Transjugular renal biopsy in a case of nephrotic syndrome with extrahepatic portal venous obstruction

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ABSTRACT

Renal biopsy in patients with nephrotic syndrome helps to establish the pathological diagnosis and subsequent treatment. In certain circumstances, biopsies are difficult to obtain because of the risk of bleeding. We report a case where renal biopsy was obtained through the transjugular route in a patient who had nephrotic syndrome with extrahepatic portal venous obstruction.

Key words: Nephrotic syndrome, renal biopsy, transjugular

Introduction

Considerable literature exists regarding the presence of renal disease in patients with liver disease. Most of these studies are in patients with hepatitis virus infection, and in established cases of cirrhosis of liver. In majority of these studies, renal pathological evaluation is often limited because of the presence of hypersplenism/ bleeding abnormalities. The presence of hypersplenism and its attendant pancytopenia makes biopsies difficult. We report the safe use of the transjugular approach for a simultaneous renal biopsy in a patient with extrahepatic portal venous obstruction and nephrotic syndrome who was offered partial splenic embolization for his portal hypertension.

Case Report

A 25-year-old male patient, a known case of extrahepatic portal venous obstruction and portal hypertension,

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was detected to have nephrotic syndrome during his pre-anesthetic evaluation for hydrocele surgery. He was told that his surgery was deferred till his kidney disease was evaluated and treated. It was felt that the worsening ascites and subsequent scrotal swelling was aggravated by the presence of his nephrotic state, and hence correction of his nephrotic state will reduce post-operative complications and recurrence of his scrotal swelling. He was subsequently referred to our hospital for his renal evaluation and management.

At the age of 12 years, he had an episode of hematemesis and was evaluated and found to have esophageal varices. Evaluation of his portal hypertension revealed the presence of extrahepatic portal venous obstruction. He was managed with oral propranolol. He stayed asymptomatic except for minimal ascites. His liver functions on regular follow-up stayed normal.

Three months prior to this admission, he started developing scrotal swelling, abdominal distension and pedal edema for which he consulted a surgeon. Physical examination revealed pallor, pedal oedema, tense ascites, splenomegaly and bilateral hydrocele. Laboratory investigations showed evidence of anemia, thrombocytopenia, renal insufficiency hypoalbuminemia, and normal liver enzymes. His coagulation work up was normal. His cardiac evaluation did not reveal any abnormality. His urine analysis revealed the presence of proteinuria. Ultrasonography of the abdomen showed massive splenomegaly, ascites and no evidence of liver cirrhosis. Liver biopsy showed normal liver architecture with no abnormality. Adiagnosis of nephrotic syndrome was made and the bilateral hydrocele repair was

postponed, and the patient was referred to the Nephrology services of our hospital for the management of nephrotic syndrome and renal insufficiency.

The laboratory investigations in our hospital are shown in Table1.

A renal biopsy was advised for histopathological diagnosis. However, due to pancytopenia and tense ascitis, a percutaneous renal biopsy was deferred and open renal biopsy was considered. An open surgical biopsy was also considered risky as the anatomical position of his kidneys necessitated 12th rib resection for renal access. The case was discussed with the interventional radiologists whose opinion was sought for management of hypersplenism. It was felt that during his partial splenic embolisation via the transjugular route, a renal biopsy can be performed. Transjugular renal biopsy along with splenic embolization was thus advised and after optimisation with blood and blood products the patient underwent both the procedures in the same sitting. For the transjugular renal biopsy, 8 Fr guiding sheath was guided into the lower pole of right (R) renal vein. The long transjugular liver biopsy needle (Cook USA) was used to get a core biopsy. The needle tip was protected with a 5 Fr catheter. Under fluoroscopic control, the needle was positioned deep in the vein, allowing the tip to protrude into the renal cortex [Figure 1]. No post procedure complications like drop in hematocrit, gross hematuria or perirenal hematoma were encountered.

Post procedure his platelet counts improved to 120,000/μL (vs. 77,000) and WBC count to 16,300, three days after the splenic embolization. However, the biopsy specimen was inadequate for an opinion and a re-biopsy through the transjugular approach was done with no complications [Figure 1]. Biopsy was suggestive of minimal change disease.



