



P K R JAIN HEALTHCARE INSTITUTE

NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

☎ 0171-2532620, 8222896961 ✉ pkrajainhealthcare@gmail.com

NAME : Mrs. PARVINDER KAUR
AGE/ GENDER : 29 YRS/FEMALE **PATIENT ID** : 1650546
COLLECTED BY : **REG. NO./LAB NO.** : 122410220015
REFERRED BY : **REGISTRATION DATE** : 22/Oct/2024 04:21 PM
BARCODE NO. : 12505294 **COLLECTION DATE** : 22/Oct/2024 04:24PM
CLIENT CODE. : P.K.R JAIN HEALTHCARE INSTITUTE **REPORTING DATE** : 23/Oct/2024 12:08PM
CLIENT ADDRESS : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

Test Name	Value	Unit	Biological Reference interval
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ENDOCRINOLOGY

DUAL MARKER MATERNAL SCREENING

DUAL MARKER TEST

PATEINT SPECIFICATIONS

DATE OF BIRTH	30-10-1994		
MATERNAL AGE	30.5	YEARS	
WEIGHT	50.5	Kg	
DATE OF LMP	24-07-2024		
ETHNIC ORIGIN	ASIAN		ASIAN
H/O IVF	ABSENT		
H/O SMOKING	ABSENT		
H/O INSULIN DEPENDANT DIABETES	ABSENT		
H/O TRISOMY 21 SCREENING	ABSENT		

ULTRA SOUND SCAN DETAILS

DATE OF ULTRASOUND	22-10-2024		
by ULTRASOUND SCAN			
METHOD FOR GESTATION AGE ESTIMATION	ULTRASOUND SCAN DETAILS		
by ULTRASOUND SCAN			
FOETUS (NOS)	1		
by ULTRASOUND SCAN			
GA ON THE DAY OF SAMPLE COLLECTION	13	WEEKS	
by ULTRASOUND SCAN			
CROWN RUMP LENGTH (CRL)	66.9	mm	38 - 84
by ULTRASOUND SCAN			
GESTATIONAL AGE BY CRL	13		
by ULTRASOUND SCAN			
NUCHAL TRANSLUCENCY (NT)	2.3	mm	0.1 - 6.0
by ULTRASOUND SCAN			
NUCHAL TRANSLUCENCY (NT) MOM	1.35		
by ULTRASOUND SCAN			

DUAL MARKER - BIOCHEMICAL MARKERS

PREGNANCY ASSOCIATED PLASMA	8094.01	mIU/L
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PROTEIN A (PAPP-A) by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)			
BETA HCG - FREE; SERUM by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	21.67	ng/mL	
<u>MULTIPLE OF MEDIAN (MOM) VALUES</u>			
PAPP-A MOM by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	1.21		
BETA HCG - FREE MOM by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	0.43		
<u>TRISOMY 21 SCREENING (DOWNS SYNDROME) RISK ASSESSMENT</u>			
TRISOMY 21 SCREENING RISK RESULT by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	NEGATIVE (-ve)		NEGATIVE (-ve)
TRISOMY 21 AGE RISK by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	1: 909 NEGATIVE (-ve)		
TRISOMY 21 BIOCHEMICAL RISK by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	< 1:10000 NEGATIVE (-ve)		RISK CUT OFF 1:150
TRISOMY 21 COMBINED RISK (BIOCHEMICAL + NT) by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	< 1:10000 NEGATIVE (-ve)		RISK CUT OFF 1:150
<u>TRISOMY 18 SCREENING RISK ASSESSMENT</u>			
TRISOMY 18 AGE RISK by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	NEGATIVE (-ve)		
TRISOMY 13/18 SCREENING RISK by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)	< 1:10000 NEGATIVE (-ve)		RISK CUT OFF 1:300

INTERPRETATION:

1.Double marker test (maternal serum screen – first trimester) is a prenatal test to screen for Trisomy 21 (down's syndrome) and Trisomy 13/18 during gestational period 8 – 13 weeks.
2.Besides the biochemical markers tested – maternal pregnancy associated plasma protein a (papp-a) & maternal free beta hcg , the risk is calculated combining usg measurement of nuchal translucency (nt), gestational age at the time of sample with other maternal factors as age, weight, h/o diabetes, smoking, race, twin pregnancies, use of assisted reproductive technologies (IVF).




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NOTE:

1.This is only screening test based purely on statistical analysis which is further based on the data submitted; hence the correctness of data is vital for risk analysis.

2.A negative screen indicates a lower probability of having a baby with trisomy 21 ,trisomy 18 and neural tube defects, but does not completely exclude the possibility.

3.A positive screen on the contrary only indicates a higher probability of having a baby with trisomy 21, trisomy 18 and neural tube defects, and needs confirmation by cytogenetic studies and/or level ii scan.

4.The detection rate by this test is about 60%, with 5% false positive rate when assesment is done for only biochemical parameters and increase to 85 % with 5% false positive rate when both biochemical parameters and nt are combined for analysis.

5.Correlation with patient history, family history and detailed USG scan is required to decide further course of action in cases who have high risk statistically calculated by this test.



