, Jeffery HE. Women with gestational diabetes in Vietnam: A qualitative study to determine attitudes and health behaviours. BMC Pregnancy Childbirth 2012;12.

Hjelm K, Bard K, Berntorp K, Apelqvist J. Beliefs about health and illness postpartum in women born in Sweden and the Middle East. Midwifery 2009;25:564-75.

Hjelm K, Bard K, Nyberg P, Apelqvist J. Management of gestational diabetes from the patient's perspective - A comparison of Swedish and Middle-Eastern born women. Journal of Clinical Nursing 2007;16:168-78.

Hjelm K, Bard K, Nyberg P, Apelqvist J. Swedish and Middle-Eastern-born women's beliefs about gestational diabetes. Midwifery 2005;21:44-60.

Hjelm K, Berntorp K, Apelqvist J. Beliefs about health and illness in Swedish and Africanborn women with gestational diabetes living in Sweden. JCN 2012;21:1374-86.

Hjelm K, Berntorp K, Frid A, Aberg A, Apelqvist J. Beliefs about health and illness in women managed for gestational diabetes in two organisations. Midwifery 2008;24:168-82.

Hoedjes M, Berks D, Vogel I, et al. Motivators and Barriers to a Healthy Postpartum Lifestyle in Women at Increased Cardiovascular and Metabolic Risk: A Focus-Group Study. Hypertension in Pregnancy 2012;31(1):147-55.

Hoedjes M, Berks D, Vogel I, et al. Preferences for Postpartum Lifestyle Counseling Among Women Sharing an Increased Cardiovascular and Metabolic Risk: A Focus Group Study. Hypertension in Pregnancy 2011;30(1):83-92.

Hoffman L, Nolan C, Wilson JD, Oats JJ, Simmons D. Gestational diabetes mellitus-management guidelines. The Australasian Diabetes in Pregnancy Society. Med J Aust 1998;169(2):93-7.

Hordern MD, Dunstan DW, Prins JB, Baker MK, Singh MA, Coombes JS. Exercise prescription for patients with type 2 diabetes and pre-diabetes: a position statement from Exercise and Sport Science Australia. J Sci Med Sport 2012;15(1):25-31.

Hoy-Rosas J, Lancaster KJ. Psychosocial factors influencing lifestyle change in minority women with a history of recurrent gestational diabetes. FASEB Journal 2011; 25:770.4.

Hunsberger ML, Donatelle RJ, Lindsay K, Rosenberg KD. Physician Care Patterns and Adherence to Postpartum Glucose Testing after Gestational Diabetes Mellitus in Oregon. PLOS ONE 2012;7(10).

Institute of Medicine (US) and National Research Council (US) Committee to Reexamine IOM Pregnancy Weight Guidelines; Rasmussen KM, Yaktine AL, editors. Weight Gain During Pregnancy: Reexamining the Guidelines. Washington (DC): National Academies Press (US); 2009. Available from: http://www.ncbi.nlm.nih.gov/books/NBK32813/

International Association of Diabetes Pregnancy Study Groups Consensus Panel, Metzger BE, Gabbe SG, Persson B, Buchanan TA, Catalano PA, Damm P, Dyer AR, Leiva A, Hod M, Kitzmiler JL, Lowe LP, McIntyre HD, Oats JJ, Omori Y, Schmidt MI. International association of diabetes and pregnancy study groups recommendations on the diagnosis and classification of hyperglycemia in pregnancy. Diabetes Care 2010;33(3):676-82.

Jefferson VW, Melkus GD, Spollett GR. Health-promotion practices of young black women at risk for diabetes. Diabetes Educ 2000;26(2):295-302.

Jensen DM, Damm P, Sorensen B, Molsted-Pedersen L, Westergaard JG, Klebe J, Beck-Nielsen H. Clinical impact of mild carbohydrate intolerance in pregnancy: a study of 2904 nondiabetic Danish women with risk factors for gestational diabetes mellitus. Am J Obstet Gynecol 2001;185(2):413-9.

Jensen DM, Korsholm L, Ovesen P, Beck-Nielsen H, Molsted-Pedersen L, Damm P. Adverse pregnancy outcome in women with mild glucose intolerance: is there a clinically meaningful threshold value for glucose? Acta Obstet Gynecol Scand 2008;87(1):59-62.

Jones EJ, Appel SJ, Eaves YD, Moneyham L, Oster RA, Ovalle F. Cardiometabolic Risk, Knowledge, Risk Perception, and Self-Efficacy among American Indian Women with Previous Gestational Diabetes. JOGNN 2012;41:246-57.

Jones EJ, Appel SJ. High levels of cardiometabolic risk, knowledge, and risk perception cooccur with low self-efficacy to prevent cardiometabolic disease in American indian women with previous gestational diabetes. Circulation 2011;124.

Katreddy MV, Pappachan JM, Taylor SE, Nevill AM, Indusekhar R, Nayak AU. Hemoglobin A1c in early postpartum screening of women with gestational diabetes. World J Diabetes 2013;4(3):76-81.

Keely E, Clark H, Karovitch A, Graham I. Screening for type 2 diabetes following gestational diabetes: family physician and patient perspectives. Can Fam Physician 2010;56(6):558-63.

Keely E, Kucey W, Clark H, Karovitch A, Malcolm J. Patient Preferences for Automated Reminders Following Gestational Diabetes. Can J Diabetes 2012;36:42-3.

Keely E. An Opportunity Not to be Missed - How do we improve postpartum screening rates for women with gestational diabetes? Diabetes Metab Res Rev 2012;28(4):312-6.

Kerimoglu OS, Yalvac S, Karcaaltincaba D, Kandemir O, Altinbas SK, Dede H. Early postpartum diabetes mellitus screening rates in patients with history of gestational diabetes. Arch Gynecol Obstet 2010;282:613-6.

Kevat DA, Sinha AK, McLean AG. Lower treatment targets for gestational diabetes: is lower really better? Med J Aust 2014;201(4):204-7.

Kieffer EC, Willis SK, Arellano N, Guzman R. Perspectives of pregnant and postpartum latino women on diabetes, physical activity, and health. Health Educ Behav 2002;29(5):542-56.

Kim C, Berger DK, Chamany S. Recurrence of gestational diabetes mellitus: a systematic review. Diabetes Care 2007;30(5):1314-9.

