

PATIENT INFORMATION

First name	MI	Last name	Date of birth (MM/DD/YYYY)		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Sex	MRN (medical record number)	Ancestry			
<input type="radio"/> M <input type="radio"/> F	<input type="text"/>	<input type="radio"/> Asian <input type="radio"/> Black/African American <input type="radio"/> White/Caucasian <input type="radio"/> Ashkenazi Jewish <input type="radio"/> Hispanic <input type="radio"/> Native American <input type="radio"/> Pacific Islander <input type="radio"/> French Canadian <input type="radio"/> Sephardic Jewish <input type="radio"/> Mediterranean <input type="radio"/> Other: _____			
Email address (for report access after release by medical professional)			Phone		
<input type="text"/>			<input type="text"/>		
Address					
<input type="text"/>					
City	State	ZIP code	Country		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>		

ORGANIZATION INFORMATION

Organization name and address					
Organization name				Phone	
Address				Fax	
City	State	ZIP code	Country		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>		
Primary clinical contact					
Name				NPI	
Email address (for report access)				Phone	
Ordering physician					
<input type="radio"/> Same as primary clinical contact					
Name				NPI	
Email address (for report access)				Phone	
Additional clinical or laboratory contacts (optional)					
Name	Email address (for report access)	Name	Email address (for report access)		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>		
Name	Email address (for report access)	Name	Email address (for report access)		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>		

<input type="radio"/> INSURANCE BILLING (Please attach a copy of the patient's card.)	
We do not accept insurance for certain tests or patients outside the US. Before completing this section, confirm your test is eligible at www.invitae.com/billing.	
Primary insurance company name	Primary member ID#
<input type="text"/>	<input type="text"/>
Secondary insurance company name	Secondary member ID#
<input type="text"/>	<input type="text"/>
<input type="radio"/> Patient has Medicare and was treated as a hospital inpatient (>24 hour stay) in the last 14 days.	Prior-authorization #
<input type="text"/>	<input type="text"/>
Letter of Medical Necessity (LMN)	
<input type="radio"/> I have attached an LMN and/or other documents for insurance billing purposes.	
<input type="radio"/> I agree to allow Invitae to transfer the information from this requisition to an LMN and/or other documentation using the ordering physician's name as the signature for insurance billing.	

<input type="radio"/> INSTITUTIONAL BILLING
Invitae will send an invoice to the organization address above. Please contact Invitae if this order should be billed to a different location.

<input type="radio"/> PATIENT PAY BILLING
Invitae will send an electronic invoice to the patient email listed above

<input type="radio"/> OTHER BILLING
Invitae partner code:
<input type="text"/>

SPECIMEN INFORMATION

The specimen type for this test is trophectoderm (blastocyst) biopsy. Embryos must be frozen after biopsy, as we cannot accommodate fresh transfers.

Was ICSI performed? Yes No

**Samples derived from non-ICSI embryos may have an increased risk for false negative or false positive results.*

Does this case include a re-biopsy of previously tested embryo(s)? Yes No

If yes, please provide the original PAT# _____

PARTNER AND DONOR INFORMATION

Partner's first name

Partner's last name

Partner's sex

 M F

Partner's DOB (MM/DD/YYYY)

Partner's MRN

 Egg donor cycle? Yes No

Age of donor: _____

 Sperm donor cycle? Yes No

Age: _____

REASON FOR TESTING

Please indicate the reason for testing by selecting all that apply:

- | | |
|--|--|
| <input type="radio"/> Advanced maternal age | <input type="radio"/> Routine aneuploidy screening/patient request |
| <input type="radio"/> Repeat failed implantation/failed IVF cycles
of failed cycles: _____ | <input type="radio"/> Family balancing → MALE/FEMALE |
| <input type="radio"/> Recurrent pregnancy loss
of losses: _____ | <input type="radio"/> Male infertility |
| <input type="radio"/> Known structural chromosomal rearrangement (Please submit a copy of the PGT-SR Referral Form [available at www.invitae.com/forms] for clinical review prior to placing an order for structural chromosome rearrangements from known translocation carriers.) | <input type="radio"/> Other:
Please specify:

_____ |

TEST INFORMATION

PGT-A (aneuploidy, also known as PGS) identifies embryos with the correct number of chromosomes.

- Preimplantation genetic testing for aneuploidy (PGT-A) **WITH** sex reported
- Preimplantation genetic testing for aneuploidy (PGT-A) **WITHOUT** sex reported

PGT-SR (structural rearrangements) evaluates embryos for unbalanced chromosome rearrangements from a known translocation carrier. This test also screens for aneuploidy. If ordering PGT-SR, please select "Known structural chromosomal rearrangement" under reason for testing and submit the PGT-SR Referral Form. Once Invitae receives this form and required karyotypes, our medical team will determine within 1 to 2 business days whether your patient is a candidate for PGT-SR.

- Preimplantation genetic testing for structural rearrangements/translocations (PGT-SR) **WITH** sex reported
- Preimplantation genetic testing for structural rearrangements/translocations (PGT-SR) **WITHOUT** sex reported

To request a complimentary specimen collection kit visit
www.invitae.com/request-a-kit

SHIPPING INSTRUCTIONS

Please ship specimen overnight in insulated containers:

Attn: Invitae Client Services
1400 16th Street, San Francisco, CA 94103, USA

By signing this form, the medical professional acknowledges that the individual/family member authorized to make decisions for the individual (collectively, the "Patient") has been supplied information regarding and consented to undergo genetic testing, substantially as set forth in Invitae's Informed Consent for Genetic Testing (www.invitae.com/PGT-consent), and for orders originating outside the US, has been informed that the Patient's personal information and specimen will be transferred to and processed in the US. The Patient has further been informed and authorizes Invitae Corporation ("Invitae") and its designees to release information concerning testing to their insurer, if applicable, in order to process and/or appeal claims on behalf of the Patient. If a letter of medical necessity (LMN) has not been provided, the medical professional agrees to allow Invitae to transfer the information from this requisition to a LMN and/or other documentation using the medical professional's name as the signature for insurance billing. For amounts received directly, the Patient has agreed to remit payment to Invitae for testing services rendered. I acknowledge that I offered pre-test genetic counseling to the Patient, if required by their insurer. In addition to the above, I attest that I am the ordering physician, or I am authorized by the ordering physician to order this test, or I am authorized under applicable law to order this test.

Medical professional signature

Date