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CONTACT US

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040 35353535
 helpdesk@yodalifeline.in
 www.yodadiagnostics.com
 6-3-862/A, Lal Bungalow add on, Ameerpet, Hyderabad - 500016



Visit ID	: YOD637568	UHID/MR No	: YOD.0000615250
Patient Name	: Mr. SANJAY KUMAR LABANIA	Client Code	: YOD-DL-0021
Age/Gender	: 38 Y 0 M 0 D /M	Barcode No	: 10943621
DOB	:	Registration	: 24/Feb/2024 09:40AM
Ref Doctor	: SELF	Collected	: 24/Feb/2024 09:45AM
Client Name	: MEDI WHEELS	Received	: 24/Feb/2024 10:08AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 12:01PM
Hospital Name	:		

DEPARTMENT OF HAEMATOLOGY					
Test Name	est Name Result Unit Biological Ref. Range Method				

ESR (ERYTHROCYTE SEDIMENTATION RATE)						
Sample Type : WHOLE BLOOD EDTA						
ERYTHROCYTE SEDIMENTATION RATE	7	mm/1st hr	0 - 15	Capillary		
				Photometry		
COMMENTS: ESR is an acute phase reactant which indicates presence and intensity of an inflammatory process. It is never diagnostic of a specific disease. It is used to monitor the course or response to treatment of certain diseases. Extremely high levels are found in cases of malignancy, hematologic diseases, collagen disorders and renal diseases. Increased levels may indicate: Chronic renal failure (e.g., nephritis, nephrosis), malignant diseases (e.g., multiple myeloma,						

Hodgkin disease, advanced Carcinomas), bacterial infections (e.g., hepfinitis, hepfinitis), inflatination diseases (e.g., functional infections, acute pelvic inflatinatory diseases, syphilis, pneumonia), inflatinatory diseases (e.g. temporal arteritis, polymyalgia rheumatic, rheumaticid arthritis, rheumatic fever, systemic lupus erythematosus [SLE]), necrotic diseases (e.g., acute myocardial infarction, necrotic tumor, gangrene of an extremity), diseases associated with increased proteins (e.g., hyperfibrinogenemia, macroglobulinemia), and severe anemias (e.g., iron deficiency or B12 deficiency).

Falsely decreased levels may indicate: Sickle cell anemia, spherocytosis, hypofibrinogenemia, or polycythemia vera.

Verified By : S MD ISMAIL

Approved By :

A. Peart

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST





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DEPARTMENT OF HAEMATOLOGY				
Test NameResultUnitBiological Ref. RangeMethod				

BLOOD GROUP ABO & RH Typing						
Sample Type : WHOLE BLOOD EDTA						
ABO		В				
Rh Typing		POSITIVE				
Method : Hemagglutination Tube method by forward and reverse grouping						
COMMENTS:						
The test will detect common blood						

will not be detected by this method. Further investigation by a blood transfusion laboratory, will be necessary to identify such groups.

Disclaimer: There is no trackable record of previous ABO & RH test for this patient in this lab. Please correlate with previous blood group findings. Advsied cross matching before transfusion

Verified By : S MD ISMAIL



Approved By :

A. Paa -

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST





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Hospital Name	:		

DEPARTMENT OF HAEMATOLOGY				
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method

CBC(COMPLETE BLOOD COUNT)								
Sample Type : WHOLE BLOOD EDTA								
HAEMOGLOBIN (HB)	11.9	g/dl	13.0 - 17.0	Cyanide-free SLS method				
RBC COUNT(RED BLOOD CELL COUNT)	5.69	million/cmm	4.50 - 5.50	Impedance				
PCV/HAEMATOCRIT	37.7	%	40.0 - 50.0	RBC pulse height detection				
MCV	66.3	fL	83 - 101	Automated/Calculated				
МСН	20.9	pg	27 - 32	Automated/Calculated				
MCHC	31.6	g/dl	31.5 - 34.5	Automated/Calculated				
RDW - CV	17	%	11.0-16.0	Automated Calculated				
RDW - SD	39	fl	35.0-56.0	Calculated				
TOTAL LEUCOCYTE COUNT	4,910	cells/ml	4000 - 11000	Flow Cytometry				
DLC (by Flow cytometry/Microscopy)								
NEUTROPHIL	54.5	%	40 - 80	Impedance				
LYMPHOCYTE	35.6	%	20 - 40	Impedance				
EOSINOPHIL	2.4	%	01 - 06	Impedance				
MONOCYTE	6.7	%	02 - 10	Impedance				
BASOPHIL	0.8	%	0 - 1	Impedance				
PLATELET COUNT	1.75	Lakhs/cumm	1.50 - 4.10	Impedance				



