Papillary vs non-papillary access during percutaneous nephrolithotomy: Retrospective, match-paired case-control study

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Summary

Objective: The most crucial steps of percutaneous nephrolithotomy (PCNL) are the percutaneous access and dilation of the access route. Recent literature suggests that papillary access to renal calyx is the accepted method. Despite this rule, we do not always make papillary puncture and we puncture wherever we can to achieve stone-free status and reduce unnecessary access. In this study, we present our results with papillary vs non-papillary access in patients with a kidney stone.

Material and methods: Two hundred and seven patients with non-papillary access and 69 patients with papillary access who had similar demographics (age, body mass index (BMI), stone size) were selected with pair match analysis (3:1). Preoperative and postoperative data were collected from the patient's chart. Operative time (from starting surgery to nephrostomy tube), drop-in hematocrit level, transfusion rate, duration of hospital stay, perioperative and postoperative complications (Clavien-Dindo Classification) and stone-free status (no or < 3 mm residual stone) were also evaluated in both groups.

Results: The mean operative time was similar in between two groups. The mean hematocrit decreases not differ between the two groups (p = 0.56). In papillary group, only 2 patients (3.2%) required transfusion and only one patient (1.4%) in the non-papillary group had a transfusion with no statistically significant difference (p = 0.43). The overall complication rates were 7.1% in the papillary group and 7.2% in the non-papillary group (p = 0.89). Postoperative mean creatinine level was similar between the two groups.

Conclusions: In this study, we found that non-papillary access is a feasible option for PCNL in the terms of stone-free rate and complication rates.

KEY WORDS: Percutaneous nephrolithotomy; Access; Papillary; Non-papillary.

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INTRODUCTION

Percutaneous nephrolithotomy (PCNL) is still the standard therapy for the larger calculi in the kidney (1). The most crucial steps of PCNL are the percutaneous access and dilation of the access route. As accepted by literature, punctures must be done through the papilla of the posterior renal calyx to avoid the major vascular structures of the kidney (2). Despite this rule, access through the papilla is not always achievable. In our huge volume percutaneous surgery centre (over 400 cases per year) we do not always make papillary puncture and we puncture wherever we can to achieve stone-free status and reduce unnecessary access. Non-calyceal puncture method was recently published and the authors concluded that it is feasible and probably not as dangerous as it was stated (3). In this study, we present our results with papillary vs non-papillary access in patients with kidney stone.

MATERIAL AND METHODS

After an Ethics Committee approval was obtained, a total of 638 patients patient undergoing PCNL between January 2017 and June 2018 were analyzed. Two hundred and seven patients with non-papillary access and 69 patients with papillary access who had similar demographics (age, BMI, maximum diameter of the stone) were selected with pair match analysis (3:1). Preoperative and postoperative data were collected from the patient's chart. Operative time (from starting surgery to nephrostomy tube), drop-in hematocrit level, transfusion rate, duration of hospital stay, perioperative and postoperative complications (Clavien-Dindo Classification) and stone-free status (no or < 3 mm residual stone) were also evaluated in both groups. Preoperative and postoperative third-month creatinine level were also recorded and analyzed. Follow-up was made with low dose non-enhanced computed tomography (NECT) three months after surgery.

Patients with a solitary kidney, history of previous surgery (open or endoscopic) or Extracorporeal Shock Wave Lithotripsy (ESWL) for the same kidney, congenital anatomical variants (horseshoe kidney, ectopic kidney) were excluded.

