

# **LOOKING BEYOND THE USUAL**

By -Dr.Rishikesh Chauhan  
Department of General Medicine.

# Case 1

## PRESENTING COMPLAINTS

76 yrs old male farmer by occupation, known case of Type II diabetes mellitus since 5 years on oral medications was

### **Admitted with complaints of**

- Weight Loss of 5 kg in last 2 months,
- Diffuse abdominal pain since 15 days,
- High grade continuous fever with chills since 7 days,
- Burning micturition with increased frequency since 5 days.
- No history of nausea ,vomiting ,loss of appetite,diarrhoea
- No significant past history

## GENERAL EXAMINATION

Patient was conscious, oriented

- Pulse -98 /min regular,
- BP - 120/80 mmHg in Rt Arm supine position
- Febrile –T **101.2 F**
- RR- 16/min
- Spo2- 98% on RA
- No cyanosis, clubbing, icterus, edema,
- No lymphadenopathy

## SYSTEMIC EXAMINATION

On Per abdomen-

-Diffuse Tenderness was present on palpation, no rebound tenderness, no free fluid, no hepatosplenomegaly was present.

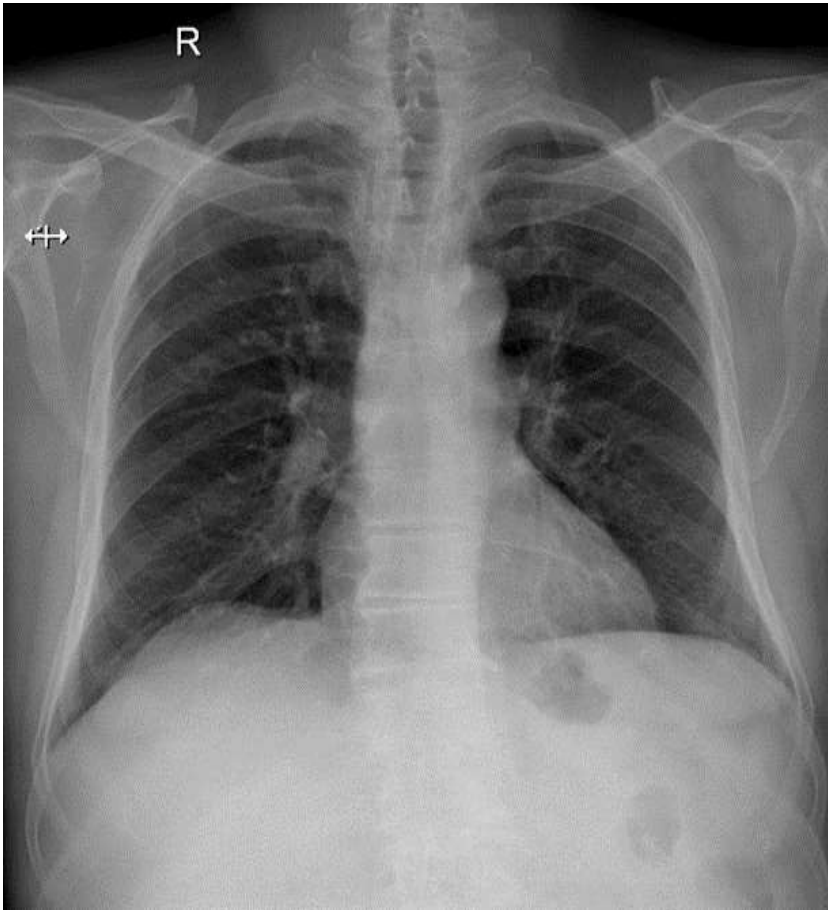
-No abdominal bruit present

- CVS , RS and CNS examination was normal.

# INVESTIGATIONS

CBC		RFT		Urine R/Ms	
Hb	10 gm%	Urea	28	PUS CELL	80-90
TLC	18000 N-78	Creat	1.06	Glucose	++
PLT	310000	<b>S.Electrolyte</b>		D-dimer	7000
PCV	28	Na/K/Cl	137/4.5/102	Procalcitonin	0.9
MCV	64	ESR	35	Lactate	20
<b>LFT</b>		CRP	15	TFT	Normal
Total bilirubin	1.15	HbA1C-	8.9	Mantoux test	Negative
Direct	0.48	UPCR-	1.01	Urine culture and blood culture was sent.	
Indirect	0.67				
SGPT	34				
SGOT	60				
ALP	118				

