



BIRTH CERTIFICATE

JAVITRI HOSPITAL & TEST TUBE BABY CENTRE

Telibagh, Lucknow- 226 002 ☎ : 7800427000, 0522-3218637.

This is to Certify that Mrs. Roohem Nigam

W/o. Manish Nigam S/o. Mr. ....

Resident of 3/310 Vrindavan Sahar, State Janki Nigam, Lko.

Delivered a Female Child on 18/4/15

At 3:22 PM Birth weight 2.800 Kg. Type of delivery .....

L.S.C.S

Dr. Mrs. Rajul Tyagi  
M.B.B.S., D.G.O., M.D., FICMCH

Self Attested  
Roohem Nigam

Past Obstetrical History

Annexure 1/2

(A) Previous Pregnancy

Antenatal      During Labour      Neonatal      Puerperium

- i
- ii
- iii

(B) Abortion in Past

Week of Preg.	Year	Treatment
i 2 1/2 month G.A	NOV. 2012	Ectopic Pregnancy 2nd. trimester ↓ Mifedolone taken
ii		
iii		

Present Pregnancy

(a) High Risk Factor

(b) Immunization Schedule

i

December  
14/12/14

ii

January  
14/1/15

iii

February

- Tetanus
- Hepatitis
- Other

Hematological Investigations

	Date	Date	Date	Date
Hb% TCL, DCL, GBP			12/2	
ABO-Rh	B + ve		11.5 mm	
HIV/HCV	-ve			
HBs Ag	-ve			
VDRL	-ve			
Blood Urea				
S. Creatinin				
BI Sugar <small>F</small>				
<small>PF</small>				
Urine R <small>R</small>				
<small>M/E</small>				
GTT				
Glycosylated Hb A1C				
TSH, T4, T3	3.28 (18/11/14)		1.39 mg/l	
LFT				
TORCH				
Rh-Antibody				
OTHERS				

H1A - (N)  
Triple Test (N)

Self Attested  
Footmark.



S.No.	Date of Visit	Anaemia	Oedema	Denial & Gums	Lymph Glands	Breast Nipples	CVS	Respiratory	Ophthalmic	Height
1.	10/12/14	-	(+)	(N)	(N)	formed				39.5"
2.	11/1/15	-	+	N	N	formed		N		40"
3.	11/1/15	-	+	N	N	formed	-	N		41.5"
4.	11/2/15	-	N	N	N	formed		N		42.0"
5.	11/3/15	-	N	N	N	formed		N		42.5"
6.	11/3/15	-	(+)	(N)	(N)	formed	-	(+)		43.0"

S.No.	Date of Visit	Any Complaint	B/P	PR	WT	Fundal Height	FHS	Presentation
1.	10/12/14	Pain in abdomen (on & off)	97/85	105/min	61.0 kg	17 wks 5 days	136 bpm	
2.	11/1/15	Pain in upper abd.	99/70	106/min	64 kg	21 wks 4 days		
3.	11/2/15	Backache Ticking over abd.	112/79	109/min	66 kg	26 wks 5 days		
4.	11/3/15	Indigestion	116/82	96/min	68 kg	28 wks 27 wks 30 wks		
5.	28/03/15	Backache	132/96	80/min	68 kg	32 wks 2 days 33 wks 0 days		
6.	04/04/15	Backache	123/92	92/min	69.0 kg	31 weeks 5 days		
7.	27/4/15	Fade slow fetal heart, nooch	140/98					

Ultrasonology Findings

	Date 11/11/14	Date	Date	Date
Presentation	TRIPLET unstable lie			
FHS	TEST (+) 144 bpm			
Placental Position	TIFA			
Maturity				
Amniotic Fluid				
Gestational Age				
Fetal Weight				
IUGR				
Fetal Anomaly				
Comments If Any	Tri-amniotic Tri-zygotic & Bi-chorionic triplet pregnancy 13 weeks 3 days 13 weeks 5 days 14 weeks ± 1 day			

Self Attended  
Koonayy



**IVF RECORD SHEET**

**INDIRA IVF**  
FERTILITY & IVF CENTRE

**DEMOGRAPHICS**

Date of first Consultation : 23/02/2020

Name POONAM Age 36 Yrs. Reg. No. EDDUPY-756  
 Husband Name MANISH Age 40 Yrs. Address H/4 H3 Mumfordganj  
 City Ald. State U.P. Contact No. 7007899031  
 Height 4'11" Ft. Weight 63 Kg. BMI 28.0 BP 110/70 Pulse 72/min  
 Infertility 1°/2° 1 Married life 10yr. Duration of Infertility 5yr.

