The Use of Simple Metabolic Evaluation in Determining the Etiology of Recurrent Stone Formers

Abstract

Objectives: Recurrence rates in urolithiasis may reach to 50% after 5-10 years. Simple metabolic evaluation (SE) is commonly used for evaluating patients with low risk for recurrence. In the present study, patients with high risk to develop for stone recurrence were evaluated by single sample of urine collected in 24 hours urination with standard diet.

Methods: Two hundred and sixty eight cases who were supposed to be patients with high risk for stone recurrence were evaluated. Fifty three patients who had routine follow up for 5 years were investigated. Calcium, sodium, potassium, magnesium, creatinine, BUN, uric acid, parathormone in blood and calcium, uric acid, magnesium, citrate, urine volume and oxalate in the sample of urine collected in 24 hours urination with standard diet were checked.

Results: Metabolic abnormalities were determined in 87% of the patients. During the five years, more than 3 parameters corrupted in 9 (17%) and less than three parameters corrupted in 12 (23%) patients. In 5 (1%) patients the same parameters never changed. In 14 (26%) patients all of the parameters improved whereas at least one parameter recovered in 13 (25%) patients. Stone recurrence was seen in 2 (4%) patients.

Conclusion: Metabolic evaluation and preventive medical therapy are supplementary steps in the treatment of urolithiasis. We conclude that, although SE seems to be safe, inexpensive and easy to perform in the evaluation of patients with high risk to develop stone recurrence, the compliance of the patients may be the restrictive parameter for the success.

Abbreviations: SE: Simple Metabolic Evaluation; SWL: Shock Wave Lithotripsy; URS: Ureterorenoscopic Modalities; PNL: Percutaneous Nephrolithotomy; Ca: Calcium; Na: Sodium; K: Potassium; Mg: Magnesium; CIT: Citrate; OX: Oxalate; HCU: Hypocitraturia; HCall: Hypercalciuria; CE: Comprehensive Evaluation; SF: Stone Formers

Introduction

Urolithiasis is one of the most common disorders in urology practice with an incidence of 0.4-1% in industrialized countries [1]. Innovations in the therapeutic applications, such as extracorporeal shock wave lithotripsy (SWL), minimally invasive ureterorenoscopic modalities (URS) and percutaneous nephrolithotomy (PNL), have made most of the stones treatable. On the other hand, even for children, recurrence rates may reach to 50% in 5 to 10 years and 75% in 20 years without routine follow up and medical treatment [2-4]. Patients carrying risk factors such as developmental urinary anomalies, genetic predisposition, excessive excretion of promoters or reduced excretion of inhibitors for crystallization are at high risk for new stone formation [5]. These patients should be evaluated to reveal the underlying pathology for preventive medical therapy [6]. Metabolic evaluation in patients with calculous disease reveals some form of abnormality in over 90% of patients [6]. However, comprehensive metabolic evaluation is often neglected due to meticulous attention needed for urine collection, costs and dietary restriction. In this study, the efficacy of simple metabolic evaluation, commonly used for low risk stone formers, is evaluated in the management of high risk stone formers.

