	Title	CODE UK_M_F_033	VERSION 1.0
	Pre-PGT-M Test Requisition Form	Author (Name): Araz Raberi & Roy Pascal Naja	Date of issue: 28/05/2019
		Authorized by (Name): Roy Pascal Naja	Date of next review: 28/05/2021
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Pre-Preimplantation Genetic Testing for Monogenic Disorders (PGT-M) Test Requisition Form

Samples will only be processed if all mandatory fields (in blue and italics) have been completed

PATIENT INFORMATION

Patient Surname
Patient Name

Patient date of birth
Unique patient ID

Partner Surname
Partner Name

Partner date of birth
Unique partner ID

Address

City/Town Country Postcode

Telephone Number Email address

FERTILITY CLINIC INFORMATION

Fertility clinic name
Date of test request

Clinician ordering test
Clinician email

- Complete the following fields **only** if not already registered with Igenomix UK

Fertility clinic address


City/Town Country Postcode

Telephone Number Email address

CYCLE INFORMATION


Confirm cycle type: Frozen cycle *Date of expected oocyte retrieval/thaw*

Confirm specimen type: Day 5/6 (Trophectoderm)


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REASON FOR REFERRAL

Patient	Last name/ Name <input type="text"/>		
	Date of Birth: <input type="text"/>	Relation to embryo <input type="text"/>	
	Please specify the presence (if any) of a blood borne transmitted disease <input type="text"/>		
Genetic disorder	Gene	Mutation	
Disorder 1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>	
Disorder 2 (If available): <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>	
Partner	Last name/ Name <input type="text"/>		
	Date of Birth <input type="text"/>	Relation to embryo <input type="text"/>	
	Please specify the presence (if any) of a blood borne transmitted disease <input type="text"/>		
Genetic disorder	Gene	Mutation	
Disorder 1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>	
Disorder 2 (If available): <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>	

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Family member 1	Last name/ Name <input style="width: 100%;" type="text"/>		
	Date of Birth <input style="width: 200px;" type="text"/>	Relation to embryo <input style="width: 150px;" type="text"/>	
	Please specify the presence (if any) of a blood borne transmitted disease <input style="width: 150px;" type="text"/>		
Genetic disorder	Gene	Mutation	
Disorder 1: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	1: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	1: <input style="width: 100%;" type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input style="width: 100%;" type="text"/>	
Disorder 2 (If available): <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	2: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	2: <input style="width: 100%;" type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input style="width: 100%;" type="text"/>	
Family member 2 (If available)	Last name/ Name <input style="width: 100%;" type="text"/>		
	Date of Birth <input style="width: 200px;" type="text"/>	Relation to embryo <input style="width: 150px;" type="text"/>	
	Please specify the presence (if any) of a blood borne transmitted disease <input style="width: 150px;" type="text"/>		
Genetic disorder	Gene	Mutation	
Disorder 1: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	1: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	1: <input style="width: 100%;" type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input style="width: 100%;" type="text"/>	
Disorder 2 (If available): <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	2: <input style="width: 100%;" type="text"/> OMIM No. <input style="width: 100%;" type="text"/>	2: <input style="width: 100%;" type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input style="width: 100%;" type="text"/>	
Family member 3 (If available)	Last name/ Name <input style="width: 100%;" type="text"/>		
	Date of Birth <input style="width: 200px;" type="text"/>	Relation to embryo <input style="width: 150px;" type="text"/>	
	Please specify the presence (if any) of a blood borne transmitted disease <input style="width: 150px;" type="text"/>		

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Genetic disorder	Gene	Mutation
Disorder 1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> OMIM No. <input type="text"/>	1: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>
Disorder 2 (If available): <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> OMIM No. <input type="text"/>	2: <input type="text"/> <i>Genetic status (e.g. Unaffected):</i> <input type="text"/>

CLINICIAN AUTHORIZATION

It is the referring clinician's responsibility to ensure that the patient/carer knows the purpose of the test and that the sample(s) may be stored for future testing related to specific diagnosis for the patient. In signing this form, the clinician confirms that they have obtained consent for testing and storage. The patient should be advised that the clinical information/sample(s) may be used anonymously for audits, quality assurance, research, and training purposes. Please advise of any restrictions.

<i>Clinician signature:</i> <input type="text"/>	<i>Date:</i> Click or tap to enter a date. <input type="text"/>
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FOR IGENOMIX UK USE ONLY

<i>Please fill sample reception table:</i>		
<i>Received by:</i> <input type="text"/>	<i>Time:</i> <input type="text"/>	<i>Date:</i> <input type="text"/>
<i>Sample accepted/rejected:</i>		
<i>If rejected, state the reason:</i>		