

# ಭಾರತ ಸರ್ಕಾರ Government of India

ಆರ್ ಕುಮಾರ್ R Kumar ಜನ್ಮ ದಿನಾಂಕ / DOB : 21/01/1976 ಪುರುಷ / Male

9502 2458 1166 ನನ್ನ ಆಧಾರ್, ನನ್ನ ಗುರುತು

PID No. : MED111028055 **Register On** : 21/03/2022 10:01 AM : 922017444 SID No. Collection On : 21/03/2022 10:04 AM Age / Sex : 46 Year(s) / Male

Report On : 22/03/2022 3:37 PM

: 23/03/2022 4:21 PM **Printed On** 

Ref. Dr : MediWheel

: OP

Type



Investigation  HAEMATOLOGY	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Complete Blood Count With - ESR			
Haemoglobin (EDTA Blood/Spectrophotometry)	15.4	g/dL	13.5 - 18.0
Packed Cell Volume(PCV)/Haematocrit (EDTA Blood/Derived from Impedance)	49.1	%	42 - 52
RBC Count (EDTA Blood/Impedance Variation)	5.91	mill/cu.mm	4.7 - 6.0
Mean Corpuscular Volume(MCV) (EDTA Blood/Derived from Impedance)	83.0	fL	78 - 100
Mean Corpuscular Haemoglobin(MCH) (EDTA Blood/Derived from Impedance)	26.1	pg	27 - 32
Mean Corpuscular Haemoglobin concentration(MCHC) (EDTA Blood/Derived from Impedance)	31.3	g/dL	32 - 36
RDW-CV (EDTA Blood/Derived from Impedance)	15.1	%	11.5 - 16.0
RDW-SD (EDTA Blood/Derived from Impedance)	43.87	fL	39 - 46
Total Leukocyte Count (TC) (EDTA Blood/Impedance Variation)	5700	cells/cu.mm	4000 - 11000
Neutrophils (EDTA Blood/Impedance Variation & Flow Cytometry)	51.5	%	40 - 75
Lymphocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	37.7	%	20 - 45
Eosinophils (EDTA Blood/Impedance Variation & Flow Cytometry)	2.3	%	01 - 06



**VERIFIED BY** 



Name : Mr. KUMAR R
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Monocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	8.0	%	01 - 10
Basophils (EDTA Blood/Impedance Variation & Flow Cytometry)	0.5	%	00 - 02
Absolute Neutrophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	2.94	10^3 / μl	1.5 - 6.6
Absolute Lymphocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	2.15	10^3 / μl	1.5 - 3.5
Absolute Eosinophil Count (AEC) (EDTA Blood/Impedance Variation & Flow Cytometry)	0.13	10^3 / μl	0.04 - 0.44
Absolute Monocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.46	10^3 / μl	< 1.0
Absolute Basophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.03	10^3 / μl	< 0.2
Platelet Count (EDTA Blood/Impedance Variation)	245	10^3 / μl	150 - 450
MPV (EDTA Blood/Derived from Impedance)	8.5	fL	7.9 - 13.7
PCT (EDTA Blood/Automated Blood cell Counter)	0.21	%	0.18 - 0.28
ESR (Erythrocyte Sedimentation Rate) (EDTA Blood/Modified Westergren)	8	mm/hr	< 15



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: Mr. KUMAR R Name

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<b>BIOCHEMISTRY</b>			
Liver Function Test			
Bilirubin(Total) (Serum/Diazotized Sulfanilic Acid)	0.8	mg/dL	0.1 - 1.2
Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid)	0.2	mg/dL	0.0 - 0.3
Bilirubin(Indirect) (Serum/Derived)	0.6	mg/dL	0.1 - 1.0
Total Protein (Serum/Biuret)	7.0	gm/dL	6.0 - 8.0
Albumin (Serum/Bromocresol green)	4.22	gm/dL	3.5 - 5.2
Globulin (Serum/Derived)	2.8	gm/dL	2.3 - 3.6
A : G Ratio (Serum/Derived)	1.5		1.1 - 2.2
SGOT/AST (Aspartate Aminotransferase) (Serum/IFCC Kinetic)	22	U/L	5 - 40
SGPT/ALT (Alanine Aminotransferase) (Serum/IFCC / Kinetic)	28	U/L	5 - 41
Alkaline Phosphatase (SAP) (Serum/IFCC Kinetic)	68	U/L	53 - 128
GGT(Gamma Glutamyl Transpeptidase) (Serum/SZASZ standarised IFCC)	36.8	U/L	< 55



