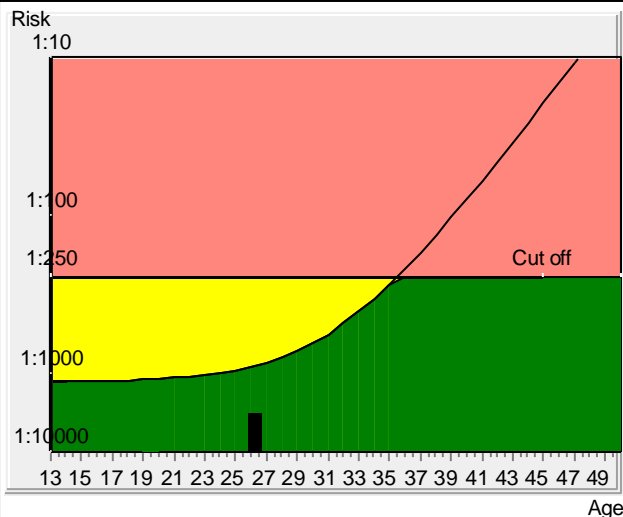


JITM Diagnostics

Patient data				
Name	MRS. ASHPREET KAUR		Patient ID	
Birthday	4/01/1999		Sample ID	2503040386/NOD
Age at sample date	26.2		Sample Date	20/03/2025
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	70.5	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	4.65 mIU/ml	1.23	Method	CRL Robinson
fb-hCG	118 ng/ml	2.87	Scan date	19/03/2025
Risks at sampling date			Crown rump length in mm	65.21
Age risk		1:915	Nuchal translucency MoM	0.79
Biochemical T21 risk		1:639	Nasal bone	present
Combined trisomy 21 risk		1:3489	Sonographer	DR. PREETY SHARMA AGNIHOTRI
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	M.D
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 3489 women with the same data, there is one woman with a trisomy 21 pregnancy and 3488 women with not affected pregnancies.

The free beta HCG level is high.

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