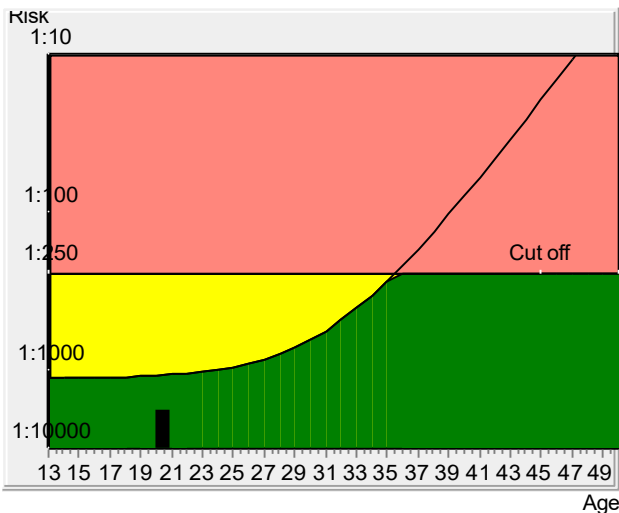


JITM Diagnostics

Patient data			
Name	MRS. POOJA SHARMA	Patient ID	
Birthdate	11/06/2004	Sample ID	2411001642/NOD
Age at sample date	20.4	Sample Date	1/11/2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.26 mIU/ml	1.83	Gestational age 12 + 4
fb-hCG	162 ng/ml	3.90	Method CRL Robinson
Risks at sampling date			Scan date 30/10/2024
Age risk		1:1095	Crown rump length in mm 64
Biochemical T21 risk		1:673	Nuchal translucency MoM 0.74
Combined trisomy 21 risk		1:3462	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. PULKIT SONI
			Qualifications in measuring NT M.D
			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 3462 women with the same data, there is one woman with a trisomy 21 pregnancy and 3461 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
			Trisomy 13/18 + NT
			<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>



Sign of Physician