

Patient data			
Name	MRS. HARJINDER KAUR	Patient ID	1807220769/AMB
Birthdate	11-11-1982	Sample ID	1807220769/AMB
Age at sample date	35.7	Sample Date	28-07-2018
Gestational age	11 + 4		
Correction factors			
Fetuses	1 IVF	no	Previous trisomy 21 no
Weight	68 diabetes	no pregnancies	
Smoker	no	Origin Asian	
Biochemical data	Ultrasound data		
Parameter	Value	Corr. MoM	
PAPP-A	4.28 mIU/ml	2.14	Gestational age 11 + 2
fb-hCG	140 ng/ml	3.49	Method CRL Robinson
Risks at sampling date	Scan date 26-07-2018		
Age risk	1:223	Crown rump length in mm 46.8	
Biochemical T21 risk	1:215	Nuchal translucency MoM 1.33	
Combined trisomy 21 risk	1:262	Nasal bone present	
Trisomy 13/18 + NT	<1:10000	Sonographer .	
Risk	1:10	Qualifications in measuring NT MD	
	1:1000	Trisomy 21	
	1:250	<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 262 women with the same data, there is one woman with a trisomy 21 pregnancy and 261 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</p>	
	Cutoff		
	1:10000		

1315 1719 212325 2729 313335 3739 414345 4749  
Age

on the risk assessment ! Calculated risks have no diagnostic value!

Trisomy 13/18 + NT

 below cut off **The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.**  above cut off

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**Sign of Physician**

