Give her the difference of Harmony

SUPERIOR PERFORMANCE TO FTS^{1,3}

with exceptionally low false-positive rates for

1.4 MILLION

tests performed¹⁰

10 WEEKS

Clear results in

3-5 DAYS

from sample

Validated for use

ANY AGE OR RISK CATEGORY*

SUPPORT

and your patients

BARBADOS REFERENCE LABORATORY LTD.

Mayfield Medical Centre, Belleville | Manor Lodge, Warrens Sandy Crest Medical Centre | Maxwell Main Rd., Top Rock

Tel: (246) 436-3472 | Email: info@brllimited.com website: www.brllimited.com

SERVING BARBADOS AND THE EASTERN CARIBBEAN **FOR OVER 35 YEARS**





Answers that matter

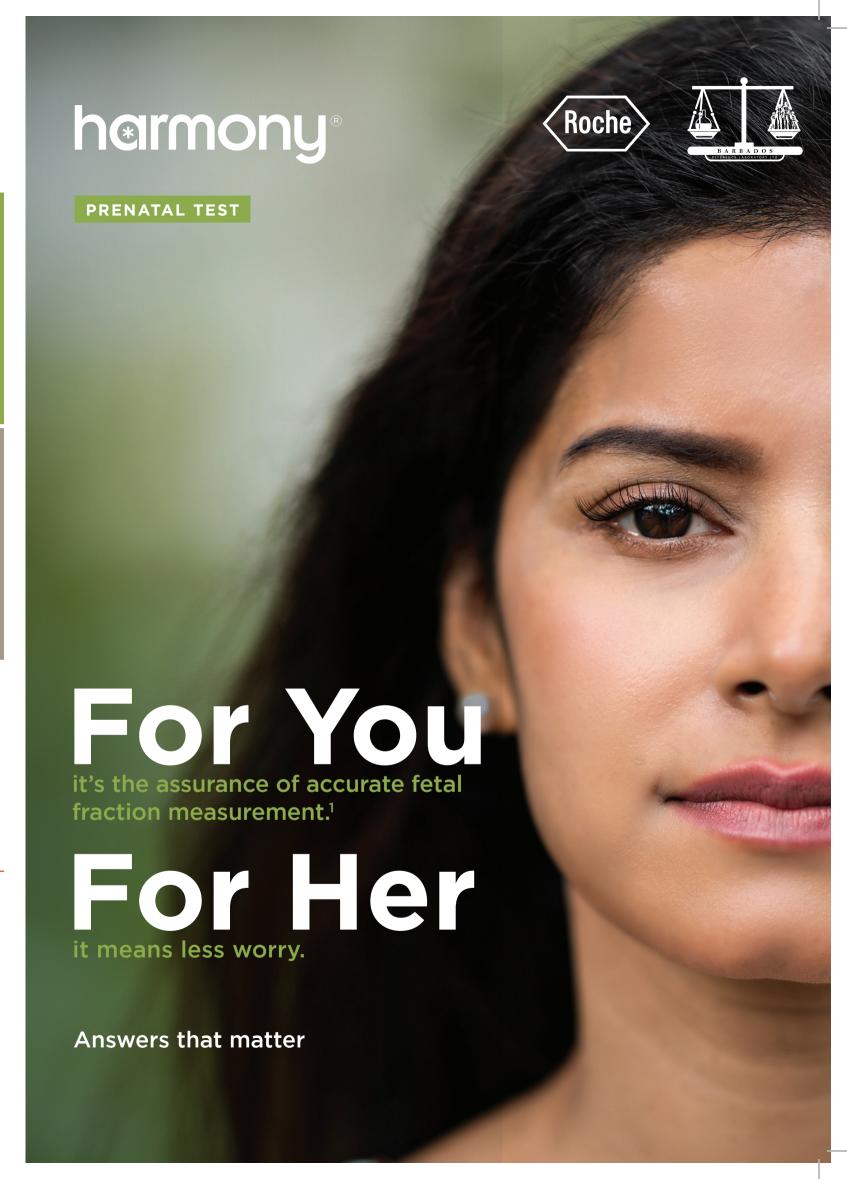
*Any risk refers to the average risk population (under age 35) and high risk population (over age 35). Pregnancies with more than two fetuses, a history of vanishing twin, maternal organ transplant or maternal aneuploidy are not eligible for the Harmony test.