**INDICATION**

- Female
- 1) Unexplained
  - 2) Diminished Ovarian Reserve / PCOS / Endometriosis
  - 3) Tubal Factor
  - 4) Uterine Factor (Fibroids, Adenomyosis, Polyp, Endometrial Adhesions, Congenital Malformation etc.)
  - 5) RPL, Genetic
- Male
- 1) Normal
  - 2) Oligospermia, Asthenospermia, OAS, OATS
  - 3) Azoospermia
  - 4) Others \_\_\_\_\_

**MENSTRUAL HISTORY**

LMP 10/2/2020 Day of Cycle 14 days. Duration 3-4 days. Cycle length 28-30  
 Hysterosalpingogram: NO Date \_\_\_\_\_ Findings \_\_\_\_\_  
 REG/IRR  NF/SCANTY/HEAVY  PL/PP

**OBSTETRIC HISTORY**

S.No.	Mode of conception Natural / IUI / IVF	Weeks	Outcome-Abortion / Ectopic / Preterm / ND / LSCS	Any Surgical Intervention	Any Complication	Comments
1 <sup>st</sup> Pregnancy	<u>Natural</u>	<u>8wk</u>	<u>Rd ectopic.</u>			
2 <sup>nd</sup> Pregnancy	<u>IVF</u>	<u>9w.</u>	<u>Lscs.</u>			<u>Fen-syn 110</u>
3 <sup>rd</sup> Pregnancy						
4 <sup>th</sup> Pregnancy						
5 <sup>th</sup> Pregnancy						
6 <sup>th</sup> Pregnancy						

Past History mother-Dm.  
 Hospital Admission for any reason. ~~hysterectomy~~ 2015, Laparoscopy/ 2011  
 Allergies not known Ucs  
 DM / HTN / Thyroid / Asthma / Epilepsy / Skin Disease / Jaundice / MI / TIA / DVT Thyroid

**MEDICAL HISTORY (WIFE)**

S.No.	Medical Problem	Current Medications	Previous Treatment
1	<u>hypertension</u> <u>(shelley)</u>	<u>thiazide</u>	<u>foosung</u>

AKT: Yes / No yes Months 6w Date started 2011 Indication 2-P.B.

Self Attested  
Poonam

**SURGICAL HISTORY (WIFE)**

S.No.	Surgery Done	Date	Place	Details / Findings
1	Laparoscopy	2011	Kanpur	
2	Laparotomy	2011	Lucknow	LSCS - FCH - My H/O

**MEDICAL HISTORY (HUSBAND) NO**

S.No.	Medical Problem	Current Medications	Previous Treatment
/			

**SURGICAL HISTORY (HUSBAND) NO**

S.No.	Surgery Done	Date	Place	Details / Findings
/				

**PREVIOUS ART TREATMENT / OVULATION STUDY / OVULATION INDUCTION**

S.No.	Treatment--	Attempts	Drug Used	Protocol	Result	Comments
1	IVF (2015)	1			Self (H/O)	Jointly (L/O)

**BLOOD TESTS : FEMALE**

Haemoglobin..... 10.5  
 Blood Group (ABO/RH)..... B+ve  
 Random Blood Sugar (RBS)..... 120.1  
 HIV.....  
 HBsAg.....  
 VDRL.....  
 HCV.....  
 TSH..... 3.70  
 PRL..... 25.83 AMH..... 1.72  
 FSH..... LH..... E<sub>2</sub>.....  
 Karyotyping..... RPL Panel.....  
 SGOT..... 18  
 SGPT..... 18  
 Sr. CREATININE..... 0.60  
 BUN..... 16

**BLOOD TESTS : MALE**

HIV..... HBsAg.....  
 VDRL.....  
 HCV.....  
 Blood Group (ABO/RH).....