Approved By :

A. Pea-th

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



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Visit ID	: YOD637568	UHID/MR No	: YOD.0000615250
Patient Name	: Mr. SANJAY KUMAR LABANIA	Client Code	: YOD-DL-0021
Age/Gender	: 38 Y 0 M 0 D /M	Barcode No	: 10943621
DOB	:	Registration	: 24/Feb/2024 09:42AM
Ref Doctor	: SELF	Collected	: 24/Feb/2024 09:45AM
Client Name	: MEDI WHEELS	Received	: 24/Feb/2024 10:19AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 11:19AM
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DEPARTMENT OF BIOCHEMISTRY
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	NT
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1 C31	

Unit

Result **Biological Ref. Range** 

#### **THYROID PROFILE (T3,T4,TSH)**

#### Sample Type : SERUM

T3	1.05	ng/ml	0.60 - 1.78	CLIA
T4	8.36	ug/dl	4.82-15.65	CLIA
TSH	1.45	ulU/mL	0.30 - 5.60	CLIA

#### INTERPRETATION:

1. Serum T3, T4 and TSH are the measurements form three components of thyroid screening panel and are useful in diagnosing various disorders of thyroid gland function.

2. Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values along with depressed TSH levels.

3. Primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels. 4. Normal T4 levels accompanied by high T3 levels are seen in patients with T3 thyrotoxicosis. Slightly elevated T3 levels may be found in pregnancy and in estrogen therapy while depressed levels may be encountered in severe illness, malnutrition, renal failure and during

therapy with drugs like propanolol and propylthiouracil. 5. Although elevated TSH levels are nearly always indicative of primary hypothyroidism, rarely they can result from TSH secreting pituitary tumors (secondary hyperthyroidism).

6. Low levels of Thyroid hormones (T3, T4 & FT3, FT4) are seen in cases of primary, secondary and tertiary hypothyroidism and sometimes in non-thyroidal illness also.

7. Increased levels are found in Grave's disease, hyperthyroidism and thyroid hormone resistance.

8. TSH levels are raised in primary hypothyroidism and are low in hyperthyroidism and secondary hypothyroidism. 9 REFERENCE RANGE

9.	HEILINGE HANGE .	
	PREGNANCY	TSH in uIU/ mL
	1st Trimester	0.60 - 3.40
	2nd Trimester	0.37 - 3.60

( References range recommended by the American Thyroid Association)

Comments:

3rd Trimester

1. During pregnancy, Free thyroid profile (FT3, FT4 & TSH) is recommended.

0.38 - 4.04

2. TSH levels are subject to circadian variation, reaches peak levels between 2-4 AM and at a minimum between 6-10 PM. The

variation of the day has influence on the measured serum TSH concentrations.

Verified By : S MD ISMAIL



Approved By :

Method





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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method	

	LIVER FUNCTION TEST(LFT)					
Sample Type : SERUM						
TOTAL BILIRUBIN	1.58	mg/dl	0.3 - 1.2	JENDRASSIK & GROFF		
CONJUGATED BILIRUBIN	0.29	mg/dl	0 - 0.2	DPD		
UNCONJUGATED BILIRUBIN	1.29	mg/dl		Calculated		
AST (S.G.O.T)	28	U/L	< 50	KINETIC WITHOUT P5P- IFCC		
ALT (S.G.P.T)	42	U/L	< 50	KINETIC WITHOUT P5P- IFCC		
ALKALINE PHOSPHATASE	106	U/L	30 - 120	IFCC-AMP BUFFER		
TOTAL PROTEINS	7.3	gm/dl	6.6 - 8.3	Biuret		
ALBUMIN	4.8	gm/dl	3.5 - 5.2	BCG		
GLOBULIN	2.5	gm/dl	2.0 - 3.5	Calculated		
A/G RATIO	1.92			Calculated		



Approved By :