Kim C, Brawarsky P, Jackson RA, Fuentes-Afflick E, Haas JS. Changes in health status experienced by women with gestational diabetes and pregnancy-induced hypertensive disorders. J Women's Health 2005;14(8):729-36.

Kim C, Herman WH, Cheung NW, Gunderson EP, Richardson C. Comparison of hemoglobin A1c with fasting plasma glucose and 2-h postchallenge glucose for risk stratification among women with recent gestational diabetes mellitus. Diabetes Care 2011;34(9):1949-51.

Kim C, McEwen LN, Kerr EA, et al. Preventive Counseling among women with histories of gestational diabetes Mellitus. Diabetes Care 2007;30(10):2489-95.

Kim C, McEwen LN, Kieffer EC, Herman WH, Piette JD. Self-efficacy, social support, and associations with physical activity and body mass index among women with histories of gestational diabetes mellitus. Diabetes Educator 2008;34(4):719-28.

Kim C, McEwen LN, Piette JD, Goewey J, Ferrara A, Walker EA. Risk perception for diabetes among women with histories of gestational diabetes mellitus. Diabetes Care 2007;30:2281-6.

Kim C, Newton KM, Knopp RH. Gestational diabetes and the incidence of type 2 diabetes: a systematic review. Diabetes Care 2002;25(10):1862-8.

144

Kim C, Sinco B, Kieffer EA. Racial and ethnic variation in access to health care, provision of health care services, and ratings of health among women with histories of gestational diabetes mellitus. Diabetes Care 2007;30:1459-65.

Kim C, Vahratian A. Self-Rated Health and Health Care Use Among Women With Histories of Gestational Diabetes Mellitus. Diabetes Care 2010;33(1):41-42.

Kirke AB, Evans SF, Walters B. Gestational diabetes in a rural, regional centre in south Western Australia: predictors of risk. Rural Remote Health 2014;14:2667.

Knowler WC, Barrett-Connor E, Fowler SE, Hamman RF, Lachin JM, Walker EA, Nathan DM, Diabetes Prevention Program Research G. Reduction in the incidence of type 2 diabetes with lifestyle intervention or metformin. N Engl J Med 2002;346(6):393-403.

Ko JY, Dietz PM, Conrey EJ, Rodgers L, Shellhaas C, Farr SL, Robbins CL. Gestational diabetes mellitus and postpartum care practices of nurse-midwives. J Midwifery Womens Health 2013;58(1):33-40.

Koh D, Miller YD, Marshall AL, Brown WJ, McIntyre D. Health-enhancing physical activity behaviour and related factors in postpartum women with recent gestational diabetes mellitus. J Sci Med Sport 2010;13(1):42-45.

Korpi-Hyovalti E, Laaksonen DE, Schwab U, Heinonen S, Niskanen L. How can we increase postpartum glucose screening in women at high risk for gestational diabetes mellitus? Int J Endocrinol 2012;2012:519267.

Kristman V, Manno M, Cote P. Loss to follow-up in cohort studies: how much is too much? Eur J Epidemiol 2004;19(8):751-60. Kwong S, Mitchell RS, Senior PA, Chik CL. Postpartum diabetes screening: adherence rate and the performance of fasting plasma glucose versus oral glucose tolerance test. Diabetes Care 2009;32:2242-4.

Landon MB, Spong CY, Thom E, Carpenter MW, Ramin SM, Casey B et al. A multicenter, randomized trial of treatment for mild gestational diabetes. N Engl J Med 2009;361(14):1339-48.

Langer O, Umans JG, Miodovnik M. The proposed GDM diagnostic criteria: a difference, to be a difference, must make a difference. J Matern Fetal Neonatal Med 2013;26(2):111-5.

Lapolla A, Di Cianni G, Di Benedetto A, et al. Quality of Life, Wishes, and Needs in Women with Gestational Diabetes: Italian DAWN Pregnancy Study. Int J Endocrinol 2012; 2012(2012): Article ID 784726. http://dx.doi.org/10.1155/2012/784726

Lawrence JM, Black MH, Hsu JW, Chen W, Sacks DA. Prevalence and timing of postpartum glucose testing and sustained glucose dysregulation after gestational diabetes mellitus. Diabetes Care 2010;33:569-76.

Lawson EJ, Rajaram S. A transformed pregnancy: the psychosocial consequences of gestational diabetes. Sociol Health Illn 1994;16:536-62.

Lee AJ, Hiscock RJ, Wein P, Walker SP, Permezel M. Gestational diabetes mellitus: clinical predictors and long-term risk of developing type 2 diabetes: a retrospective cohort study using survival analysis. Diabetes Care 2007;30(4):878-83.

Lega IC, McLaughlin H, Coroneos M, Handley-Derry F, Donovan N, Lipscombe LL. A physician reminder to improve postpartum diabetes screening in women with gestational diabetes mellitus. Diabetes Res Clin Pract 2012;95(3):352-7.

Lie M, Hayes L, Barned NL, May C, White M, Bell R. Preventing type 2 diabetes after gestational diabetes: qualitative studies with postnatal women to inform intervention development. Ann Hum Biol 2011;38(4):466-67.

Limmer K, LoBiondo-Wood G, Dains J. Predictors of cervical cancer screening adherence in the United States: a systematic review. J Adv Pract Oncol 2014; 5(1): 31-41.

Lindmark A, Smide B, Leksell J. Perception of healthy lifestyle information in women with gestational diabetes: A pilot study before and after delivery. Eur Diab Nursing 2010;7: 16–20. doi: 10.1002/edn.150.

Liu B, Jorm L, Banks E. Parity, breastfeeding, and the subsequent risk of maternal type 2 diabetes. Diabetes Care 2010;33(6):1239-41.

Macintosh MC, Fleming KM, Bailey JA, Doyle P, Modder J, Acolet D, Golightly S, Miller A. Perinatal mortality and congenital anomalies in babies of women with type 1 or type 2 diabetes in England, Wales, and Northern Ireland: population based study. BMJ 2006;333(7560):177.

Malcolm J, Lawson ML, Gaboury I, Keely E. Risk perception and unrecognized type 2 diabetes in women with previous gestational diabetes mellitus. Diabetes Care 2009;2(3):107-10.

