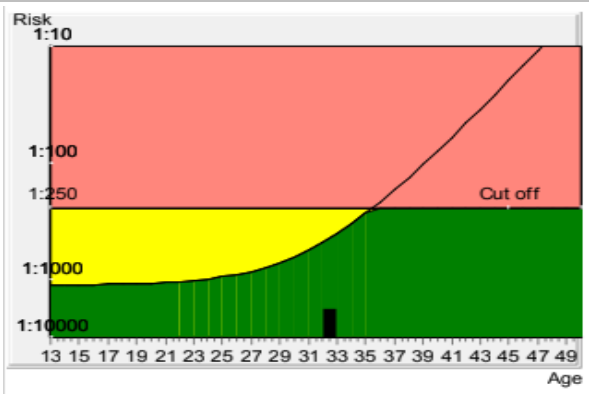



Date of Report 2/6/2022  
PRISCA 5.1.0.17

Patient Data					
Name	<b>SARASWATI (Surrogate)</b>	Patient ID	012206010055		
Birthday	31/12/1989 (Donor)	Sample ID	11447542		
Age at term	32.1	Sample Date	1/6/2022		
Gestational age	12+4				
Correction factors					
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	49	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+2	
PAPP-A	4.17 mIU/ml	0.72	Method	CRL (<>Robinson)	
fb-hCG	59.3 ng/ml	1.19	Scan date	30/05/2022	
Risks at sampling date			Nuchal Translucency	1.1	
Age Risk	1:452		Nuchal Translucency MoM	0.74	
Biochemical T21 risk	1:901		Nasal Bone	Present	
Combined T21 risk	1:5004				
Trisomy 13/18+NT	<1:10000				
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
			<p><b>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5004 women with the same data, there is one woman with a trisomy 21 pregnancy and 5003 women with not affected pregnancies.</p> <p>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 approaches and have no diagnostic value!</p>		
Trisomy 13/18+NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
<p><b>The calculated risk for Trisomy 13/18 (with NT) is &lt;1:10000 , which indicates a low risk</b></p>					

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk