Priska	5.0.2.37
Date of report:	**24-06-2019**

Patient data			
Name	MRS. PARNEET KAUR	Patient ID	
Birthday	08-02-1998	Sample ID	1906221093/AMB
Age at sample date	21.4	Sample Date	22-06-2019
Gestational age	12 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	unknown
Weight	65	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 0
PAPP-A	2.8 mIU/ml	0.87	Method	CRL Robinson
fb-hCG	145.1 ng/ml	3.66	Scan date	18-06-2019

Risks at sampling date		Crown rump length in mm	56.82
Age risk	1:1069	Nuchal translucency MoM	0.95
Biochemical T21 risk	1:192	Nasal bone	unknown
Combined trisomy 21 risk	1:910	Sonographer	.
Trisomy 13/18 + NT	<1:10000	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 910 women with the same data, there is one woman with a trisomy 21 pregnancy and 909 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

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