In surgical technique: after insertion of open-end 4 F ureteral catheter in lithotomy position, patients were set to the prone position. All patients were treated with combined fluoroscopic and ultrasound guided PCNL. In papillary access group, papilla of the calyx was punctured. In the other group, the puncture was made through infundibulum of the calyx. The puncture site was corrected under ultrasound control. After insertion of hydrophilic guidewire all cases were dilated up to 16 F with Amplatz dilators and then balloon dilatation was made up to 30 F. Rigid nephroscope and pneumatic lithotripter were used to remove calculi. Fragments were

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removed using basket catheter and forceps. At the end of the procedure, 14 F nephrostomy catheter was inserted in all cases.
Statistical analysis was performed with the IBM SPSS version 20 (IBM Corp., Armonk, NY, USA). Fischer’s exact test, Pair-Match analysis, Mann-Whitney U, t-test and were used for the analysis of the data and statistical significance was accepted as p-value < 0.05.

RESULTS
Patient age, BMI and stone size were similar (nearly the same) in both groups according to case match analysis (Table 1). The mean operative time was also similar. The average of hematocrit decrease was 3.45 ± 2.2 in the papillary group and 3.89 ± 3.3 in the non-papillary group with no difference between the two groups (p = 0.56). In the papillary group, only 7 patients (3.8%) required transfusion and only one patient (1.4%) in the non-papillary group had a transfusion with no statistically significant difference (p = 0.43).
The overall complication rates were similar being 7.1% in the papillary group and 7.2% in the non-papillary group (p = 0.89). According to Clavien-Dindo Classification, only one patient in the non-papillary group and three patients in non-papillary group had Class IIIa (required selective angiobolization because of uncontrolled bleeding) complication (p = 0.87).
The mean duration of hospital stay was also similar. Postoperative mean creatinine level was similar between the two groups. All statistical analysis are showed in Table 2.

**Table 1.**
**Patient characteristics.**

<table>
<thead>
<tr>
<th></th>
<th>Papillary group (n: 207)</th>
<th>Non-papillary group (n: 89)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>51.6 ± 13.5</td>
<td>52.2 ± 12.43</td>
<td>0.94</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>28.3 ± 5.2</td>
<td>27.0 ± 3.6</td>
<td>0.93</td>
</tr>
<tr>
<td>Stone size (max. diameter, cm)</td>
<td>2.46 ± 4.6</td>
<td>2.38 ± 5.1</td>
<td>0.85</td>
</tr>
<tr>
<td>Gender f/M, n</td>
<td>66/141</td>
<td>24/45</td>
<td></td>
</tr>
<tr>
<td>Location</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Upper</td>
<td>23</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Middle</td>
<td>25</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Pelvis</td>
<td>82</td>
<td>26</td>
<td></td>
</tr>
<tr>
<td>Lower</td>
<td>77</td>
<td>23</td>
<td></td>
</tr>
</tbody>
</table>

Values are presented as mean values ± standard deviation. BMI = body mass index; M = male; F = female.

<table>
<thead>
<tr>
<th></th>
<th>Papillary group (n: 207)</th>
<th>Non-papillary group (n: 89)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Operation time, min</td>
<td>58.3 ± 14.3</td>
<td>56.8 ± 15.3</td>
<td>0.56</td>
</tr>
<tr>
<td>Drop in hematocrit level</td>
<td>3.45 ± 2.2</td>
<td>3.45 ± 2.7</td>
<td>0.93</td>
</tr>
<tr>
<td>Length of hospital stay, days</td>
<td>4.45 ± 1.9</td>
<td>4.51 ± 1.8</td>
<td>0.42</td>
</tr>
<tr>
<td>Stone Free Status (%)</td>
<td>86.4%</td>
<td>85.5%</td>
<td>0.66</td>
</tr>
<tr>
<td>Mean change in Creatinine, mg/dl</td>
<td>0.06 ± 0.29</td>
<td>0.05 ± 0.41</td>
<td>0.68</td>
</tr>
<tr>
<td>Overall complications (%)</td>
<td>7.1</td>
<td>7.2</td>
<td>0.89</td>
</tr>
<tr>
<td>&gt; Clavien-Dindo Class III</td>
<td>1.4</td>
<td>1.4</td>
<td>0.87</td>
</tr>
<tr>
<td>Transfusion rate (%)</td>
<td>3.8</td>
<td>1.4</td>
<td>0.16</td>
</tr>
</tbody>
</table>

Values are presented as mean values ± standard deviation.