**SEMEN ANALYSIS**

Count..... 85 mil/ml Motility..... 55%  
 Morphology..... 3 Vitality..... 60

**DNA Fragmentation :**

Remark..... Spermatogenesis with karyogenesis.  
 TRUS..... FNAC.....

Self Attested  
 [Signature]





Patients Name Poonam - MAMSH Age 36 Yrs. Reg. No. EPDUPV-756  
 Address: SULTANPUR, U.P. Mobile No. 9007899031

23.2.20

WIFE

Haemoglobin..... 10.5  
 Blood Group (ABO/RH)..... B+ve  
 Random Blood Sugar (RBS)..... 120.1  
 HIV..... NEG  
 HBsAg..... NEG  
 VDRL..... NEG  
 HCV..... NEG  
 TSH..... 5.70  
 PRL..... 25.83  
 AMH..... 1.73  
 SGOT..... 18  
 SGPT..... 18  
 Sr.CREATININE..... 0.60  
 BUN..... 16  
 Rubella IgG..... IgM  
 Thalassemia screen.....  
 Pap Test.....  
 Karyotype.....

HUSBAND

HIV..... NEG  
 HBsAg..... NEG  
 VDRL..... NEG  
 HCV..... NEG  
 Blood Group (ABO/RH)..... B+ve  
 Thalassemia screen.....  
 Karyotype.....

SEMEN ANALYSIS

Count..... 85 ..... ml/ml  
 Motility..... 55 ..... %  
 Remarks..... TERATOZOOSPERMIA WITH  
 DFI..... %

PREVIOUS IVF ATTEMPTS

IVF 2015 (1) / JAYTRI ESORTAC (1) / SELF (1)

Height..... 4'11" Ft. Weight..... 63 Kg. BMI..... 28.0  
 Pulse rate..... 72 /Min. Blood Pressure..... 110/70 mmhg  
 Galactorrhea..... NO H/o T.B..... 9-12-2011 / 6 months  
 Allergy..... NOT KNOWN  
 Diabetes (Yes / No)..... NO Hypertension (Yes / No)..... NO  
 Surgical / Medical History..... LAPAROSCOPY / 2011, LAPAROTOMY / 2011 -  
HYPOTHYROIDISM SINCE 11/2  
 Obstetrics History..... 2012 / 1st / 2nd / 3rd / 4th / 5th / 6th / 7th / 8th / 9th / 10th / 11th / 12th  
 P/SV..... 2015 / IVF / 1st / 2nd / 3rd / 4th / 5th / 6th / 7th / 8th / 9th / 10th / 11th / 12th

Self Attested  
 Poonam



Date: 21-Feb-20

Name: Poonam Manish/36yrs

Day: 13 of cycle

Reg. No-EADUPV756

### TRANSVAGINAL SCAN REPORT

- Uterus is normal in size measuring 7.0cm X 4.0cm X 4.6cm is anteverted and anteflexed in nature.
- Myometrium is heterogeneous in texture with no evident focal lesion.
- Endometrium is homogeneous and measures 9.5mm in thickness and is centrally placed.
- Right ovary measures 1.7cm X 1.4cm X 1.3cm in size with evidence of few small follicles
- Left ovary not visualised.
- Left Adnexa normal with no evidence of any abnormality.
- Right adnexa shows evidence of tubular cystic structure adjacent to the ovary suggestive of hydrosalpinx.
- No Free fluid in POD.

#### Impression:

1. Adenomyotic Uterus
2. Poor Ovarian Reserve
3. ~~Left ovary not visualised~~
4. Right hydrosalpinx

Laparo Salpingect

(P): OPU/ICSI

23.2.20

PREPARED BY  
Neha

Tab Iron-1

X CHECKED BY

Tab Folinine 5mg OD x 30 days

Tab Thyronorm 12.5µg BBF / 4 weeks.

Tab C'blin (0.5) weekly x 1

Follow on D<sub>8</sub> for lap. Salpingectomy.