McCambridge J, Kypri K, Elbourne D. In randomization we trust? There are overlooked problems in experimenting with people in behavioral intervention trials. J Clin Epidemiol 2014; 67:247-253.

McCloskey L, Bernstein J, Winter M, et al., Follow-up of gestational diabetes mellitus in an urban safety net hospital: missed opportunities to launch preventive care for women, J Women's Health 2014;23:327-334.

McCulloch DK. Overview of medical care in adults with diabetes mellitus. In: UpToDate, Waltham, MA (Accessed on 4th Feb 2015).

McGovern A, Butler L, Jones S, van Vlymen J, Sadek K, Munro N, Carr H, de Lusignan S. Diabetes screening after gestational diabetes in England: a quantitative retrospective cohort study. Br J Gen Pract 2014;64(618):e17-23.

Megia A, Naf S, Herranz L, Serrat N, Yanez RE, Simon I, Vendrell J. The usefulness of HbA1c in postpartum reclassification of gestational diabetes. BJOG 2012;119(7):891-4.

Meltzer S, Leiter L, Daneman D, Gerstein HC, Lau D, Ludwig S, Yale JF, Zinman B, Lillie D: Clinical practice guidelines for the management of diabetes in Canada. Canadian Diabetes Association. CMAJ 1998, 159 (Suppl 8):1–29.

Metzger BE, Buchanan TA, Coustan DR, de Leiva A, Dunger DB, Hadden DR, Hod M, Kitzmiller JL, Kjos SL, Oats JN, et al: Summary and recommendations of the Fifth International Workshop-Conference on Gestational Diabetes Mellitus. Diabetes Care 2007; 30(Suppl 2):S251–260.

Middleton P and Crowther CA. Reminder systems for women with previous gestational diabetes mellitus to increase uptake of testing for type 2 diabetes or impaired glucose tolerance. Cochrane Database Syst Rev 2014 Mar 18;3:CD009578. doi: 10.1002/14651858.CD009578.pub2.

Ministry of Health. 2014. Screening, Diagnosis and Management of Gestational Diabetes in New Zealand: A clinical practice guideline. Wellington: Ministry of Health.

Moher D, Liberati A, Tetzlaff J, Altman DG, Group P. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. BMJ 2009;339:b2535.

Morrison MK, Collins CE, Lowe JM. Postnatal testing for diabetes in Australian women following gestational diabetes mellitus. Aust N Z J Obstet Gynaecol 2009;49(5):494-98.

Morrison MK, Koh D, Lowe JM, et al. Postpartum diet quality in Australian women following a gestational diabetes pregnancy. Eur J Clin Nutr 2012;66(10):1160-65.

Morrison MK, Lowe JM, Collins CE. Perceived risk of Type 2 diabetes in Australian women with a recent history of gestational diabetes mellitus. Diabet Med 2010;27(8):882-6.

Nankervis A and Conn J. Gestational diabetes mellitus--negotiating the confusion. Aust Fam Physician 2013;42(8):528-31.

Nankervis A, McIntyre HD, Moses R, Ross G, Callaway L, Porter C, Jeffries W, Boorman C, De Vries B, McElduff A. Australasian Diabetes In Pregnancy Society (ADIPS) Consensus guidelines for the testing and diagnosis of gestational diabetes mellitus in Australia. Online, available from: http://www.adips.org/downloads/ADIPSConsensusGuidelinesGDM-03.05.13VersionACCEPTEDFINAL.pdf, 2013.

Nankervis A, McIntyre HD, Moses R, Ross G, Callaway L, Porter C, Jeffries W, Boorman C, De Vries B, McElduff A. Australasian Diabetes in Pregnancy Society (ADIPS) Consensus Guidelines for the Testing and Diagnosis of Hyperglycaemia in Pregnancy in Australia and New Zeal modified November 2014 2014. Online, available from: http://adips.org/downloads/2014ADIPSGDMGuidelinesV18.11.2014.pdf. National Diabetes Data Group. Classification and diagnosis of diabetes mellitus and other categories of glucose intolerance. Diabetes 1979;28(12): 1039-1057.

National Diabetes Services Scheme, National Gestational Diabetes Register, in, Australia. Last accessed 10 Dec 2014. Available online: http://www.ndss.com.au/GD/Home/Headline-Containers-for-L2/National-Gestational-Diabetes-Register/

National Institute for Health and Care Excellence. Diabetes in pregnancy: Management of diabetes and its complications [CG63]. London: National Institute for Health and Care Excellence. 2008.

Neufeld HT. Food perceptions and concerns of aboriginal women coping with gestational diabetes in Winnipeg, Manitoba. J Nut Educ Behav 2011;43:482-91.

Nicklas JM, Seely EW, Zera CA, Abdul-Rahim ZS, Rudloff ND, Levkoff SE. Using Focus Groups and Informant Interviews To Identify Novel Approaches To Preventing Type 2 Diabetes in Women with a History of Gestational Diabetes. Endocrine Reviews 2010;31.

Nicklas JM, Zera CA, Seely EW, Abdul-Rahim ZS, Rudloff ND, Levkoff SE. Identifying postpartum intervention approaches to prevent type 2 diabetes in women with a history of gestational diabetes. BMC Pregnancy Childbirth. 2011;11:23.

Nielsen KK, Kapur A, Damm P, de Courten M, Bygbjerg IC. From screening to postpartum follow-up - the determinants and barriers for gestational diabetes mellitus (GDM) services, a systematic review. BMC Pregnancy Childbirth. 2014;14:41.

Noctor E, Crowe C, Carmody LA, Avalos GM, Kirwan B, Infanti JJ, O'Dea A, Gillespie P, Newell J, McGuire B, O'Neill C, O'Shea PM, Dunne FP, investigators AD. ATLANTIC DIP: simplifying the follow-up of women with previous gestational diabetes. Eur J Endocrinol 2013;169(5):681-7.

Nolan JA, McCrone S, Chertok IRA. The maternal experience of having diabetes in pregnancy. J Am Pract 2011;23(11):611-18.

O'Sullivan J and Mahan C. Criteria for the oral glucose tolerance test in pregnancy. Diabetes 1964;13:278-185.

Ogonowski J, Miazgowski T. The prevalence of 6 weeks postpartum abnormal glucose tolerance in Caucasian women with gestational diabetes. Diabetes Res Clin Pract 2009;84:239-44.

