



Name : Mrs.VINODHA
Age/Gender : 24 Y 0 M 0 D /Female
Ref Doctor : TRIVENI MS OBG
Client Code : NCL006
PUP Name : TRIVENI HOSPITAL

Barcode No : 10003756
Visit No : NCD3104
Collection On : 30/Dec/2024 07:26PM
Received On : 30/Dec/2024 07:26PM
Authorized On : 06/Jan/2025 01:21PM


DEPARTMENT OF HORMONE ASSAYS

Test Name	Result	Unit	Bio. Ref. Range	Method
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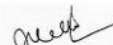
Double Marker (SERUM)

PATIENT DEMOGRAPHICS	:			
Origin	Asian			
Weight	60 kg			
Date of Birth	10/06/2000			
H/O Diabetes	Unknown			
H/O IVF	Unknown			
No. of Fetus	1			
Date of Scan	30/12/2024			
Crown-Rump Length (CRL)	59.80	MM		
PAPP-A	4.27	mIU/mL	<0.1:Non-Pregnant 0.1-19.5:Pregnant	CLIA
Mom for PAPP-A	1.36			
Free Beta Hcg	159.00	ng/mL	<2:Non-Pregnant 5.4-393.4:Pregnant	CLIA
Mom For Free Beta Hcg	4.32			
Nuchal Translucency (NT)	2.30	MM		
MoM for NT	1.480			
Down syndrome (T21) Biochemical Risk	1:330	Risk Cutoff	High Risk:< 1:250 Low Risk: > 1:250	
Down syndrome (T21) Combined Risk	1:216	Risk Cutoff	High Risk:< 1:250 Low Risk: > 1:250	
Combined Trisomy 13/18 Risk Estimate.	<1:10000	Risk Cutoff	High Risk : < 1:250 Low Risk : > 1:250	
Down syndrome(T21) Maternal Age Risk:	1:971			

Reviewed By:


Mr. SK. Rasheed
(Biochemist)




Dr. P. H. Latha MD
Consultant Pathologist



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Interpretation :

NOTE:

Multiple of the Median (MoM): Analyte values are compared to median values at a given gestational age and multiple of the median (MoM) results obtained. The MoM results are used in a multivariate algorithm that includes the mother's age to derive risk factors for Down syndrome and Trisomy 18. An interpretive report is provided.

- Double marker screen or first-trimester screen is performed by measuring analytes in maternal serum that are produced by the fetus and the placenta. Additionally: the Nuchal translucency (NT) measurement is a sonographic marker shown to be effective in screening fetuses for Down syndrome.

A mathematical model is used to calculate a risk estimate by combining the analyte values. NT measurement and maternal demographic information.

- All serum marker multiple of medians are adjusted for maternal weight (to account for dilution effects in heavier mothers). The estimated risk calculations and screen results are dependent on accurate information for gestation, maternal age: and weight. Inaccurate information can lead to significant alterations in the estimated risk.

- PAPP-A:** is highly expressed in first-trimester trophoblasts, participating in regulation of fetal growth. Levels in maternal serum increase throughout pregnancy. Low PAPP-A levels before the 14th week of gestation are associated with an increased risk for Down syndrome and Trisomy 18.


- Nuchal translucency (NT):** The NT measurement, an ultrasound marker, is obtained by measuring the fluid-filled space within the Nuchal region (back of the neck) of the fetus. While fetal NT measurements obtained by ultrasonography increase in normal pregnancies with advancing gestational age. Down syndrome fetuses have larger NT measurements than gestational age-matched normal fetuses.

Increased fetal NT measurements can therefore serve as an indicator of an increased risk for Down syndrome.

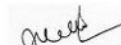
Page 3 PDF Graph attached

*** End Of Report ***

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Mr. SK. Rasheed
(Biochemist)




Dr. P. H. Latha MD
Consultant Pathologist

Patient data			
Name	VINODHA	Patient ID	NCD3104
Birthday	10/06/00	Sample ID	10003756
Age at sample date	24.6	Sample Date	30/12/24
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	unknown
Weight	60	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.27 mIU/ml	1.36	12 + 2
fb-hCG	159 ng/ml	4.32	Method
			CRL Robinson
			Scan date
			30/12/24
Risks at sampling date		Crown rump length in mm	
		59.8	
Age risk	1:971	Nuchal translucency MoM	
Biochemical T21 risk	1:330	1.48	
Combined trisomy 21 risk	1:216	Nasal bone	
Trisomy 13/18 + NT	<1:10000	present	
		Sonographer	
		DR.MD DR.MD	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 216 women with the same data, there is one woman with a trisomy 21 pregnancy and 215 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician