SAGEPATH LABS PVT LTD.

Prisca 5.1.0.17

Date of report: 21-07-2023

ASHRA KANWAL

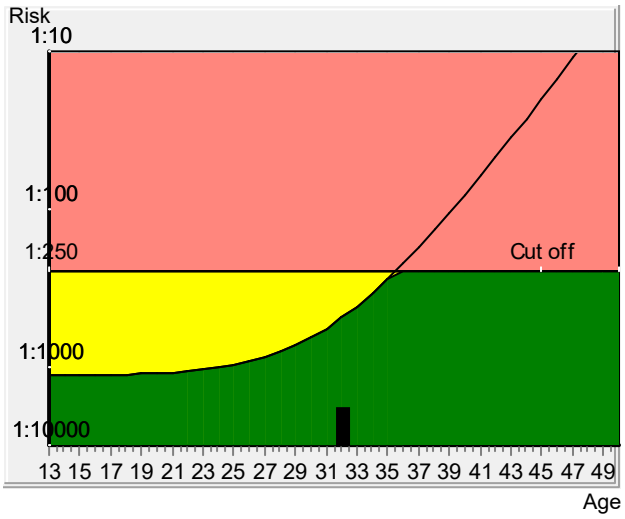
Patient data			
Name	Mrs. B KUMARI	Patient ID	0012307200070
Birthday	11-06-1991	Sample ID	24499814
Age at sample date	32.1	Sample Date	20-07-2023
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	57.8	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	2.3 mIU/mL	0.46
fb-hCG	35.74 ng/mL	0.95

Risks at sampling date	
Age risk	1:484
Biochemical T21 risk	1:513
Combined trisomy 21 risk	1:3105
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	12 + 5
Method	CRL Robinson
Scan date	18-07-2023
Crown rump length in mm	65
Nuchal translucency MoM	0.66
Nasal bone	present
Sonographer	ASHRA KANWAL
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.

After the result of the Trisomy 21 test (with NT) it is expected that among 3105 women with the same data, there is one woman with a trisomy 21 pregnancy and 3104 women with not affected pregnancies.

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

below cut off | Below Cut Off, but above Age Risk | above cut off