S K. Deeptri Dr.S.K.DEEPTHI FFM, FDM MD BIOCHEMISTRY





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Hospital Name	:		

DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method	

LIPID PROFILE							
124	mg/dl	Refere Table	Below	Cholesterol oxidase/peroxidase			
42	mg/dl	> 40		Enzymatic/ Immunoinhibiton			
43.8	mg/dl	Refere Table	Below	Enzymatic Selective Protein			
191	mg/dl	See Tab	le	GPO			
38.2	mg/dl	< 35		Calculated			
2.95		Refere Table	Below	Calculated			
4.55	Ratio	< 2.0		Calculated			
82	mg/dl	< 130		Calculated			
TOTAL CHOLESTEROL	TRI GLYCER	I DE LDL CHOLESTEROL	NON HDI CHOLESTEF				
<200	<150	<100	<130	-			
-	-						
-							
L Ratio							
	124 42 43.8 191 38.2 2.95 4.55 82 TOTAL CHOLESTEROL	124       mg/dl         42       mg/dl         43.8       mg/dl         191       mg/dl         38.2       mg/dl         2.95	124       mg/dl       Refere Table         42       mg/dl       > 40         43.8       mg/dl       Refere Table         191       mg/dl       See Table         191       mg/dl       See Table         38.2       mg/dl $< 35$ 2.95       Refere Table         4.55       Ratio $< 2.0$ 82       mg/dl $< 130$ TOTAL CHOLESTEROL         200       <150	124       mg/dl       Refere Table Below         42       mg/dl       > 40         43.8       mg/dl       Refere Table Below         43.8       mg/dl       Refere Table Below         191       mg/dl       See Table         38.2       mg/dl       < 35			

Note:

1. Measurements in the same patient can show physiological& analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL& LDL Cholesterol

2. NLA-2014 identifies Non HDL Cholesterol(an indicator of all atherogenic lipoproteins such as LDL, VLDL, IDL, Lpa, Chylomicron remnants)along with LDL-cholesterol as co- primary target for cholesterol lowering therapy. Note that major risk factors can modify treatment goals for LDL &Non HDL.

3.Apolipoprotein B is an optional, secondary lipid target for treatment once LDL & Non HDL goals have been achieved

4. Additional testing for Apolipoprotein B, hsCRP, Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement

Verified By : S MD ISMAIL

Approved By :

SK. Deepthi Dr.S.K.DEEPTHI FFM, FDM MD BIOCHEMISTRY





Visit ID	: YOD637568	UHID/MR No	: YOD.0000615250
Patient Name	: Mr. SANJAY KUMAR LABANIA	Client Code	: YOD-DL-0021
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Client Name	: MEDI WHEELS	Received	: 24/Feb/2024 10:19AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 10:49AM
Hospital Name	:		

DEPARTMENT OF BIOCHEMISTRY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

HBA1C Sample Type : WHOLE BLOOD EDTA					
ESTIMATED AVG. GLUCOSE	100	mg/dl			

Note:

1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled .

2. Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targeting a goal of < 7.0 % may not be appropriate. 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 .

Verified By : S MD ISMAIL







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Hospital Name	:		

DEPARTMENT OF BIOCHEMISTRY					
Test NameResultUnitBiological Ref. RangeMethod					

	FBS (GLUC	OSE FASTING)				
Sample Type : FLOURIDE PLASMA						
FASTING PLASMA GLUCOSE	85	mg/dl	70 - 100	HEXOKINASE		
INTERPRETATION:						
Increased In						
Diabetes Mellitus						
<ul> <li>Stress (e.g., emotion, burns, shock</li> </ul>	, anesthesia)					
Acute pancreatitis						
<ul> <li>Chronic pancreatitis</li> </ul>						
Wernicke encephalopathy (vitamin	B1 deficiency)					
<ul> <li>Effect of drugs (e.g. corticosteroids</li> </ul>	, estrogens, alcoho	l, phenytoin, thiazio	les)			
Decreased In						
Pancreatic disorders						
<ul> <li>Extrapancreatic tumors</li> </ul>						
<ul> <li>Endocrine disorders</li> </ul>						
Malnutrition						
<ul> <li>Hypothalamic lesions</li> </ul>						
Alcoholism						
<ul> <li>Endocrine disorders</li> </ul>						



Approved By :