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<u>Investigation</u>	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
<u>Lipid Profile</u>			
Cholesterol Total (Serum/Cholesterol oxidase/Peroxidase)	155	mg/dL	Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240
Triglycerides (Serum/Glycerol phosphate oxidase / peroxidase)	152	mg/dL	Optimal: < 150 Borderline: 150 - 199 High: 200 - 499 Very High: >= 500

**INTERPRETATION:** The reference ranges are based on fasting condition. Triglyceride levels change drastically in response to food, increasing as much as 5 to 10 times the fasting levels, just a few hours after eating. Fasting triglyceride levels show considerable diurnal variation too. There is evidence recommending triglycerides estimation in non-fasting condition for evaluating the risk of heart disease and screening for metabolic syndrome, as non-fasting sample is more representative of the "usual" circulating level of triglycerides during most part of the day.

HDL Cholesterol (Serum/Immunoinhibition)	38	mg/dL	Optimal(Negative Risk Factor): >= 60 Borderline: 40 - 59 High Risk: < 40
LDL Cholesterol (Serum/Calculated)	86.6	mg/dL	Optimal: < 100 Above Optimal: 100 - 129 Borderline: 130 - 159 High: 160 - 189 Very High: >= 190
VLDL Cholesterol (Serum/Calculated)	30.4	mg/dL	< 30
Non HDL Cholesterol (Serum/Calculated)	117.0	mg/dL	Optimal: < 130 Above Optimal: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very High: >= 220

**INTERPRETATION:** 1. Non-HDL Cholesterol is now proven to be a better cardiovascular risk marker than LDL Cholesterol. 2. It is the sum of all potentially atherogenic proteins including LDL, IDL, VLDL and chylomicrons and it is the "new bad cholesterol" and is a co-primary target for cholesterol lowering therapy.



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The results pertain to sample tested.

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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Total Cholesterol/HDL Cholesterol Ratio (Serum/Calculated)	4.1		Optimal: < 3.3 Low Risk: 3.4 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 High Risk: > 11.0
Triglyceride/HDL Cholesterol Ratio (TG/HDL) (Serum/Calculated)	4		Optimal: < 2.5 Mild to moderate risk: 2.5 - 5.0 High Risk: > 5.0
LDL/HDL Cholesterol Ratio (Serum/Calculated)	2.3		Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0



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Investigation	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Glycosylated Haemoglobin (HbA1c)  HbA1C	6.6	%	Normal: 4.5 - 5.6
(Whole Blood/HPLC)	0.0	, c	Prediabetes: 5.7 - 6.4 Diabetic: >= 6.5

INTERPRETATION: If Diabetes - Good control: 6.1 - 7.0 %, Fair control: 7.1 - 8.0 %, Poor control >= 8.1 %

Estimated Average Glucose 142.72 mg/dL

(Whole Blood)

#### **INTERPRETATION: Comments**

HbA1c provides an index of Average Blood Glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.

Conditions that prolong RBC life span like Iron deficiency anemia, Vitamin B12 & Folate deficiency,

hypertriglyceridemia, hyperbilirubinemia, Drugs, Alcohol, Lead Poisoning, Asplenia can give falsely elevated HbAlC values. Conditions that shorten RBC survival like acute or chronic blood loss, hemolytic anemia, Hemoglobinopathies, Splenomegaly, Vitamin E ingestion, Pregnancy, End stage Renal disease can cause falsely low HbAlc.



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-	<u>Value</u>		Reference Interval

# **IMMUNOASSAY**

#### THYROID PROFILE / TFT

T3 (Triiodothyronine) - Total 1.35 ng/mL 0.7 - 2.04

(Serum/CMIA)

#### INTERPRETATION:

#### **Comment:**

Total T3 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T3 is recommended as it is Metabolically active.

T4 (Thyroxine) - Total 9.79 μg/dL 4.2 - 12.0

(Serum/CMIA)

#### INTERPRETATION:

#### Comment:

Total T4 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T4 is recommended as it is Metabolically active.

TSH (Thyroid Stimulating Hormone) 3.54 µIU/mL 0.35 - 5.50

(Serum/Chemiluminescent Microparticle

Immunoassay(CMIA))

#### INTERPRETATION:

Reference range for cord blood - upto 20

1 st trimester: 0.1-2.5 2 nd trimester 0.2-3.0 3 rd trimester: 0.3-3.0

(Indian Thyroid Society Guidelines)

#### **Comment:**

- 1.TSH reference range during pregnancy depends on Iodine intake, TPO status, Serum HCG concentration, race, Ethnicity and BMI.
- 2.TSH Levels are subject to circadian variation, reaching peak levels between 2-4am and at a minimum between 6-10PM. The variation can be of the order of 50%, hence time of the day has influence on the measured serum TSH concentrations.
- 3. Values&amplt 0.03 µIU/mL need to be clinically correlated due to presence of rare TSH variant in some individuals.