Oza-Frank R. Resources to improve care among women with a history of gestational diabetes: A provider perspective. Diabetes 2012;61:A642.

Pan XR, Li GW, Hu YH, Wang JX, Yang WY, An ZX, Hu ZX, Lin J, Xiao JZ, Cao HB, Liu PA, Jiang XG, Jiang YY, Wang JP, Zheng H, Zhang H, Bennett PH, Howard BV. Effects of diet and exercise in preventing NIDDM in people with impaired glucose tolerance. The Da Qing IGT and Diabetes Study. Diabetes Care 1997;20(4):537-44.

Parsons J, Ismail K, Amiel S, Forbes A. Perceptions among women with gestational diabetes. Qual Health Res 2014;24:575-85.

Persily CA. Relationships between the perceived impact of gestational diabetes mellitus and treatment adherence. J Obstet Gynecol Neonatal Nurs 1996;25(7):601-7.

Persson M, Hornsten A, Winkvist A, Mogren I. "Mission impossible"? Midwives' experiences counseling pregnant women with gestational diabetes mellitus. Patient Educ Couns 2011;84(1):78-83.

Persson M, Hornsten A, Winkvist A, Mogren I. 'Dealing with ambiguity'—the role of obstetricians in gestational diabetes mellitus. Acta Obstet Gynecol Scand 2012;91:439–46.

Persson M, Winkvist A, Mogren I. 'From stun to gradual balance'- women's experiences of living with gestational diabetes mellitus. Scand J Caring Sci 2010;24:454-62.

Peticca P, Shah BR, Shea A, Clark HD, Malcolm JC, Walker M, et al. Clinical predictors for diabetes screening in the first year postpartum after gestational diabetes. Obstet Med 2014;7:116-20.

Pettitt DJ and Knowler WC. Long-term effects of the intrauterine environment, birth weight, and breast-feeding in Pima Indians. Diabetes Care 1998;21 Suppl 2:B138-41.

Pettitt DJ, Knowler WC, Baird HR, Bennett PH. Gestational diabetes: infant and maternal complications of pregnancy in relation to third-trimester glucose tolerance in the Pima Indians. Diabetes Care 1980;3(3):458-64.

Picon MJ, Murri M, Munoz A, Fernandez-Garcia JC, Gomez-Huelgas R, Tinahones FJ. Hemoglobin A1c versus oral glucose tolerance test in postpartum diabetes screening. Diabetes Care 2012;35(8):1648-53.

Pierce M, Modder J, Mortagy I, Springett A, Hughes H, Baldeweg S. Missed opportunities for diabetes prevention: Post-pregnancy follow-up of women with gestational diabetes mellitus in England. Br J Gen Pract 2011;61(591):e611-e19.

Power ML, Wilson EK, Hogan SO, Loft JD, Williams JL, Mersereau PW, Schulkin J. Patterns of preconception, prenatal and postnatal care for diabetic women by obstetriciangynecologists. J Reprod Med 2013;58(1-2):7-14. Ratner RE, Christophi CA, Metzger BE, Dabelea D, Bennett PH, Pi-Sunyer X, Fowler S, Kahn SE, Diabetes Prevention Program Research G. Prevention of diabetes in women with a history of gestational diabetes: effects of metformin and lifestyle interventions. J Clin Endocrinol Metab 2008;93(12):4774-9.

Rautio N, Jokelainen J, Korpi-Hyovalti E, et al., Lifestyle Intervention in Prevention of Type 2 Diabetes in Women with a History of Gestational Diabetes Mellitus: One-Year Results of the FIN-D2D Project, J Women's Health 2014;23:506-512.

Razee H, van der Ploeg HP, Blignault I, Smith BJ, Bauman AE, McLean M, et al. Beliefs, barriers, social support, and environmental influences related to diabetes risk behaviours among women with a history of gestational diabetes. Health Promot J Austr 2010;21:130-7.

Reinblatt SL, Morin L, Meltzer SJ. The importance of a postpartum 75 g oral glucose tolerance test in women with gestational diabetes. J Obstet Gynaecol Can 2006;28(8):690–694.

Remsberg KE. Women with Gestational Diabetes Informed, but not Convinced of Power to Prevent CVD Epidemiology and Prevention/Physical Activity, Nutrition and Metabolism 2012 San Deigo, CA, United States Circulation; 2012.

Rhoads-Baeza ME, Reis J. An exploratory mixed method assessment of low income, pregnant Hispanic women's understanding of gestational diabetes and dietary change. Health Education Journal 2012;71(1):80-89.

Risa CF, Lide'n E, Friberg F. Communication patterns in antenatal diabetes care: an explorative and descriptive study of midwife-led consultations. J Clin Nurs 2011;20:2053–63.

Rumbold AR, Crowther CA. Women's experiences of being screened for gestational diabetes mellitus. Aust N Z J Obstet Gynaecol 2002;42(2):131-37.

Sacks DA, Greenspoon JS, Abu-Fadil S, Henry HM, Wolde-Tsadik G, Yao JF. Toward universal criteria for gestational diabetes: the 75-gram glucose tolerance test in pregnancy. Am J Obstet Gynecol 1995;172(2 Pt 1):607-14.

Salomon IM, Soares SM. Understanding the impact of gestational diabetes diagnosis. Nurs J Minas Gerais 2004;4:349-57.

Scheil W, Scott J, Catcheside B, Sage L, Kennare R. Pregnancy outcome in South Australia 2011: Pregnancy Outcome Unit, SA Health, Government of South Australia, 2013.

Schoen C, Osborn R, Squires D, Doty MM. Access, affordability, and insurance complexity are often worse in the United States compared to ten other countries. Health Affairs 2013;32:2205–15.

Schumann I, Schneider A, Kantert C, Lowe B, Linde K. Physicians' attitudes, diagnostic process and barriers regarding depression diagnosis in primary care: a systematic review of qualitative studies. Fam Pract 2012;29:255-63.

Segall-Gutierrez P, Liu X, Xiang AH, Toro MM, Jurow R, Reyes C, et al. Beneficial impact of promatoras on compliance with postpartum glucose tolerance testing in latina women with recent gestational diabetes mellitus (GDM). Diabetes 2011;60:A350.