S K. Deeptri Dr.S.K.DEEPTHI FFM, FDM MD BIOCHEMISTRY





Visit ID	: YOD637568	UHID/MR No	: YOD.0000615250
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Ref Doctor	: SELF	Collected	: 24/Feb/2024 12:01PM
Client Name	: MEDI WHEELS	Received	: 24/Feb/2024 12:17PM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 01:06PM
Hospital Name	:		

DEPARTMENT OF BIOCHEMISTRY					
Test Name Result Unit Biological Ref. Range Meth					

PPB	PPBS (POST PRANDIAL GLUCOSE)				
Sample Type : FLOURIDE PLASMA					
POST PRANDIAL PLASMA GLUCOSE	107	mg/dl	<140	HEXOKINASE	
INTERPRETATION:					
Increased In Diabetes Mellitus Stress (e.g., emotion, burns, shock, anesthes Acute pancreatitis Chronic pancreatitis Wernicke encephalopathy (vitamin B1 deficien Effect of drugs (e.g. corticosteroids, estrogen	ncy)	ytoin, thiazides)			
<u>Decreased In</u>					
Pancreatic disorders					
<ul> <li>Extrapancreatic tumors</li> </ul>					
Endocrine disorders					
Malnutrition					
Hypothalamic lesions					
<ul> <li>Alcoholism</li> <li>Endocrine disorders</li> </ul>					
Endocrine disorders					



Approved By :

S K. Deeptri Dr.S.K.DEEPTHI FFM, FDM MD BIOCHEMISTRY





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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test NameResultUnitBiological Ref. RangeMethod					

SERUM CREATININE				
Sample Type : SERUM				
SERUM CREATININE	0.78	mg/dl	0.70 - 1.30	KINETIC-JAFFE
Increased In:				
<ul> <li>Diet: ingestion of creatinine (ro</li> <li>Impaired kidney function.</li> </ul>	ast meat), Muscle disea	se: gigantism, acro	omegaly,	
Decreased In:				
<ul> <li>Pregnancy: Normal value is 0.4 diagnostic evaluation.</li> <li>Creatinine secretion is inhibited</li> </ul>				e clinician to further









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DEPARTMENT OF BIOCHEMISTRY					
Test NameResultUnitBiological Ref. RangeMethod					

SERUM UREA						
Sample Type : SERUM						
SERUM UREA	15	mg/dL	13 - 43	Urease GLDH		
Interpretation		<u> </u>				
Determination of blood urea is the most widely used screening test for renal function. When used in conjunction with serum creatinine determinations it can aid in the differential diagnosis of the three types of azotemia: prerenal, renal and postrenal.						

Elevations in blood urea concentration are seen in inadequate renal perfusion, shock, diminished blood volume (prerenal causes), chronic nephritis, nephrosclerosis, tubular necrosis, glomerular nephritis (renal causes) and urinary tract obstruction (postrenal causes). Transient elevations may also be seen during periods of high protein intake. Unpredictable levels occur with liver diseases.

Verified By : S MD ISMAIL







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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test NameResultUnitBiological Ref. RangeMethod					

ELECTROLYTES SERUM							
Sample Type : SERUM							
SERUM SODIUM	136	mEq/L	136-145	ISE			
SERUM POTASSIUM	4.2	mEq/L	3.5 - 5.1	ISE			
SERUM CHLORIDE 104 mEq/L 98 - 107 ISE							

#### **USEFUL FOR**

Identifying a suspected imbalance in electrolytes or acid/base imbalance

CLINICAL INFORMATION

The electrolytes is ordered to identify electrolyte, fluid, or pH imbalance. Electrolyte concentrations are evaluated to assist

in investigating conditions that cause electrolyte imbalances such as dehydration, kidney disease, lung diseases, or heart

conditions. Repeat testing of the electrolyte or its components may be used to monitor the patients response to treatment of any condition that may be causing the electrolyte, fluid or pH imbalance.

Electrolyte and acid-base imbalances can often be indicative of many acute and chronic illnesses. For this reason, the

electrolyte panel is often used in the hospital and emergency settings to evaluate patients.