Materials and Methods

With the approval of Institutional Board and Local Ethic Committee (KA/0250), 53 patients (28 male, 25 female) of 268 patients (149 males, 119 females) regarded as high risk cases for stone recurrences were evaluated for 5 years, prospectively. These 53 patients attended routine controls in the 5 year period. The mean age of the patients was 46 (3-71) years. The patients underwent SWL, URS, PNL, open surgery or passed the stone spontaneously. One month later, all of the patients were checked for stone-free status by performing non-contrast spiral computerized tomography or intravenous urography. Stones were analysed by using X-ray diffractometry method. At the initial visit the patients were queried about the stone episodes, history of any surgery (gastrointestinal or genitourinary surgery), systemic diseases such as osteoporosis, sarcoidosis, etc., family history for urolithiasis, dietary methods and fluid intake, stone analysis, the age at the first episode and medication (Vit-D, calcium, etc).
Urine was analysed for pH, glucose, density, microscopy for erythrocyte, leucocyte, crystallization (cystine) and culture. Calcium (Ca), sodium (Na), potassium (K), magnesium (Mg), creatinine, urate (UA), parathormone in blood and Ca, Na, K, Mg, UA, citrate (CIT), urine volume and oxalate (OX) in the sample of urine collected in 24 hours urination with standard diet were checked. The urine sample was collected in a dry and clean plastic container. No preservative material, such as hydrochloric acid, was used. The samples were delivered to the laboratory as soon as possible. In the laboratory urinary volume and pH were measured immediately, and then a part of it was acidified for urinary OX, Ca, Na, K and CIT whereas the remaining was alkalinized for urinary UA calculation. The patients were treated according to the metabolic disorder determined by SE and were re-assessed in 6 months’ period. The changes in blood and urine parameters after the treatment were examined.

**Results**

The total numbers of the stones were 1.34 [1-3]. Initial stone burdens were 28.5 (10-300) mm². The stone analysis of the patients was as seen in Figure 1. In the beginning, 87% (233/268) of the patients revealed metabolic abnormalities. Hypocitraturia (HCU) was detected in 60% of the patients as the most common pathology (Table 1). The hypocitraturia was treated by potassium citrate of 30-60 mEq/day and urine pH was evaluated in every visit. Despite the treatment, another pathology accompanied to HCU in 1 (1/53) patient. In the end of the study, hypocitraturia was completely improved in 40% (21/53) of the patients. New stone formation was not reported (Table 2).

Hypercalciuria (HCAU) was established in 49% (26/53) of the subjects (Table 1). The hypercalciuric patients were treated by indapamide of 2.5 miligrams/day and potassium citrate (30-60 mEq/day). Potassium citrate was used to prevent the HCU secondary to indapamide treatment. During the follow up, stone recurrence occurred in only one patient in whom HCAU never improved. More than 50% of the patients with HCAU reached normal level by this treatment. Interestingly, when detected as the sole pathology, HCAU improved in all patients during the 5 years period (Table 2). Hyperoxaluria was determined in 32% (17/53) of the subjects (Table 1). It was treated by correcting the dietary habits and encouraging high fluid intake. It recovered in 41% of the patients. However, new stone formation appeared in 1 patient while hyperoxaluria persisted in 8 patients (Table 2).

During the five years, more than 3 parameters became abnormal in 9 (17%) and less than three parameters became abnormal in 12 (23%) patients. Interestingly, 5 of 7 patients whose parameters were normal initially, revealed various abnormalities during the follow up. In 5 (1%) patients the same abnormal parameters never changed. In 14 (26%) patients all of the parameters improved whereas at least one parameter recovered in 13 (25%) patients (Table 2). Stone recurrence was seen in 2 (4%) patients whose parameters were always abnormal. Primary hyperparathyroidism was detected in 5 (9%) patients. These patients were treated by surgically. Urine volumes lower than 2000 ml were accepted as a risk factor and were detected in 6 (13%) patients. These cases were treated by adequate fluid intake to have a daily urine volume of 2000 ml. Urine volumes were turned to normal in 3 patients (50%). New stone formation occurred in 1 patient (Table 1 & 2).

![Figure 1](image_url)

*Figure 1: The types of the stones revealed by stone analysis.*

Com: calcium oxalate monohydrate  
Cod: calcium oxalate dehydrate  
Ua: uric acid

