Prisca 5.2.0.13

Date of report: 18/01/2025

JITM Diagnostics

Patient data

Name	MRS. POOJA (A)	Patient ID	
Birthday	1/01/1993	Sample ID	2501026640/NOD (A)
Age at sample date	32.0	Sample Date	18/01/2025
Gestational age	13 + 2		

Correction factors

Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	no
Weight	44.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

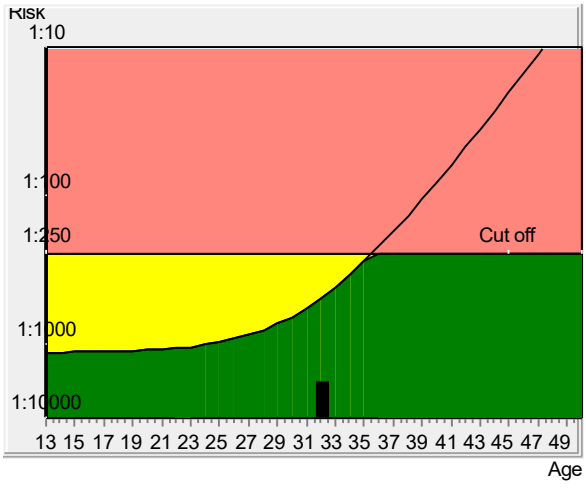
Parameter	Value	Corr. MoM
PAPP-A	10.1 mIU/ml	0.71
fb-hCG	84.1 ng/ml	0.85

Risks at sampling date

Age risk	1:494
Biochemical T21 risk	1:2001
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 1
Method	CRL Robinson
Scan date	17/01/2025
Crown rump length in mm	71
Nuchal translucency MoM	0.73
Nasal bone	present
Sonographer	DR. KIRTI SHARMA
Qualifications in measuring NT	M.D



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!

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

Prisca 5.2.0.13

Date of report: 18/01/2025

JITM Diagnostics

Patient data

Name	MRS. POOJA (B)	Patient ID	
Birthday	1/01/1993	Sample ID	2501026640/NOD (B)
Age at sample date	32.0	Sample Date	18/01/2025
Gestational age	13 + 3		

Correction factors

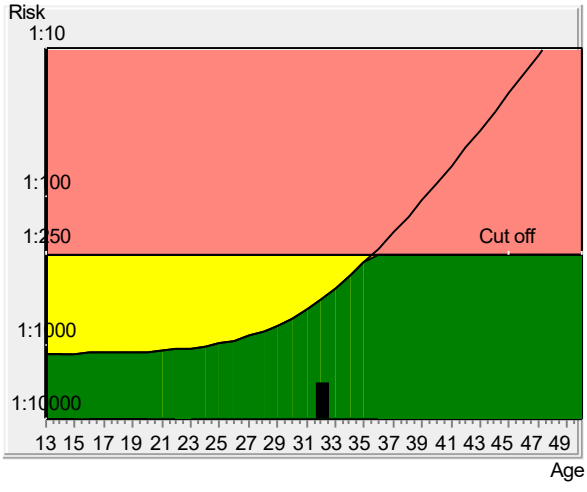
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	no
Weight	44.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	10.1 mIU/ml	0.67
fb-hCG	84.1 ng/ml	0.86

Risks at sampling date

Age risk	1:496
Biochemical T21 risk	1:1704
Combined trisomy 21 risk	1:4747
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 + 2
Method	CRL Robinson
Scan date	17/01/2025
Crown rump length in mm	74
Nuchal translucency MoM	1.15
Nasal bone	present
Sonographer	DR. KIRTI SHARMA
Qualifications in measuring NT	M.D

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 4747 women with the same data, there is one woman with a trisomy 21 pregnancy and 4746 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