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Maternity 21

Blood Test Cpt

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Blood Test

Cpt Code

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The following

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information must be provided with the test request form: patient's date of birth, gestational age, additional patient demographic information: pregnancy type (singleton or multiple), donor egg status and the clinical indications (including advanced maternal age, abnormal ultrasound, history suggestive of increased risk for

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aneuploidy, positive serum screen, or other indications).

451927: MaterniT21 PLUS Core (chr21,18,13,sex) | LabCorp

The following information must be provided with the test request form: patient's date of birth, gestational age, additional patient demographic information, pregnancy

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type (singleton or multiple), donor egg status and the clinical indications (including advanced maternal age, abnormal ultrasound, history suggestive of increased risk for aneuploidy, positive serum screen, or other indications).

**451934: MaterniT21
PLUS Core + SCA |
LabCorp**

Prenatal Genetic

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Screening MaterniT 21

Non-Invasive Prenatal
Testing - CPT: 81420 •

Analyzes genetic
information that enters
the bloodstream from
the placenta. It screens
for certain

chromosomal
abnormalities that
could affect your
baby's health and
development such as
Trisomy 21 (Downs),
18, and 13. It can also
detect the sex of the
fetus.

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Prenatal Genetic Screening MaterniT 21 Non-Invasive ...

A noninvasive blood test. There are many ways to get this genetic information, including methods such as serum screens and diagnostic procedures such as amniocentesis. As a noninvasive prenatal test, MaterniT 21 PLUS is different from both.

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**MaterniT 21 PLUS
Test | Integrated
Genetics**

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Title: Sample ...

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Booklection.com**

Myriad Prequel™
Prenatal Screen: Non-
invasive prenatal
screen (cell-free fetal
DNA). Conditions
covered: CPT code:
Common aneuploidies
including Trisomy 21,
Trisomy 18, Trisomy
13: 81420:
Microdeletions
including 22q11.2
deletion syndrome,

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1p36 deletion
syndrome, ...

Which CPT codes could be used for the Prequel Prenatal Screen?

Noninvasive prenatal test A noninvasive prenatal test (NIPT) is a single blood test performed any time at or after 10 weeks.

During pregnancy, 3-13% of the DNA in your blood stream is circulating cell free

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fetal DNA that comes from the placental cells. A NIPT works by evaluating the amount of cell free DNA in your blood.

Prenatal Testing Information and Codes

MaterniT 21 PLUS performs in key areas that ensure your time is spent wisely, delivering fast, reliable, and effective prenatal screening results.

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Integrated Genetics has run over 1.5 million non-invasive prenatal tests (NIPT) since pioneering the technology in 2011, including more than 30,000 twins and more than 50,000 genome-wide tests; 5 the depth of experience you should expect from an NIPT.

**Prenatal MaterniT
21 Plus | Integrated
Genetics**

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The Harmony prenatal test is a screening test. Results should be confirmed with diagnostic testing such as amniocentesis and chorionic villus sampling (CVS). Read more about the accuracy of the Harmony prenatal test. References. Stokowski et al. Prenat Diagn. 2015 Dec;35(12):1243-6. ACOG Practice Bulletin 163 Obstet Gynecol.

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127(5):815-816, May ...

Code

**Harmony Prenatal
Test for Expecting
Parents**

The test analyzes the relative amount of 21, 18, 13; X and Y chromosome material in circulating cell-free DNA from a maternal blood sample. This test can be performed at any time after 10 weeks gestation. If you elect this test, you will also have an AFP

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(alpha-fetoprotein)
blood test to test for
open neural tube
defects.

GUIDE TO GENETIC TESTING | Obstetrics and Gynecology ...

Test Name: PANORAMA

PRENATAL TEST* Test

Code: 2191746: Alias:

NIPD NIPS NIPT

Prenatal NIPT: CPT

Code(s): 81420 : Test

Includes: Screening
chromosomes 13, 18,

21, X

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Health Lab

Trisomy 21, 18 or 13 means there are three copies of either chromosome 21, 18 or 13, instead of two. Noninvasive prenatal laboratory testing analyzes circulating cell-free DNA from a maternal blood sample which detects increases of chromosome material associated with trisomy 21 (Down syndrome),

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trisomy 18 (Edwards syndrome), or trisomy 13

**Medical Necessity
Guidelines: Genetic
Testing: Maternal ...**

Confirmation of
Pregnancy Office Visit:
(Your first
appointment). The
Diagnosis code will be
V72.42--confirmation of
pregnancy. Prenatal
Panel (blood-work)
(CPT Codes 85025,
86900, 86901, 86850,

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87340, 86780, 86762)

HCG, Quant (CPT Code

84702) HIV testing

(CPT 87389)

Hemoglobin (CPT Code

85025) Pap Smear

(CPT Code 88175)

**2015 Prenatal Billing
Information - Clinic
Sofia**

The MaterniT21® test is a prenatal screening blood test which is designed to detect increased amounts of material found from

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chromosome 21 (Down syndrome), chromosome 13 (trisomy 13) and chromosome 18 (trisomy 18).

How Much Does the Sequenom MaterniT21® Test Cost ...

Non-invasive prenatal testing (NIPT)

Panorama is a blood-based genetic, prenatal screening test of the pregnant mom that

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Chrom

screens for common chromosomal conditions that affect a baby's health.

Panorama uses unique SNP*-based technology to deliver the most accurate NIPT on the market.

Panorama Prenatal Screening Test | Fetal Genetic Testing

Aetna considers noninvasive prenatal testing (NIPT) using

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measurement of cell-free fetal nucleic acids in maternal blood (e.g., MaterniT21, MaterniT21 PLUS, Verifi Prenatal Test, Harmony Prenatal Test, Panorama Prenatal Test, QNatal Advanced) medically necessary for testing for fetal aneuploidy (trisomy 13, 18 and 21) in pregnant women with single gestations who meet any of the following

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**Serum and Urine
Marker Screening
for Fetal Aneuploidy**

...

Specimen: 5 - 7 mL
Whole Blood. Transport
Temperature: Room
Temperature If the sex
chromosomes option is
selected, results for
Monosomy X, XXX, XXY
and XYY. will be
included. Note: Test
requires the following
information for testing.
1) Estimated Due Date

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(EDD) 2) Pregnancy

Type (Single, Twin) 3)

Is gender required on
report (Yes/No)

**Test Code ClariNIPT
ClariTest Non-
invasive Prenatal ...**

It consists of a blood
test to check levels of
pregnancy-associated
proteins and hormones
in the mother's blood
and an ultrasound to
look for ... the
marker of the

MaterniT 21 Plus test,

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New Prenatal Blood Tests Come With High Hopes And Some ...

test), maternal serum screening (a blood test) and a high-resolution ultrasound evaluation in the second trimester.

These tests may identify women with an increased risk of having a child with trisomy 21, 18 or 13,

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but they cannot
diagnose, confirm or
exclude the possibility
of a chromosomal
disorder.

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