Self attested  
Neha





CLIENT CODE : C000067890

**CLIENT'S NAME AND ADDRESS :**

OPD - ALLAHABAD SRL LIMITED  
 INDIRA IVF HOSPITAL PVT. LTD., 41/2, MOHALLA CIVIL STATION, ELGIN ROAD,  
 NEAR HANUMAN MANDIR,  
 ALLAHABAD 211001  
 UTTAR PRADESH INDIA  
 9956666714

**SRL LIMITED**

43/1, Sardar Patel Marg, Civil Lines,  
 Allahabad, 211001  
 Uttar Pradesh, INDIA  
 Tel : 0532-2260438, 8601433777, Fax : CIN -  
 U74899PB1995PLC045956

**PATIENT NAME : POONAM EADUPV 756**

**PATIENT ID :**

ACCESSION NO : **0201TB004422** AGE : 36 Years SEX : Female

DATE OF BIRTH :

DRAWN : 21/02/2020 00:00

RECEIVED : 21/02/2020 17:54

REPORTED : 21/02/2020-19:25

REFERRING DOCTOR : DR. ANJALI SHARMA

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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**INFERTILITY PANEL - F (WITH AMH)**

**HIV 4TH GEN ASSAY (P24AG + HIV AB), SERUM**

HIV 4TH GEN ASSAY (P24AG + HIV AB) NON REACTIVE NON REACTIVE

**HEPATITIS B SURFACE ANTIGEN, SERUM**

HEPATITIS B SURFACE ANTIGEN NON REACTIVE NON REACTIVE  
 PATIENT VALUE 0.51 Ref. ranges for Electrochemiluminescence  
 < 0.90 (Non Reactive)  
 > or = 1.00 (Reactive) IU/mL

**VDRL, SERUM**

VDRL NONREACTIVE NONREACTIVE TITER

METHOD : NON TREPONEMAL FLOCCULATION TEST

**HEPATITIS C ANTIBODIES, SERUM**

HEPATITIS C ANTIBODIES NON REACTIVE NON REACTIVE  
 PATIENT VALUE 0.06 Ref. ranges for Electrochemiluminescence  
 < 0.90 (Non Reactive)  
 > or = 1.00 (Reactive) IU/mL

**TSH 3RD GENERATION ULTRA( TSH3 - UL), SERUM**

TSH 3RD GENERATION 3.700 0.27 - 4.20 µIU/mL

**PROLACTIN, SERUM**

PROLACTIN 25.83 High 4.79 - 23.3 ng/mL

**GLUCOSE RANDOM, PLASMA**

GLUCOSE RANDOM, PLASMA 120.1 Non-Diabetic: < 200  
 Diabetic: > or = 200  
 "In individuals with symptoms of hyperglycemia or hyperglycemic crisis." mg/dL

**ABO GROUP & RH TYPE, EDTA WHOLE BLOOD**

ABO GROUP TYPE B  
 RH TYPE POSITIVE

**ANTI MULLERIAN HORMONE**

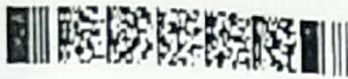
ANTI MULLERIAN HORMONE 1.73 0.777 - 5.240 ng/ml

**BLOOD COUNTS**

HEMOGLOBIN 10.5 Low 12.0 - 15.0 g/dL  
 METHOD : SPECTROPHOTOMETRY AUTOMATED HEMATOLOGY ANALYSER  
 RED BLOOD CELL COUNT 3.76 Low 3.8 - 4.8 mil/µL  
 WHITE BLOOD CELL COUNT 4.0 - 10.0 thou/µL

*Self-attested  
 Poonam*

**AGNOSTIC REPORT**  
**DIAGNOSTIC REPORT**



Annexure no 2/7  
SRL  
Diagnostics  
SRL  
Diagnostics

CLIENT CODE: C000067890

CLIENT'S NAME AND ADDRESS:

OPD - ALI AHABAD SRL LIMITED  
INDRA JYU HOSPITAL PVT. LTD., 41/2, MOHALLA CIVIL STATION, ELGIN  
ROAD,  
NEAR HANUMAN MANDIR,  
ALAHABAD 211001  
UTTAR PRADESH INDIA  
955666714

SRL LIMITED  
43/1, Sardar Patel Marg, Civil Lines,  
Allahabad, 211001  
Uttar Pradesh, INDIA  
Tel - 0532-2260438, 8601433777, Fax : CIN -  
U74899PB1995PLC045956

PATIENT NAME: POONAM EADUPV 756

PATIENT ID :

ACCESSION NO: 0201TB004422

AGE: 36 Years

SEX: Female

DATE OF BIRTH :

