

Prisca 5.1.0.17  
**Date of report:** 11-04-2025

N A

Patient data								
Name	Mrs. K ANITHA TWIN A	Patient ID	0042504110003					
Birthday	02-01-1997	Sample ID	23168743					
Age at sample date	28.3	Sample Date	10-04-2025					
Gestational age	13 + 2							
Correction factors								
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	101	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	9.62 mIU/mL	1.88	Gestational age 13 + 2					
fb-hCG	35.01 ng/mL	0.54	Method CRL Robinson					
Risks at sampling date								
Age risk	1:797		Scan date 10-04-2025					
Biochemical T21 risk	<1:10000		Crown rump length in mm 75					
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.76					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
Risk								
1:10			Sonographer N A					
1:100			Qualifications in measuring NT MD					
1:250			Trisomy 21					
1:1000			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.					
1:10000			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.					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	Age		The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.					
			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!					
			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).					
			The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!					
Trisomy 13/18 + NT								
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.								

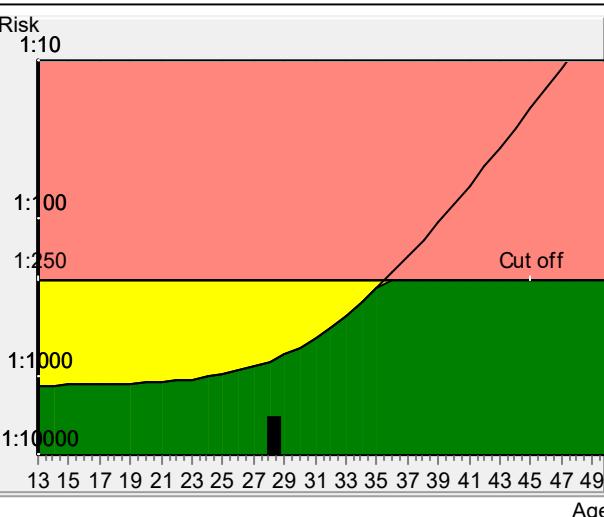
Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off

N A

Patient data								
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Combined trisomy 21 risk		<1:10000	Nuchal translucency MoM 0.87					
Trisomy 13/18 + NT		<1:10000	Nasal bone present					
Sonographer N A								
Qualifications in measuring NT MD								
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> <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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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								
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off