

LABORATORY REPORT

NAME : MISS.BC0874 REFERRED BY : SELF VISIT NO : VAMP26148185
AGE : 40Y 0M 0D ZERO TARIFF CLIENT CODE COLLECTED ON : 21-04-2026 10:00
GENDER : Female **LAB MR#** : AAMP01479464 RECEIVED ON : 21-04-2026 19:51
OP / IP / DG # : APPROVED ON : 22-04-2026 17:55
REPORT STATUS : Final Report



BIOCHEMISTRY

Maternal Marker Enhanced (1T QUAD) (Serum)

Results

Risk for Down's syndrome (Trisomy 21)	1:100000	Low Risk
Risk for Edward's syndrome (Trisomy 18)	1:100000	Low Risk
Risk for Patau's syndrome (Trisomy 13)	1:100000	Low Risk

Risk for Pre-Eclampsia

Risk for Preeclampsia < 32 weeks	1:100000	Low Risk
Risk for Preeclampsia < 34 weeks	1:33360	Low Risk
Risk for Preeclampsia < 37 weeks	1:2714	Low Risk

Sanjeeta

Dr. Sanjeeta
MBBS,MD (Biochemistry)
Consultant Biochemist

Disclaimer:

1. All results released pertain to the specimen as received by the lab for testing and under the assumption that the patient indicated or identified on the bill/test requisition form is the owner of the specimen.
2. Clinical details and consent forms, especially in Genetic testing, histopathology, as well as wherever applicable, are mandatory to be accompanied with the test requisition form. The non-availability of such information may lead to delay in reporting as well as misinterpretation of test results. The lab will not be responsible for any such delays or misinterpretations thereof.
3. Test results are dependent on the quality of the sample received by the lab. In case the samples are preprocessed elsewhere (e.g., paraffin blocks), results may be compromised.

Generated On 25-Apr-2026 11:17:51

This is an electronically authenticated laboratory report.

Page 1 of 2

Sin No: 20385481



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- 4. Tests are performed as per the schedule given in the test listing and in any unforeseen circumstances, report delivery may be affected.
- 5. Test results may show inter-laboratory as well as intra-laboratory variations as per the acceptable norms.
- 6. Genetic reports as well as reports of other tests should be correlated with clinical details and other available test reports by a qualified medical practitioner. Genetic counselling is advised in genetic test reports by a qualified genetic counsellor, medical practitioner or both.
- 7. Samples will be discarded post processing after a specified period as per the laboratory's retention policy. Kindly get in touch with the lab for more information.
- 8. If accidental damage, loss, or destruction of the specimen is not attributable to any direct or negligent act or omission on the part of Ampath Labs or its employees, Ampath shall in no event be liable. Ampath lab's liability for a lack of services, or other mistakes and omissions, shall be restricted to the amount of the patient's payment for the pertinent laboratory services.



PATIENT REPORT Miss.BC0874 -

22/04/2026

Requestor: -, -

REQUESTOR FIRST NAME: RAJITHA	REQUESTOR SURNAME: DR
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Patient 20385481: Miss.BC0874, -

PATIENT ID: 20385481	BIRTH DATE: 01/01/2000	ETHNICITY: South Asian
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Pregnancy, Calculated EDD: 06/11/2026 (MAEDD: 26.85)

MAEDD: 26.85	CALCULATED EDD: 06/11/2026	GEST. DATE: 30/01/2026
SELECTED GEST. METHOD: CRL	LMP DATE: 11/01/2026	SMOKING STATUS: Non smoker
INSULIN DEP. DIABETIC: No	NO. OF FETUSES: 1	MONOZYGOUS: No
MATERNAL WEIGHT [KG]: 50	HEIGHT [CM]: 170	PAST T21 - DOWN'S SYNDROME: No
PAST T18 - EDWARDS' SYNDROME: No	PAST T13 - PATAU'S SYNDROME: No	RISK ASSESSED: At term
SCREENING PROTOCOL: Screening_4.0		

Ultrasound

SCAN DATE: 22/04/2026	CRL: 50	BPD: -	GEST. AT SAMPLE DATE (W + D): 11 w 5 d
CRL (#2): -	BPD (#2): -	HC (#2): -	WEIGHT [KG]: 50