ORDERN 21/02/2020 00:00

RECEIVED: 21/02/2020 17:54

REPORTED: 21/02/2020 19:25

REFERRING DOCTOR: DR. ANJALI SHARMA

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
PLATELET COUNT		150	150 - 410	thou/ $\mu$ L
RBC AND PLATELET INDICES				
HEMATOCRIT		33.2	Low 36 - 46	%
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
MEAN CORPUSCULAR VOL		88.0	83 - 101	fL
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
MEAN CORPUSCULAR HGB.		28.0	27.0 - 32.0	pg
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION		31.8	31.5 - 34.5	g/dL
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
RED CELL DISTRIBUTION WIDTH		16.9	High 11.6 - 14.0	%
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
MEAN PLATELET VOLUME		11.3	High 6.8 - 10.9	fL
METHOD: CALCULATED (HEMATOLOGY ANALYSER)				
WBC DIFFERENTIAL COUNT				
SEGMENTED NEUTROPHILS		67	40 - 80	%
ABSOLUTE NEUTROPHIL COUNT		4.76	2.0 - 7.0	thou/ $\mu$ L
EOSINOPHILS		04	1 - 6	%
ABSOLUTE EOSINOPHIL COUNT		0.28	0.02 - 0.50	thou/ $\mu$ L
LYMPHOCYTES		26	20 - 40	%
ABSOLUTE LYMPHOCYTE COUNT		1.85	1.0 - 3.0	thou/ $\mu$ L
MONOCYTES		03	2 - 10	%
ABSOLUTE MONOCYTE COUNT		0.21	0.2 - 1.0	thou/ $\mu$ L
BASOPHILS		00	< 1 - 2	%
ABSOLUTE BASOPHIL COUNT		0	Low 0.02 - 0.10	thou/ $\mu$ L
DIFFERENTIAL COUNT PERFORMED ON:		EDTA SMEAR		
METHOD: AUTOMATED ANALYZER / MICROSCOPY				
DISCLAIMER: THE ABSOLUTE WHITE CELL COUNTS ARE OUTSIDE THE NABL ACCREDITED SCOPE OF THE LABORATORY.				
ASPARTATE AMINOTRANSFERASE, SERUM				U/L
ASPARTATE AMINOTRANSFERASE (AST/SGOT)		18	0 - 32	
ALANINE AMINOTRANSFERASE, SERUM				U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)		18	0 - 33	U/L
SERUM BLOOD UREA NITROGEN				mg/dL
BLOOD UREA NITROGEN		16	6 - 20	
CREATININE, SERUM				mg/dL
CREATININE		0.60	0.50 - 0.90	

Self Affected  
Smt Poonam



**DIAGNOSTIC REPORT**

**DIAGNOSTIC REPORT**

Annexure no. 278



CLIENT CODE : C000067890

**CLIENT'S NAME AND ADDRESS :**

ALLAHABAD SRI LIMITED  
 ANJALI HOSPITAL PVT. LTD., 41/2, MOHALLA CIVIL STATION, ELGIN

ANJUMAN MANDIR,  
 ALLAHABAD 211001  
 UTTAR PRADESH INDIA  
 996666714

SRL LIMITED  
 43/1, Sardar Patel Marg, Civil Lines,  
 Allahabad, 211001  
 Uttar Pradesh, INDIA  
 Tel : 0532-2260438, 8601433777, Fax : CIN -  
 U74899PB1995PLC045956

**PATIENT NAME : POONAM EADUPV 756**

PATIENT ID :

ADMISSION NO : 0201TB004422

AGE : 36 Years

SEX : Female

DATE OF BIRTH :

DRAWN : 21/02/2020 00:00

RECEIVED : 21/02/2020 17:54

REPORTED : 21/02/2020 19:25

REFERRING DOCTOR : DR. ANJALI SHARMA

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
LIQUID-BASED CYTOLOGY		RESULT PENDING		
LETTER		RESULT PENDING		
HEMOGLOBIN VARIANT ANALYSIS, BLOOD		RESULT PENDING		
RUBELLA IGG & IGM, SERUM		RESULT PENDING		
RUBELLA IGG AVIDITY, SERUM		RESULT PENDING		

**Interpretation(s)**

HIV 1/2 ELISA ASSAY (P24AG + HIV AB), SERUM-Acquired Immunodeficiency syndrome (AIDS) is caused by 2 types of human immunodeficiency viruses, collectively designated HIV. HIV is transmitted by sexual contact, exposure to blood or blood products, and prenatal infection of a fetus or perinatal infection of a newborn.

