

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. MANPREET	Patient ID	
Birth day	01-07-2000	Sample ID	2206220371/AMB
Age at sample date	22.0	Sample Date	14-06-2022
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	50.8	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	12.1 mIU/ml	1.57	Gestational age
fb-hCG	321 ng/ml	7.74	Method
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:1097	67
Biochemical T21 risk		1:482	Nuchal translucency MoM
Combined trisomy 21 risk		>1:50	1.18
Trisomy 13/18 + NT		<1:10000	Nasal bone
			absent
			Sonographer
			.
			Qualifications in measuring NT
			MD
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</b></p> <p>After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy.</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>			
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>			

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**Sign of Physician**