

Sample Collection date: **10062024** am/pm
 Sample Collection time: _____
 Sample Collection from: _____
 Sample Collected By: **Ketan Darvi**

Arkhana wani Kaper Khairne
Requisition form for Prenatal Screening

Patient Details:
 Name: **Mrs. Manju Mali**
 Address: **Ambe bhawan Room no. 1 Plot nos 5 Sec-12 Vashi**
 Pin Code: _____

3000) - Gapy

City: _____ State: _____
 DOB: **19061989** Hospital ID: _____
 Weight: **60** kg Height: **5.1** cm
 Ethnicity: South Asian East Asian Caucasian African Other Smoking status: Yes No

Ultrasound History
 First trimester ultrasound details
 USG date: **08062024**
 CRL: **67.8** mm NT: **1.6** mm
 NB: Present Absent
 DCDA MCDA MCMA
 Twin pregnancy ultrasound details
 USG date: _____
 Twin A: _____ mm Twin B: _____ mm
 CRL: _____ mm NT: _____ mm
 NB: Present Absent Present Absent
 BPD: _____ mm
 FL: _____ mm
 HC: _____ mm

Requesters Information:
 Name of Hospital/Collection Centre: _____
 City: _____
 Name of Ordering Physician: **Dr. Rakul Wani**
 Name of Sonographer: **Dr. Shlok S. Jogte**
 FMF code (if available): _____

Dual Plac
Important - USG report is mandatory to avoid discrepancies in the information through human error or missing out some important findings other than basic parameters required to perform the test.

Pregnancy Details:
 LMP: **15032024** USG/Corr EDD: _____
 LMP certainty: Regular Irregular Unknown
 Obstetric History: Partly (pregnancy at ≥ 24 weeks) Gravida Abortion Live
 Details of last pregnancy at ≥ 24 weeks
 PE: Yes No Date of delivery: _____
 GA at delivery: _____ Weeks _____ Days

TEST REQUESTED
First Trimester Screening (FTS) (10 weeks to 13 weeks 6 days)
 EVICO Duo : Dual Marker
 EVICO Duo Plus : Dual Marker with Macrosonia
 EVICO Duo PE : Dual Marker with Macrosonia & PE
 EVICO Duo PE+ : Dual Marker with Macrosonia, SGAI/UGR & PE (with PLGF)
 EVICO FT Enhanced PE : Enhanced FTS with SGAI/UGR, Macrosonia & PE
 Only Biochemical Values
Second Trimester Sc (15 weeks to 21 week)
 STS cannot be performed
 EVICO Quad : Quad
 EVICO Trio : Triple Marker Test
 Integrated Screeni
Pre-eclampsia Prog (After 20 weeks)
 EVICO PE-Pro (sFlt-1/PlGF Ratio)

Present pregnancy: Singleton Twin Vanishing Twin
 Type of Conception: Natural Assisted Ovulation drugs
 If assisted reproduction, kindly mention the type of procedure: _____
 Extraction date: _____ Transfer date: _____
 Egg source: Self/ donor. If donor, then donor's age/DOB: _____
 Diabetes: Yes No If Yes, Type: Gestational Type 1 Type 2
 Treatment method: No treatment / Insulin / Metformin / Insulin+Metformin / Diet Control
 If on Insulin, Insulin start date: _____
 Patient on hCG: Yes No If yes, latest date of hCG intake: _____
 Bleeding/Spotting in last two weeks: Yes No

Fill this section for Pre-eclampsia screening:
 BP measurement date: **10062024**

Markers	Left arm	Right arm	MAP
Blood pressure (mm/Hg)	Systolic BP	Diastolic BP	Systolic BP
First reading	110	70	
Second reading			

 The difference should not be more than 10 mm/Hg in first and second reading
 Family History of Pre-eclampsia: Not Known No
 Chronic Hypertension: Not known No
 Uterine Artery pulsative index (UAD-PI): Right PI: _____ Left PI: _____
 Previous small baby: Yes No
Thalassemia Screening Hb-HPLC Test Iron Therapy: Yes No
 Blood Transfusion History: Yes No