**DISCUSSION**
After Fernstrom and Johansson reported the first percutaneous nephrolithotomy in 1976 and Allen et al. published the first series of percutaneous interventions for removing renal stones, PCNL started to become widely used for renal stones (4, 5). Operative technique and endoscopic equipment are still evolving to increase success and decrease morbidity.
Recent literature suggests that preferred puncture site is papillary access on the avascular line to avoid the risk of bleeding. Sampaio et al. were studied to determine the best route to puncture in 1992 and they found that, in infundibular access, upper site was injured in 67.6% (41.1% venous and 26.5% arterial), mid site of kidney in 61.5% (38.4% venous and 23.1% arterial) and lower site in 68.2% (54.6% venous and 13.6% arterial).
In the direct puncture of pelvis 33.2% injuries were recorded but in calycal fornix overall injuries were recorded only 7.7% and they were all venous injuries (6). This study had three major concerns; firstly, it was cadaveric study and renal-related anatomical tissue was not evaluated, secondly renal and cortical system was native and anatomical changes due to stone was not considered, thirdly no renal functional evaluation was made, and recovery was not assessed.
Kallidonis et al. published a prospective randomized trial that compared papillary vs non-papillary access in PCNL, and they found that access to infundibulum is the feasible and safe procedure and it is not associated with higher blood loss and transfusion rate (7).
Kallidonis et al. evaluated infundibulum of the middle calyx approach technique by 99mTc-dimercapto-succinie acid SPECT/CT renal scintigraphies and/or computerized tomographies perfusion (CTP), they found that the punctures to the mid calyceal papilla fornix and infundibulum as well as pelvis have similar angles of approach and that effects on parenchyma involved in the tract dilatation are similar (8).
They concluded that infundibular puncture can be an option to puncture in the performance of PCNL.
In another study Kyriazis et al. investigated the feasibility and safety of PCNL with non-calycal access track; they operated 137 patients consecutively, including 10 cases with anatomical variations, under fluoroscopic guidance (3). Stone free status was 89.2% for a single stone, 80.4% with multiple and 66.7% for staghorn stones. The overall complication rate was 10.2% and the major complication rate was 3.6%. The authors concluded that calycal access is feasible and safe with stone-free status and low complications.
The length of hospitalization is a new investigation era for PCNL. The technology continues to improve, and postoperative complications are decreasing. “outpatient” procedure is an option now for highly selected patients (9). However, we still routinely admitted patients to follow-up. In terms of hospital stay our results are comparable with a recent literature (10).
In this study, the overall complication rates were 7.1% for the papillary group and 7.2% for the non-papillary group. Transfusion rates were 3.8% and 1.4%.
In a study by Wiesenthal et al. which compared shockwave lithotripsy, ureteroscopy, and PCNL for renal cal-

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culi between 100-300 mm²; they found that overall complication in PCNL was 14% (11). In a recent study comparing retrograde intrarenal surgery versus PCNL, in PCNL group bleeding rate was 6.87% and it was higher than our results (10). This study has several limitations; firstly, the study has a retrospective nature. Secondly, control patients were not randomized and selected for analysis. The surgeons (three surgeons) are very experienced – over 300 cases – and that may cause low complication rates. Patients are not consecutively enrolled in this study and anatomical variations are not evaluated. Multiple access and long-term complications are not also evaluated. Low number of patients because of the wide exclusion criteria is another limitation of this study.

**CONCLUSIONS**

In this study, we found that non-papillary access is a feasible option for PCNL in terms of stone-free status and complication rates. More anatomical and radiological, prospective randomized studies involving a great number of patients may be needed to determine the safety and efficacy of non-papillary access.

**REFERENCES**


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