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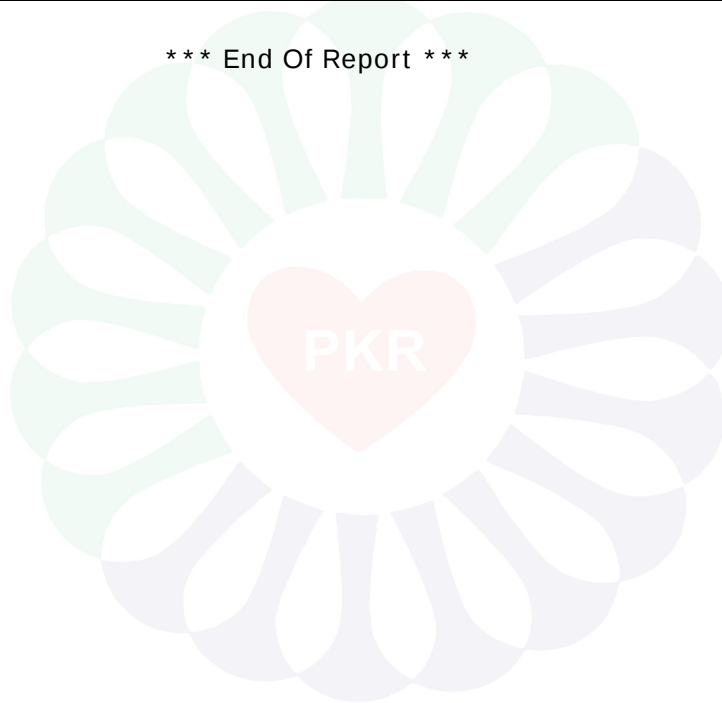
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*** End Of Report ***




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Basic Information

Name: PARVINDER KAUR Contact: Gender: Female
Weight: 50.50 Kg Birthdate: 1994-10-30 Age of EDC: 30.50 Year
Race: Asian Twins: No GA calc method: CRL Robinson
LMP Day: Sender:

Sample information

Send time: 2024-10-23 Sample NO.: 202410230002 Scan Date: 2024-10-22
Lab: Sample Date: 2024-10-22 GA: 13+0
BPD: -- mm CRL length: 66.90 mm NT length: 2.30 mm

Assay

NO.	Item abbr	Result	Unit	MOM	Reference range
1	free-β-HCG	21.67	ng/ml	0.43	
2	PAPP-A	8094.01	mIU/L	1.21	
3	NT	2.30	mm	1.35	

Risk calculate

Age risk: 1:909

Parameter: Trisomy21

Risk: 1:27149

Cut Off: (< 1:150)

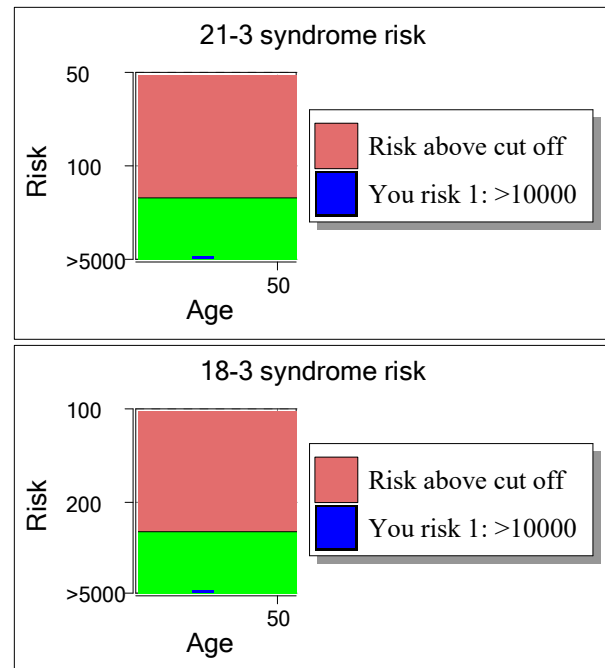
Screening Result: Negative

Parameter: Trisomy18/13

Risk: 1:205554

Cut Off: (< 1:300)

Screening Result: Negative



Parameter:

Cut Off:

Screening Result:

Advice:Diagnostic results with less risk

Note: *The basic information on the basis of Down's risk assessment in this report is provided at the time of your onsite. When you get this report, please first check whether your relevant information is correct. If there is any discrepancy, please contact your doctor in time, so as to feedback us the correct information and documents, then obtain the correct report.

*The high risk and borderline risk of trisomy 21 or trisomy 18 requires further interventional prenatal diagnosis (from fetuses such as villus, amniotic fluid, cord blood, etc.); high risk of neural tube defect (NTD), please go to ultrasound prenatal diagnosis qualified hospitals use ultrasound to exclude.

*The risk of NTD is only calculated at 14-22 weeks.

*The screening result with low risk only shows that the chance of this kind of congenital abnormality in your fetus is less, and the possibility of this kind of abnormality or other abnormalities cannot be completely ruled out. Please consult a doctor in time after you get the report, and the doctor will follow your Risks and other conditions (whether you are older than 35 years old, whether you have had more than one child with other deformities, or have other diseases such as tumors) are comprehensively considered to suggest whether you need to take further examination to confirm the diagnosis.

****This report only can be reference and assistant for doctor, cannot directly give conclusion by this ****

Doctor:

Checked by :

Print date: 2024-10-23 12:02:06