EVICScreen™ Evidence Based Comprehensive Screening

EVICDUO™ Dual Marker Screening

To: **Dr.Manju Gupta-New Delhi**
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Delhi
Delhi - 110091
Contact: 9873146127
Report Of: Mrs. SHREYA SENGAR
Pt. Contact: 9755531390

Sample ID	2460009603
Patient ID	1602417774
Received on	21/11/2024 16:27
Registered on	21/11/2024 16:35
Reported on	-
Referred by	**Dr. Manju Gupta**
Sonography by	**Dr. ARCHIT SINGHAL**

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EVICOSCREEN - EVIDENCE BASED COMPREHENSIVE PRENATAL SCREENING REPORT

Patient Name: Mrs. SHREYA SENGAR **Patient DOB: 29/01/1997**

EVICScreen™ is an evidence based prenatal screening program curated by Lilac Insights in accordance with the Fetal Medicine Foundation (UK) guidelines for First Trimester Screening to determine the probality of most common chromosomal aneuploidies in a pregnancy. It utilizes:

- Hormonal values from the pregnancy measured on Fetal Medicine foundation (UK) accredited analyzers and reagents
- Robust indigenous medians from over 7 lac+ pregnancies for different gestation ages
- Risk calculations from evidence based algorithms validated through large international studies

UKNEQAS: United Kingdom National External Quality Assessment Service

RIQAS: Randox International Quality Assessment Scheme

ASSURED QUALITY ASSURED

The Risk Assessment Performed Using CE-Marked Antenatal Risk Evaluation Software Certified by the British Standards Institute (BSI)- ISO 13485:2016

RISK ASSESSMENT

T21 (Down syndrome)	**1: 2100**	Low Risk	LOW INTERMEDIATE HIGH
T18 (Edwards' syndrome)	**1: 100000**	Low Risk	LOW HIGH
T13 (Patau syndrome)	**1: 100000**	Low Risk	LOW HIGH

MULTIPLE OF MEDIAN (MoM)

Free ß-hCG	7.34
PAPP-A	1.17

INTERPRETATION

The First Trimester Screening for the given sample is found **SCREEN NEGATIVE.**

SUGGESTIONS AND OTHER FINDINGS

In view of free bHCG MoMs observed in the mother, kindly consider correlation with fetal growth and well being scan at 28 - 30 weeks.

NABL MC-6043

UK NEQAS International Quality Expertise Lab Reg. No. 90968

Verified by
Mr. Pradip Kadam
Incharge Biochemistry
(FMF ID: 147760)

Verified by
Dr. Suresh Bhanushali
MD (Path), Consultant Pathologist

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Patient name : Mrs. SHREYA SENGAR **Sample ID** : 2460009603

Sample Type: Serum **Risk assessment:** Algorithm validated by SURUSS 2003, N.J Wald

Method: Electrochemiluminescence

PREGNANCY DETAILS

No. of fetuses	: 1	**EDD**	: 29/05/2025	**Age at Term**	: 28.3 Years
GA is Based on	: CRL 66mm at 20/11/2024	**LMP Date**	: 24/08/2024	**LMP Certainty**	: Regular
Smoking : None	**Parity** :	**Height**	:	**Weight**	: 72.40 Kg
Ethinicity :Asian	**FHR** :				

Previous pregnancy history		Pre-eclampsia history		Other findings	
☐ Down syndrome	☐ Edwards' syndrome	☐ PE in previous pregnancy		☐ Insulin dependent diabetes	
☐ Patau syndrome	☐ NTD syndrome	☐ Pat. mother had PE		☐ Chronic hypertension	

EDD: Estimated Due Date | GA: Gestation Age | LMP: Last Menstrual Period | FHR: Fetal Heart Rate | NTD: Neural Tube Defect | PE: Pre-eclampsia | DOB: Date of Birth

SPECIMEN DETAILS

Sample ID	: 2460009603	**CRL**	: 66 mm
Collection Date	: 21/11/2024	**CRL2**	:
Scan Date	: 20/11/2024	**BPD**	:
GA at Coll Date	: 13 Weeks 0 Days	**BPD2**	:
GA at Scan Date	: 12 Weeks 6 Days	**HC**	:
Received on	: 21/11/2024	**HC2**	:

Test Name	Conc.	Unit	Corr. Mom
Free-ß-hCG	224.10	ng/mL	7.34
NB	Present		
NT	1.5	mm	1.02
PAPP-A	4691.00	mIU/L	1.17

GA: Gestation Age | CRL: Crown Rump Length | BPD: Bi-parietal Diameter | HC: Head Circumference | free-ß-hCG: free-Beta Human Chorionic Gonadotropin NT: Nuchal Translucency | PAPP-A: Pregnancy-associated Plasma Protein-A

RISKS

Disorder: Down Syndrome **Result:** Low Risk

Final risk: 1:2100 Age risk: 1:1200

Cutoff 1:250 Risk type Risk At Term

Disorder: Edwards' Syndrome **Result:** Low Risk

Final risk: 1:100000 Age risk: 1:8100

Cutoff 1:100 Risk type Risk At Term

Disorder: Patau Syndrome **Result:** Low Risk

Final risk: 1:100000 Age risk: 1:12000

Cutoff 1:100 Risk type Risk At Term

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Verified by **Mr. Pradip Kadam** Incharge Biochemistry (FMF ID: 147760)