Previous pregnancy History:
 History of Down Syndrome: Yes No Edwards' Syndrome: Yes No Patau Syndrome: Yes No ONTD: Yes No
 History of Systemic Lupus Erythematosus: Yes No History of Anti Phospholipid Syndrome (APLA): Yes No
Patient Consent: I have read & understood the Requisition Form for Prenatal screening & Pre-eclampsia. I consent that my sample shall be the sole exclusive property of LILAC INSIGHTS PVT LTD & I transfer all my sample rights to LILAC INSIGHTS for its research and/or commercial use. I agree to be contacted by Lilac Insights for information regarding their tests and updates.

Important: 1) This form has to be completely filled up for us to process the sample. 2) Sample(s) accepted are subject to verification at our Laboratory. If sample found unfit for processing, the healthcare professional will be notified. 3) Kindly attach relevant copy(s) of diagnostic report(s).
 Signature of the Patient: _____
 Signature of Ordering Physician: _____
For Lilac Insights
 PAPP-A:
 β -hCG:
 AFP:
 PIGF:
 UE3:
 Inhibin A:
 sFlt-1:





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First Trimester Screening Report

MANJU MALL
 Date of birth : 19 June 1989, Examination date: 08 June 2024

Referring doctor: Dr Rahul Wani

Maternal / Pregnancy Characteristics:

Racial origin: South Asian (Indian, Pakistani, Bangladeshi).
 Parity: 0.
 Maternal weight: 60.0 kg; Height: 152.0 cm.
 Smoking in this pregnancy: no; Diabetes Mellitus: no; Chronic hypertension: no; Systemic lupus erythematosus: no; Antiphospholipid syndrome: no; Patient's mother had preeclampsia: no.
 Method of conception: Spontaneous;
 Last period: 15 March 2024

EDD by dates: 20 December 2024

First Trimester Ultrasound:

US machine: Philips Affiniti 70 G. Probe: C5-1 Pure wave. Visualisation: good.
 Gestational age: 13 weeks + 0 days from CRL
 EDD by scan: 14 December 2024

Findings
 Alive fetus visualised
 Fetal heart activity visualised
 Fetal heart rate 151 bpm
 Crown-rump length (CRL) 67.8 mm
 Nuchal translucency (NT) 1.6 mm
 Biparietal diameter (BPD) 21.7 mm
 Ductus Venosus PI 1.100
 Placenta anterior high
 Amniotic fluid normal
 Cord 3 vessels

Chromosomal markers:

Nasal bone: present; Tricuspid Doppler: normal.

Fetal anatomy:

Skull/brain: appears normal; Spine: appears normal; Heart: left sided heart; Abdominal wall: appears normal; Stomach: visible; Bladder / Kidneys: visible; Hands: both visible; Feet: both visible; Intracranial translucency ~ 2.3 mm.

Uterine artery PI: 1.60

equivalent to 1.010 MoM

Endocervical length: 3.4 mm

Risks / Counselling:

Patient counselled and consent given.