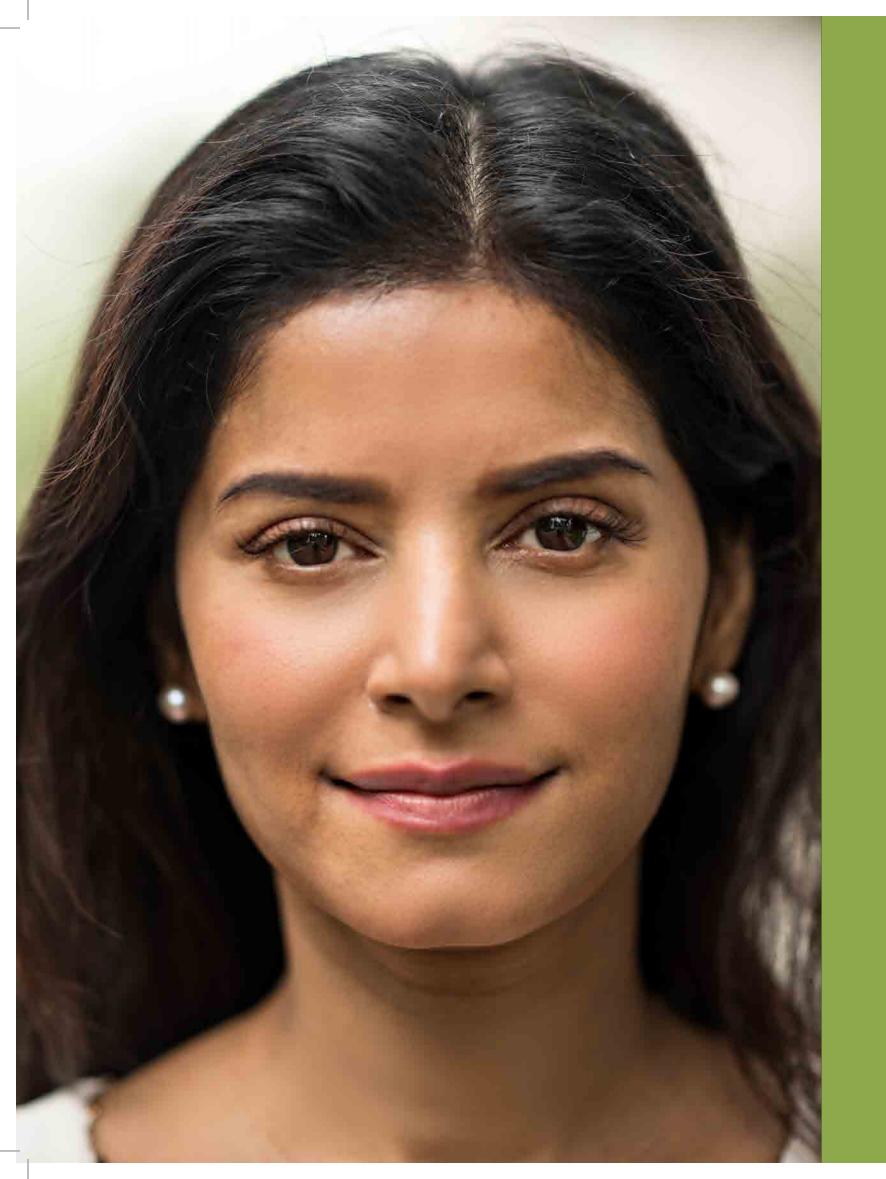
- Stokowski et al. Prenat Diagn. 2015 Dec;35(12):1243-6.
- Demonstrated by 48 peer-reviewed published studies using the Harmony prenatal test as of Jan 2018. For the entire 48 references, please go to harmonytest.com/references
- Norton ME et al. N Engl J Med. 2015;372(17):1589-1597.
- Wax et al. J Clin Ultrasound. 2015 Jan;43(1):1-6.
- Takoudes et al. Ultrasound Obstet Gynecol. 2015 Jan;45(1):112. Dondorp et al. Eur J Hum Genet. 2015 Nov:23(11):1592.
- Benn et al. Prenat Diagn. 2015 Aug;35(8):725-34.
- Gregg et al. Genetics in Medicine 2016 Oct;18(10):1056-65.
- Schmid et al. Ultrasound Obstet Gynecol. DOI: 10.1002/uog.19036.
- Data on file.

The Harmony non-invasive prenatal 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

The Harmony prenatal test was developed and its performance characteristics determined by Ariosa Diagnostics, Inc. a CLIA-certified and CAP-accredited clinical laboratory in San Jose, CA USA. This testing service has not been cleared or approved by the US Food and Drug Administration (FDA).

