

Prisca 5.1.0.17

Date of report: 09/08/24

Patient data			
Name	Mrs. PRANALI UTANE	Patient ID	0372408080041
Birthday	01/09/01	Sample ID	A0231282
Age at sample date	22.9	Sample Date	08/08/24
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	18.7 mIU/mL	3.14	Gestational age 12 + 4
fb-hCG	44.25 ng/mL	1.00	Method CRL Robinson
Risks at sampling date			
Age risk	1:1039		Scan date 07/08/24
Biochemical T21 risk	<1:10000		Crown rump length in mm 64.3
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 1.09
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer N A			
Qualifications in measuring NT MD			
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.			
<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 PAPP-A 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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off