Verified by **Dr. Suresh Bhanushali** MD (Path), Consultant Pathologist

Patient name : Mrs. SHREYA SENGAR **Sample ID** : 2460009603

PRENATAL SCREENING BACKGROUND

Every pregnant woman carries a certain degree of risk that her fetus/baby may have certain chromosomal defect/ abnormalities. Diagnosis of these fetal chromosomal abnormalities requires confirmatory testing through analysis of amniocytes or Chorionic Villous Samples (CVS). However, amniocentesis and CVS procedures carry some degree of risk for miscarriage or other pregnancy complications (Tabor and Alfirevic, 2010). Therefore in routine practice, prenatal screening tests are offered to a pregnant woman to provide her a personalised risk for the most common chromosomal abnormalities (T21-Down syndrome, T18- Edwards' syndrome, T13- Patau syndrome) using her peripheral blood sample. Based on this risk assessment, if the risk is high or intermediate, you can take informed decision of opting for invasive procedure such as amniocentesis or CVS followed by confirmatory diagnostic test(s), as per discussion with your clinician.

PRENATAL SCREENING TESTS ARE NOT CONFIRMATORY TESTS. THEY ARE LIKELIHOOD ASSESSMENT TESTS.

You may get your prenatal screening result as either of the following:-

High Risk

High Risk or Screen Positive Result: A High Risk Result does not mean that the pregnancy is affected with the condition. It means that the likelihood of the pregnancy having a condition is higher than the cut-off (Most commonly used cut-off is 1:250 and this represents the risk of pregnancy loss from confirmatory testing through CVS or amniocentesis).

Low Risk

Low Risk or Screen Negative Result: A Low Risk result does not mean that the pregnancy is not affected with a condition. It means that the likelihood of the pregnancy having a condition is lower than the cut-off.

Intermediate Risk

Intermediate Risk result: An intermediate Risk result means that the pregnancy has an equivocal or a borderline risk of being affected with a condition. In this case, you may want to choose a second stage screening modality like an Integrated Screening Test that is done between 16 to 20 weeks of pregnancy or a Non-invasive Prenatal Screening Test between 12 to 20 weeks of pregnancy before taking a decision on an invasive confirmatory testing. This will help you improve the sensitivity of the screening test keeping an invasive test a last option were you to come as a high risk in the second stage screening test.

SIGNIFICANCE OF MULTIPLE OF MEDIANS (MoMs)

Prenatal Screening determines the likelihood of the pregnancy being affected with certain conditions by analysing levels of certain hormones. These hormones are Feto placental products (released by Fetus or placenta). Their levels not only indicate propensity of the fetus being affected with certain chromosomal conditions, they also provide indication of placental insufficiency that can potentially lead to pregnancy complications like Pre-Eclampsia or Intra-Uterine Growth Restriction. It is therefore important to take cognisance of the Reported MoMs alongside the Risk results.

For more information, visit our website at: www.lilacinsights.com/faq-pns

DISCLAIMERS

Limitations of the Test:

As prenatal screening tests are not confirmatory diagnostic tests, the possibility of false positive or false negative results can not be denied. The results issued for this test does not eliminate the possibility that this pregnancy may be associated with other chromosomal or sub- chromosomal abnormalities, birth defects and other complications.

Nuchal Translucency is the most prominent marker in screening for Trisomy 13, 18, 21 in the first trimester and should be measured in accordance with the Fetal Medicine Foundation (UK) guidelines. Nuchal Translucency or Crown Rump Length measurement, if not performed as per FMF (UK) imaging guidelines may lead to erroneous risk assessments and Lilac Insights bears no responsibility for errors arising due to sonography measurements not performed as per these criteria defined by international bodies such as FMF (UK), ISUOG.

It is assumed that the details provided along with the sample are correct. The manner in which this information is used to guide patient care is the responsibility of the healthcare provider, including advising for the need for genetic counselling or additional diagnostic testing like amniocentesis or Chorionic Villus Sampling. Any diagnostic test should be interpreted in the context of all available clinical findings. As with any medical test, there is always a chance of failure or error in sample analysis though extensive measures are taken to avoid these errors.

Note:

- Quality of the Down syndrome screening program (Biochemical values, MoMs and Risk assessments) is monitored by UKNEQAS on an ongoing basis.
- This interpretation assumes that patient and specimen details are accurate and correct.
- Lilac Insights does not bear responsibility for ultrasound measurements like CRL,NT,NB etc. We strongly recommend that ultrasound measurements are performed as per FMF (UK)/ISUOG practice guidelines.
- It must be clearly understood that the results represent risk and not diagnostic outcomes. Increased risk does not mean that the baby is affected and further tests must be performed before a firm diagnosis can be made. A Low Risk result does not exclude the possibility of Down's syndrome or other abnormalities, as the risk assessment does not detect all affected pregnancies.
- Each sample received at Lilac Insights' processing centre is handled with the utmost sensitivity and care. All samples received on Sundays and National holidays are stored as per specific guidelines for the respective specimens and processed on the next day.

END OF REPORT