## Sarita Vihar

Prisca 5.1.0.17

Date of report: 09-06-2022

Patient data			
Name N	MRS. GURJOT KAUR		2206220204/AMB
Birthday	17-04-1997	Sample ID	2206220204/AMB
Age at sample date	25.1	Sample Date	e 08-06-2022
Gestational age	11 + 5		
Correction factors			
Fetuses 1	IVF	no	Previous trisomy 21 no
Weight 64.9	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data	Ultrasound data		ata
Parameter Value	Value Corr. MoM Gestational age 11 +		
PAPP-A 1.9 mIU/m	I 0.76	Method CRL Robinson	
fb-hCG 154 ng/ml	3.21	Scan date 07-06-2022	
Risks at sampling date		Crown rump length in mm 49.98	
Age risk	1:927	Nuchal translucency MoM 1.33	
Biochemical T21 risk	1:176	Nasal bone present	
Combined trisomy 21 risk	1:225	Sonographer .	
Trisomy 13/18 + NT	<1:10000	1 3	
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal
1:1000 1:10	Age /18 (with nuchal	translucency) is above the cut off, which indicates an increased risk.  After the result of the Trisomy 21 test (with NT) it is expected that among 225 women with the same data, there is one woman with a trisomy 21 pregnancy and 224 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!	

Sign of Physician

below cut off