

*Free Home Sample Collection 9999 778 778



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Date of Report 14-02-2025 PRISCA 5 2 0 13

5.2.0.13	PRISCA					
						Patient Data
2502130149	125	Patient ID	KI KHATUN	ARS. SILI	1	Name
11925792		Sample ID	08-03-2005			Birthday
13-02-2025	1	Sample Date	19.9			Age at Sample date
			13+3			Gestational age
						Correction factors
unknown	Previous trisomy 21	unknown		IVF	1	Fetuses
unknown	Pregnancies	NO		Diabetes	52	Weight in kg
		Asian		Origin	NO	Smoker
	ıta	Ultrasound Da				Biochemical Data
13+3		Gestational age	Corr Mom		Value	Parameter
<>Robinson)	CRL (<>	Method	0.66	mIU/ml	5.9	PAPP-A
13-02-2025	J	Scan date	1.3	ng/ml	39.1	fb-hCG
71.6	ength in mm	Crown rump le			e	Risks at sampling date
0.67	cency MoM	Nuchal translu	1:1122			Age Risk
PRESENT	I	Nasal bone	1:1502			Biochemical T21 risk
DR.MONIKA	1	Sonographer	1:8622		risk	Combined trisomy 21 ris
MD	n measuring NT	Qualifications	<1:10000			Trisomy 13/18 + NT
Down's Syndrome Risk (Trisomy 21 Screening)						Risk
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 (with NT) it is expected that among 8622 women with the same data, there is one woman with a trisomy 21 pregnancy and 8621 women with not affected pregnancies.			1:10 1:100 1:250 Gtoff			
The laboratory cannot be hold responsible for their impact on the						1:1000 1:1000 13 15 17 19 21 23 25 Trisomy 13/18+NT The calculated risk for Tri
trisomy 21 pregnancy and 8621 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note the the risk calculations are statistical aapproaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-521998).			414345474		Trisomy 13/18	1:20 1:1000 1:1000 1:10101

which indicates a low risk