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	Value		Reference Interval

# **CLINICAL PATHOLOGY**

## PHYSICAL EXAMINATION

Colour (Urine) Pale yellow	
Appearance Clear Clear (Urine)	
Volume 20 mL (Urine)	
<u>CHEMICAL EXAMINATION(Automated-Urineanalyser)</u>	
pH 5.0 4.5 - 8.0 (Urine/AUTOMATED URINANALYSER)	)
Specific Gravity 1.025 1.002 -	)35
Ketones Negative Negative (Urine)	e
Urobilinogen 0.2 0.2 - 1.0 (Urine/AUTOMATED URINANALYSER)	)
Blood Negative (Urine/AUTOMATED URINANALYSER)	e

(Urine/AUTOMATED URINANALYSER)

Protein Negative (Urine)

Negative

Negative

Glucose Negative Negative

(Urine)

Bilirubin



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(Urine/AUTOMATED URINANALYSER)



Negative

Negative

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Leukocytes (Urine)  MICROSCOPY(URINE DEPOSITS)	Negative	leuco/uL	Negative
Pus Cells (Urine/Flow cytometry)	3-4	/hpf	3-5
Epithelial Cells (Urine)	3-4	/hpf	1-2
RBCs (Urine/Flow cytometry)	Nil	/hpf	2-3
Others (Urine)	Nil		Nil
Casts (Urine/Flow cytometry)	Nil	/hpf	0 - 1
Crystals (Urine)	Nil		NIL

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Investigation BIOCHEMISTRY	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
BUN / Creatinine Ratio	11		6 - 22
Glucose Fasting (FBS) (Plasma - F/GOD - POD)	86	mg/dL	Normal: < 100 Pre Diabetic: 100 - 125 Diabetic: >= 126

**INTERPRETATION:** Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level.

Glucose Postprandial (PPBS) 102 mg/dL 70 - 140

(Plasma - PP/GOD - POD)

#### INTERPRETATION:

Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level. Fasting blood glucose level may be higher than Postprandial glucose, because of physiological surge in Postprandial Insulin secretion, Insulin resistance, Exercise or Stress, Dawn Phenomenon, Somogyi Phenomenon, Anti- diabetic medication during treatment for Diabetes.

Glucose Postprandial - Urine (Urine - PP)	Negative		Negative
Blood Urea Nitrogen (BUN) (Serum/Urease-GLDH)	10	mg/dL	7.0 - 21
Creatinine (Serum/Jaffe Kinetic)	0.9	mg/dL	0.9 - 1.3

**INTERPRETATION:** Elevated Creatinine values are encountered in increased muscle mass, severe dehydration, Pre-eclampsia, increased ingestion of cooked meat, consuming Protein/ Creatine supplements, Diabetic Ketoacidosis, prolonged fasting, renal dysfunction and drugs such as cefoxitin, cefazolin, ACE inhibitors, angiotensin II receptor antagonists, N-acetylcyteine, chemotherapeutic agent such as flucytosine etc.

Uric Acid **8.0** mg/dL 3.5 - 7.2

(Serum/Uricase/Peroxidase)



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Prostate: > 10.0

<u>Investigation</u>	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
<u>IMMUNOASSAY</u>			
Prostate specific antigen - Total(PSA) (Serum/Chemiluminescent Microparticle Immunoassay(CMIA))	0.464	ng/mL	Normal: 0.0 - 4.0 Inflammatory & Non Malignant conditions of Prostate & genitourinary system: 4.01 - 10.0 Suspicious of Malignant disease of

#### INTERPRETATION: Analytical sensitivity: 0.008 - 100 ng/mL

PSA is a tumor marker for screening of prostate cancer. Increased levels of PSA are associated with prostate cancer and benign conditions like bacterial infection, inflammation of prostate gland and benign hypertrophy of prostate/ benign prostatic hyperplasia (BPH).

Transient elevation of PSA levels are seen following digital rectal examination, rigorous physical activity like bicycle riding, ejaculation within 24 hours.

PSA levels tend to increase in all men as they age.

Clinical Utility of PSA:

- •In the early detection of Prostate cancer.
- •As an aid in discriminating between Prostate cancer and Benign Prostatic disease.
- •To detect cancer recurrence or disease progression.