# INVESTIGATIONS



Chest X-ray- Normal,

- ECG-Normal sinus rhythm;
- Echocardiography- EF- 60%; NO RWMA;

# INVESTIGATIONS

## ➤ **USG ABDOMEN PELVIS**

- Left renal non obstructive calculi. (17mm in size)
- B/L renal minimal simple exophytic cyst
- Few air foci in lower pole calyces-  
likely s/o **Emphysematous pyelonephritis.**

## ➤ **Urine Culture & sensitivity-**

s/o **E.coli** sensitive to

- 1) Piperacillin tazobactam ,
- 2) Trimethoprim sulphomethoxazole

Resistant to nitrofurantoin and ciprofloxacin.

# TREATMENT

- Patient was started on Inj. piperacillin tazobactam 3.375gm 6 hourly for 14 days .
- **Patient became afebrile and was symptomatically better ,but was constantly complaining of deep seated abdominal pain.**
- So to investigate the constant abdominal pain,**CECT (abdo+pelvis)** was done.

# CECT ABDOMEN

## PELVIS

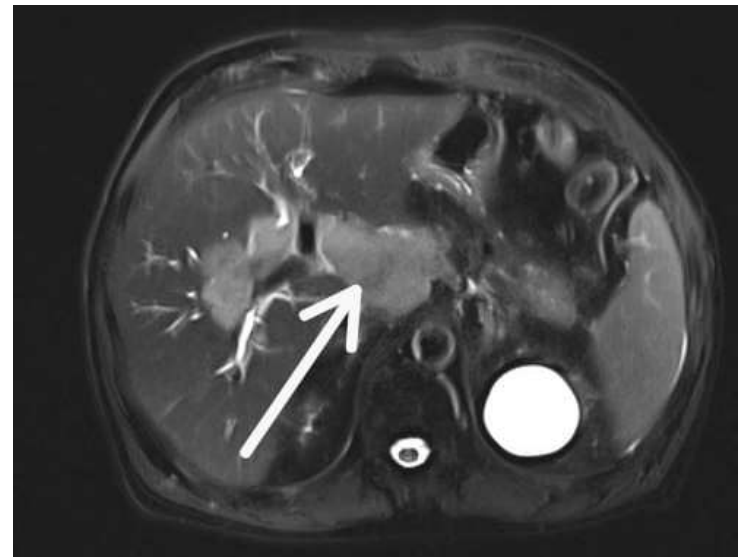
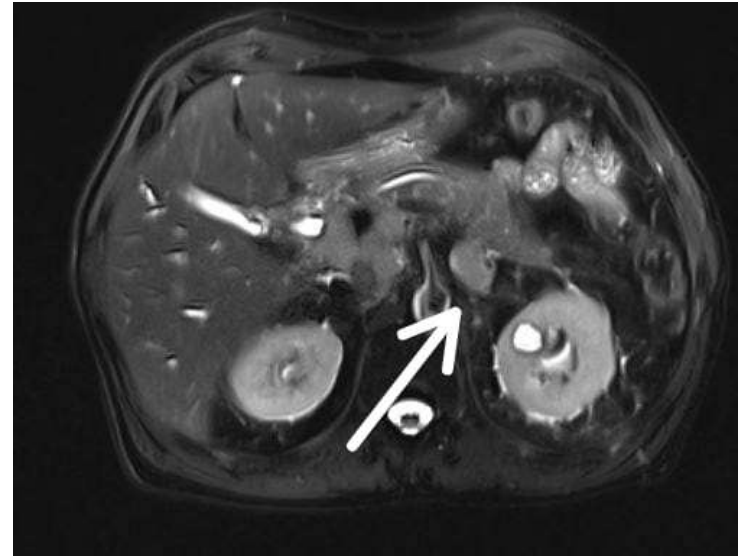
- Left renal calculi (**16\*11 mm in size**) with air foci in the collecting system- s/o Emphysematous pyelonephritis
- Retro-peritoneal necrotic lymphadenitis in peripancreatic, periportal, paraaortic region with largest being **4\*5 cm**, with mass effect and compression of lower CBD and mild IHBR dilatation. - likely Koch's infection/necrotic metastasis.





# MRI ABDOMEN

- Well defined solid lesion with lobulated outlines of 5.5cm\*4cm\*4.3cm in periportal space adjacent to right Portal vein in Rt hepatic lobe  
?Neoplastic aetiology-  
?infiltrating cholangiocarcinoma or  
?Lymphoma.
- Multiple LN at porta hepatis, peripancreatic, periportal, paraaortic and aorto caval region.



