



	Total Study Population (n)
Women in study	405
Pregnancies	418
Multiples	8
Total babies	426
Early fetal losses < 20 weeks	30
Late fetal losses 20-24 weeks	6
Pregnancies ongoing >24 weeks	382
Live births delivered > 24 weeks	373
Live births delivered < 24 weeks	0
Total live births	373
Neonatal deaths	3 *
Total live at 28 days	370
Stillbirths	17
All births registered	390
Congenital malformations	32

		N	umber of pregnanci	es
		Type 1	Type 2	All
		n=269	n=149	n=418
	Median age of			
	mother at entry to			
	study, Years	29 (9)	33 (8)	31 (9)
	(IQR)			
	Ethnicity n (%)			
	European	249 (92.6)	64 (43.0)	313 (74.9)
	Black African	0	4 (2.7)	4 (1.0)
	Black Caribbean	5 (1.9)	12 (8.1)	17 (4.1)
	Indian	7 (2.6)	11 (7.4)	18 (4.3)
	Pakistani	3 (1.1)	45 (30.2)	48 (11.5)
	Bangladeshi	0	7 (4.7)	7 (1.7)
	Other	5 (1.9)	6 (4.0)	11 (2.6)
	Parity n (%)			
	Primigravida	94 (34.9)	21 (14.1)	115 (27.5)
	Multigravida	175 (65.1)	128 (85.9)	303 (72.5)
	Median age at			
	onset of diabetes,	13 (13.25)	28 (9)	20 (16)
	years (IQR)			
(iii	Median duration	44 (40.05)		0 (10)
4B	of diabetes, years	14 (13.25)	4 (4)	9 (13)







(Referenced separately at the end of the document)

- 1 Preconception counselling
- 2 Prepregnancy folic acid.
- 3 First trimester ultrasound scan for accurate dating.
- 4 Corticosteroids for early delivery.
- 5 HbA1C < 7% (standard reported on in chapter IV).
- 6 Hypoglycaemia events / Glucagon use.
- 7 Retinal screening in first trimester.
- 8 Neonatal facilities.
- 9 Mode and timing of delivery discussed.
- 10 Continuous electronic fetal heart monitoring.
- 11 IV glucose and insulin for delivery.

Preconception counselling:

perinatal

Standard: A preconception clinic should be run jointly by the adult diabetes service and the maternity service for women with diabetes wishing to become pregnant.

Table 4.1a Documentation of pre pregnancy counselling and control

	Type 1 (%) n = 269	Type 2 (%) n = 149	All women (%) n = 418
Pre-pregnancy counselling documented	82 (30.4%)	36 (24.2%)	118(28.2%)
Pre-pregnancy HbA1C	125 (46.4%)	48 (32.2%)	173 (41.4%)
Pre-pregnancy HbA1C < 7% in those that had a	31 (24.8%)	22 (45.8%)	53 (30.6%)

	Pre- pregnancy	Test closest to 10 wks	Test Closest to 20 wks	34 weeks onwards
Type 1 (n=269)	125 (46.5%)	236 (87.7%)	226 (83.3%)	220 (81.8%)
Type 2 (n=149)	48 (32.2%)	111 (74.5%)	122 (84.0%)	116 (77.9%)
All (n = 418)	173 (41.4%)	347 (83%)	348 (81.9%)	336 (80.4%)

Γ		Pre-	10 weeks	20 weeks	34 weeks
	All Women: Result <7% n(%) ª	pregnancy 53 (30.6)	155 (44.7)	244 (70.1)	230 (68.5)
	Women with Type 1 diabetes: Result <7% n(%) ^a	31 (24.8)	95 (40.3)	149 (65.9)	141 (64.1)
	Women with Type 2 diabetes: Result <7% n(%) ^a	22 (45.8)	60 (51.4)	95 (77.9)	89 (76.7)

	Table 6.3: Mo general materi	des of deliver nity populatio	y for all won n.	nen compared	to National	dataset and
	MODE	Type 1 (n = 247)	Type 2 (n = 135)	All (n = 382)	National Dataset	Normal Population
	Vaginal Spontaneous Instrumental	58 (23.5%) 44(17.8%) 14 (5.7%)	49 (36.3%) 45 (33.3%) 4 (3.0%)	107 (28%) 89 (23.3%) 18 (4.7%)	32.1% 24.4% 7.7%	67% 11%
	C Section Emergency Elective	188 (76%) 107 (43%) 81 (33%)	86 (60%) 41 (29%) 45 (30%)	274 (71.7%) 148 (38.7%) 12 (33%)	67.4% 37.6% 29.8%	22% 13% 9%
	Not vaginal delivery	1 (%)	0	1 (%)	0.2%	0%
Ger				I		I

