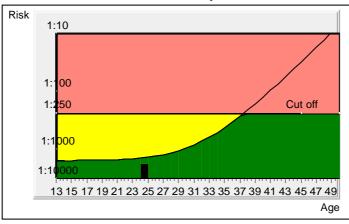
KOS DIAGNOSTIC LAB 6349/1 NICHOLSON ROAD, AMBALA CANTT

Results for:Sample noDate of report:MRS. SHARANPREET KAUR2209220623/AMB27-09-2022

Referring Doctors

KOS DIAGNOSTIC LAB

Summary



	Patient data
Age at delivery	24.5
WOP	22 + 0
Weight	74.0 kg
Patient ID	
Ethnic origin	Asian

Risks at term				
Biochemical risk for Tr.21	1:2155			
Age risk:	1:1405			
Neural tube defects risk	1:1008			

For MRS. SHARANPREET KAUR, born on 08-08-1998, a screening test was performed on the 22-09-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	_	_	-
	Valu	ıe	Corr. MoMs
AFP	114	ng/ml	1.61
HCG	32304	miu/ml	2.55
uE3	2.1	ng/ml	1.05
Gestation ag	ae	22+0	

Gestation age 22+0

Method BPD Hadlock

The MoMs have been corrected according to: maternal weight ethnic origin

TRISOMY 21 SCREENING

The calculated risk for Trisomy 21 is below the cut off which represents a low risk.

After the result of the Trisomy 21 test it is expected that among 2155 women with the same data, there is one woman with a trisomy 21 pregnancy and 2154 women with not affected pregnancies.

The 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!

TRISOMY 18 SCREENING

The calculated risk for trisomy 18 is < 1:10000, which indicates a low risk.

NEURAL TUBE DEFECTS (NTD) SCREENING

The corrected MoM AFP (1.61) is located in the low risk area for neural tube defects.





