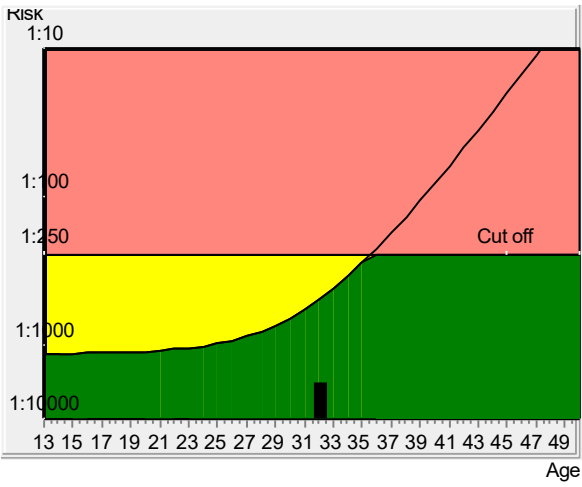


JITM Diagnostics

Patient data			
Name	MRS. HARPREET KAUR	Patient ID	
Birthday	23/10/1992	Sample ID	2411001891/NOD
Age at sample date	32.0	Sample Date	2/11/2024
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.46 mIU/ml	0.91	Gestational age 13 + 2
fb-hCG	27.1 ng/ml	0.66	Method CRL Robinson
Risks at sampling date			Scan date 1/11/2024
Age risk		1:498	Crown rump length in mm 73.3
Biochemical T21 risk		1:6190	Nuchal translucency MoM 0.72
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. NEERJA CHOPRA
			Qualifications in measuring NT M.D
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>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.</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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			



**Sign of Physician**