Sage Path Labs Pvt Ltd
Plot No. 564, First Floor, Buddanagar, Peerzadiguda, Boduppal, Hyderabad
Telangana - 500092

Prisca 5.1.0.17

Date of report: 18/04/25

N A

Patient data			
Name	W/O SHIV KUMAR TWIN A	Patient ID	0692504170029
Birthday	05/05/93	Sample ID	B2762425
Age at sample date	32.0	Sample Date	17/04/25
Gestational age	13 + 0		

Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	49	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

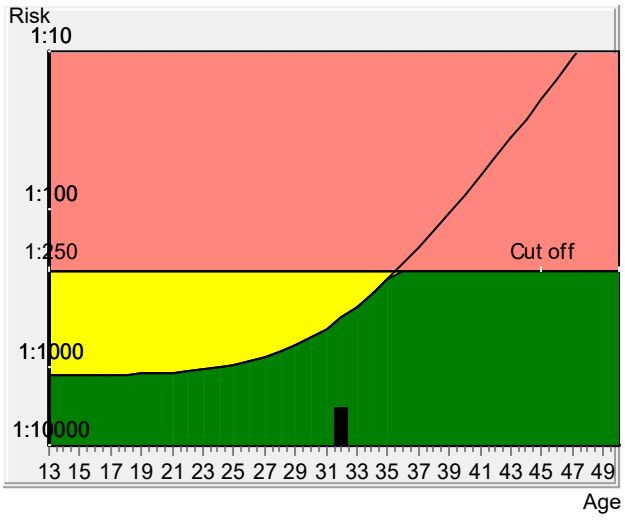
Parameter	Value	Corr. MoM
PAPP-A	8.57 mIU/mL	0.76
fb-hCG	37.85 ng/mL	0.44

Risks at sampling date

Age risk	1:497
Biochemical T21 risk	1:9307
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 5
Method	CRL Robinson
Scan date	15/04/25
Crown rump length in mm	65
Nuchal translucency MoM	1.02
Nasal bone	present
Sonographer	N A
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 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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Patient data			
Name	W/O SHIV KUMAR TWIN B	Patient ID	0692504170029
Birthday	05/05/93	Sample ID	B2762425
Age at sample date	32.0	Sample Date	17/04/25
Gestational age	13 + 0		

Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	49	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

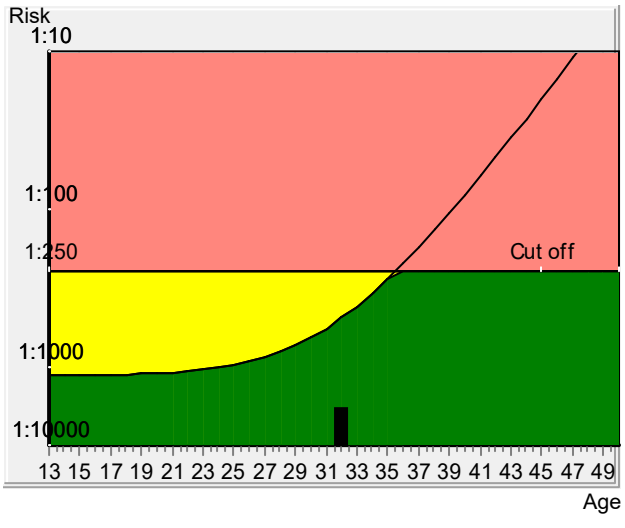
Parameter	Value	Corr. MoM
PAPP-A	8.57 mIU/mL	0.76
fb-hCG	37.85 ng/mL	0.44

Risks at sampling date

Age risk	1:497
Biochemical T21 risk	1:9307
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 5
Method	CRL Robinson
Scan date	15/04/25
Crown rump length in mm	65
Nuchal translucency MoM	0.84
Nasal bone	present
Sonographer	N A
Qualifications in measuring NT	MD



Trisomy 13/18 + NT

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

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 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!
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