© 2018 Roche Diagnostics, Inc. All Rights Reserved. HARMONY is a trademark of Roche SEQ100040 07/18



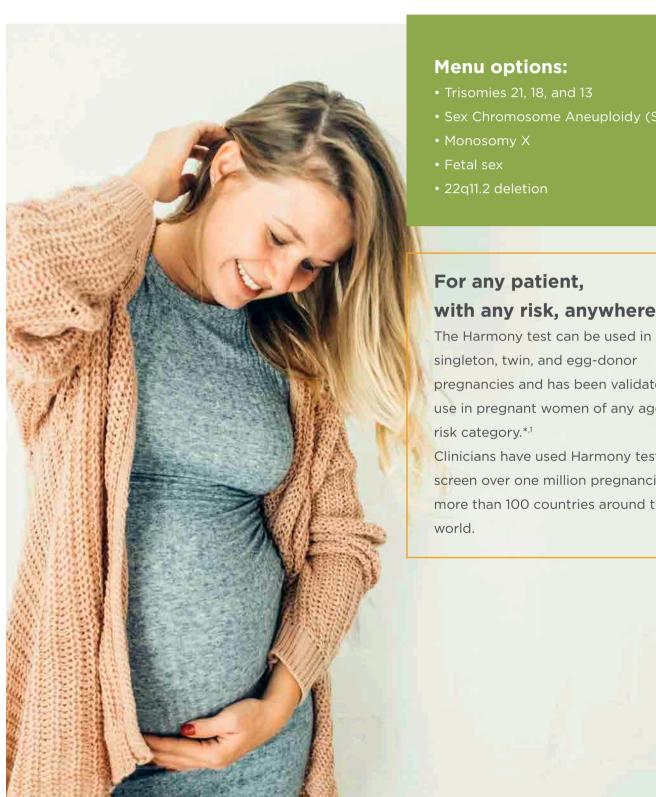


The reliability you want, and the accuracy you need.

THE HARMONY TEST EXPERIENCE

Most proven

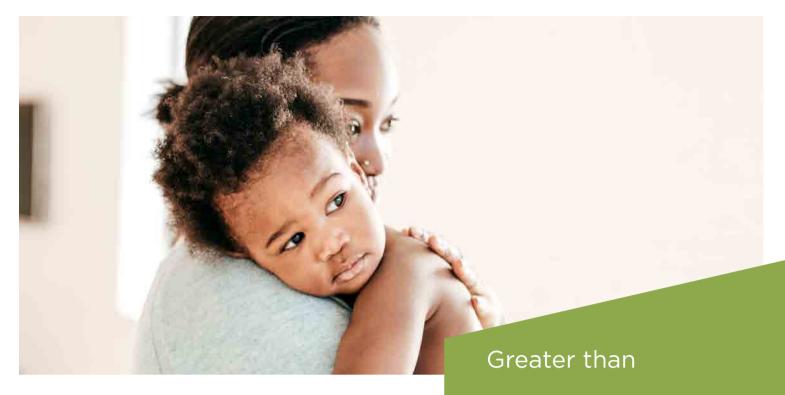
The Harmony test is the most proven cell-free DNA-based prenatal blood screen.²



with any risk, anywhere.*

singleton, twin, and egg-donor pregnancies and has been validated for use in pregnant women of any age or

Clinicians have used Harmony test to screen over one million pregnancies in more than 100 countries around the



Reliable results for all tested conditions

The Harmony prenatal test delivers consistent, industry-leading performance across clinical studies¹

	Detection Rate	False-Positive Rate
Trisomy 21	99.3% (418/421)	<0.1% (10/23,155)
Trisomy 18	97.4% (147/151)	<0.1% (5/22,399)
Trisomy 13	93.8% (30/32)	<0.1% (3/14,243)

99% detection rate for trisomy 21.1

Less than false-positive rate.1

A LANDMARK STUDY

Clinical evidence

Cell-free DNA analysis for Non-invasive Examination of Trisomy (NEXT) Study³

The Harmony test, as a first-line screen, is supported by extensive clinical evidence.³

NEXT study design

- Maternal age range: 18 to 48 years old (mean: 31 years old)
- Gestational age range: 10.0 to 14.3 weeks (mean: 12.5 weeks)

All patients in this study received both first-trimester screening (FTS) and the Harmony prenatal test. Pregnancy outcome data was obtained on each pregnancy by genetic testing or newborn examination.

15,841 women

were included in the primary analysis population.³



Significantly outperforms

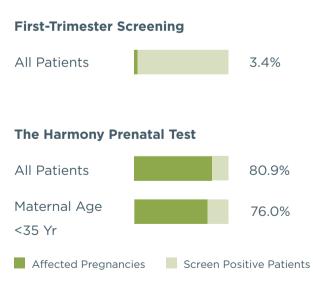
In the largest blinded, prospective study of its kind, the Harmony test significantly outperformed first-trimester screening (FTS[†]) for trisomy 21 in both detection and false-positive rates.³

NEXT study results³

	Harmony Prenatal Test	FTS	P-value
DETECTION RATE (affected pregnanices identified as high risk)	100% (38/38)	79% (30/38)	0.008
FALSE-POSITIVE RATE (unaffected pregnancies incorrectly identified as high risk)	0.06% (9/15803)	5.4% (854/15803)	<0.001

The Harmony prenatal test demonstrates superior positive predictive value (PPV) for trisomy 21 compared to FTS.

