

| For Juno Genetics internal use only | Juno Genetics number | Date and time of reception | Received by | Case Status | State the reason for rejection |
|-------------------------------------|----------------------|----------------------------|-------------|---|--------------------------------|
| | | | | <input type="checkbox"/> Accepted <input type="checkbox"/> Rejected | |

**The sections marked with (*) are mandatory to fill in to request the test*

| REFERRING CLINIC DETAILS | | | |
|--------------------------|--|-----------------------|--|
| Referring Clinic * | | Referring Clinician * | |
| Clinician's email * | | | |

| FEMALE PATIENT CLINICAL INFORMATION | | | |
|--|--|----------------|--|
| Surname/Name * | | DOB * | DD/MM/YYYY |
| Clinic ID * | | Gamete donor * | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Sample type * | <input type="checkbox"/> Blood <input type="checkbox"/> Saliva/ buccal swab <input type="checkbox"/> DNA <input type="checkbox"/> Other: _____ | | |
| 1 st | Genetic Disorder * | Gene * | Mutation * |
| | | | |
| | OMIM# | OMIM# | |
| | Genetic Status * | | |
| <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |
| 2 nd | Genetic Disorder * | Gene * | Mutation * |
| | | | |
| | OMIM# | OMIM# | |
| | Genetic Status * | | |
| <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |

| MALE PATIENT CLINICAL INFORMATION | | | |
|--|--|----------------|--|
| Surname/Name * | | DOB * | DD/MM/YYYY |
| Clinic ID * | | Gamete donor * | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Sample type * | <input type="checkbox"/> Blood <input type="checkbox"/> Saliva/ buccal swab <input type="checkbox"/> DNA <input type="checkbox"/> Other: _____ | | |
| 1 st | Genetic Disorder * | Gene * | Mutation * |
| | | | |
| | OMIM# | OMIM# | |
| | Genetic Status * | | |
| <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |
| 2 nd | Genetic Disorder * | Gene * | Mutation * |
| | | | |
| | OMIM# | OMIM# | |
| | Genetic Status * | | |
| <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |

CLINICAL INFORMATION OF FAMILY MEMBERS
(if applicable; also, in case of additional family members, reuse this page)

| 1st family member as reference | | | | |
|--|--|---|--------|--|
| Surname/Name * | | DOB * DD/MM/YYYY | | |
| Gender * | | <input type="checkbox"/> Male <input type="checkbox"/> Female | | |
| Sample type * | | <input type="checkbox"/> Blood <input type="checkbox"/> Saliva/ buccal swab <input type="checkbox"/> DNA <input type="checkbox"/> Other: _____ | | |
| Relationship to pre-embryos for PGT-M* | | <input type="checkbox"/> Grandparent <input type="checkbox"/> Sibling <input type="checkbox"/> Uncle/ aunt <input type="checkbox"/> Other: _____ | | |
| | | Select one or both options: <input type="checkbox"/> Maternal affiliation <input type="checkbox"/> Paternal affiliation | | |
| 1 st | Genetic Disorder * | | Gene * | |
| | OMIM# | | OMIM# | |
| | Genetic Status * | | | |
| | <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |
| 2 nd | Genetic Disorder * | | Gene * | |
| | OMIM# | | OMIM# | |
| | Genetic Status * | | | |
| | <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |

| 2nd family member as reference | | | | |
|--|--|---|--------|--|
| Surname/Name * | | DOB * DD/MM/YYYY | | |
| Gender * | | <input type="checkbox"/> Male <input type="checkbox"/> Female | | |
| Sample type * | | <input type="checkbox"/> Blood <input type="checkbox"/> Saliva/ buccal <input type="checkbox"/> DNA <input type="checkbox"/> Other: _____ | | |
| Relationship to pre-embryos for PGT-M * | | <input type="checkbox"/> Grandparent <input type="checkbox"/> Sibling <input type="checkbox"/> Uncle/ aunt <input type="checkbox"/> Other: _____ | | |
| | | Select one or both options: <input type="checkbox"/> Maternal affiliation <input type="checkbox"/> Paternal affiliation | | |
| 1 st | Genetic Disorder * | | Gene * | |
| | OMIM# | | OMIM# | |
| | Genetic Status * | | | |
| | <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |
| 2 nd | Genetic Disorder * | | Gene * | |
| | OMIM# | | OMIM# | |
| | Genetic Status * | | | |
| | <input type="checkbox"/> Unaffected <input type="checkbox"/> Carrier <input type="checkbox"/> Affected <input type="checkbox"/> Not tested | | | |

Health Professional Authorised To Request The prePGT-M TEST

I certify that, to the best of my knowledge, the patients' and clinical information provided in this form are correct. Based on the clinical indication and my professional expertise, I have requested this test for the patient(s). The limitations of the test, including the fact that PGT-M is not 100% accurate and that prenatal testing is needed to confirm the test result in any pregnancy obtained after PGT, have been explained to the patients and all relevant questions have been answered. I agree to provide any additional information requested by Juno Genetics with regard to this particular test.

| | |
|---------------------------------|---|
| Authorised referrer Signature * | Date * DD/MM/YYYY |
|---------------------------------|---|