Immunologic analysis classifies HIV-1 into groups M (major), N (non-M, non-O), and O (outlier). HIV-2 is similar to HIV-1 in its structural morphology, genomic organization, in vitro cytopathogenicity, transmission routes, and ability to cause AIDS. However, HIV-2 is less pathogenic than HIV-1. HIV-2 infections have a longer latency period, slower progression to disease, lower viral titers, and lower rates of vertical and horizontal transmission. HIV-2 is endemic to West Africa but HIV-2 infections, at a lower frequency compared to HIV-1, have been identified in the USA, Europe, Asia, and other regions of Africa. India predominantly has HIV-1M subtype C.

**Use Utility**

The test is used as an aid in the diagnosis of HIV-1/HIV-2 infection.

If HIV reactive result is obtained, confirmation of HIV antibody status is done using 2 more antibody tests (as per NACO guidelines-Strategy III algorithm). If indicated HIV serostatus may be confirmed by repeating antibody test on fresh specimen or HIV-1 Western Blot (Immunoblot) Assay (SRL test code #3012).

**Limitations:**

Antibody tests may give false negative during the window period, an interval of 3 weeks to 6 months between the time of HIV infection and the production of measurable antibodies to HIV seroconversion. Most people develop detectable antibodies approximately 30 days after infection, although some seroconvert later. The vast majority of people (97%) have detectable antibodies by three months after HIV infection. A 6-month window is extremely rare with modern antibody testing. Early antiretroviral therapy during the window period may alter antibody responses. This does not apply to individuals undergoing treatment with post-exposure prophylaxis.

Antibody tests may yield false negative results in patients with X-linked agammaglobulinemia.

A positive HIV result in an infant <18 months of age may not reflect the infant's HIV infection status. HIV antibodies persist in the sera of infants up to 18 months of age, due to transplacentally acquired maternal antibodies. HIV PCR testing is recommended in this age group for diagnosis.

HEPATITIS B SURFACE ANTIGEN, SERUM-Hepatitis B is caused by infection with HBV, an enveloped DNA agent that is classified as hepadnavirus. This test detects the presence of surface antigen (HbsAg) in serum sample and is indicative of an active HBV infection, either acute or chronic.

**Use Utility**

HbsAg is the first serologic marker appearing in the serum 6-16 weeks following hepatitis B viral infection. In typical HBV infection, HbsAg will be detected 2-4 weeks before the serum enzyme levels (ALT) become abnormal and 3-5 weeks before patient develops jaundice. In acute cases HbsAg usually disappears 1-2 months after the onset of symptoms. Persistence of HbsAg for more than 6 months indicates development of either a chronic carrier state or chronic liver disease. The presence of HbsAg is frequently associated with infectivity. HbsAg when accompanied by Hepatitis Be antigen and/or hepatitis B viral DNA almost always indicates infectivity.

**Limitations:**

For diagnostic purposes, results should be used in conjunction with patient history and other hepatitis markers for diagnosis of acute or chronic infection. If the antibody results are inconsistent with clinical evidence, additional testing is suggested to confirm the result.

HbsAg detection will only indicate the presence of surface antigens in the serum and should not be used as the sole criteria for diagnosis, staging or monitoring of HBV infection. This test may be negative during "window period" i.e. after disappearance of anti-HBs.

The current assay being a highly sensitive test, may yield a small percentage of false positive reports. Hence all HbsAg positive specimens should be confirmed with an assay based upon Neutralisation of Human anti Hepatitis B Surface antibody.

**THE SERUM:**

VDRL (RPR/TRUST) are Non-Treponemal screening tests for syphilis. They achieve a sensitivity of 100% in the detection of secondary syphilis. After the secondary stage, titres drop and about a third of patients with late syphilis are seronegative. The titres may rise in patients developing cardiovascular, neurological or gummatous lesions. The test usually becomes negative 6 to 18 months after effective treatment of syphilis, depending on the stage at which treatment is initiated. If treatment is started late, tests may remain positive in low titres.