Sermer M, Naylor CD, Gare DJ, Kenshole AB, Ritchie JW, Farine D, Cohen HR, McArthur K, Holzapfel S, Biringer A, et al. Impact of increasing carbohydrate intolerance on maternalfetal outcomes in 3637 women without gestational diabetes. The Toronto Tri-Hospital Gestational Diabetes Project. Am J Obstet Gynecol 1995;173(1):146-56.
Shea AK, Shah BR, Clark HD, Malcolm J, Walker M, Karovitch A, Keely EJ. The effectiveness of implementing a reminder system into routine clinical practice: does it increase postpartum screening in women with gestational diabetes? Chronic Dis Can 2011;31(2):58-64.

Shea BJ, Hamel C, Wells GA, Bouter LM, Kristjansson E, Grimshaw J, Henry DA, Boers M. AMSTAR is a reliable and valid measurement tool to assess the methodological quality of systematic reviews. J Clin Epidemiol 2009;62(10):1013-20.

Shera AS, Jawad F, Basit A. Diabetes related knowledge, attitude and practices of family physicians in Pakistan. J Pak Med Assoc 2002;52:465–70.

Shih ST, Davis-Lameloise N, Janus ED, Wildey C, Versace VL, Hagger V, Asproloupos D, O'Reilly S, Phillips PA, Ackland M, Skinner T, Oats J, Carter R, Best JD, Dunbar JA, Group MR. Mothers After Gestational Diabetes in Australia Diabetes Prevention Program (MAGDA-DPP) post-natal intervention: study protocol for a randomized controlled trial. Trials 2013;14:339.

Simmons DS, Walters BN, Wein P, Cheung NW. Guidelines for the management of gestational diabetes mellitus revisited. Med J Aust 2002;176(7):352.

Smith BJ, Cheung NW, Bauman AE, Zehle K, McLean M. Postpartum physical activity and related psychosocial factors among women with recent gestational diabetes mellitus. Diabetes Care 2005;28:2650-4.

Smith-Morris CM. Diagnostic controversy: Gestational diabetes and the meaning of risk for Pima Indian women. Med Anthropol 2005;24(2):145-77.

Soares S, Santos D. Prevention of diabetes mellitus type 2 in women with history of gestational diabetes. A descriptive study [Portuguese]. Online Brazilian Journal of Nursing 2006;5.

South Australian Perinatal Practice Guidelines (2012). Chapter 65 Diabetes mellitus and abnormal glucose tolerance. Available from:

http://www.sahealth.sa.gov.au/wps/wcm/connect/Public+Content/SA+Health+Internet/Clinic al+resources/Clinical+topics/Perinatal+practice+guidelines/#D.

Stage E, Ronneby H, Damm P. Lifestyle change after gestational diabetes. Diabetes Res Clin Pract 2004;63:67-72.

Stasenko M, Cheng YW, McLean T, Jelin AC, Rand L, Caughey AB. Postpartum follow-up for women with gestational diabetes mellitus. Am J Perinatol 2010;27:737-42.

Stasenko M, Liddell J, Cheng YW, Sparks TN, Killion M, Caughey AB. Patient counseling increases postpartum follow-up in women with gestational diabetes mellitus. Am J Obstet Gynecol 2011;204(6):522 e1-6.

Sterne V, Logan T, Palmer M. Factors affecting attendance at postpartum diabetes screening in women with gestational diabetes mellitus. Practical Diabetes International 2011;28(2):64-8.

Stuebe A, Ecker J, Bates DW, Zera C, Bentley-Lewis R, Seely E. Barriers to follow-up for women with a history of gestational diabetes. Am J Perinatol 2010;27(9):705-10.

Stuebe AM, Rich-Edwards JW, Willett WC, Manson JE, Michels KB. Duration of lactation and incidence of type 2 diabetes. JAMA 2005;294(20):2601-10.

Subramanian S, Klosterman M, Amonkar M, Hunt T. Adherence with colorectal cancer screening guidelines: a review. Prev Med 2004; 38(5): 536–50.

Swan W, Kilmartin G, Liaw ST. Assessment of readiness to prevent type 2 diabetes in a population of rural women with a history of gestational diabetes. Rural Remote Health. 2007;7:802.

Swan WE, Liaw ST, Dunning T, Pallant JF, Kilmartin G. Diabetes risk reduction behaviours of rural postpartum women with a recent history of gestational diabetes. Rural Remote Health 2010;10(4):1461.

Teh WT, Teede HJ, Paul E, Harrison CL, Wallace EM, Allan C. Risk factors for gestational diabetes mellitus: implications for the application of screening guidelines. Aust N Z J Obstet Gynaecol 2011;51(1):26-30.

The American College of Obstetricians and Gynecologists. Practice Bulletin: Clinical Management Guidelines for Obstetrician-Gynecologists. Gestational Diabetes Mellitus. Obstet Gynecol 2013;122(2): 406-416.

Thomas J, Harden A. Methods for the thematic synthesis of qualitative research in systematic reviews. BMC Med Res Methodol 2008;8:45.

Thompson D, Berger H, Feig D, Gagnon R, Kader T, Keely E, Kozak S, Ryan E, Sermer M, Vinokurof C. Diabetes and Pregnancy. Can J Diabetes 2013;37(supple 1):S168-83.

Tong A, Morton R, Howard K, Craig JC. Adolescent experiences following organ transplantation: a systematic review of qualitative studies. J Pediatr 2009;155:542–9.

Torloni MR, Betran AP, Horta BL, Nakamura MU, Atallah AN, Moron AF, et al. Prepregnancy BMI and the risk of gestational diabetes: a systematic review of the literature with meta-analysis. Obes Rev 2009; 10:194-203. Tovar A, Chasan-Taber L, Eggleston E, Oken E. Postpartum screening for diabetes among women with a history of gestational diabetes mellitus. Prev Chronic Dis 2011;8(6):A124.

Trewin D. Australian Standard Classification of Education (ASCED). Canberra, Australia: Australian Bureau of Statistics; 2001.

Trutnovsky G, Dorfer M, Panzitt T. Gestational diabetes: Patients reactions to diagnosis and treatment satisfaction. J Psychosom Obstet Gynecol 2010;31:124.

