



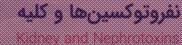




AKI & rare etiology (a case report)

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Case Presentation: (on admission, 1th day)

- A 59yrs-old F.
- Acute skin rush & jaundice since 12 hrs ago.
- Maculo-pupular pruritic ,generalized lesions.
 Concomitant red urine.
- Through recent 12 hrs, UOP decreased and progressive edema.progressive weakness.(cr: 1.8mg/dl).







Second day:

Pt is anuric and may be uremic. LOC is normal.

Exam: only RUQ tenderness.
BP: 90/60. T: 37.8
Percent data:

Recent data:....





Hematology data	WBC=17.2 Hb=13.5 Plt =118000 MCV=90 PTT=53 PT=33.5 INR= 5.9 EOS: 3% Neut: 78% Retic count : 1.1%
Liver tests	Bill T= 14.5 Bill D: 8 Alp= 396 AST=271 ALT=81
Kidney tests	Urea=131 Cr=3.52 U/A: Pr: +++ ، RBC: 30-34 ، WBC: 18-20 ، Glc: + ، Granular cast:1-2
Electrolytes & ABG	PH=7.28 Pco2=36.6 Po2=35 Hco3=17.5 Na=132 Ca=8.35 Alb: 3.5 Ph=5.1 K=5
Other data	CPK=1311CK_MB=65Amylase=148LDH=1540Lipase=109CRP=+Troponin=0.15ESR: 25COVIDPCR: negESR:25
Serology finding	HbsAg : (?) HIVab: (?) HCVab: (?) HAVab :(?) ANA: (?) ANCAc : (?) ANCAp: (?) ,C3,C4,CH50: ????
	نفروتوکسینها و کلیه

Kidney and Nephrotoxins

Additional data:

- Abdominal-pelvic Ultrasonography, Doppler's st : OK
 Echocardiography: OK
 CXR: OK
- DD: SEPSIS.....LEPTOSPIROSIS......VIRAL HEPATITIS....VASCULITIS.....DRUG IND AKI.
- Management : initiation of Ab & short course hemodialysis.





Following days:

Anuria : continue for next days.

Hemodialysis : continue for uremia

 5th day: Cr: 5mg/dl
 First culture :(-)....other serologic test:(-) A course of methylprednisolone tx was done





Key points:

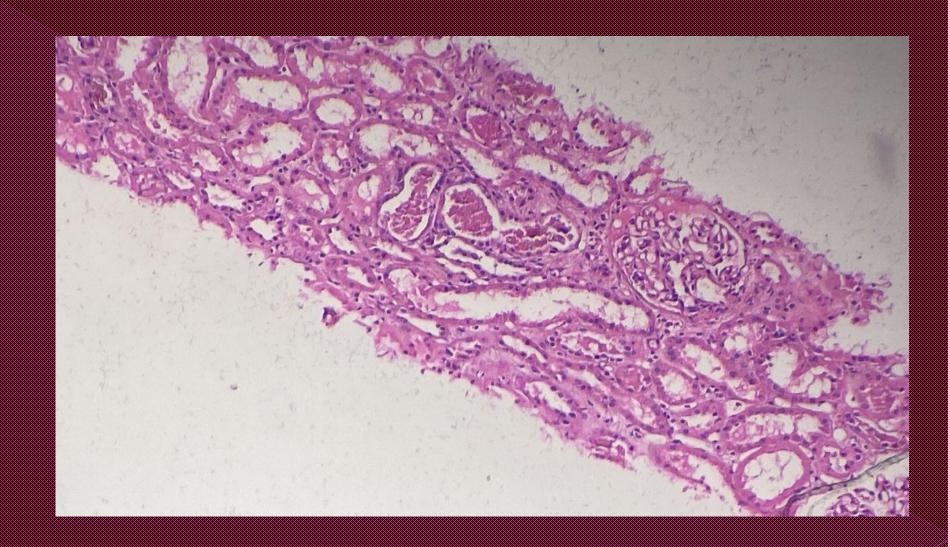
 History of one dose of rifampin cap 1 day before admission.

She had history of brucellosis some years ago and partial tx due to allergic reaction.

Kidney biopsy was done.?????