PPV: Proportion of screen positive results that accurately reflect an affected pregnancy



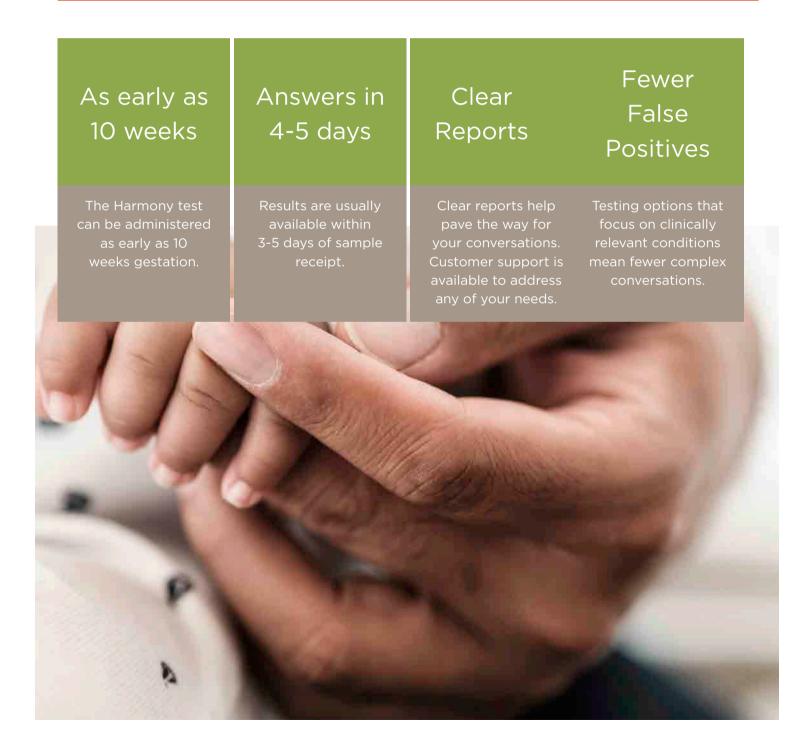
A superior first-line screen for Trisomy 213

 $^{^{\}dagger}$ Serum PAPP-A, total of free $\beta\text{-hCG}$ & Nuchal Translucency

Clarity is one simple step away

Your patients who are expecting may have many questions. That's why it's important for you to be equipped with the right knowledge at the right time. With the Harmony test, results are clear and focused, so that your conversations can be too.

It's about time



It's about confidence

Superior performance

to help minimize anxiety and invasive procedures due to false-positive results.^{3,4}

Unparalleled evidence

as demonstrated in studies of over 148,000 patients in more than 48 peerreviewed publications.²

Trusted results

the Harmony test will only provide a result when there is sufficient cell-free fetal DNA for an accurate analysis.⁵

Minimize patient anxiety

caused by false-positive results.⁴



Our commitment to your care

Information and support

The Harmony test was developed with the belief that all women should have access to the best quality information about their pregnancies.

Provider portal – Track results and records easily, so you can give patients the answers they need when they need them.

Patient education - We're committed to developing resources to help your patients understand more about their pregnancy, test results, and more.

Market access initiative – Our team works at the health-system level to provide the clinical evidence necessary to bring testing to your population.

First line screening test - The Harmony test makes superior prenatal screening available to any woman, regardless of age or risk status.*,3



Professional medical societies

NIPT as a first-line screening test for trisomy 21.

Recent positioning statements recommend using NIPT as a first-tier test, which significantly decreases the number of women who will receive an initial false alarm.⁶

66

Performance of cfDNA screening is considerably greater than conventional screening...

- ISPD Statement 2015

66

informing all pregnant women that NIPS [noninvasive prenatal screening] is the most sensitive screening option for T21, 18, 13...

- ACMG Statement 2016

All labs should include clearly visible fetal fraction on all reports...[and] establish clinical validation of fetal fraction...

- ACMG Statement 2016⁸

The Harmony test has been demonstrated to provide highly accurate and reproducible assessment of fetal fraction.⁹

EXPERIENCE

the Harmony difference.
Visit www.brllimited.com
Tel: 436-3472
Email: info@brllimited.com