Trutnovsky G, Panzitt T, Magnet E, Stern C, Lang U, Dorfer M. Gestational diabetes: Women's concerns, mood state, quality of life and treatment satisfaction. J Matern Fetal Neonatal Med 2012;25:2464-6.

Tuomilehto J, Lindstrom J, Eriksson JG, Valle TT, Hamalainen H, Ilanne-Parikka P, Keinanen-Kiukaanniemi S, Laakso M, Louheranta A, Rastas M, Salminen V, Uusitupa M, Finnish Diabetes Prevention Study G. Prevention of type 2 diabetes mellitus by changes in lifestyle among subjects with impaired glucose tolerance. N Engl J Med 2001;344(18):1343-50.

Van Ryswyk E, Middleton P, Hague W, Crowther C. Clinician views and knowledge regarding healthcare provision in the postpartum period for women with recent gestational diabetes: a systematic review of qualitative/survey studies. Diabetes Res Clin Pract 2014;106(3):401-11.

Vandorsten JP, Dodson WC, Espeland MA, Grobman WA, Guise JM, Mercer BM, Minkoff HL, Poindexter B, Prosser LA, Sawaya GF, Scott JR, Silver RM, Smith L, Thomas A, Tita AT. NIH consensus development conference: diagnosing gestational diabetes mellitus. NIH Consens State Sci Statements 2013;29(1):1-31.

Vesco KK, Dietz PM, Bulkley J, Bruce FC, Callaghan WM, England L, Kimes T, Bachman DJ, Hartinger KJ, Hornbrook MC: A system-based intervention to improve postpartum diabetes screening among women with gestational diabetes. Am J Obstet Gynecol 2012; 207(4):281–286.

Walker JD. NICE guidance on diabetes in pregnancy: management of diabetes and its complications from preconception to the postnatal period. NICE clinical guideline 63. London, March 2008. Diabet Med 2008;25(9):1025-7.

Weaver SP. New research gestational diabetes indicates risk later in life. Family Medicine 2004;36(3):159-60.

Wilkinson SA, Lim SS, Upham S, Pennington A, O'Reilly SL, Asproloupos D, et al. Who's responsible for the care of women during and after a pregnancy affected by gestational diabetes? Med J Aust 2014;201:S78-81.

World Health Organization (2013). Diagnostic Criteria and Classification of Hyperglycaemia First Detected in Pregnancy. Geneva, World Health Organisation.

Wylie J, Millward A, Stenhouse E. P473: Gestational diabetes a window of opportunity for the prevention of Type 2 diabetes. Diabet Med 2011;28:32-203.

Wylie J, Millward A, Stenhouse E. Pregnant women's understanding and knowledge of gestational diabetes and the impact of diagnosis on their pregnancy experience. Diabet Med 2011;28:175.

Wylie J, Tomlinson J, Pinkney J, Letherby G, Stenhouse E. Cardiovascular risk related to gestational diabetes and polycystic ovary syndrome: Are women aware? Diabet Med 2012;29:102.

159

Zehle K, Smith BJ, Chey T, McLean M, Bauman AE, Wah Cheung N. Psychosocial factors related to diet among women with recent gestational diabetes opportunities for intervention. Diabetes Educ 2008;34:807-14.

### Appendix A/1: Search Strategy (27<sup>th</sup> Feb 2013)

EMBASE (301 results)							
• Emtree	was	used to identify search terms.					
• Extensi	ive se	arching (mapping, explosion, as keyword) un-tick	ed.				
• Used A	ND ł	between columns.					
('pregnancy diabetes   'help seeking behavior'/syn OR 'patient   'qualitative							
mellitus'/syn OR		attitude'/syn OR 'reminder system'/syn OR	resea	research'/syn OR			
'gestational		reminder*:ti,ab OR 'child care' OR (risk*	inter	interview/syn OR (focus			
diabetes':ti,ab)		NEAR/10 perce*) OR 'follow up':ti,ab OR	NEX	NEXT/3 group*):ti,ab			
		followup:ti,ab OR 'attitude to health'/syn OR	OR s	survey*:ti,ab			
		'knowledge':de,ti,ab OR barrier*:ti,ab OR					
		facilitator*:ti,ab OR 'practice guideline'/syn					
		OR 'oral glucose tolerance test'/syn OR					
		'hemoglobin A1c'/syn OR 'haemoglobin					
		A1c':ti,ab OR 'glycated haemoglobin':ti,ab					
		OR 'glycosylated haemoblobin':ti,ab OR (fast*	*				
		NEXT/2 glucose) OR 'non insulin dependent					
		diabetes mellitus'/syn					
WEB OF SCIEN	ICE (	301 results)					
• Used T	opic	search with lemmatization.					
Used A	ND ł	between columns.					
"gestational	risk	* NEAR/10 perce* OR Patient* NEAR/2 attitude*	• OR	qualitative OR			
diabetes"	Atti	tude* OR "Reminder Systems" OR Reminder* OI	2	interview* OR			
	"chi	ld care" OR childcare OR follow up OR followup	OR	"focus group" OR			
	Barı	ier* OR Facilitator* OR knowledge OR postnatal	OR	focus NEAR			
postpartum OR "Practice Guidelines" OR guideline*				group* OR			
"Glucose Tolerance Test" OR "Hemoglobin A,			survey*				
	Glycosylated" OR glycated hemoglobin OR glycated						
	haemoglobin OR glycosylated haemoglobin OR						
	glycosylated hemoglobin OR HbA1c OR Haemoglobin						
	A1c	OR Hemoglobin A1c OR Fasting near/2 glucose	OR				
"type 2 diabetes" OR diabetes							

#### PUBMED (375 results)

Used AND between columns.

