



HRN: Site: DOB: yyyy/mon/dd

Last Name: First and Additional Names:

PHN: Gender: Age in Years:
Admitting Physician: Encounter #:

Address: Street, City, Province, Postal Code

Telephone Number:
Date of Admission: yyyy/mon/dd Family Physician:

PHYSICIAN REFERRAL FORM

Please type directly into the form. Where indicated, required referral information may be attached.

Date ^(yyyy/Mon/dd) :	Refer to: Maternal Genomics	Fax: +1 (866) 566-7683
Referring Physician:		Phone:
Address:		Fax:
Family Physician:		
Referring Physician Practice ID:		Referring Physician License ID:

Referral Information

Reason for Referral: Non-invasive prenatal testing (NIPT)	
Type of referral: <input type="checkbox"/> New Referral <input type="checkbox"/> Re-referral <input type="checkbox"/> 2 nd Opinion <input type="checkbox"/> Urgent Referral <input type="checkbox"/> Service/consultant is aware of urgent referral Reason for urgency:	
Test Indications (choose at least one): <input type="checkbox"/> Advanced maternal age (≥35 years) <input type="checkbox"/> Positive serum screen <input type="checkbox"/> Abnormal ultrasound <input type="checkbox"/> History suggestive of increased risk for the specified aneuploidies <input type="checkbox"/> Low risk/maternal anxiety <input type="checkbox"/> Other:	
Past medical history (if relevant)	<input type="checkbox"/> Attached
Current medication/Allergies	<input type="checkbox"/> Attached
Gestational age: Weeks _____ Days _____ As estimated on ^{yyyy-Mon-dd} :	
Dating Method: <input type="checkbox"/> LMP <input type="checkbox"/> Date of implantation <input type="checkbox"/> CRL (Please attach ultrasound(s), if applicable.) <input type="checkbox"/> Other (please specify):	

NIPT Selection

<p>NIPT Test Chosen:</p> <p><input type="checkbox"/> Singleton Verifi Prenatal Test (chromosomes 21, 18, 13)</p> <p style="padding-left: 20px;"><input type="checkbox"/> Additional option – Sex chromosome aneuploidies (MX, XXX, XXY, and XYY)</p>	<p><input type="checkbox"/> Verifi Plus Prenatal Test (chromosomes 21, 18, 13) (Singleton only)</p> <p style="padding-left: 20px;"><input type="checkbox"/> Additional option – Microdeletions (1p36 deletion, 4p- (Wolf-Hirschhorn syndrome), 5p- (Cri-du chat syndrome), 15q11.2 (Prader-Willi syndrome), Angelman syndrome), 22q11. Deletion (DiGeorge syndrome)</p> <p style="padding-left: 20px;"><input type="checkbox"/> Additional option – Sex chromosome aneuploidies (MX, XXX, XXY, and XYY)</p> <p style="padding-left: 20px;"><input type="checkbox"/> Additional option – All chromosome aneuploidies (MX, XXX, XXY, and XYY)</p>
<p><input type="checkbox"/> Twin Verifi Prenatal Test (chromosomes 21, 18, 13)</p> <p style="padding-left: 20px;"><input type="checkbox"/> Additional option – Presence of Y chromosome</p>	
<p>Factors that may affect consultation/care</p>	<p><input type="checkbox"/> Language:</p> <p style="padding-left: 20px;"><input type="checkbox"/> Interpreter needed</p> <p><input type="checkbox"/> Physical limitations</p> <p><input type="checkbox"/> Social/psychological</p> <p><input type="checkbox"/> Specific patient request:</p>

Maternal Genomics use only

Referral received by:	Signature:
Received date ^{yyyy-Mon-dd} :	