Name: N	MAANVI	C	amta atı			Gender: Female
Weight: 73			ontact: rthdate: 2003-0	7-03	Λ -	
	Sian Kg		Twins: No	, 03		ge of EDC: 21.28 Year c method: CRL Robinson
LMP Day:			ender:		GA cal	c method: CRL Robinson
Sample info	rmation					
Send time: 2024-04-09		Sar	nple NO.: 115024	14	Scan Date:	2024-04-08
Lab:		Sample Date: 2024-04-08			GA:	13+3
BPD:	mm	CR	RL length: 74.01	mm	NT length:	1.70 mm
Assay –						
NO.	Item abbr	Result	Unit	MOM	Refere	ence range
1 fr	ree-ß-HCG	41.80	ng/ml	1.18		
2	PAPP-A	3937.00	mIU/L	0.80		
3	NT	1.70	mm	0.93		
sk calculate —						
Age risk: 1:1512					21-3 s	yndrome risk
				50		
Paramet	ter: Trisomy21			~	Risk above cut off	
Risk: 1:14649				<u> </u>	You risk 1: >10000	
Cut Off: ( < 1:150 )						
Screaning Result: Negative				>5000		50
Screaming Res	suit. Tregutive				Age	
					18-3 s	yndrome risk
Parameter: Trisomy18/13  Risk: 1:4175057  Cut Off: ( < 1:300 )				100		
				×	Risk above cut off	
				호 200		You risk 1: >10000
Screening Res				>5000		
~ 51 5511111g 1000	1,05411,0			/3000	F	50

Advice: Diagnostic results with less risk

Parameter:

Note: \*The basic information on the basis of Down's risk assessment in this report is provided at the time of your onsite. When you get this report, please first check whether your relevant information is correct. If there is any discrepancy, please contact your doctor in time, so as to feedback us the correct information and documents, then obtain the correct report.

\*The high risk and borderline risk of trisomy 21 or trisomy 18 requires further interventional prenatal diagnosis (from fetuses

Screening Result:

Cut Off:

Doctor: Checked by :

Print date: 2024-04-09 13:14:09

<sup>\*</sup>The high risk and borderline risk of trisomy 21 or trisomy 18 requires further interventional prenatal diagnosis (from fetuses such as villus, amniotic fluid, cord blood, etc.); high risk of neural tube defect (NTD), please go to ultrasound prenatal diagnosis qualified hospitals use ultrasound to exclude.

<sup>\*</sup>The risk of NTD is only calculated at 14-22 weeks.

<sup>\*</sup>The screening result with low risk only shows that the chance of this kind of congenital abnormality in your fetus is less, and the possibility of this kind of abnormality or other abnormalities cannot be completely ruled out. Please consult a doctor in time after you get the report, and the doctor will follow your Risks and other conditions (whether you are older than 35 years old, whether you have had more than one child with other deformities, or have other diseases such as tumors) are comprehensively considered to suggest whether you need to take further examination to confirm the diagnosis.

<sup>\*\*</sup>This report only can be reference and assistant for doctor, cannot directly give conclusion by this \*\*