Used AND between columns.							
Diabetes, gestationa	Reminder Systems [Mesh] OR Reminder* [TIAB]	"Qua	alitative				
[MH] OR gestationa	OR "child care" [Mesh] OR "follow up" [TIAB] OR		Research"[Mesh] OR				
diabetes [TIAB]	Followup [TIAB] OR Barrier* [TIAB] OR		"Interviews as				
	Facilitator* [TIAB] OR "Health Knowledge,	Topic"[Mesh] OR					
	Attitudes, Practice"[Mesh] OR "postpartum	"Hea	alth				
	period"[MESH] OR Practice Guidelines as Topic	Surv	Surveys"[Mesh] OR				
	[MESH] OR "Glucose Tolerance Test" [Mesh] OR	"focus groups"[Mesh]					
	Glucose tolerance test* [TIAB]] OR "Hemoglobin A,	OR i	OR interview*				
	Glycosylated" [Mesh] OR glycated hemoglobin		[TIAB] OR				
	[TIAB] OR glycated haemoglobin [TIAB] OR	qual	itative [TIAB]				
	glycosylated haemoglobin [TIAB] OR glycosylated	OR s	survey				
	hemoglobin [TIAB] ORHbA1c [TIAB] OR						
	Haemoglobin A1c [TIAB] OR Hemoglobin A1c						
	[TIAB] OR ((Risk [TIAB]] OR risks [TIAB]]) AND						
	perce* [TIAB] OR (Patient [TIAB] OR patients						
	[TIAB] OR patient's [TIAB]) AND attitude*						
	[TIAB]) OR Fasting blood glucose [TIAB] OR						
	Fasting glucose [TIAB] OR "Diabetes Mellitus, Type						
	2"[Mesh]						
CINAHL (272 resu	llts)						
Advanced search, ne	o changes to default.						
MH "Diabetes	MH "Reminder Systems" OR TX Reminder* OR TX Patie	ent*	MH "Qualitative				
Mellitus,	N2 attitude* OR TX risk* N10 perce* OR MH "Child Care	e+"	Studies+" OR				
Gestational" OR	OR TX "child care" OR MH "after care" OR TX "after care"		TX qualitative				
TX "gestational	OR TX "follow up" OR TX "followup" OR MH "Attitude	of	OR MH				
diabetes"	health personnel" OR MH "attitude to health" OR MH		"Interviews+"				
	"attitude to illness" OR TX Barrier* OR TX Facilitator* O	R	OR TX				
	MH "health knowledge" OR TX "health knowledge" MH		Inteview* OR				
	"Postnatal Period+" OR TX postnatal OR TX postpartum M	ИН	MH "Focus				
"Diabetes Mellitus, Type 2/EP" OR MH "Practice Guidelines"		nes"	Groups" OR TX				
OR TX guideline* MH "Glucose Tolerance Test" OR TX			focus NEXT/3				
"Glucose Tolerance Test" OR MH "Hemoglobin A,			group* OR MH				
	Glycosylated" OR TX glycated hemoglobin OR TX glycat	ted	surveys TX				
	haemoglobin OR TX glycosylated haemoglobin OR TX		survey*				
	glycosylated haemoglobin OR TX HbA1c OR TX						
	Haemoglobin A1c OR TX Hemoglobin A1c OR TX Fastin	g					
	NEXT/2 glucose						

	Exclusion Reason			
Study ID	Results from women with hGDM not reported separately	Did not report our pre-specified outcomes	Women did not have GDM	
Ali et al 2010	•			
Coffman and Ray 2002		•		
Cosson et al 2012		•		
Dahlberg et al 1981			•	
Daniells et al 2003		•		
Doran and Davis 2011		•		
Kieffer et al 2002	•			
Kim et al 2005		•		
Kim et al 2007		•		
Kim et al 2008		•		
Hoedjes et al 2011		•		
Hoedjes et al 2012	•			
Jefferson et al 2000	•			
Lapolla et al 2012		•		
Kim and Vahratian 2010		•		
Koh et al 2010		•		
Lie et al 2011		•		
Morrison et al 2009		•		
Morrison et al 2012		•		
Nolan et al 2011	•			
Persily 1996		•		
Rhoads-Baeza and Reis 2012	•			
Rumbold and Crowther 2002		•		
Smith-Morris 2005	•			
Swan et al 2010		•		
Trutnovsky et al 2010	•			
Wylie et al 2011		•		

#### Appendix 2: Excluded Studies Table with Reasons

#### **Appendix 3: Included Studies Records (refer to Bibliography for full references)**

#	Study ID	Article type
	Australia	
1	Bandyopadhyay 2011 (Bandyopadhyay et al 2011)	Full-text
2	Carolan 2013 (Carolan 2013;Carolan, et al. 2012)	Full-text
3	Graco 2009 (Graco et al 2009)	Full-text
4	Doran 2008 (Doran 2008)	Full-text
5	Morrison 2010 (Morrison, et al. 2010)	Full-text
6	Carolan 2010 (Carolan et al 2010;Carolan et al 2010)	Full-text
7	Razee 2010 (Razee, et al. 2010)	Full-text
8	Smith 2005 (Smith et al 2005)/Zehl 2008(Zehle et al 2008)	Full-text
9	Sterne 2011 (Sterne, et al. 2011)	Full-text
10	Swan 2007 (Swan et al 2007)	Full-text
	United States	
11	Jones 2012 (Jones and Appel 2011;Jones et al 2012)	Full-text
12	Hoy-Rosas 2011 (Hoy-Rosas and Lancaster 2011)	Abstract
13	Bennett 2011 (Bennett et al 2010;Bennett et al 2010;Bennett et al 2011)	Full-text
14	Collier 2011 (Collier, et al. 2011)	Full-text
15	Downs 2006 (Downs and Ulbrecht 2006)	Full-text
16	Bieda 2009 (Bieda 2009)	Thesis (full-
		text)
17	Kim 2007 (Kim, et al. 2007) (Risk = "a")	Full-text
18	Kim 2007 (Kim, et al. 2007) (Racial = "b")	Full-text
19	Lawson 1994 (Lawson and Rajaram 1994)	Full-text
20	Nicklas 2011 (Nicklas et al 2010;Nicklas et al 2011)	Full-text
21	Remsberg 2012 (Remsberg 2012)	Abstract
22	Segall-Gutierrez 2011 (Segall-Gutierrez et al 2011)	Abstract
	Canada	
23	Evans 2005 (Evans and O'Brien 2005)	Full-text
24	Evans 2010 (Evans et al 2010)	Full-text
25	Feig 1998 (Feig et al 1998)	Full-text
26	Gaudreau 2012 (Gaudreau and Michaud 2012)	Full-text
27	Keely 2010 (Keely, et al. 2010)	Full-text
28	Keely 2012 (Keely 2012)	Commentary
29	Neufeld 2011 (Neufeld 2011)	Full-text
	Europe/UK	
30	Stage 2004 (Stage et al 2004)	Full-text
31	Hjelm (Hjelm et al 2009;Hjelm, et al. 2007;Hjelm, et al. 2005;Hjelm et al 2012;Hjelm	Full-text
	et al 2011;Hjelm et al 2008)	
32	Lindmark 2010 (Lindmark, et al.)	Full-text