Figure 1: Transjugular renal biopsy

Table 1: Biochemical, hematological, radiological and endoscopic findings

endoscopic findings	
Investigations	Results
Laboratory	
Hematology	
Hemoglobin	7.8 g/dl
Total leucocyte count	2800/µL
Platelet count	77000/µL
Prothrombin time	11.4 s
INB	1.08
APTT	38.9 s (test)
ALLI	26.0 (control)
Biochemistry	20.0 (00/11/01)
Creatinine	1.5
Urea	59.9
Uric acid	7.73
BUN	28
BUN/Creatinine ratio	22.2
Albumin (A)	2.4
Globulin (G)	3.5
Total protein	5.86
A/G ratio	0.7
Sodium	136.6
Potassium	4.9
Chloride	110
Calcium	8.2
Phosphorus	4.52
Alkaline phosphatase	118.6
Cholesterol	278
LDL Cholesterol	166
24 h urine protein	10.2 g/24 h
Urine protein/creatinine ratio	11.02
Microbiology	
Urine Routine	4 plus protein, 2-4 pus cells/hpf,
	80-100 RBCs, No casts
Ascitic fluid analysis	
Appearance	Slightly hazy
Color	Colorless
Total proteins	1.81 g/dl
Albumin	0.69 g/dl
Glucose	96.8 mg/dl
Total nucleated cells	20 cells
Neutrophils	15%
Lymphocytes	85%
RBCs	+
Microscopic examinations	No organisms seen
Immunology	
C3	Normal
C4	Normal
ANA	Negative
dsDNA	Negative
Antiphospholipid antibody IgG	12.69(0-15)
Antiphospholipid antibody IgM	3.81(0-15)
Endoscopy/liver scan	
Upper GI endoscopy	Esophageal varices
Liver scan	Massive splenomegaly, ascites,
	normal liver size
Radiology	
CT abdomen with contrast	Splenomegaly. Multiple
	peripancreatic, periportal,
	perisplenic, splenorenal, gastric
	and esophageal varices. No
	radiological changes of liver
	cirrhosis. Extrahepatic portal
	obstruction, tense ascites

He was started on oral steroids. His condition gradually improved with resolution of pallor and pedal edema and ascites. Pre-steroid urine protein/creatinine ratio was 11.67 which improved to 3.02 after 1week of steroids and ACE inhibitors. His haemoglobin and albumin levels improved subsequently. He underwent bilateral hydocele repair successfully with no complications. Predischarge he received his pneumococcal vaccination. He is now asymptomatic and under regular follow-up. At 6-months post follow-up, he was in clinical and biochemical remission for his nephrotic syndrome and had normal renal functions.

Discussion

Renal glomerular changes are well recognized in patients with liver disease. Membranous and membranoproliferative glomerulonephritis have been described with hepatitis B and Hepatitis C infection.[1,2] There are case reports of association of minimal change nephropathy with chronic hepatitis C infection.[3] However, glomerular lesions are not as common as ATN/AKI in cirrhosis of liver.[4] In patients with non-cirrhotic portal hypertension, IgA nephropathy appears to be common.^[5] Interestingly, there is an increase in incidence of glomerulonephritis in patients with non-cirhhotic portal hypertension who have undergone splenorenal shunt surgery.[6] This study had reported minimal change nephropathy as an uncommon renal lesion (incidence of 3%). Our patient had minimal change nephrotic syndrome on biopsy.

Most patients with liver disease don't undergo indicated renal histological evaluation because of the presence of bleeding abnormalities. In such situations, transjugular renal biopsy may be attempted if the expertise is available. The first report of transjugular renal biopsy appeared in literature 20years back.[7] This procedure is practiced commonly in continental Europe and its experience is limited in USA. However, literature from across the Atlantic consider it safe and effective in high-risk patients.[8-10] The effectiveness of the procedure in the hands of interventional nephrologists is also reported.[11] In certain centers, this procedure is attempted with simultaneous insertion of internal jugular hemodialysis catheter in patients with acute renal failure (ARF).[12] This early availability of renal histology altered the management in 50% of the patients with ARF. The procedure is considered relatively safe with minor complications like renal hematoma, macroscopic hematuria and capsular perforation reported in literature. [8,13,14] Contrast nephrotoxicity has also been reported. Major complications of arterio-pelvicalyceal fistula and renal vein thrombosis are rare.[15] Overall, complication rates are similar

to that of percutaneous biopsy. Our patient had no bleeding complications.

The diagnosis of minimal change disease and the consequent treatment with steroids allowed his surgeons to perform the required surgery after 3 months. Six months later he was back to his desk job with no major medical problems.

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