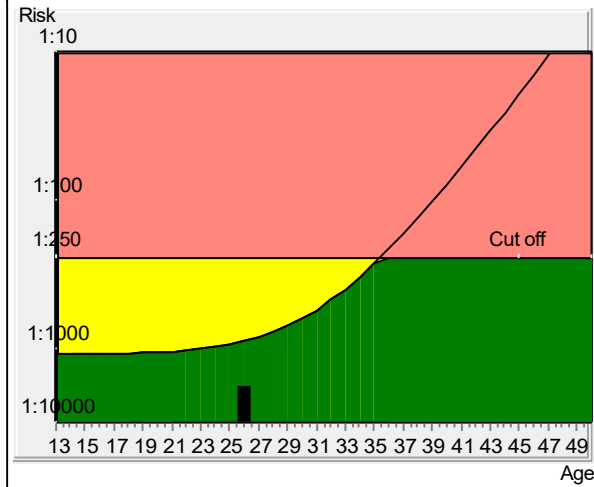


Patient data			
Name	MRS. CHERCHI HELEN	Patient ID	
Birthday	27/07/1998	Sample ID	2408015627/NOD
Age at sample date	26.0	Sample Date	9/08/2024
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	87	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.6 mIU/ml	1.22	11 + 5
fb-hCG	118 ng/ml	2.80	Method
			CRL Robinson
			Scan date
			6/08/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:900	51.65
Biochemical T21 risk		1:663	Nuchal translucency MoM
Combined trisomy 21 risk		1:3618	0.58
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..



Trisomy 21
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
 After the result of the Trisomy 21 test (with NT) it is expected that among 3618 women with the same data, there is one woman with a trisomy 21 pregnancy and 3617 women with not affected pregnancies.
 The free beta HCG level is high.
 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 hold 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