## Sarita Vihar

Prisca 5.2.0.13

Date of report: 8/06/2024

Patient data						
Name		MR	S. MEHAK	Patient ID		
Birthday	27/10/1999			Sample ID		2406013111/NOD
Age at sample date				Sample Date 7/06/2024		
Gestational age 12 + 5						
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	57.6	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound da	ata	
Parameter	Value		Corr. MoM	Gestational	age	12 + 5
PAPP-A	4.15 mIU/m	nl	0.91	Method		CRL Robinson
fb-hCG	46.5 ng/ml 1.05			Scan date		7/06/2024
Risks at sampling date			Crown rump length in mm 66.7			
Age risk	1:983			Nuchal translucency MoM 0.77		
Biochemical T21 risk				Nasal bone		present
Combined trisomy 21 risk <1:10000			Sonographe			
•				Qualifications in measuring NT		
Risk 1:10				Trisomy 21	ated risk for Trisomy 2	
1:1000 1:10				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 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!		
translucency) is < 1:						

Below Cut Off, but above Age Risk

below cut off

Sign of Physician

above cut off