

Prisca

5.1.0.17

Date of report:

10/07/24

Patient data				
Name	Mrs. POOJA SINGH		Patient ID	0662407090335
Birthday	05/10/97		Sample ID	A0779764
Age at sample date	26.8		Sample Date	09/07/24
Gestational age	13 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	42.9	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 4
PAPP-A	16.15 mIU/mL	1.79	Method	CRL Robinson
fb-hCG	38.99 ng/mL	1.13	Scan date	08/07/24
Risks at sampling date			Crown rump length in mm	78.4
Age risk	1:908		Nuchal translucency MoM	0.95
Biochemical T21 risk	<1:10000		Nasal bone	present
Combined trisomy 21 risk	<1:10000		Sonographer	NA
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD
Risk			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 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>				

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off