Complications of Kidney Biopsy in a Tertiary Hospital of Central Nepal, Chitwan

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ABSTRACT

Background: Kidney biopsy is an important diagnostic tool in Nephrology and is said to be relatively a safe procedure. There are limited studies in the complications of kidney biopsy from this region. We therefore thought of looking into the complications of kidney biopsy and its risk factors.

Methods: A hospital based analytical cross sectional study was carried out over a period of 3 years. Kidney biopsies were done under ultrasonography guidance. The complications and its risk factors were recorded and were analyzed using mean, standard deviation, ratio, percentage and chi square.

Results: A total of 210 patients were analysed. The mean± standard deviation of 210 patients was 35.7±14.9 years. The commonest minor complication was biopsy site pain not requiring analgesics 136 (64.8%) and the commonest major complication was biopsy site pain requiring analgesics 18(8.6%) followed by perinephric hematoma 10 (4.8%). There was a significant association between low platelet count and the development of complications like pain requiring analgesic (p value 0.04), perinephric hematoma (p value 0.022) and gross hematuria (p value 0.011).

Conclusions: Kidney biopsy is a safe procedure and low platelet count is a significant risk factor for complications (p value <0.05).

Keywords: complications; kidney biopsy; perinephric hematoma; platelet count.

INTRODUCTION

Kidney biopsy is an important invasive procedure in Nephrology. It is useful not only in diagnosis but also in the prognosis of the disease.1 It is known that invasive procedure has some inherent complications however the kidney biopsy is said to be a relatively safe and effective procedure with life-threatening complications occurring in less than 0.1% of biopsies.2,3 Previous study from Chitwan had also shown kidney biopsy to be a safe procedure.4 Kidney biopsies are regular procedure in various center but there are limited studies in the complications of kidney biopsy from this region. We therefore thought of looking into the complications of kidney biopsy procedure. This study was done with an objective to know the nature of the complications and the risk factors associate associated with the complication of the kidney biopsy in our center.

METHODS

This was a hospital based analytical cross sectional study carried out College of Medical Sciences-Teaching Hospital, Chitwan over a period of 3 years from May 2016 to April 2019. All the consecutive kidney biopsies were included in the study. The approval to conduct the study was given by the Institutional Review Committee of the hospital and a written consent was taken from each patient prior to the biopsy procedure, after explaining the risk and benefit of the procedure and the possible complications of the procedure. The indication of kidney biopsies were the standard indication based on clinical presentation and investigations. Kidney biopsies were done by a nephrologist who had an experience of kidney biopsy for > 7 years. A self-adjustable, automated, spring loaded gun biopsy needle of 16 to 18 gauge (Bard Monotopy USA 16-18 G) was used for kidney biopsy. Kidney biopsies were performed under ultrasonography (USG) guidance with the help of radiology residents or consultants. The lower pole of the kidney was identified for biopsy. Two cores of renal tissue were removed with maximum number of pricks not being more than four times. After the procedure, patients lay in bed flat on their back till they void their first urine and then remained in bed for 24 h of observation. Patients were monitored closely after biopsy for signs or symptoms of complications, such as flank pain requiring and not requiring analgesics, gross hematuria, need of blood transfusion or hypotension. The vital signs were checked every 15 min for 2 h, every hour for 4 h, every 2 h for 6
RESULTS
A total of 210 kidney patients were analyzed over a period of 3 years. The mean ±SD age of the patient was 35.7 ± 14.9 years. The age ranges of patients were from 9 years to 76 years. The mean ±SD size of biopsy tissue obtained was 0.9 ± 0.3 cm. The minimum and the maximum size was 0.3 and 1.8 cm (Table 1).

The average ±SD number of glomeruli obtained was 35.7 ± 14.9 years. The age ranges of patients were from 9 years to 76 years. The mean ±SD size of biopsy tissue obtained was 0.9 ± 0.3 cm. The minimum and the maximum size was 0.3 and 1.8 cm (Table 1).

The average ±SD number of glomeruli obtained was 23.4 ± 11.03. The minimum number was 2 and the maximum was 76. Of total glomeruli < 20 were seen in 94 (44.8%), glomeruli 20-30 were seen in 70 (33.3%) and >30 were seen in 46 (21.9%) (Table 2).

The average serum creatinine was 2.2 ± 2.5 mg/dl. The minimum serum creatinine was 0.3 mg/dl and maximum serum creatinine was 18.1 mg/dl (Table 3). None of the patients developed hypotension or need of blood transfusion post biopsy and none of had mortality related to kidney biopsy (Table 4).