Key Findings : Labour and Delivery 28% vaginal delivery rate in WM – 14% spontaneous labour 72% CS rate – primarily due to obstetric reason - maternal diabetic complications/maternal request are rare 1 in 3 women are induced – 60% vaginal del rate 32% preterm delivery rate

	Table 8.3: Neor delivery (percen number of babi 'missing')	natal admissio tages are pro es with valid	n at any time fo portion of babie response, i.e.	llowing delivery, es in category excluding 'not	by gestation a out of the tot applicable'an
	G	iestation at d	elivery, complet	ed weeks n (%)
	Type of Neonatal Care	<32 (n=16)	32-36 (n=98)	37+ (n=259)	All (n=373)
	Special Care	4 (25.0%)	39 (39.8%)	54 (20.8%)	97 (26.0%)
	High Dependency Care	5 (31.3%)	28 (28.6%)	45 (17.4%)	78 (20.9%)
	Intensive Care	6 (37.5%)	16 (16.3%)	22 (8.5%)	44 (11.8%)
	Other Specialist Care	0	0	3 (1.2%)*	3 (0.8%)
	Postnatal ward normal care with Mother	0	15 (15.3%)	133 (51.4%)	148 (39.7%)
\sim	Not Known	1 (6.3%)	0	2 (0.8%)	3 (0.8%)

	Table 9.3: Reasons for admission to information is given:	o special care in 222 babies where
	Reason for Admission	All infants (n = 222)
	Routine/observation/infant of diabetic mother/maternal/not known	54 (24.3%)
	Hypoglycaemia/feeding problem	85 (38.3%)
	Prematurity	52 (23.4%)
	Respiratory /Ventilated	23 (10.4%)
	Other infant medical problems	8 (3.6%)
perinatal		



	Early fetal loss<20/40	Late feta Loss >20	Stillbirth	Neonatal Death	Alive at 28 days	Grand Total
		23+6				
Cases	2	6	15	3	13	39
Type 1 Type 2	1 1	5 1	8 7	1** 2	8 5	23 16
Congenital Anomalies	2	4	3	2	11	22
Controls	0	0	0	0	38	38
Type 1 Type 2	0	0 0	0	0	20	

	Cases (n=39)	Controls (n=38)
First contact with health professional (From EDD) Days //	QR 47 (25.5)	48.5 (17.75
First hospital appointment (from EDD) Days / IQR	62 (19)	67 (31)
Health professional involved with care (%)		
DNS	33 (84.6)	31 (81.6)
Physician	37 (94.9)	37 (97.4)
Dietician	19 (48.7)	18 (47.4)
Midwife	5 (12.8)	5 (13.2)
Obstetrician	37 (94.9)	36 (94.7)

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Specific diabetes complication	Women in the enqui n/N (%)
Recurrent hypoglycaemia during pregnancy	37/71 (52.1)
Hypoglycaemia requiring external help	12/56 (21.4)
Retinal assessment performed in1st trimester	43/65 (66.2)
Evidence of Retinopathy in pregnancy	22/55 (40.0)
Pre-existing retinopathy	9/55 (16.4)
Pre-existing retinopathy - deteriorating	4/55 (7.3)
New retinopathy	7/55 (12.7)
Nephropathy	3/71 (4.2)

	Cases n= 39		Controls n= 38	
	N	3 7 %	N	1= 38 %
Diabetic care during pregnancy (other than glycaemic control)	23/37	62	25/38	66
Maternity Care - antenatal	23/39	59	15/38	40
Maternity care - delivery	13/37	35	14/37	38

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Pai	nel comments on antepa	rtum c	are	
		Cases (n23)	Controls (n15)	
	Issues relating to provision of health service Clinical practice - Fetal survelliance -Management of pregnancy complications -Mode / timing of delivery -Use of steroids -Need for senior Obstetric input -Poor / no management plan -Blank	7 4 2 0 3 3	5 10 2 1 1 2 2	
	Resources -In-appropriate appointments -Use of interpreter -Neonatal cots	3 1 0	0 0 1	
	Communication -Between professionals -Between professionals & woman	3 0	1 0	
perinatal	Issues relating to woman / family Maternal actions - Smoking Maternal compliance Poor attendance	1 1 1	1 0 0	
institute	Total Comments	31	26	



