

Prisca 5.1.0.17
Date of report: 08-08-2019

Patient data			
Name	MRS. HARPAL KAUR	Patient ID	1908220432/AMB
Birthday	08-01-1986	Sample ID	1908220432/AMB
Age at delivery	34.1	Sample Date	07-08-2019
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.5 mIU/ml	0.61	Gestational age 12 + 0
fb-hCG	103 ng/ml	3.30	Method CRL Robinson
			Scan date 30-07-2019
Risks at term		Crown rump length in mm 56	
Age risk		1:513	Nuchal translucency MoM 0.54
Biochemical T21 risk		1:53	Nasal bone present
Combined trisomy 21 risk		1:354	Sonographer .
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 354 women with the same data, there is one woman with a trisomy 21 pregnancy and 353 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.</p> <p>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>	
		<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>	
Trisomy 13/18 + NT			

Sign of Physician

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