Operator: Shlok Loige, FMF Id: 166441

Condition Background risk Adjusted risk

DR SHLOK'S DIAGNOSTIC CENTRE



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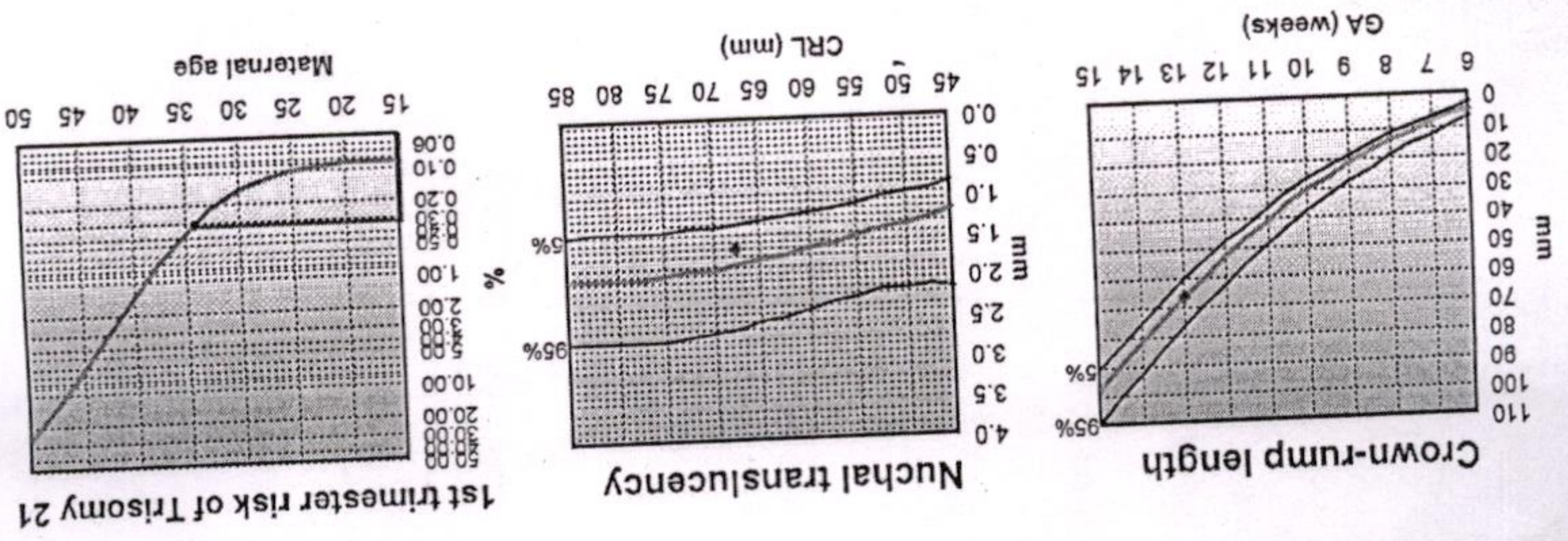
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First Trimester Screening Report

Trisomy 21	1: 280	1: 2955
Trisomy 18	1: 696	1: 3022
Trisomy 13	1: 2180	<1: 20000

The background risk for aneuploidies is based on maternal age (34 years). The adjusted risk is the risk at the time of screening, calculated on the basis of the background risk and ultrasound factors (fetal nuchal translucency thickness, nasal bone, fetal heart rate).
 All biophysical markers are corrected as necessary according to several maternal characteristics including racial origin, weight, height, smoking, method of conception and parity.

The estimated risk is calculated by the FMF-2012 software (version 2.81) and is based on findings from extensive research coordinated by the Fetal Medicine Foundation (UK Registered charity 1037116). The risk is only valid if the ultrasound scan was performed by a sonographer who has been accredited by the Fetal Medicine Foundation and has submitted results for regular audit (see www.fetalmedicine.com).



Comments - Single live intrauterine gestation corresponding to 13 weeks 0 day of gestational age

May I suggest: Detailed anomaly scan between 19 - 20 weeks.

Dr. Shlok J. Loige.
 MD (Radiology)
 Consultant Radiologist

Thanks for reference, with regards.
 (Please note: Detailed fetal anatomy may not always be visible due to technical difficulties related to embryonic size, and abdominal wall thickness. Therefore all embryonic / fetal anomalies may not necessarily be detected at every examination. All measurements including estimated fetal weight are subject to statistical variations)
 I Dr Shlok J Loige declare that while performing sonography of, I have neither detected nor disclosed the sex of fetus to anybody in any manner. Signature.....



