

Prisca 5.0.2.37  
Date of report: 04-06-2019

Patient data			
Name	MRS. HARPREET	Patient ID	
Birthday	01-05-1996	Sample ID	211906030008
Age at sample date	23	Sample Date	03-06-2019
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58.8	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.8 mIU/ml	1.51	13 + 0
fb-hCG	75.1 ng/ml	2.08	Method
			CRL Robinson
			Scan date
Risks at sampling date		Crown rump length in mm	
Age risk	1:888	Nuchal translucency MoM	
Biochemical T21 risk	1:2146	Nasal bone	
Combined trisomy 21 risk	1:4321	Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	
		MD	
		Trisomy 21	
Risk		<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 4321 women with the same data, there is one woman with a trisomy 21 pregnancy and 4320 women with not affected pregnancies.</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

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