**Limitations:**

Non-treponemal tests lack sensitivity in primary and late syphilis. False negative reactions can occur in stages of the disease where there is minimal tissue damage, particularly in early infection and in latent stages.

Biologic false positive reactions are common in a variety of other infections (Leprosy, Malaria, Relapsing fever, Infectious mononucleosis, hepatitis), Rheumatic diseases and Auto-Immune disorders. More specific Treponemal tests, such as Treponema pallidum Hemagglutination assay (TPHA) test are recommended for confirmation.

HEPATITIS C ANTIBODIES, SERUM-Hepatitis C Virus (HCV) is a blood borne flavivirus. It is one of the most important causes of post-blood transfusion as well as community acquired non-A non-B hepatitis and chronic liver failure. Although the majority of infected individuals may be asymptomatic, HCV infection may develop into chronic hepatitis and/or increased risk of hepatocellular carcinoma.

**Notes & Limitations:**

HCV antibody is typically not detected until approximately 14 weeks after infection (or 5 weeks after appearance of the first biochemical marker of illness) and is almost

Self Attested  
 Page 3 Of 6



**DIAGNOSTIC REPORT**  
**DIAGNOSTIC REPORT**

*Anneure*  
*no. 2/19*  
**SRL**  
**DIAGNOSTICS**  
**SRL**  
Diagnostics



CLIENT CODE : C000067890

CLIENT'S NAME AND ADDRESS :  
OPD - ALLAHABAD SRL LIMITED  
INDIRA IVF HOSPITAL PVT. LTD., 41/2, MOHALLA CIVIL STATION, ELGIN  
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PATIENT NAME : POONAM EADUPV 756

PATIENT ID :

ACCESSION NO : D201TB004422 AGE : 36 Years SEX : Female

DATE OF BIRTH :

DRAWN : 21/02/2020 00:00

RECEIVED : 21/02/2020 17:54

REPORTED : 21/02/2020 19:25

REFERRING DOCTOR : DR. ANJALI SHARMA

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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always detectable by the late convalescent stage of infection. - A negative result may also be observed due to loss of HCV antigen, years following resolution of infection. Infants born to hepatitis C infected mothers may have delayed seroconversion to anti-HCV. Hence a negative result should be evaluated cautiously with respect to clinical findings. It is to be noted that absence of HCV antibodies after 14 weeks of exposure is strong evidence against HCV infection.  
- Presence of HCV antibodies does not imply an active Hepatitis C infection but is indicative of both past and/or recent infection. It has been reported that as many as 90% of individuals receiving intravenous commercial immunoglobulin test falsely positive for HCV antibody. Also, patients with autoimmune liver disease may show a false positive HCV antibody result. Hence it is advisable to confirm a positive antibody result with a supplemental test. A positive result when followed by a positive supplemental test (i.e. HCV-RNA-PCR) suggests active hepatitis C infection.  
TSH 3RD GENERATION ULTRA (TSH3 - UL), SERUM-Comment: The Biological Reference Interval of TSH-3rd Generation Ultra (TSH3-UL) is not established for age less than 2 years.

Below mentioned are the guidelines for Pregnancy related reference ranges for TSH.

Levels in Pregnancy	TSH (µIU/mL)
1st Trimester	0.1 - 2.5
2nd Trimester	0.2 - 3.0
3rd Trimester	0.3 - 3.0

PROLACTIN, SERUM-Reference Ranges of Prolactin for Pregnant and Post-Menopausal Females:

PREGNANT:  
9.7 - 208.5 ng/mL  
POSTMENOPAUSAL:  
1.8 - 20.3 ng/mL

GLUCOSE RANDOM, PLASMA-GLUCOSE RANDOM, PLASMA

As per ADA Guidelines 2012, Diabetic- Random plasma glucose = 200 mg/dL in a patient with classic symptoms of hyperglycemia or hyperglycemic crisis.  
ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A, B, O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.  
ANTI MULLERIAN HORMONE-

Anti Mullerian hormone (AMH) or Mullerian inhibiting substances (MIS) is a glycoprotein dimer composed of two 72 kDa monomers linked by disulfide bonds. AMH belongs to the transforming growth factor β (TGF - β) superfamily. AMH is a hormone marker for quantitative prediction of ovarian reserve, ovarian aging, ovarian dysfunction and ovarian responsiveness. The levels of AMH decrease in pre-menopausal women as the quality and number of ovarian follicles decline with age.

