Namai	DANHIZATID					Gender: Female	
	RANI KAUR		ontact: thdate: 1998-	07.04	24		
_	53.90 Kg			07-04	_	e of EDC: 26.17 Ye	
Race: LMP Day:	Asian		Twins: No ender:		GA calc	method: CRL Robi	nson
Sample inf	formation		ender.				
Send time: 2024-02-17		San	mple NO.: Y0076	5514	Scan Date:	2024-02-13	
Lab:		Sample Date: 2024-02-16			GA:	11+4	
BPD: mm		CRL length: 44.60 mm			NT length:	3.60 mm	
Assay	]				- TVI length.		
NO.	Item abbr	Result	Unit	MOM	Refere	nce range	
1	free-ß-HCG	50.70	ng/ml	0.76			
2	PAPP-A	6431.00	mIU/L	1.77			
3	NT	3.60	mm	2.92			
k calculate .							
Age risk: 1:1309					21-3 syndrome risk		
				50 -	,		
Param	eter: Trisomy21						
Risk: 1:35				<u>×</u> 100 -	पूर्व 100 - Risk above cur You risk 1:35		
				<u> </u>			5
Cut	Off: ( < 1:150 )			>5000			
Screaning R	Lesult: Positive				Age	50	
					18-3 sy	rndrome risk	
Parameter: Trisomy18/13  Risk: 1:791  Cut Off: ( < 1:300 )				100 -			
					Risk above cut off You risk 1:791	ut off	
				호 200			
				_		1 ou risk 1:79	71
. D	esult: Negative			>5000			

Advice: Trisomy 21 High 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:

Doctor: Checked by:

Cut Off:

Print date: 2024-02-17 11:31:53

<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 \*\*