DISCUSSION
There are limited data on short term and long term complications of kidney biopsy from Nepal. In
most of the studies done in complications of kidney biopsy from Nepal, the overall complications rate are low. The most common complication observed in our study was pain not requiring analgesics (minor complication) in 136 (64.7%) patients. However the most common major complication was moderate to severe flank pain requiring analgesics in 18 (8.6%) patients. Burnstein et al., reported complications in 91 (14.3%) patients, out of which 6.6% were minor complications and rest were major. A study by Mendelsson and Cole in their series found an overall complication rate of kidney biopsy to be 5.3%. Similarly a study done by Tuladhar et al., showed the overall complication to be three (4%). Other studies from Nepal done by Maskey et al., Manandhar et al., and Ghimire et al., showed gross hematuria as the commonest complication of kidney biopsy in 15 (8.5%), six (8%) and five (6.7%) patients respectively. Perinephric hematoma was seen in 10 (4.8%) patients in our study. However not all patients had routinely undergone USG evaluation to look for perinephric hematoma, which was one of the limitation of our study. Only those patients who had severe flank pain requiring analgesics in 18 (8.6%) had undergone USG evaluation to look for development of perinephric hematoma and of them (n=18) 10 were found to have perinephric hematoma. We adopted this approach to minimize the overall cost of kidney biopsy. Because of this, we could not establish the association between flank pain and the perinephric hematoma. This will be an area of future research.

In some other studies done by in Manandhar et al., Tuladhar et al., Ghimire et al., and Maskey et al., perinephric hematoma was seen in two (2.67%) patients, three (4%), four (5.3%) patients and eight (4.5%) respectively. According to a study done by Ralls et al., clinically significant perinephric hematomas was seen in 6% cases. We tried to look for the association between low e-GFR, hemoglobin level and platelet count and complications. In our study, > 50% had chronic renal insufficiency (e-GFR <60ml/min) and we tried to see the association between low e-GFR and complications, however, there was no significant association between the low e-GFR <60ml/min and the complications like pain requiring the analgesia (P value =0.543), perinephric hematoma (p value 0.860) and gross hematuria (p value 0.569). However previous studies showed low e-GFR to be the predisposing factor for complications and the risk of bleeding increases with worsening renal function.

In our study none of the patient needed any surgical or radiological intervention and none of the patient had hypotension or mortality. Similarly Maskey et al., Pandey et al., Ghimire et al., Manandhar et al., and Tuladhar et al., also did not report any cases of hypotension or biopsy related death. In a review done by Kobert SM in 9595 kidney biopsies over last 50 years, only 0.3% required major surgical or radiographic intervention and the mortality associated with the procedure was in < 0.1% of cases. Requirement of blood transfusion post procedure was not seen in our study. Similar observations were made by studies done by Ghimire et al., Manandhar et al. and Tuladhar et al. However three (2.5%) patients required blood transfusion post biopsy in the study done by Pandey et al. and in 1 (0.6%) patient in the study done by Maskey et al.

The average number of glomeruli and the average size of the biopsy tissue obtained in our study were 23.4±11.03 and 0.9±0.3cm respectively. A study from Romania done by Covic et al., showed that serious complications were more frequent when a large tissue fragments were harvested (>20 glomeruli). However we had not tried to see the association between the sizes of the tissue recovered and the number of pricks applied with the complications. This could be an area of future research. In our institute we do not recommend pricking more than 4 times. There was no significant association between the Hb level and the complications in our study however, there was significant association between low platelet count and the development of complications like pain requiring analgesics (p value 0.04), perinephric hematoma (p value 0.022) and gross hematuria (p value 0.011). Similar association between low platelet count and complication was seen in a study carried out over the 5-year period in 3,577 native kidney biopsies, where multivariable logistic regression analysis demonstrated low platelet count to be independently associated with the risk of complications. Likewise significant association between low platelet count and the risk of bleeding was seen in a study done by Monahan et al. (p = 0.002).

With the existing studies within and across the country and our study, it seems kidney biopsy is a safe procedure. However this was an observational study, so all the inherent limitations of observational studies were there in the study. This was a single center study so the results could not be generalized to the whole region or the country. We had not done the post biopsy USG in all the cases to minimize the cost, so we could not look for perinephric hematoma in all cases and the true prevalence of post biopsy perinephric hematoma was not represented and its association with the complications couldn’t be established. We had not seen the association of age, sex, blood pressure, size of the biopsy needle (16 Vs. 18 gauge), number of pricks applied and the size of the
biopsy tissue with the complication variables. We had also not seen the long term complication of kidney biopsy. These limitations demand another well structured, larger and multicentric study to better define the complications of kidney biopsy in our center and the country as a whole.

CONCLUSIONS
The kidney biopsy was a safe procedure and the low platelet count was a significant risk factor for complications (p value <0.05). The overall complication rate was 18 (8.6%). Kidney biopsy seems a safe procedure, so one should not hesitate to perform it whenever indicated. However, we need a large and multicentric study to understand the true prevalence of complications in the country.

REFERENCES

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