Clinical Utility:

- Evaluating Fertility Potential - Serum AMH levels correlate with the number of early antral follicles with greater specificity than Inhibin B, Oestradiol, Follicle Stimulating Hormone and Luteinizing Hormone on cycle day 3. Thus, Day 3 AMH may reflect ovarian follicular status better than these hormone markers.
- Measuring Ovarian Aging - Diminished ovarian reserve, associated with poor response to IVF, is signaled by reduced baseline serum AMH concentrations. AMH would appear to be a useful marker for predicting ovarian aging and the potential for successful IVF.
- Predicting Onset of Menopause - The duration of the menopausal transition can vary significantly in individuals and reproductive capacity may be seriously compromised prior to clinical diagnosis. AMH can predict the occurrence of the menopausal transition.
- Assessing Polycystic Ovary Syndrome - Serum AMH levels are elevated in patients with polycystic ovary syndrome and may be useful as a marker for the extent of the disease.

Interpretation:

AMH levels do not change significantly throughout the menstrual cycle and decrease with age. Healthy women, below 38 years old, with normal follicular status at day 3 of the menstrual cycle, have AMH levels of 2.0 - 6.8 ng/ml (14.28 - 48.55 pM).

Ovarian Fertility Potential	pmol/L	ng/mL
Optimal Fertility	28.6 - 48.5	4.0 - 6.8
Satisfactory Fertility	15.7 - 28.6	2.2 - 4.0
Low Fertility	2.2 - 15.7	0.3 - 2.2
Very Low / undetectable	0.0 - 2.2	0.0 - 0.3
High Level	> 48.5	> 6.8

*Self Attested*  
*Poonam*

The interpretation guide provided above are only suggestions which are based upon examination of multiple published studies. It is expected in the near future that refinement of these ranges may occur.

References:

1. Durlinger ALL, Visser JA, Themmen APN. Regulation of ovarian function: the role of anti-Müllerian hormone. Reproduction 2002 124:601-609.



CLIENT CODE : CD00067890

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**PATIENT NAME :** POONAM EADUPV 756

PATIENT ID :

ACCESSION NO : **0201TB004422** AGE : 36 Years SEX : Female DATE OF BIRTH :

DRAWN : 21/02/2020 00:00 RECEIVED : 21/02/2020 17:54 REPORTED : 21/02/2020 19:25

REFERRING DOCTOR : DR. ANJALI SHARMA

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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2. Ficioglu C, Kutlu T, Baglam E, Bakacak Z. Early follicular antimüllerian hormone as an indicator of ovarian reserve. Fertility and Sterility 2006 85:592-6.  
 3. Human Reproduction 2007 22(9):2414-2421 doi:10.1093/humrep/dem204.  
 4. Fertil Steril. 2005 83(4):979-87 (ISSN: 1556-5653)

**BLOOD COUNTS**-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

**RBC AND PLATELET INDICES**-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

**ASPARTATE AMINOTRANSFERASE, SERUM**-Aminotransferase (AST) is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile ducts, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous exercise.

**ALANINE AMINOTRANSFERASE, SERUM**-Alanine aminotransferase (ALT) test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine the cause of liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis.

**SERUM BLOOD UREA NITROGEN**-Causes of Increased levels

Pre renal

- High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal
- Renal Failure
- Post Renal
- Malignancy, Nephrolithiasis, Prostatism

Causes of decreased levels

- Liver disease
- SIADH.

**CREATININE, SERUM**-Higher than normal level may be due to:

- Blockage in the urinary tract
- Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
- Loss of body fluid (dehydration)
- Muscle problems, such as breakdown of muscle fibers
- Problems during pregnancy, such as seizures (eclampsia), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

- Myasthenia Gravis
- Muscular dystrophy

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

Please visit [www.srlworld.com](http://www.srlworld.com) for related Test Information for this accession.

*Akanksha*

Dr. Akanksha Singh  
 M. D. (PATH)

*Self affected  
 Poonam*