#### **INTERPRETATION**

With an imbalance of a single electrolyte, such as sodium or potassium, repeat testing may be ordered of that particular electrolyte, can be used to monitor the imbalance until remedied. With an acid-base imbalance, blood gases may be ordered, which will measure the oxygen, carbon dioxide, and pH levels in the arterial blood. These tests assist in evaluating the acuteness of the imbalance and monitoring the response to treatment. https://www.mayocliniclabs.com/test-catalog/overview/113632#Clinical-and-Interpretive









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Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 01:13PM
Hospital Name	:		

DEPARTMENT OF CLINICAL PATHOLOGY					
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method	





Approved By :

A. Pea-

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



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Hospital Name	:		

#### DEPARTMENT OF CLINICAL PATHOLOGY

**Test Name** 

Result

Unit

**Biological Ref. Range** 

Method

С	UE (COMPLETE U	RINE EXAMIN	NATION)	
Sample Type : SPOT URINE				
PHYSICAL EXAMINATION				
TOTAL VOLUME	20	ml		
COLOUR	Pale yellow			
APPEARANCE	Clear			
SPECIFIC GRAVITY	1.003		1.003 - 1.035	Bromothymol Blue
CHEMICAL EXAMINATION				
pH	5		4.6 - 8.0	Double Indicator
PROTEIN	Negative		NEGATIVE	Protein - error of Indicators
GLUCOSE(U)	Negative		NEGATIVE	Glucose Oxidase
UROBILINOGEN	0.1	mg/dl	< 1.0	Ehrlichs Reaction
KETONE BODIES	Negative		NEGATIVE	Nitroprasside
BILIRUBIN - TOTAL	Negative		Negative	Azocoupling Reaction
BLOOD	Negative		NEGATIVE	Tetramethylbenzidine
LEUCOCYTE	Negative		Negative	Azocoupling reaction
NITRITE	Negative		NEGATIVE	Diazotization Reaction
MICROSCOPIC EXAMINATION				·
PUS CELLS	2-3	cells/HPF	0-5	
EPITHELIAL CELLS	1-2	/hpf	0 - 15	
RBCs	Nil	Cells/HPF	Nil	
CRYSTALS	Nil	Nil	Nil	
CASTS	Nil	/HPF	Nil	
BUDDING YEAST	Nil		Nil	
BACTERIA	Nil		Nil	
OTHER	Nil			

\*\*\* End Of Report \*\*\*

Verified By : S MD ISMAIL



Approved By :

A. Pea

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST





Visit ID	: YOD637568	UHID/MR No	: YOD.0000615250
Patient Name	: Mr. SANJAY KUMAR LABANIA	Client Code	: YOD-DL-0021
Age/Gender	: 38 Y 0 M 0 D /M	Barcode No	: 10943621
DOB	:	Registration	: 24/Feb/2024 09:40AM
Ref Doctor	: SELF	Collected	: 24/Feb/2024 09:45AM
Client Name	: MEDI WHEELS	Received	: 24/Feb/2024 11:24AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 24/Feb/2024 01:13PM
Hospital Name	:		

DEPARTMENT OF CLINICAL PATHOLOGY					
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method	





Approved By :

A. Pea-

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



# **yoda** DIAGNOSTICS

DEPARTMENT OF RADIOLOGY							
Patient Name	Mr. SANJAY KUMAR LABANIA	Visit ID	YOD637568	Barcode	10943621		
Age / Gender	38/MALE	UHID	YOD.0000615250	Registration Date	24-02-2024 09:35 AM		
Ref Doctor	SELF	Client Name	MEDI WHEELS	Collection Date	24-02-2024 09:35 AM		
Hospital Name		Client Code	YOD-DL-0021	Received Date			
Sample Type		Client Add	F-701, Lado Sarai, Mehravli, New Delhi	Reported Date	24-02-2024 01:57 PM		

## **X-RAY CHEST PA VIEW**

### **FINDINGS:**

Trachea is midline.

Mediastinal outline, and cardiac silhouette are normal.

Bilateral lung fields show normal vascular pattern with no focal lesion.

Bilateral hila are normal in density.

Bilateral costo-phrenic angles and domes of diaphragms are normal.

The rib cage and visualized bones appear normal.

# **IMPRESSION:**

• No significant abnormality detected.

\*\*\* End Of Report \*\*\*

Suggested clinical correlation & follow up







Dr. G PRITHVI RANI MD, CONSULTANT RADIOLOGIST, FELLOW NEURORADIOLOGY