Dynacare®

harmony®
performed in Canada

Harmony®
Prenatal Test Requisition

Patient Information

Last name: _____

First name: _____

Date of birth: YYYY MM DD

Health insurance #: _____

Sex: ☐ M ☐ F Weight: _____ ☐ kg ☐ lbs

Address: _____
No. Street name Apt/Unit

_____
City Province Postal code

Telephone: _____

Test Menu Options

☒ **Harmony® Prenatal Test** (T21, T18, T13)

Additional options:

☐ Fetal Sex
☐ Monosomy X[1,2]
☐ Sex Chromosome Aneuploidy Panel[1,2]
☐ 22q11.2[1]

[1] Singletons only. [2] Fetal sex not reported.

☐ Please contact this patient for genetic counselling related to this test/clinical indication.

Important: Patients must be of at least 10 weeks gestational age at the time of collection.

Blood Draw Information

Collection date: YYYY MM DD

Is this a redraw? ☐ Yes ☐ No

Collection centre: _____

Collected by: _____

Collection account #: _____

Prescriber Information

Client #: _____

Last name: _____

First name: _____

Clinic: _____

Address: _____
No. Street name Office

_____
City Province Postal code

Telephone: _____

Fax: _____

Copy results to: _____
Last name, First name

cc. Fax: _____

Clinical Information

Gestation age: complete A or B

A Gestational age at date of ultrasound: _____ **weeks** _____ **days**

Date of ultrasound: YYYY MM DD

B ☐ LMP date; **or**
☐ IVF transfer date: YYYY MM DD

No. of fetuses: ☐ 1 ☐ 2

IVF pregnancy:
☐ No ☐ Yes → Egg donor is: ☐ Self ☐ Non-self
Donor age at retrieval: _____ **years**

Clinician Signature

I attest that my patient has been fully informed about details, capabilities, and limitations of the test(s).

The patient has given full consent for this test.

Clinician signature: _____

Licence #: _____ Date: YYYY MM DD

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Patient Informed Consent

The Harmony® Prenatal Test is a prenatal screening test that analyzes cell-free DNA (cfDNA) in maternal blood. The test provides a risk assessment, not a diagnosis, of fetal chromosomal or genetic conditions, and fetal sex determination. Consider Harmony results in the context of other clinical criteria. Follow up confirmatory testing based on Harmony results for Trisomy 21, 18, 13, sex chromosome aneuploidy, or 22q11.2 could reveal maternal chromosomal or genetic conditions in some cases. Results from the Harmony® Prenatal Test should be communicated in a setting designated by your healthcare provider that includes the availability of appropriate genetic counselling.

The Harmony non-invasive prenatal test is licensed in accordance with Health Canada regulation requirements for a class III license. The Harmony test is based on cell-free DNA analysis and is considered a prenatal screening test, not a diagnostic test. Harmony does not screen for potential chromosomal or genetic conditions other than those expressly identified in this document. All women should discuss their results with their healthcare provider who can recommend confirmatory diagnostic testing where appropriate.

Who is eligible for the Harmony® Prenatal Test?

Patients must be of at least 10 weeks gestational age for any of the Harmony Test offerings. Patients with a twin pregnancy are not eligible for monosomy X, sex chromosome aneuploidy or 22q11.2 options. The Harmony® Prenatal Test is not for patients with a history of or active malignancy; a pregnancy with fetal demise; a pregnancy with more than two fetuses; or a history of bone marrow or organ transplants.

What are the limitations of the Harmony® Prenatal Test for Trisomies 21, 18, and 13, sex chromosome aneuploidy, and fetal sex determination?

The Harmony® Prenatal Test is not validated for use in pregnancies with more than two fetuses, fetal demise, mosaicism, partial chromosome aneuploidy, translocations, maternal aneuploidy, transplant, malignancy, or in women under the age of 18. Harmony does not detect neural tube defects. Certain rare biological conditions may also affect the accuracy of the test. For twin pregnancies, HIGH RISK test results apply to at least one fetus; male test results apply to one or both fetuses; female results apply to both fetuses.

Not all trisomic fetuses will be detected. Some trisomic fetuses may have LOW RISK results. Some non-trisomic fetuses may have HIGH RISK results. False negative and false positive results are possible. A LOW RISK result does not guarantee an unaffected pregnancy due to the screening limitations of the test. Harmony provides a risk assessment, not a diagnosis, and results should be considered in the context of other clinical criteria. It is recommended that a HIGH RISK result and/or other clinical indications of a chromosomal abnormality be confirmed through fetal karyotype analysis such as amniocentesis. It is recommended that results be communicated in a setting designated by your healthcare provider that includes appropriate counselling. For a variety of reasons, including biological, the test has a failure rate. As such, you may be requested to redraw a new sample. In a small number of cases, a result for fetal sex and/or sex chromosome aneuploidy determination may not be obtained. This can be due to biological and technical factors influencing sex chromosome analysis that did not impact trisomy analysis. In these cases, we do not retest or redraw a new sample.

Note: Options for Fetal Sex, Monosomy X, and Sex Chromosome Aneuploidy Panel can only be added up to a maximum of 30 days following initial reporting.

What are the limitations of the Harmony® Prenatal Test for 22q11.2?

In addition to the limitations discussed above, the 22q11.2 option is not validated for use in pregnancies with more than one fetus or for women with a 22q11.2 duplication or deletion.

A 22q11.2 deletion may not be detected in all fetuses. Due to the limitations of the test, a LOW PROBABILITY result does not guarantee that a fetus is unaffected by a chromosomal or genetic condition. Some fetuses with a 22q11.2 deletion may receive a test result of LOW PROBABILITY. Some fetuses without the 22q11.2 deletion may receive a test result of HIGH PROBABILITY. In cases of HIGH PROBABILITY results and/or other clinical indications of a chromosomal condition, confirmatory testing is necessary for diagnosis.

Note: The 22q11.2 option can only be added up to a maximum of 30 days following initial reporting.

What is done with my sample after testing is complete?

No additional clinical testing will be performed on your blood sample other than those authorized by your healthcare provider. Dynacare will disclose the test results only to the healthcare provider(s) listed on the front of this form, or to his or her agent, unless otherwise authorized by you or as required by laws, regulations, or judicial order. Details on Dynacare's policies and procedures governing patient privacy and health information, including patient rights regarding such information, can be found at **https://www.dynacare.ca/privacy-policy.aspx**.

Harmony® Prenatal Test is a trademark of Roche.