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 Date of Report
 15/1/2022

 PRISCA
 5.2.0.13

Name MRS. DIVYA GOYAL		Patient ID		012201140001
	31/10/1998	Sample ID		11307530
	23.2	Sample Date		14/1/2022
	13+1			
1 IVF		unknown	Previous trisomy 21	unknown
64 Diabetes		NO	Pregnancies	unknown
NO Origin		Asian		
Biochemical Data		Ultrasound Data		
Value	Corr Mom	Gestational age	2	12+6
$2.54~\mathrm{mIU/ml}$	0.49	Method		CRL(<>Robinson
12.9 ng/ml	0.3	Scan date		12/1/2022
Risks at sampling date		Nuchal translucency 1.5		
	1:1046	Nuchal translucency MoM 0.89		
	<1:10000	Nasal bone presen		
	<1:10000			
	<1:10000			
Risk		Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:100000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000000		The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
	1 IVF 64 Diabetes NO Origin Value 2.54 mIU/ml 12.9 ng/ml	31/10/1998 23.2 13+1 1 IVF 64 Diabetes NO Origin Value Corr Mom 2.54 mIU/ml 0.49 12.9 ng/ml 0.3 1:1046 <1:10000 <1:10000 <1:10000 <1:10000	1 IVF unknown 64 Diabetes NO NO Origin Asian Value Corr Mom 2.54 mIU/ml 0.49 Method 12.9 ng/ml 0.3 Scan date Nuchal translu 1:1046 Nuchal translu 41:10000 Nasal bone 1:10000 1:10000 Down's Syndre The calculated cut off, which is asame data, ther pregnancy and The calculated the information note that the rihave no diagnor on the risk asset.	31/10/1998 Sample ID 23.2 Sample Date 13+1 IVF unknown Previous trisomy 21 64 Diabetes NO Origin Asian Value Corr Mom 2.54 mIU/ml 0.49 Method 12.9 ng/ml 0.3 Scan date Nuchal translucency Nuchal translucency MoM 41:10000 41:10000 1:10000 Down's Syndrome Risk (Trisomy 21 String and the represents a low risk. After the result of the Trisomy 21 (witcut off, which represents a low risk.) After the result of the Trisomy 21 test if expected that among more than 10000 same data, there is one woman with a translucency and 9999 women with not after the result of the Trisomy 21 test if expected that among more than 10000 same data, there is one woman with a translucency and 9999 women with not after the result of the Trisomy 21 test if expected that among more than 10000 same data, there is one woman with a translucency and 9999 women with not after the result of the Trisomy 21 test if expected that among more than 10000 same data, there is one woman with a translucency and 9999 women with not after the result of the Trisomy 21 test if expected that among more than 10000 same data, there is one woman with a translucency and 10000 same data, there is one woman with a translucency and 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 same data, there is one woman with a translucency are represented that among more than 10000 sam