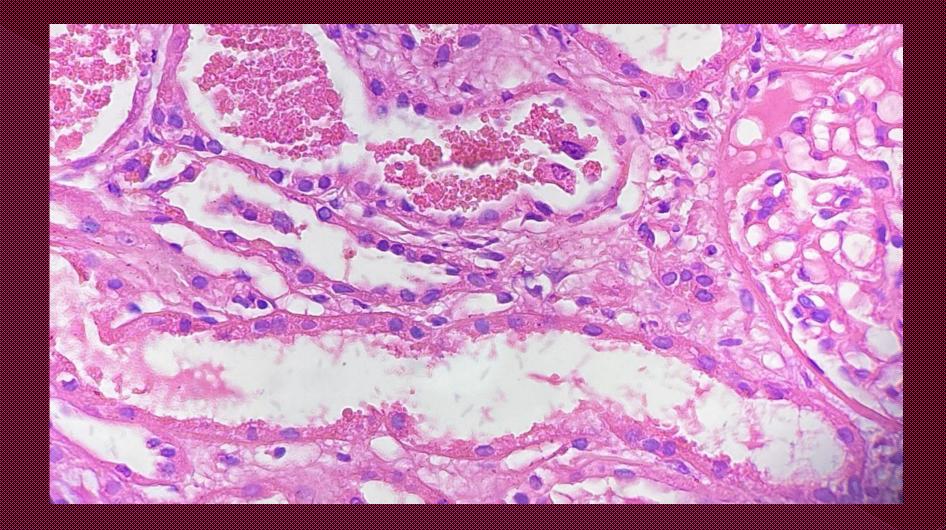










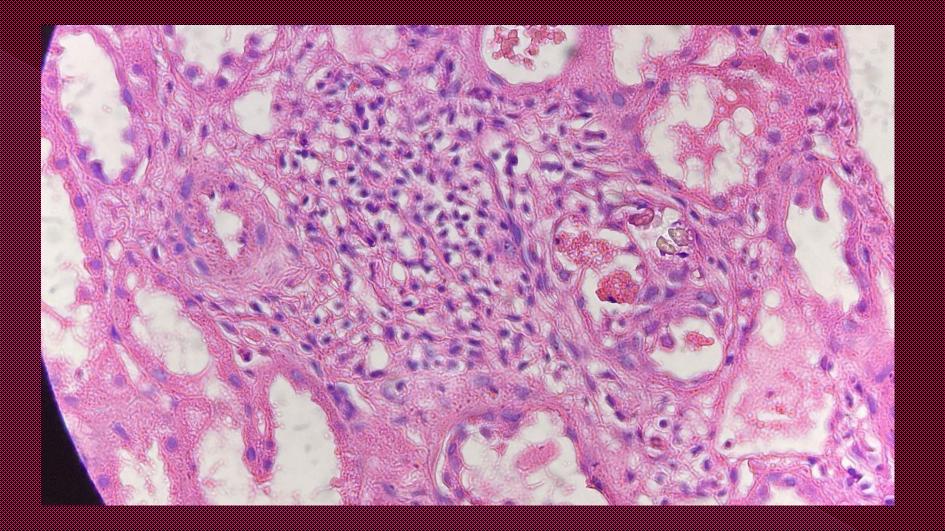




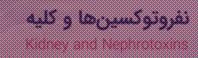




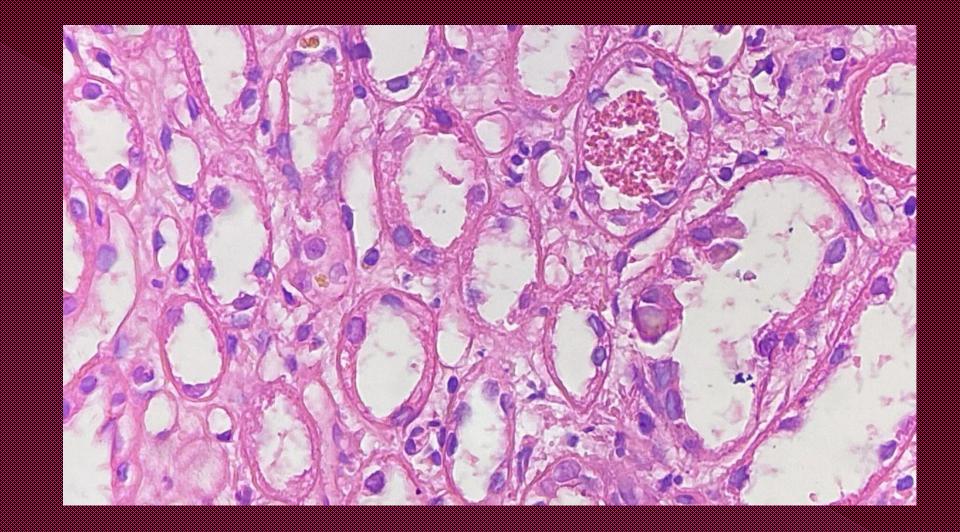




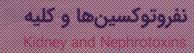
















Final dx:

- Acute interstitial nephritis related to rifampin.
- After discharge, on prednisolone 60 mg/d and IHD
 During next 14days , anuria resolved
- On 3thw, prednisolone tx was tapered and no need IHD.
- On 4th w , pt is well. Cr:1.1. UOP:OK







Discussion:

 Acute interstitial nephritis (AIN) is a common cause of acute kidney injury.

 The most common etiology of AIN is drug-induced disease, which is thought to underlie 60-70% of cases







• **IFAMPICIN** remains a key antibiotic in the treatment of tuberculosis or brucellosis and is used increasingly for the treatment of severe staphylococcal infections.

 Hepatotoxicity is the most frequently observed side effect. The present report focuses on a less known form of toxicity, acute kidney injury.







•most often when the drug is used intermittently.

 is commonly associated with <u>oliguric AK</u> with evidence of hemolytic anemia, thrombocytopenia and <u>hepolitic</u>.

- antibodies against cells expressing blood group antigen I which harms RBCs, platelets, and renal tubular epithelial cells
- type II and type IV hypersensitivity reactions

proximal tubular injury manifests as renal glycosuria
approximately two-thirds of patients require RRT.





BX finding:

- Histologically, rifampin nephrotoxicity is associated with acute tubulo-interstitial nephritis (AIN), tubular necrosis, papillary necrosis, acute cortical necrosis, and minimal change disease.
- Of these, All and tubular necrosis are the most common and frequently develop upon reintroduction of the drug or during intermittent therapy.





- In patients treated with an interrupted regimen, the rifampin-dependent antibody produces acute tubular necrosis requiring dialysis.
- In contrast, the continuous administration of ritampin has been described as progressing more insidiously..







Conclusion

- This presentation aimed to present a case of type II hypersensitivity causing renal injury following a single dose of ritampin.
- Although the occurrence of severe reactions to rifampin is rare, clinicians should be informed of this condition due to its drastic outcomes and potential mortality.





 Renal failure developed on administration of a single dose of rifampicin after a variable drug-free period after continuous ingestion in most patients.

 No variable at presentation predicted the severity of renal failure. Prognosis was excellent, with complete recovery of renal function in all patients.