# DIFFERENTIAL DIAGNOSIS

- Type II Diabetes Mellitus with urinary tract infection with B/L Emphysematous Pyelonephritis with
- Peri-portal, Peri-pancreatic lymphadenopathy under evaluation-

Tuberculosis / Lymphoma.

- For the diagnosis we decided to get a biopsy for HPE.
- CT Guided biopsy was not possible due to deep seated lymph node.
- Gastroenterologist was requested for Endoscopic USG guided biopsy.

# ENDOSCOPIC USG Guided Biopsy



**EUS showed large necrotic LN between PV and IVC.**



**Fine Needle Aspiration Biopsy (FNAB) was taken and sample was sent for histopathological investigation.**

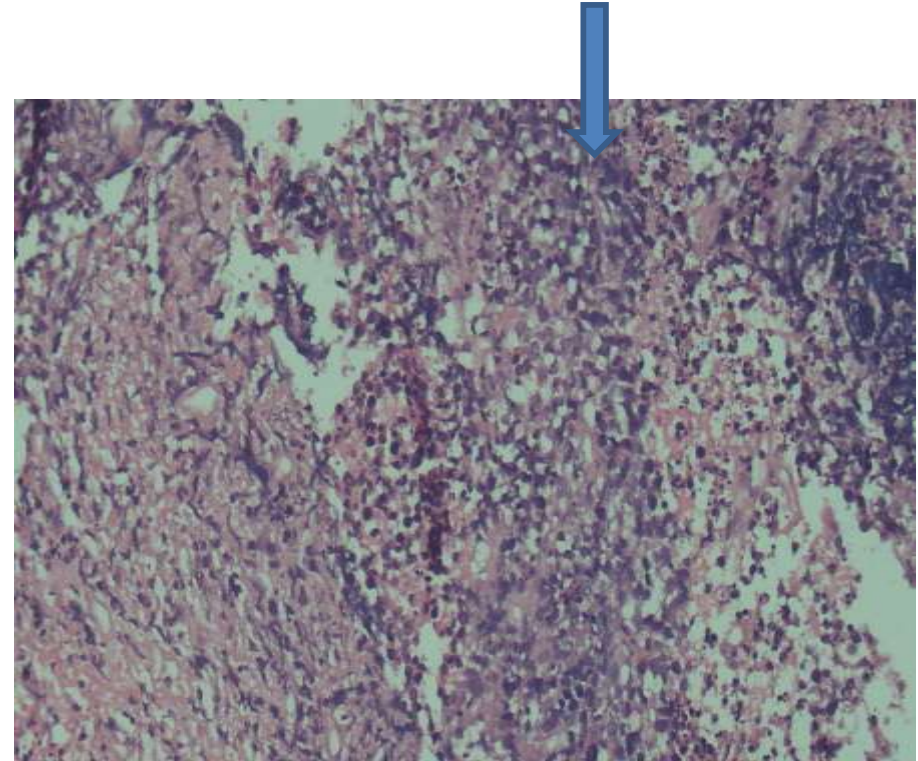
# HISTOPATHOLOGICAL EXAMINATION

Shows a poorly differentiated malignant tumor cells arranged in different sheets.

Tumor cells were large in size and have hyperchromatic nuclei, prominent nucleoli and scanty cytoplasm with increased mitoses.

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s/o **High grade B cell NHL.**



# IMMUNOHISTOCHEMISTRY MARKERS

Neoplastic cells were

-Positive for

- ☐ CD 20,
- ☐ BCL2,
- ☐ C- Myc

-Negative for CD3, CD10, Bcl-6

-MiB-1 proliferative index was 90%.

# FINAL DIAGNOSIS

## HIGH GRADE B CELL NON HODGKINS LYMPHOMA WITH DIABETES MELLITUS TYPE 2

- After the discussion with the Medical oncologist,
- Patient is advised to complete 6 cycles of R-CHOP chemo regimen therapy at an interval of every 21 days.
- Patient is on regular follow up, has completed 5 cycles of chemotherapy till date and is tolerating well.
  