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Name: MRS MANJU MALL

Date: 08/06/2024

RISK ASSESSMENT FOR PREECLAMPSIA

Report date 08-06-2024
Examination date 08-06-2024
Gestational age 13⁺ weeks

Obstetric history
Parity Nulliparous

Biophysical measurements
Uterine artery PI 1.6 (0.998 MoM)
Measurement date 08-06-2024

Preeclampsia risk from history only
< 37 weeks: 1 in 81

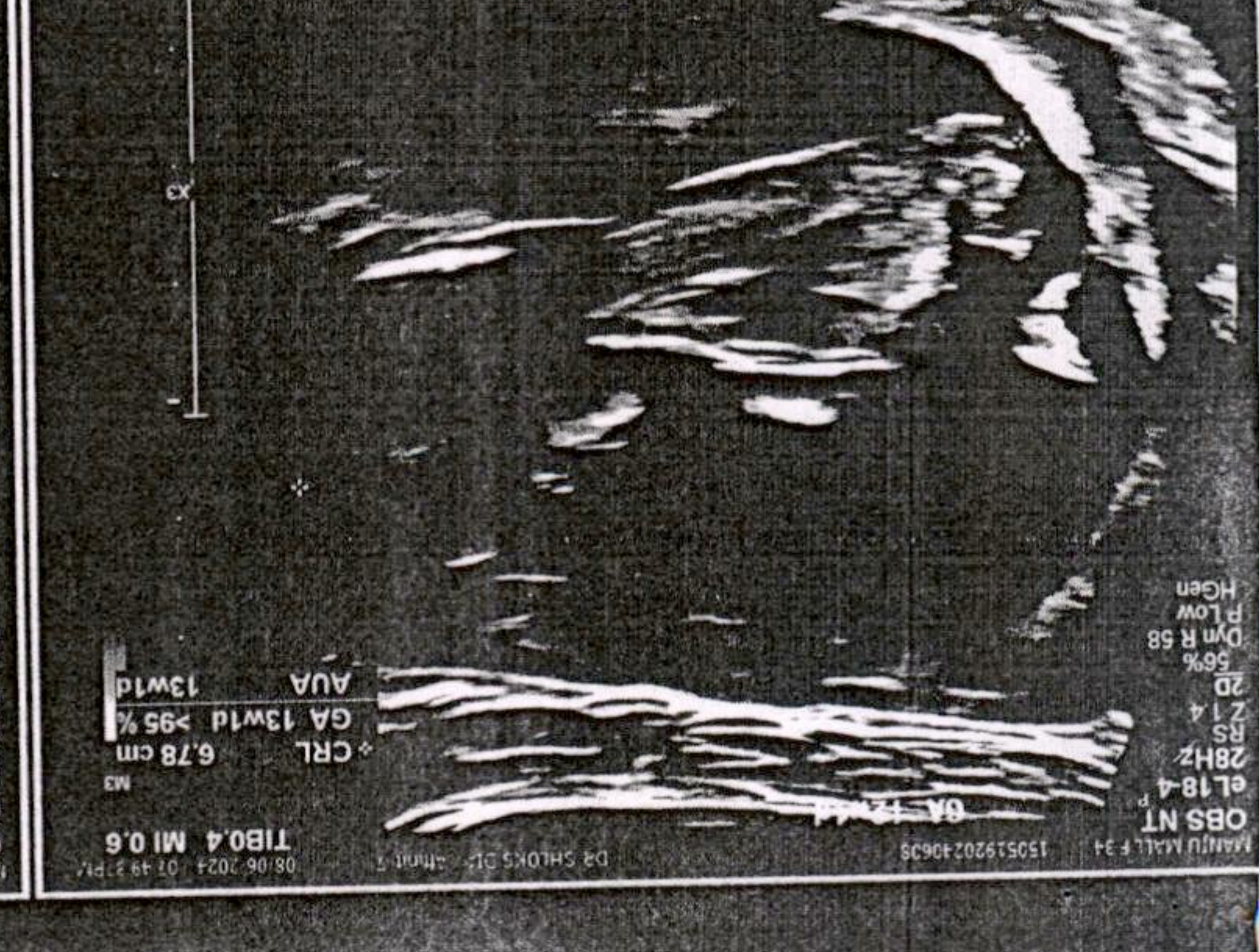
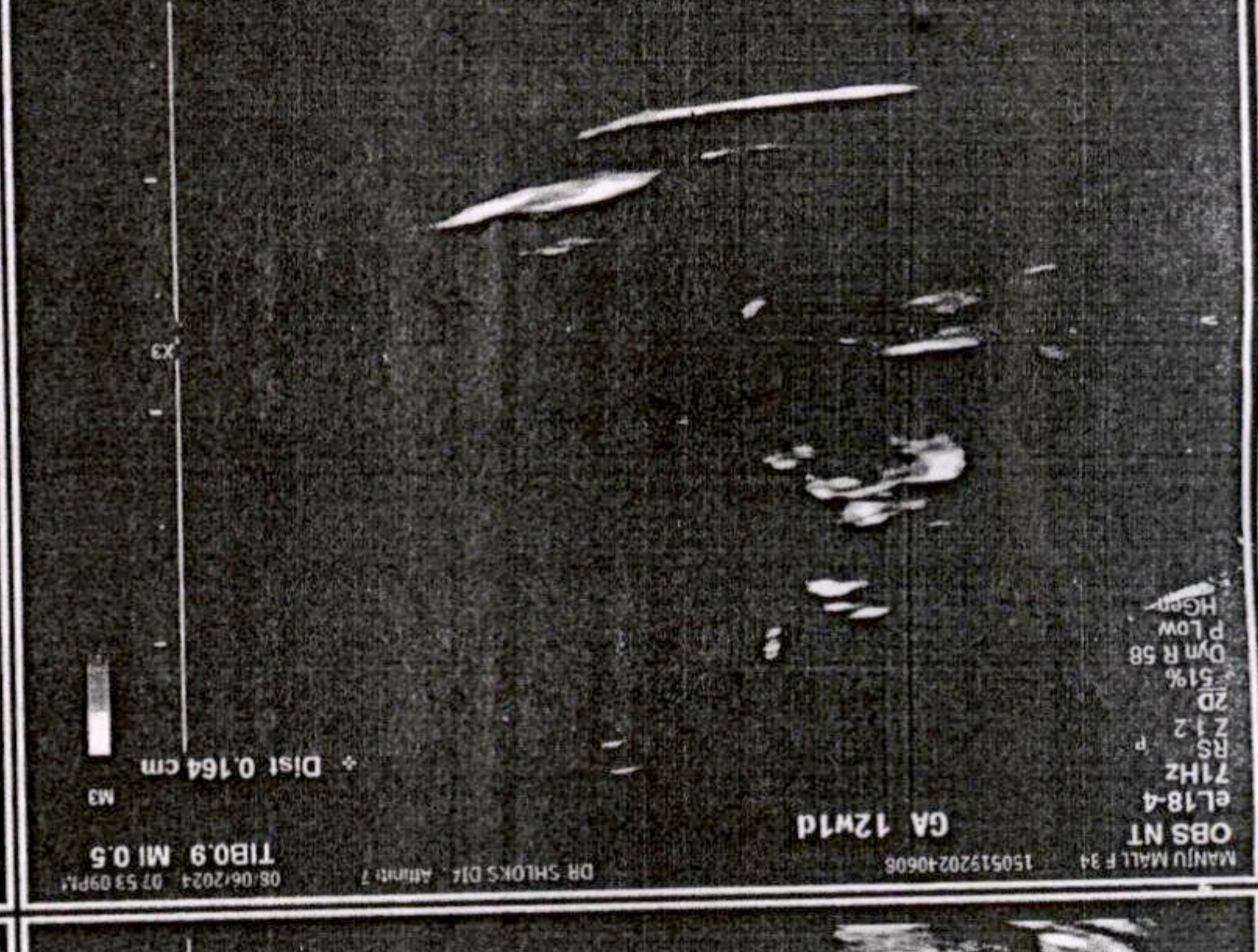
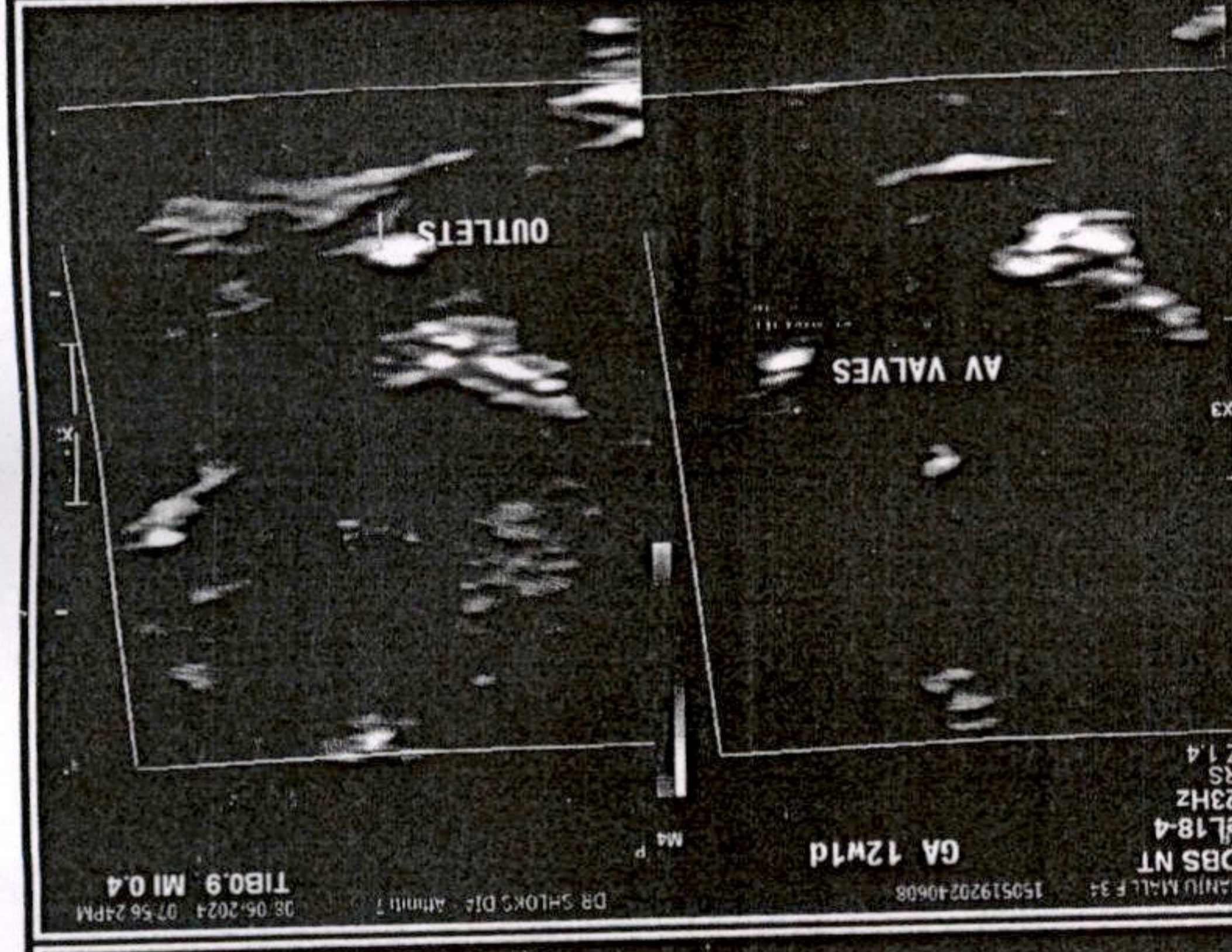
