SAGEPATH LABS PVT LTD.

Prisca 5.1.0.17

Date of report: 29/11/24

NA

Patient data			
Name	Mrs. MEENA RANSINGE TWIN A	Patient ID	0372411280058
Birthday	06/06/96	Sample ID	A0422038 TWIN A
Age at sample date	28.5	Sample Date	28/11/24
Gestational age	13 + 0		

Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	75	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

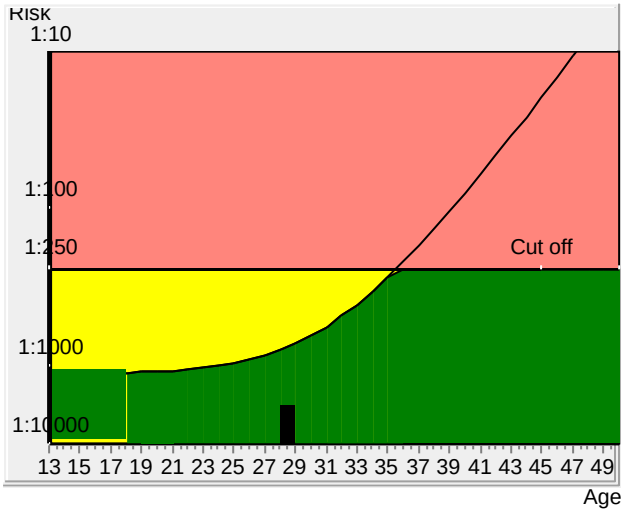
Parameter	Value	Corr. MoM
PAPP-A	4.41 mIU/mL	0.57
fb-hCG	82.45 ng/ml	1.15

Risks at sampling date

Age risk	1:774
Biochemical T21 risk	1:941
Combined trisomy 21 risk	1:5529
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	28/11/24
Crown rump length in mm	69
Nuchal translucency MoM	0.69
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 5529 women with the same data, there is one woman with a trisomy 21 pregnancy and 5528 women with not affected pregnancies.
The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.
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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Prisca 5.1.0.17

Date of report: 29/11/24

NA

Patient data

Name	Mrs. MEENA RANSINGE TWIN B	Patient ID	0372411280058
Birthday	06/06/96	Sample ID	A0422038 TWIN B
Age at sample date	28.5	Sample Date	28/11/24
Gestational age	13 + 1		

Correction factors

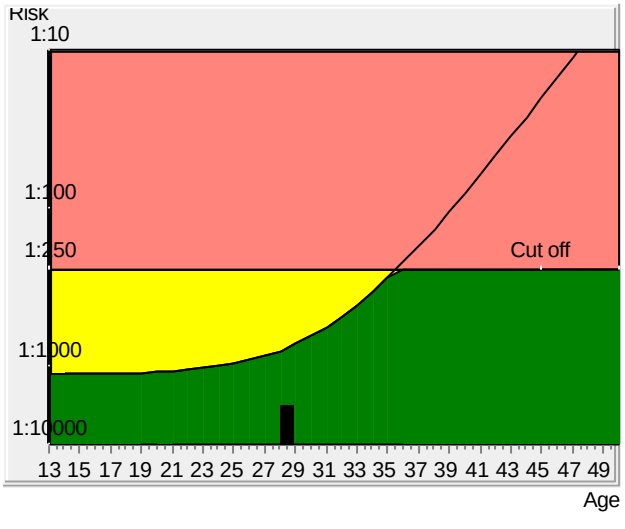
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	75	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	4.41 mIU/mL	0.54
fb-hCG	82.45 ng/ml	1.19

Risks at sampling date

Age risk	1:778
Biochemical T21 risk	1:747
Combined trisomy 21 risk	1:4481
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	13 + 1
Method	CRL Robinson
Scan date	28/11/24
Crown rump length in mm	72
Nuchal translucency MoM	0.61
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 4481 women with the same data, there is one woman with a trisomy 21 pregnancy and 4480 women with not affected pregnancies.

The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.

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