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**MEDALL** 

InvestigationObservedUnitBiologicalValueReference Interval

# **IMMUNOHAEMATOLOGY**

BLOOD GROUPING AND Rh TYPING 'B' 'Positive'

(EDTA Blood/Agglutination)





**APPROVED BY** 

-- End of Report --



Name	MR.KUMAR R	ID	MED111028055
Age & Gender	46Y/MALE	Visit Date	21/03/2022
Ref Doctor	MediWheel		

## **ABDOMINO-PELVIC ULTRASONOGRAPHY**

**LIVER** is normal in shape, size and has increased echopattern. No evidence of focal lesion or intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

**GALL BLADDER** show normal shape and has clear contents.

Gall bladder wall is of normal thickness. CBD is of normal calibre.

**PANCREAS** has normal shape, size and uniform echopattern. No evidence of ductal dilatation or calcification.

**SPLEEN** show normal shape, size and echopattern.

No demonstrable Para -aortic lymphadenopathy.

KIDNEYS move well with respiration and have normal shape, size and echopattern.

Cortico- medullary differentiations are well madeout.

No evidence of calculus or hydronephrosis.

The kidney measures as follows

e e	Bipolar length (cms)	Parenchymal thickness (cms)
Right Kidney	9.7	1.4
Left Kidney	10.0	1.7

URINARY BLADDER show normal shape and wall thickness.

It has clear contents. No evidence of diverticula.

**PROSTATE** shows normal shape, size (wt - 13.7gms) and echopattern.

No evidence of ascites.

Impression: Increased hepatic echopattern suggestive of fatty infiltration.

**CONSULTANT RADIOLOGISTS:** 

DR. PRAJNA SHENOY

DR. MAHESH. M. S.

DR. RADHA KRISHNA. A.

DR. HIMA BINDU.P Ms/pu

DR. H. K. ANAND





Name	KUMAR R	Customer ID	MED111028055
Age & Gender	46Y/M	Visit Date	Mar 21 2022 9:56AM
Ref Doctor	MediWheel		y y

# X - RAY CHEST PA VIEW

Bilateral lung fields appear normal.

Cardiac size is within normal limits.

Bilateral hilar regions appear normal.

Bilateral domes of diaphragm and costophrenic angles are normal.

Visualised bones and soft tissues appear normal.

Impression: Essentially normal study.

DR. H.K. ANAND

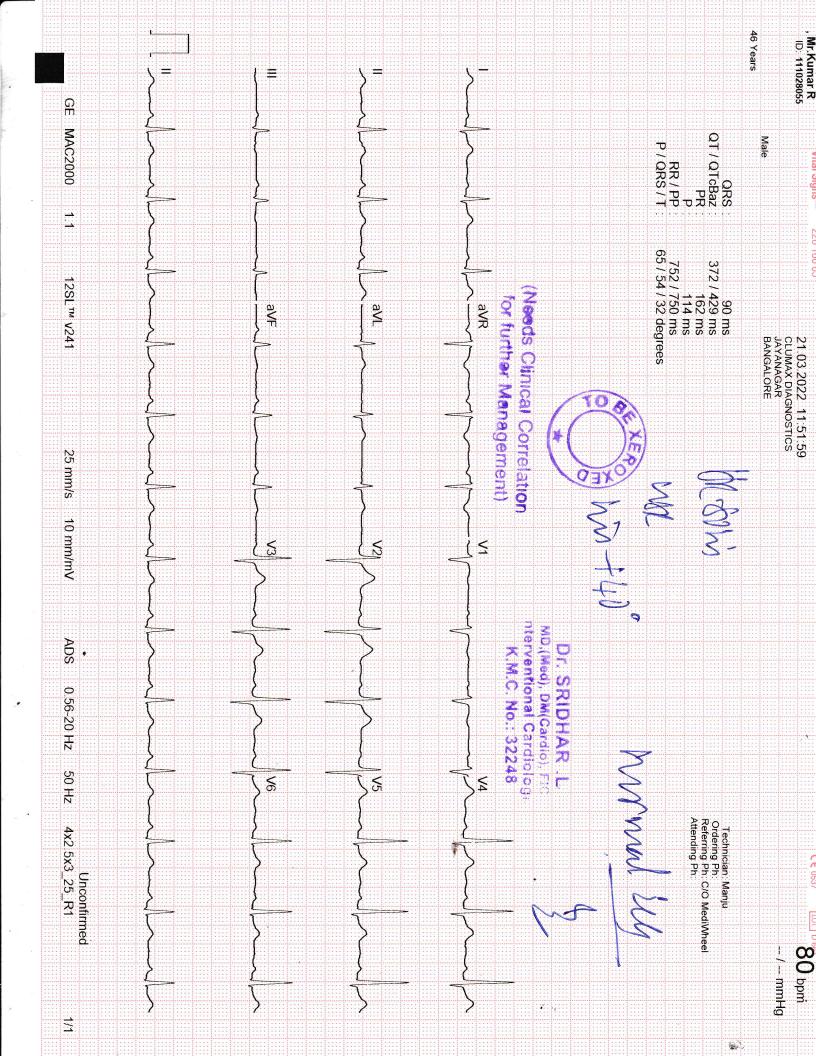
DR. POOJA B.P

DR. SHWETHA S

CONSULTANT RADIOLOGISTS

DR. PRAJNA SHENOY







# **Prabha Eye Clinic & Research Center**

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email:info@prabhaeyeclinic.com

