



Reference No.

: - 2410035632

: 19-Oct-2024

Age/Gender

: 28 Yrs/Female

Pt's Name

: Mrs. VISHAKHA KUMARI 10279446

**NOD-JIT** 

Referred By

: NA

Date

:19-Oct-2024

Sample Collection Date/Time
Sample Receiving Date/Time

: 19-Oct-2024 02:16AM

Approved Date

:19-Oct-2024 04:53AM

Sample From

: JITM Diagnostics

Report Print Time

:19-Oct-2024 11:35AM

### **SEROLOGY**

**Test Description** 

Observed Value

**Biological Reference Interval** 

#### Double Marker, Serum\*

## **DOUBLE MARKER**

## RESULT OF DOWNS SYNDROME SCREENING TEST

BIRTH DAY: 14/03/1995

AGE AT TERM: 29.6

WEIGHT: 58.7 KG LAST

**MENSTRUAL PERIOD:** 

GESTATIONAL AGE BY CRL: 13 WEEKS + 6 DAYS

CROWN RUMP LENGTH IN MM: 73.48 NUCHAL TRANSLUCENCY: 1.8 MM, NT MOM: 0.99

#### RESULTS OF MEASURED SERUM VALUE AND RISK VALUE:

PAPPA : 10.7 mIU/ml MoM : 1.56

Fb-HCG: 32.6 ng/ml MoM: 0.84

## RISK FACTOR

BIOCHEMICAL T21 RISK: < 1:10000

AGE RISK 1:709

COMBINED TRISOMY 21 RISK < 1:10000

TRISOMY 18 + NT < 1:10000

RISK CUT OFF 1:250, HIGH RISK CUT OFF 1:200

#### TRISOMY 21 (DOWN SYNDROME) RISK ASSESSMENT :SCREEN IS NEGATIVE.

THE CALCULATED RISK FOR TRISOMY 21(WITH NUCHAL TRANSLUCENCY)

IS BELOW THE CUT OFF, WHICH INDICATES A LOW RISK.

TRISOMY 13/18 SCREENING: SCREEN IS NEGATIVE.

THE RISK FOR TRISOMY 18 (WITH NUCHAL TRANSLUCENCY) IS < 1:10000. THERE IS NO

STATISTICAL INDICATION FOR TRISOMY 18 RISK.

PLEASE NOTE THAT RISK CALCULATION ARE STATISICAL APPROACHES AND HAVENO DIAGNOSTIC VALUE!

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Page 1 of 2

This test is done by Immuno Diagnostics Pvt Ltd





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\*\*\* End Of Report \*\*\*

Prisca 5.2.0.13

Date of report: 19/10/2024

# J.I.T.M. DIAGNOSTICS

Patient data					
Name MRS	MRS. VISHAKHA KUMARI		Patient ID		
Birthday		14/03/1995	Sample ID	2410035632/NOD	
Age at sample date		29.6	Sample Date	e 18/10/2024	
Gestational age		13 + 6			
Correction factors					
Fetuses 1	IVF		no	Previous trisomy 21 no	
Weight 58.7	diabetes		no	pregnancies	
Smoker no	Origin		Asian		
Biochemical data			Ultrasound data		
Parameter Value	Value Corr. MoM		.  ``		
PAPP-A 10.7 mIU/r	nl	1.56	Method	CRL Robinson	
fb-hCG 32.6 ng/ml		0.84	Scan date	14/10/2024	
Risks at sampling date			Crown rump length in mm 73.48		
Age risk	1:709		1		
Biochemical T21 risk			Nasal bone	present	
Combined trisomy 21 risk <1:10000		Sonographer DR. (MRS.) VIPULA VERMA			
•			Qualifications in measuring NT MD		
1:10  1:100  1:250  Cut off  Pl ap Th wa 1:10000				The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.  After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.  The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!  The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).  The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!	