	Early Fetal loss <20/40	Late fetal Loss >20-23+6	Stillbirth	Neonatal Death	Alive at 28 days	Grand Total
Cohort	1	1	1	1	I	
Poor Outcomes	13	1	8	2	9	33
Congenital Anomalies	1	0	1	1	9	12
Good Outcomes	0	0	0	0	116	116
Total						149



<u>1ype 2 –</u>	glycaem	nic control	
		Type 2 Cohort poor outcomes n33	Type 2 Cohort - live excl. NND and anomaly n116
HbA1c Values		n=33	n=116
Pre-pregnancy	Mean	9.0% (n10)	7.4% (n38)
10 week	Mean	7.7% (n24)	6.7% (n87)
20 week	Mean	6.4% (n18)	5.9% (n103)
34 week	Mean	6.8% (n12)	6.2% (n104)



<u>Type 2 r</u>	node of c	delivery	
Mode of delivery	European	Non- European	Total
SVD	12 (21)	33 (43)	45 (33)
Instrumental	2 (3)	2 (3)	4 (3)
Emergency CS	15 (26)	26 (34)	41 (30)
Elective CS	29 (50)	16 (21)	45 (33)
Fetal losses	6	8	14
Total	64	85	149



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	Panel Enquiry comments	
	no diabetic input in first trimester, referred to wrong consultant	
	no information prior to pregnancy	
	No folic acid. V poor communication, poor use of interpreters.	
	poor GP care	
	not sent for earlier assessment	
	no folic acid, metformin not stopped	
	incorrect dose of folic acid No folic acid, no pre pregnancy care	
	very poor glycaemic control pre conception.Not interested in weight loss & no clear targets	
	not sufficient urgency to refer	
	not advised not to get pregnant	
NE	Fructosaminine difficult to interpret, no home monitoring records, unusual insulin regime	
porinatal	No dietician. No targets	
institute		





Ethnicity	Type 1 (n=20) (%)	Type 2 (n=12) (%)	All (n=32) (%)
European	18 (90.0)	7 (58.3)	25 (78.1)
Black African	0	1 (8.3)	1 (3.1)
Black Caribbean	1 (5.0	0	1 (3.1)
Indian	1 (5.0)	0	1 (3.1)
Pakistani	0	2 (16.7)	2 (6.3)
Bangladeshi	0	1 (8.3)	1 (3.1)
Other	0	1 (8.3)	1 (3.1)

	All (n=297)	Type 1	Type 2	vs. WM	National CEMACH	vs National
1		(n=188)	(n=109)	region ¹ RR	data RR	maternity pop RR ^
10	38.0	35.7	42.1	6.0	1.4	6.7
3	11.9	6.2	22.0	2.5	1.3	3.3
12*	45.6	41.7	52.6	4.5	1.4	5.4
omprises 1 /est Midlan	0 stillbirths a ds Perinatal	and 2 early Mortality	/ neonatal de (Update Marc	aths; RR = rel ch 2006)	ative risk	
	3 12* omprises 1 /est Midlan	3 11.9 12* 45.6 omprises 10 stillbirths a /est Midlands Perinatal	3 11.9 6.2 12* 45.6 41.7 omprises 10 stillbirths and 2 early /est Midlands Perinatal Mortality	3 11.9 6.2 22.0 12* 45.6 41.7 52.6 omprises 10 stillbirths and 2 early neonatal de lest Midlands Perinatal Mortality (Update Marce)	3 11.9 6.2 22.0 2.5 12* 45.6 41.7 52.6 4.5 omprises 10 stillbirths and 2 early neonatal deaths; RR = relivest Midlands Perinatal Mortality (Update March 2006) RR = relivest Midlands Perinatal Mortality (Update March 2006)	3 11.9 6.2 22.0 2.5 1.3 12* 45.6 41.7 52.6 4.5 1.4 omprises 10 stillbirths and 2 early neonatal deaths; RR = relative risk //est Midlands Perinatal Mortality (Update March 2006) 1.4



	SB/NND (no anomaly) A n=13	Panel Anomaly* B n=22	Panel control C n=38	Cohort - live excl. NND and anomaly D <i>n=350</i>
Pre-preg	13.6	9.1	7.6	7.7
10 week	7.8	7.9	7.2	7.1
20 week	6.6	6.1	6.3	6.2
34 week	7.1	6.8	6.3	6.3
⊙Signific	cant Stillbirth/ perina ⊙ethnic minoi ⊙history of pr ⊙1st and 3rd tr	tal mortality as rities evious preterm la imester subopti	sociations : abour mal glycaemic o	control