- **R-CHOP regimen :**
- Inj. Rituximab 500mg iv infusion
- Inj. Cyclophosphamide 900 mg iv infusion
- Inj. Vincristine 2 mg iv push over 10 mins.
- Tab Prednisolone 50 mg BD for 5 days

# CASE 2



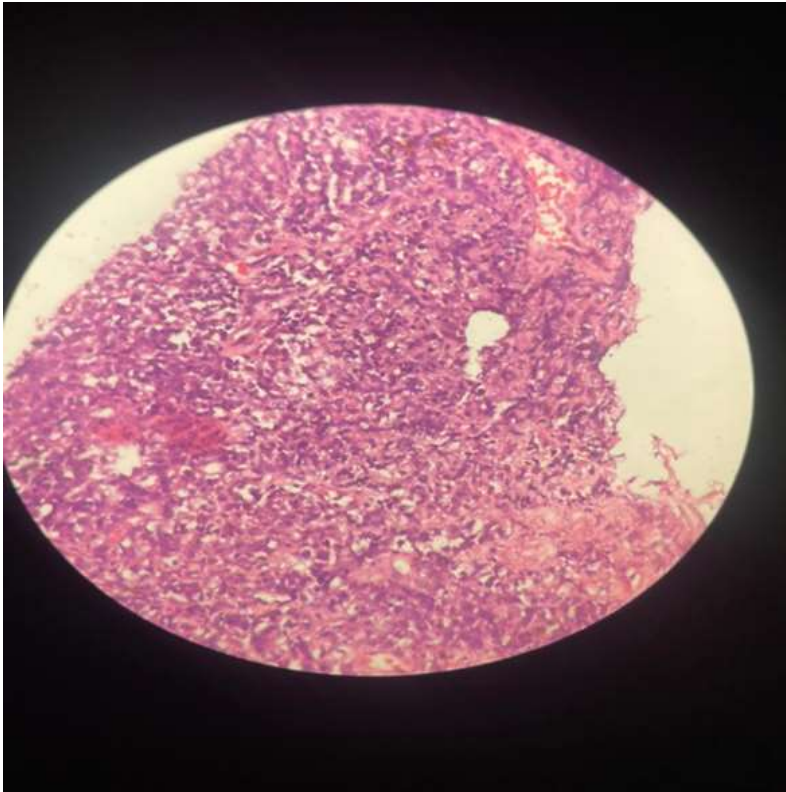
Similar case of 70 yrs old male

Chest x-ray s/o **mediastinal widening** and **cardiomegaly**.

-2d Echocardiography-s/o **moderate pericardial effusion**

**HRCT CHEST** -s/o-  
Homogenously enhancing mass lesion with epicenter in the left anterior-superior part of mediastinum; Moderate Pericardial Effusion

**CT guided biopsy of Mediastinal mass** was done and sample was sent for the HPE.



•**HPE:** shows large pleomorphic nuclei and pale cytoplasm **s/o Diffuse large B cell Lymphoma**

**IHC:**

**CD 20 positive in large cells**

**CD 3 positive in other cells**

**Ki-67- 80%**

**PAN CK- negative**

**Final Diagnosis-DIFFUSE LARGE B CELL LYMPHOMA (DLBCL) WITH PERICARDIAL EFFUSION.**



# DISCUSSION

## Usual presentation of patient with Lymphoma

### THORACIC PRESENTATION

- Cough
- Chest discomfort
- Chest pain
- May present without symptoms but with an abnormal chest radiograph.

### ABDOMINAL PRESENTATION

- Chronic pain
- Abdominal fullness
- Early satiety
- Symptoms associated with
  - Visceral obstruction
  - Acute bowel perforation
  - GI hemorrhage

-An elevated LDH (77%) and B symptoms (47%) are common.

# UNUSUAL PRESENTATION

- Patient may present with pleural and pericardial effusions.
- Superior vena cava syndrome is a frequent complication.
- Relapses can occur locally or in extranodal sites, including the
  - Liver
  - GI tract
  - Kidneys
  - Ovaries and CNS.
- Patient may present with renal or any other organ abnormality as a primary presentation.

# TAKE HOME MESSAGE

- Approximately 25 to 40% of NHL arise in tissues other than lymph node (termed as extra nodal NHL), of which the most common pathological variant is DLBCL.
- If DLBCL is diagnosed early and treated with R-CHOP regimen, it has 5 years overall survival (OS) rate of 92%, so it's highly treatable condition.
- While High grade B cell NHL have very poor prognosis and if not treated timely can be fatal.
- Hence there should be high clinical suspicion to diagnose such atypical presentation of NHL, as they are treatable and have good prognosis if treated timely.

# REFERENCES

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**THANK YOU**