Tests

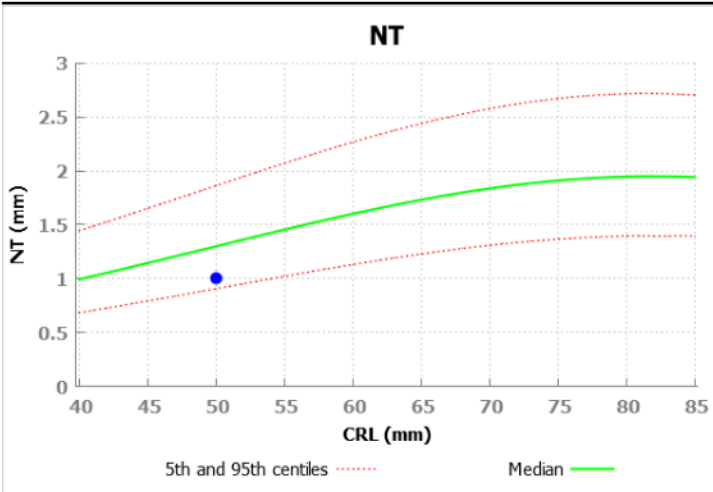
TEST	SAMPLE ID	DATE	GEST. AT SAMPLE DATE (W + D)	VALUE	UNIT	CORR. MOM	WEIGHT [KG]
AFP (Signed)	20385481	22/04/2026	11 w 5 d	34	U/mL	3.45	50
hCGb (Signed)	20385481	22/04/2026	11 w 5 d	20	ng/mL	0.38	50
PAPP-A (Signed)	20385481	22/04/2026	11 w 5 d	4000	mU/L	1.66	50
PIGF (Signed)	20385481	22/04/2026	11 w 5 d	74	pg/mL	2.41	50
NB (Signed)	-	22/04/2026	11 w 5 d	Present	-	-	50
NT (Signed)	-	22/04/2026	11 w 5 d	1	mm	0.76	50



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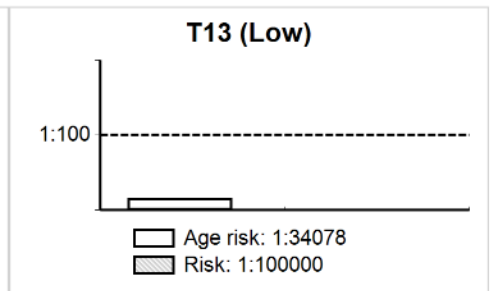
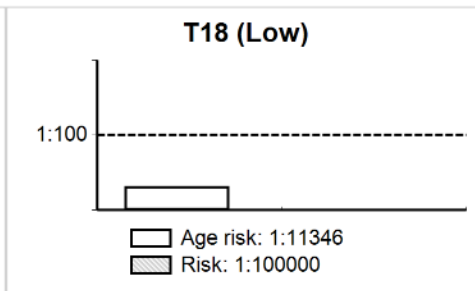
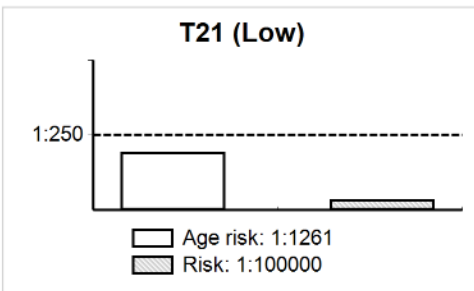
22/04/2026



Distribution median, 5th and 95th centiles obtained from publication: Wright D, Kagan KO, Molina FS, Gazzoni A, Nicolaides KH. A mixture model of nuchal translucency thickness in screening for chromosomal defects. Ultrasound Obstet Gynecol 2008;31:376-83

Risks, Risk assessed: At term

RISK NAME:	RISK RESULT:	RISK:	TWIN RISK RESULT:	TWIN RISK:	AGE RISK:	CUT-OFF:
T21 (Signed)	Low	1:100000	-	-	1:1261	1:250
T18 (Signed)	Low	1:100000	-	-	1:11346	1:100
T13 (Signed)	Low	1:100000	-	-	1:34078	1:100
PE < 32 (Signed)	Low	1:100000	-	-	1:2767	1:100
PE < 34 (Signed)	Low	1:33360	-	-	1:996	1:100
PE < 37 (Signed)	Low	1:2714	-	-	1:250	1:100

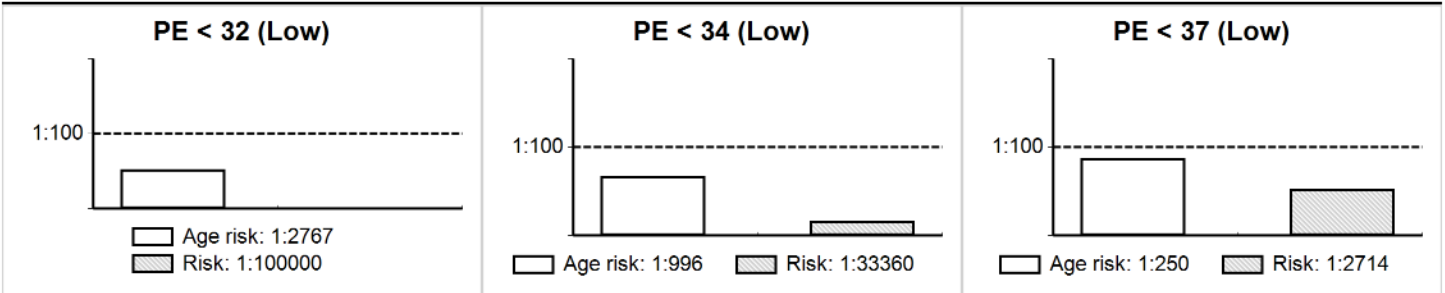


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