33	Persson 2010 (Persson et al 2010)	Full-text
34	Clarke 2012 (Clarke et al 2012)	Full-text
35	Trutnovsky 2012 (Trutnovsky et al 2012)	Full-text
36	Bell 2011 (Bell, et al. 2011)	Abstract
37	Wylie 2011 (Wylie, et al. 2011)	Abstract
38	Wylie 2012 (Wylie, et al. 2012)	Abstract
	South America	
39	Saloman 2004 (Salomon and Soares 2004)	Full-text
40	Soares 2006 (Soares and Santos 2006)	Full-text
	Asia	
41	Doran 2010 (Doran and Davis 2010)	Full-text
42	Hirst 2012 (Hirst et al 2012;Hirst et al 2012)	Full-text

Abbreviations are: not specified (NS).

Appendix 4: Questionnaire (See Overleaf)



### **'DIAMIND'**



Postpartum re<u>mind</u>ers to test for type 2 <u>dia</u>betes in women who have experienced gestational diabetes mellitus

Dear.....,

### Thank you for taking part in the DIAMIND study.

The DIAMIND study is aiming to increase attendance for oral glucose tolerance testing for type 2 diabetes or prediabetes after birth in women who have experienced gestational diabetes. Many thanks for participating in this study and completing this questionnaire. Your response is highly valued, and it will help to improve healthcare for women who have experienced gestational diabetes. You will be provided with the results of the study once it is complete.

1. In this study, a text message reminder system was used to try to encourage attendance for oral glucose tolerance testing after birth. What type of reminder system would you have preferred? (Please tick one box)

Postal (mail)	SMS text (used in this study)

Email Voice call

Unsure/Don't know

- 2. What's your current weight (in kilograms)? \_\_\_\_\_kg
- 3. For how long did you breastfeed your baby?

I did not breastfeed	First week	First month

First 3 months First 6 months Currently breastfeeding

Unsure/Don't know

#### 4. Have you had an oral glucose tolerance test since your last baby was born?

Yes  $\rightarrow$  please go to page 2 and answer questions 5-7

No  $\rightarrow$  please skip to page 3 and answer question 8 onwards

Unsure/Don't know  $\rightarrow$  please skip to page 3 and answer question 8 onwards



## **'DIAMIND'**



Postpartum re<u>mind</u>ers to test for type 2 <u>dia</u>betes in women who have experienced gestational diabetes mellitus

**<u>Page 2</u>**: Questions for women who <u>have had</u> an oral glucose tolerance test in the six months after the birth of their baby

If you have not had an oral glucose tolerance test since the birth of your baby, please skip to page 3.

5. When did you have an oral glucose tolerance test after the birth (if known)?

Date of test:		/	/
	DD	MM	YY

#### 6. Please provide your oral glucose tolerance test result (after birth), if known:

Fasting blood glucose result \_\_\_\_\_\_ OGTT two hour result\_\_\_\_\_\_

7. What did you think about taking the oral glucose tolerance test (since leaving hospital after the birth of your baby)?

It was easy to <b>fast</b> for the oral glucose tolerance test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree
(fasting is not eating or drinking anything except water for several hours)					
It was easy to find time to take the oral glucose tolerance test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree
Overall, I was happy with the oral glucose tolerance test experience	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree

The questions that follow are for women who have not had an oral glucose tolerance test since their most recent birth. If you have had an oral glucose tolerance test and completed questions 1-7, you have now completed the survey. Many thanks!



## **'DIAMIND'**



Postpartum re<u>mind</u>ers to test for type 2 <u>dia</u>betes in women who have experienced gestational diabetes mellitus

<u>Page 3:</u> Questions for women who <u>have not had</u> an oral glucose tolerance test in the six months after the birth of their baby

#### 8. Do you plan to have an oral glucose tolerance test in the near future?

Yes

No

Unsure

9. Please let us know what prevented you from having an oral glucose tolerance test in the six months after the birth of your baby. *Tick all the boxes that apply to you.* 

Concern or anxiety about being diagnosed with type 2 diabetes	Transport not available or inadequate
Baby's health	I didn't know I needed an OGTT after
Not enough time	I'm healthy or at low risk of type 2 diabetes so it wasn't necessary
Feeling down or low	I thought the test would be too unpleasant
Childcare not available or inadequate	I didn't know where to go for the test

Are there any other reasons that you did not have an oral glucose tolerance test? *Please write any other reasons in the box below:* 

10. Is there anything that would have helped you to have had an oral glucose tolerance test in the six months after the birth of your baby?



HbA1c test

# **'DIAMIND'**



Postpartum re<u>mind</u>ers to test for type 2 <u>dia</u>betes in women who have experienced gestational diabetes mellitus

<b><u>Page 4</u></b> : Questions for women who <u>have not had</u> an oral glucose tolerance test in the six months after the birth of their baby						
<b>11a. Have you had a fasting blood glucose test since leaving hospital after the birth of your baby, instead of an OGTT?</b> Sometimes a fasting blood glucose test is used to diagnose type 2 diabetes instead of an oral glucose tolerance test						
Yes	No		Unsure			
11b. If known, plea	se provide y	our fasting	blood glucose	result:		
11c. What did you	think of the	fasting bloo	d glucose test?	?		
It was easy to <b>fast</b> for the fasting blood test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree	
(fasting is not eating or drinking anything except water for several hours)						
It was easy to find time to take the fasting blood glucose test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree	
Overall, I was happy with the fasting blood glucose test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree	
<b>12a. Have you had a HbA1c (glycated haemoglobin) test since leaving hospital after the birth of your baby?</b> This test does not require fasting. It indicates the glucose levels in your blood over the last 3 months, and is sometimes used for diagnosis of type 2 diabetes.						
Yes		No	Un	sure		
12b. If known, please provide your HbA1c (glycated haemoglobin) result:						
12c. What did you	think of the	HbA1c test?	?			
It was easy to find time to take the HbA1c test	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree	
Overall, I was happy with the	Strongly agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree	

Many thanks for completing our survey!