SAGEPATH LABS PVT LTD

Prisca 5.1.0.17

Date of report: 18/11/24

NA

Patient data

Name	Mrs. PRANJYA PARIDA TWIN A	Patient ID	0652411170194
Birthday	02/04/91	Sample ID	A1583430 A
Age at sample date	33.6	Sample Date	17/11/24
Gestational age	12 + 6		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	81	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

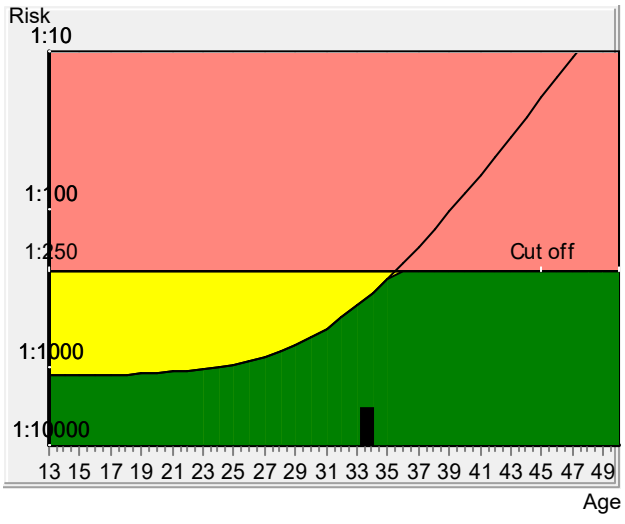
Parameter	Value	Corr. MoM
PAPP-A	4.14 mIU/mL	1.17
fb-hCG	20.56 ng/ml	0.61

Risks at sampling date

Age risk	1:366
Biochemical T21 risk	1:9046
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 6
Method	CRL Robinson
Scan date	17/11/24
Crown rump length in mm	68
Nuchal translucency MoM	0.52
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



Trisomy 21

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!

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

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NA

Patient data

Name	Mrs. PRANJYA PARIDA TWIN B	Patient ID	0652411170194
Birthday	02/04/91	Sample ID	A1583430 B
Age at sample date	33.6	Sample Date	17/11/24
Gestational age	12 + 6		

Correction factors

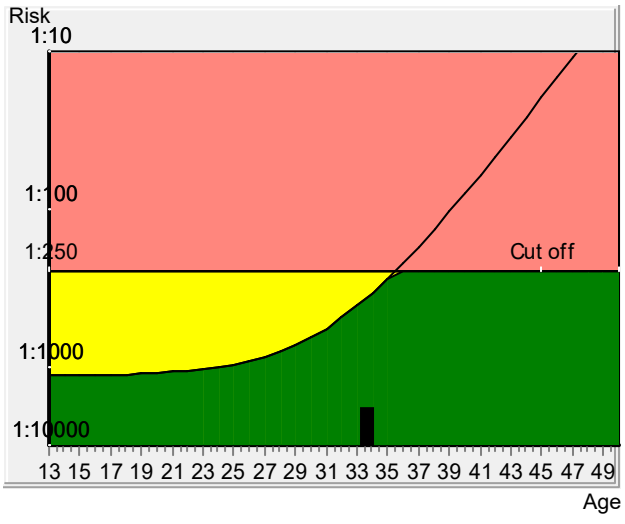
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	81	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	4.14 mIU/mL	1.17
fb-hCG	20.56 ng/ml	0.61

Risks at sampling date

Age risk	1:366
Biochemical T21 risk	1:9046
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Ultrasound data

Gestational age	12 + 6
Method	CRL Robinson
Scan date	17/11/24
Crown rump length in mm	67
Nuchal translucency MoM	0.65
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD

Trisomy 21

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!

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off