

For scanning and accuracy, affix patient label within this outlined box.

Patient Name:

MRN No:

Only use black ball point pens to ensure scanning integrity.  
**No sharpies, highlighters, or red ink please.**

## Non-Invasive Prenatal Testing Test Requisition Form

### Referring Hospital & Clinician Details

Hospital Name			
Ordering Clinician's Name		Health Authority and License No.	
Email ID		Phone Number	
<b>Assisting Clinician Details</b>			
Name		Phone Number	
Email ID			

### Patient Details

Full Name			
Date of Birth	DD/MM/YYYY	Age	
Emirates ID or Passport # (specify Citizenship in case of passport)		Phone Number	
Email ID			

### Clinical Information

Date of Blood Draw DD/MM/YYYY	Gestation Weeks/Days	Estimated Due Date DD/MM/YYYY	Maternal Weight (kg)	Maternal Height (cm)	BMI
<b>Single Pregnancy</b>	<input type="checkbox"/>	<b>Test Indications</b>		<b>Patient's Medical History</b>	
<b>Twin Pregnancy</b>	<input type="checkbox"/>	Advanced Maternal Age	<input type="checkbox"/> No <input type="checkbox"/> Yes	Received allogeneic blood transfusion	<input type="checkbox"/> No <input type="checkbox"/> Yes, when?
<b>Test Request</b>		Parental Anxiety	<input type="checkbox"/> No <input type="checkbox"/> Yes	Received heparin therapy	<input type="checkbox"/> No <input type="checkbox"/> Yes, when?
<input type="checkbox"/> 6000048 NIPT Core, with gender		Abnormal results of other prenatal screening tests	<input type="checkbox"/> No <input type="checkbox"/> Yes, Please specify	Received immunotherapy and/or human serum albumin therapy	<input type="checkbox"/> No <input type="checkbox"/> Yes, when?
<input type="checkbox"/> 6000049 NIPT Core, without gender			<input type="checkbox"/> No <input type="checkbox"/> Yes, Please specify		Diagnosed with vanishing twin syndrome
<input type="checkbox"/> 6000050 NIPT Comprehensive, with gender		Abnormal Reproductive History	<input type="checkbox"/> No <input type="checkbox"/> Yes, Please specify	Received assisted reproductive technology treatment	
<input type="checkbox"/> 6000051 NIPT Comprehensive, without gender		Previous pregnancy affected by chromosomal/genetic disorder	<input type="checkbox"/> No <input type="checkbox"/> Yes, Please specify		History of Bone Marrow Transplant
<b>Core (Common Trisomies):</b> Tests for Trisomy 21, 18, 13, and sex chromosome aneuploidies (SCAs) with the gender option.		Family history of genetic disorder/s or syndrome	<input type="checkbox"/> No <input type="checkbox"/> Yes, Please specify		
<b>Comprehensive (Genome-wide):</b> Includes all autosomal aneuploidies, rare trisomies (e.g., chromosomes 22, 16), microdeletion syndromes (e.g., DiGeorge, Prader-Willi), copy number variants (CNVs ≥7 Mb) and SCAs with gender the option.					

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### Referring Clinician's Statement

I acknowledge that the sample may not meet the acceptance criteria for the following reasons:

- Samples arrive after 48 hours.
- Gestational age below 10 weeks.
- BMI over 40.

I understand that these factors may affect the accuracy or validity of test results.

I will ensure that my patient is aware of these considerations and is willing to proceed with the testing.

Not Applicable

I verify that the patient and prescriber information in this form is complete and accurate to the best of my knowledge and that I have requested this screening test based on my professional judgement of medical necessity. I confirm that the patient understands the purpose, limitations, potential risks, scope and performance of this test and that I have answered any questions to the best of my ability. The patient has provided informed consent for this test.

Name of Physician

Signature

Date

DD/MM/YYYY

Place

### Collection Information

Collection Date DD/MM/YYYY

Collection Time

AM  PM

Sample Type  Peripheral Blood  Plasma

Collector's Signature

### Billing Details

Client Billed

Patient Billed

Insurance Billed

(Submit EID and Insurance Card)

### Receiving Lab Information

Laboratory Access Number

Date Received

DD/MM/YYYY

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