



Name : Mrs.VINODHA **Barcode No** : 10003756
Age/Gender : 24 Y 0 M 0 D /Female **Visit No** : NCD3104
Ref Doctor : TRIVENI MS OBG **Collection On** : 30/Dec/2024 07:26PM
Client Code : NCL006 **Received On** : 30/Dec/2024 07:26PM
PUP Name : TRIVENI HOSPITAL **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 ultrasound 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.

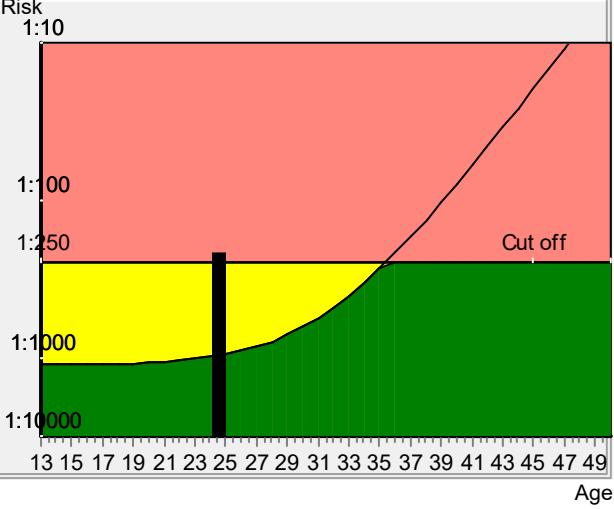
Page 3 PDF Graph attached

*** End Of Report ***

Reviewed By:
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			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	4.27 mIU/ml	1.36	Gestational age 12 + 2
fb-hCG	159 ng/ml	4.32	Method CRL Robinson
Risks at sampling date			
Age risk	1:971		Scan date 30/12/24
Biochemical T21 risk	1:330		Crown rump length in mm 59.8
Combined trisomy 21 risk	1:216		Nuchal translucency MoM 1.48
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer DR.MD DR.MD
			Qualifications in measuring NT MD
Risk			
1:10			
1:100			
1:250			
1:1000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
		Cut off	
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			
			

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off