

In conjunction with the



40000	NA CRESON																
Patient's Last Name Patient's First Name																	
Address – number, street name						Apt	/Suite/Unit										
City/Town Province					e Pos	stal Code	F	Partner's Last Name					Partner's First Name				
Telephone - Home Telephone - Work Language				ige		F	Partner's Occupation				Partner's Educational level Age			Age			
Date of birth  YYYY/MM/DD  Age Occupation E				Educati	onal level	E	Ethnic or Racial backgrounds: Mother / F				her / Father	her					
OHIP No.				Patient File No.			Marital status		E	Birth attendant		Newborn care		Family Physician		an	
Allergies or Sensitivities (describe reaction details)							ľ	Medications	s/Herba	ls							
								Pregna	ancy S	Summary							
LMP Cycle		Y/MM/DD			Certai Regula			No □ No □	EDB	EDB (by dates)			Fi	nal EDB		Dating Methor ☐ Dates	
Cycle q Contraceptive type				•			/YYY/MIV									<ul><li>□ T₁US</li><li>□ T₂US</li></ul>	
Gravida Term			Term		Premature		Abortuses		•	Living						☐ ART (e.g	g. IVF)
								Obste	etrical l	History			1				
No.	Year	Sex M/F	Gest. age (weeks)	Birth weight		gth of our	Place of birth	Type of delivery		Comments regarding pregnancy and birth							
Medical History and Physical Exam (provide details in comm							nts)	Initial Laboratory Investigations									
Current Pregnancy 1. Bleeding Y / N			Y/N	Genetic History  22. At risk population (e.g.: Ashkenazi, consanguinity, sickle cell, Tay Sachs, thalassemi			Y/N			opulation Y / N //PE, PIH/HT, pression, thyroid) amination			Test	Result		Test	Result
			Y/N Y/N									Hb			HIV		
4. Alcohol, street drugs Y /			Y/N	Family history of:			,					MC	MCV		☐ Counseled and test of		st declined
<ul><li>5. Occup/Environ. risks</li><li>6. Dietary restrictions</li><li>Y / N</li></ul>				23. Developmental delay 24. Congenital anomalies			Y/N Y/N	Y/N				ABO	ABO		Last Pap		
7. Calcium adequate Y /			Y/N	25. Chromosomal disorder			rs Y/N	1 / IN		/t		Rh			YYYY/MM/DD		
8. Preconceptual folate Y / N 26. Genetic disorders				Y/N	ВМІ	BP			Anti	body Screen		GC/C	hlamydia				
	al Histor		V (N)	Infectious Disease			N/ (NI	//N 39. Thyroid				Rubella immune			Urine C&S		
Hypertension     Endocrine			Y/N Y/N	27. Varicella susceptible 28. STDs / HSV / BV		Y/N Y/N	N 40. Chest N 41. Breasts			N / Abn N / Abn		sAg					
11. Urinary tract 12. Cardiac/Pulmonary		Y/N	29. Tuberculosis risk		Y/N				N / Abn	VDRL							
13. Liver, hepatitis, Gl			Y/N Y/N				Y/N	43. Abdomen		N / Abn Extrm. N / Abn talia N / Abn a N / Abn N / Abn N / Abn		Sick	de Cell				
,			Y/N Y/N	=			V / N	44. Varicositie Y / N 45. External g				Prenatal Genetic I			nvestigations		Result
16. Surgery Y			Y/N	32. Relationship problems			Y/N	Y / N 46. Cervix, va				a)	All ages-MSS, IPS, FTS				
17. Blood transfusion Y			Y/N Y/N	<ul><li>33. Emotional/Depression</li><li>34. Substance abuse</li></ul>			Y/N Y/N	Y / N 47. Uterus Y / N 48. Size:					Age ≥ 35 at EDB-CVS/amnio				
<ol> <li>Anaesthetic compl.</li> <li>Psychiatric</li> </ol>			Y/N	35. Family violence			Y/N	49. Adnexa			NI / Abn			o declined, or twins, then MSAFP			
20. Epilepsy/ Neurological Y /			al Y/N Y/N	•			Y/N Y/N			1	N / Abn			Counseled and test declined, or too late			
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Signature	Date	Signature	Date

## **A Guide to Pregnancy Assessment**

In the event of maternal transfer, please photocopy the front sheet and send to referral hospital.

This assessment system is intended as a basis for planning the on-going management of the pregnancy and should reflect local resources. The risk factors or problems listed below are intended as examples only.

Health	y Pregnancy, no predictable risk:		
	No pregnancy complications now or in the past  No significant maternal medical disease		No prior perinatal morbidity or mortality Fetal growth adequate
Pregna	ancy at risk:		
an app manag	ropriate specialist (obstetrician, internist, pediatric led by continuing collaborative care and birth in an e returned to the care of the referring provider with	ian, <i>etc.</i> obstetri	nancy may be necessary. In addition, consultation with may also be necessary. These patients may be ical unit with intermediate level nursing facilities OR the ested plan of management for the remainder of the
Prior pi	Diabetes, White Classes B, C, or D Chronic hypertension Other significant medical illness Obesity (BMI ≥ 35) Significant tobacco, alcohol, drug use Severe psychosocial issues Family history genetic disease or congenital anomalies Other significant family history, esp. DVT/PE and recurrent pregnancy losses regnancy history of: Preterm labour < 36 weeks Stillbirth or neonatal death Intrauterine growth restriction Previous uterine surgery including lower segment Cesarean section Cervical incompetence		Gestational hypertension Placenta previa (with or without bleeding) Other significant antepartum hemorrhage Twin pregnancy Gestational diabetes (White Class A) Abnormal fetal growth (suspected intrauterine growth restriction or large for dates) PROM 32-36 weeks Preterm labour 32-36 weeks Rh or atypical blood group sensitization Hydramnios or oligohydramnios Fetal malposition (breech, transverse) at 36 weeks Postdates ≥ 41 weeks Anemia not responding to Fe (Hb <100 g/l)
Pregna patient patient prolaps	s should be transferred to a regional perinatal cen	tre (leve vith prob	ner are obviously in danger. If at all possible, these el III) for intensive care and birth. Clearly, there are blems such as excessive antepartum bleeding, cord fetus or mother.
	High order multiple gestation (triplets or greater) Fetal congenital anomaly Diabetes beyond Class D (end-organ involvement) Renal disease with hypertension $\pm \downarrow$ function Heart disease, especially with failure Other significant severe medical illness	[ [ [	Pregnancy < 32 weeks with:  Preterm labour and/or premature rupture  Gestational hypertension with adverse conditions  Antepartum hemorrhage ongoing  Oligohydramnios  IUGR, ≤10 <sup>th</sup> %, reverse flow Doppler

**Two or more** risk problems can combine to produce a high pregnancy risk. Such a patient may need to be placed in a higher risk category