Name:	RAJINDER KAUR	Co	ntact:				Gender: Female	
Weight:		Birthdate: 1989-10-0 Twins: No Sender:				Age of EDC: 35.78 Year GA calc method: CRL Robinson		
_	Asian							
LMP Day:						OA can	memod. CKL Robinso	on
Sample int	formation							
Send time: 2024-12-30 Sample NO.: y0332974				974		Scan Date:	2024-12-31	
Lab: Sample Date: 202				1-01	-01 GA: 12+0			
BPD:	mm	CR	L length: 52.40	mm		NT length:	0.60 mm	
Assay								
NO.	Item abbr	Result	Unit	N	MOM	Reference range		
1	free-ß-HCG	23.70	ng/ml	(	0.38			
2	PAPP-A	10000.00	mIU/L		2.14			
3	NT	0.60	mm		).56			
k calculate .								
Age risk: 1:361					21-3 syndrome risk			
Damana	eter: Trisomy21				50			
	•			Risk	100 -	Risk above cut off		f
R	Risk: 1:53392			ä	100		You risk 1: >100	00
Cut Off: ( < 1:150 )					>5000			
Screaning Result: Negative				- 0000	Age 5	0		
						18-3 sv	yndrome risk	
Param	eter: Trisomy18/13				100		· 	
R	isk: 1:114318			<u>※</u> 200 -		Risk above cut off	<sub>4</sub>	
Cut	Off: ( < 1:300 )						You risk 1: >1000	00
Screening R	esult: Negative							

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 harderline risk of triscorny 21 or triscorny 18 requires first har interpretal discrepancy from fetures.

Age

Screening Result:

Doctor: Checked by:

Cut Off:

Print date: 2025-01-02 12:42:35

<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 \*\*