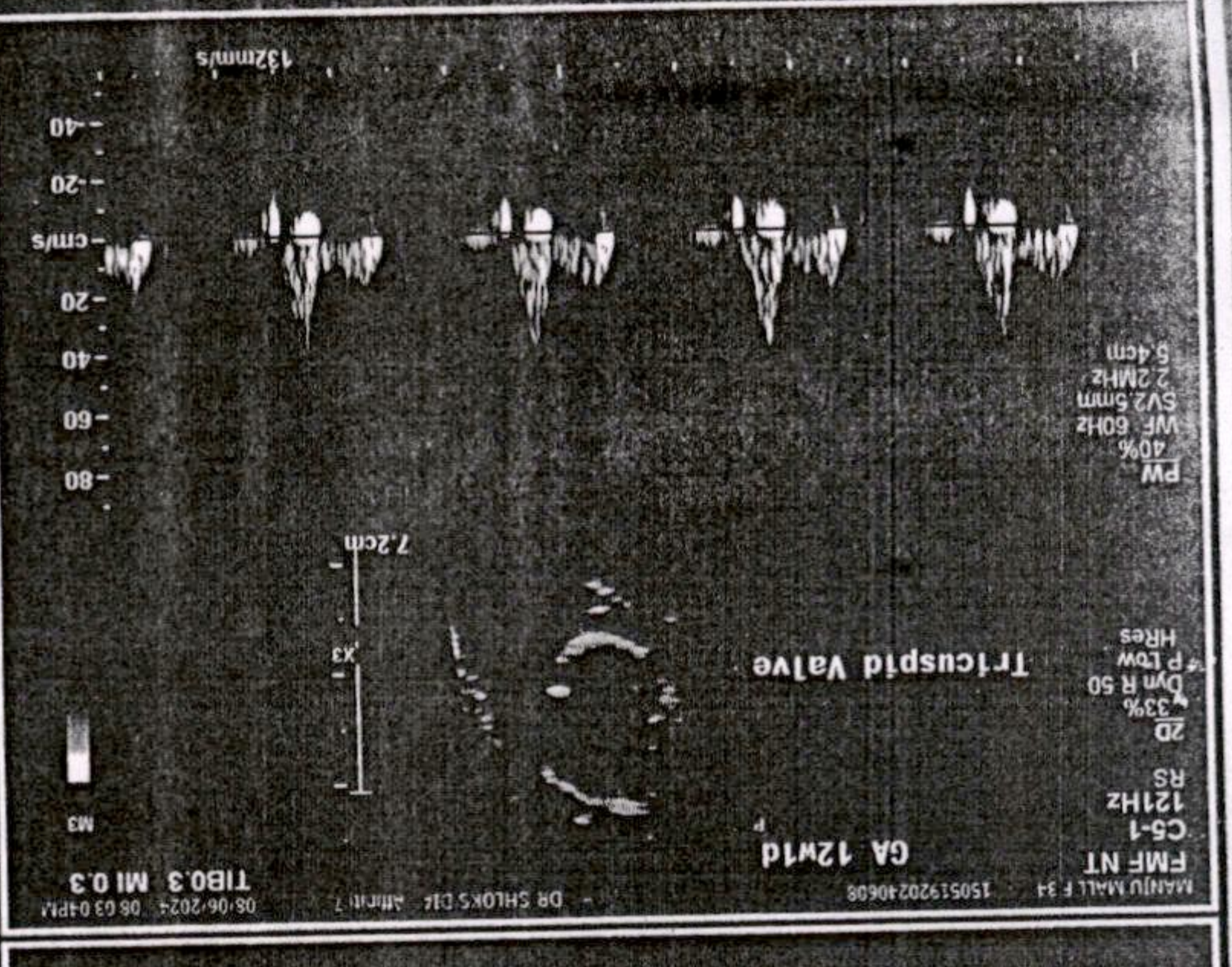
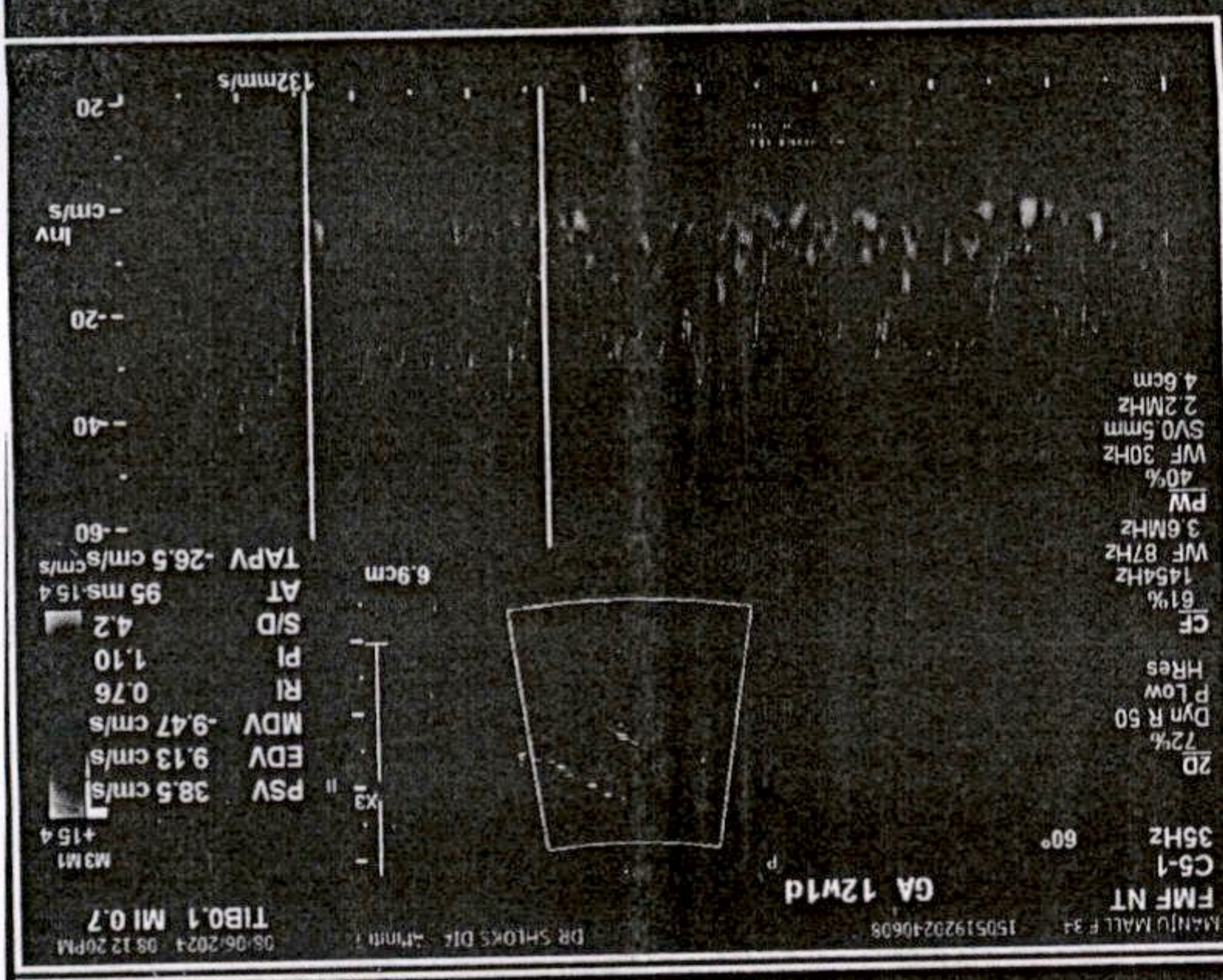
Preeclampsia risk from history plus UTP1
< 37 weeks: 1 in 106

Recommendation

The risk of preeclampsia was assessed by a combination of maternal characteristics and medical history with measurements of blood flow to the uterus.

On the basis of this assessment the patient has been classified as being at **increased risk** for developing PE before 37 weeks. The ASPRE trial has shown that in such women use of low dose aspirin (150mg/night) from now until 36 weeks reduces the incidence of PE before 32 weeks by about 90% and PE before 37 weeks by 60%.

Shlok



Dr. Shlok's Diagnostic Centre

Name : MANJU MALL F 34

08 Jun 2024 Study : Free Form