web:www.prabhaeyeclinic.com

## **PATIENT SUMMARY**

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**Patient** 

: KUMAR R - 46/Years MALE

OP Number: KA-PEC2022/347003

Address

: CLUMAX

Phone

: +919740627559

### 21/03/2022

## **OPTOMETRIST FINDINGS (-12:18:21)**

**UNAIDED VISION DIST** 

6/6 RE 6/6 LE

**UNAIDED VISION NEAR** 

N6 RE N6 LE

**COLOR VISION** 

RE Normal LE Normal

Sleeping with Contact Lens

NO

### DOCTOR ADVICE (DR.MAHESH K - 12:30:01)

PRESENTING COMPLAINTS

came for regular eye check up

LIDS & ADNEXA

RE: N; LE: N

**PUPIL** 

RE: RRR; LE: RRR

CORNEA

RERE: CLEAR; LE: CLEAR

LENS

RERE: CLEAR; LE: CLEAR

CONJUNCTIVA

RERE: N; LE: N

IRIS

RERE: N; LE: N

ANTERIOR CHAMBER

RE:ND, QUIET; LE: ND, QUIET

OTHER RESULTS

OTHER FINDINGSRE : nil intervention review after 1 year for regular eye check up

Thanking you for giving us an opportunity to provide you eye care services.



Signature:

# Sign-up & Health Assessment Form

To be filled by Customer			
ame: Mr/Ms/Mrs X V M A R	R		
ender: OMale O Female Age	: A 6. years DOB: /		
lobile:	Pincode:		
mail:			
	To be filled by Medical Hi		
	Have you been previously diagnosed with?		,
Bar code	Diabetes (Sugar)	O Yes	O No
54, 6546	Hypertension (BP)	O Yes	O No
	Cardiovascular Disease (Heart)	O Yes	O No
	Asthma/Allergies (Dust, Pollen, Food, Animals, etc.)	O Yes	O No
Vitals	Neurological Problems (Nerve)	O Yes	O No
To be filled by Technician	Are you currently taking medications for?		
Height: cms	Diabetes (Sugar)	O Yes	O No
Waist: 37 . in.	Hypertension (BP)	O Yes	O No
Hip: 4\ in.	Cardiovascular Disease (Heart)	O Yes	O No
	Liver Disease	O Yes	O No
Weight: 8 0.8 kg	Cancer Tuboroulosis (TD)	O Yes	O No
Fat: 29.5%	Tuberculosis (TB) O Yes O No  Family History		
Vice Fet.	Is there a history of below diseases in your family?		
Visc. Fat: 15. 15 %	Diabetes (Sugar)	O Yes	O No
RM: 1 7 3 4 cal	Hypertension (BP)	O Yes	O No
BMI: 28.6 kg/m²	Cardiovascular Disease (Heart)	O Yes	O No
	Cancer	O Yes	O No
Body Age: 57 years	Lifestyle Do you exercise regularly?	O Yes	O No
Sys. BP: 3 9 mmHg	Do you consume alcohol more than 2 times a week?	O Yes	O No
Dia. BP: 9 1 mmHg	Do you smoke/chew tobacco?	O Yes	O No
	Are you vegetarian?	O Yes	O No
ts pluse.	General		
	Do you see a doctor at least once in 6 months?	O Yes	O No
	Do you undergo a health checkup every year?  How would you rate your overall Health?	O Yes	O No
	· ·	O O	O O
	Women's He	alth	
	Is there a family history of Breast Cancer?	O Yes	O No
•	Is there a family history of Endometrial (Uterus) Cancer?	O Yes	O No
	Is there a family history of Ovarian Cancer?  •  Do you have irregular periods?	O Yes	O No
	Do you have heavy bleeding during periods?	O Yes	O No
	Do you have scanty periods?	O Yes	O No
	·Have you attained Menopause?	O Yes	O No
	Do you have children?	O Yes	O No O No
	Was it a normal delivery?	O Yes	O No
	Did you have diabetes/hypertension during delivery?	O yes	O No