**Table 1:** The results of simple metabolic evaluation in 53 patients who had routine follow up for 5 years.

| Pathology                      | Sole (%) | Mixed (%) | Total (%) |
|-------------------------------|----------|-----------|-----------|
| Hypercalcuiura                | 6 (11)   | 20 (37)   | 26 (49)   |
| Hyperoxaluria                 | 0        | 17 (32)   | 17 (32)   |
| Gout Diathesis                | 0        | 2 (4)     | 2 (4)     |
| Hyperuricemia                 | 0        | 1 (2)     | 1 (2)     |
| Hyperuricosuria               | 0        | 2 (4)     | 2 (4)     |
| Primary hyperparathyroidism   | 1 (2)    | 4 (8)     | 5 (9)     |
| Low urine volume              | 2 (4)    | 4 (8)     | 6 (13)    |
| Other*                        | 0        | 1 (2)     | 1 (2)     |
| Normal*: Low urine ph, cystinuria, etc. | 7 (13) | 0 | 7 (13) |

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Table 2: Results after the treatment in 53 patients who had routine follow up for 5 years.

| Parameter            | Improved (n) | Impaired (n) | New Occurrence (n) | No Change (n) |
|----------------------|--------------|--------------|--------------------|---------------|
| Hypocitraturia       | 13           | 1            | 8                  | 10            |
| Hypercalciuria       | 15           | 2            | 2                  | 7             |
| Hyperoxaluria        | 7            | 1            | 7                  | 2             |
| Gout Diathesis       | 2            | -            | -                  | -             |
| Hyperuricosuria      | 2            | -            | -                  | -             |
| Hyperuricemia        | -            | -            | -                  | -             |
| Hyperparathyroidism  | 3            | -            | 2                  | -             |
| Low Urinary Volume   | 3            | -            | 1                  | 2             |
| Other                | 1            | -            | -                  | -             |
| Normal               | -            | 5            | -                  | 2             |

Discussion

Nephrolithiasis is affecting 1-5% of the population in industrialized countries [7]. Most of these stones could be treated by SWL or endoscopic surgery (URS, PNL). However, high recurrence rates are still problem for either the patient or the clinician [4]. Therefore, metabolic evaluation is a crucial step in the treatment of stone disease. Detailed history focusing on occupation, previous use of medications, dietary and lifestyle habits, recurrent urinary tract infections and underlying disorders that can predispose to stone formation should be examined. History and physical examination are the cornerstones in evaluation of stone formers [8]. The patients should be classified as high (PHR) or low risk for recurrence (PLR). In the next step, SE or comprehensive evaluation (CE) are performed before preventive medical treatment.

Some authors stated that the stone recurrence rate was also important. Despite the high remission rates of medical prophylaxis in calcium stone formers (SF), medical prevention has been mentioned to be non cost effective in patients forming stones less frequently than once every 3 years [7,9,10]. In the present study stone formation period less than one in 3 years were excluded. Simple metabolic evaluation is recommended in patients with low risk for stone recurrence.

However, it is debate whether a single 24-hour urine collection is enough for metabolic evaluation, or not. Some authors stated that only one urine specimen was not enough in metabolic evaluation to reflect the pathology [11,12]. One should be collected under random diet and the other after restriction. This may be useful espesially for classifying the subtypes of hyper calciuria. On the other hand, Pak et al supports a single 24 hour urine collection [13]. In the study, we checked the sample of urine collected in 24 hours urination with standard diet. However, distinct from the other studies, we evaluated patients with high risk for stone recurrence. Preservation of urine seemed to be difficult by the method performed in the study, since we had only one urine sample to investigate. Ferraz et al concluded that post delivery acidification or alkalinization of urine samples did not modify the measured levels of urinary OX, Ca, Mg, creatinine and UA and change on CIT was not relevant allowing all parameters to be determined in a single urine sample [14]. We measured urinary volume, pH in the 24 hour urine sample, and the half part of the sample was acidified for OX, Ca, Mg and CIT calculation and the other half part was alkalanized for UA determination as Ferraz et al recommended [14].

During 5 years, approximately 80% of the patients who did not continue their visits were excluded from the study. When these patients were called up for visits, they refused since most of them considered the process as hopeless, time consuming, bothersome and expensive. They were acquainted about stone recurrence rates, complications and importance of metabolic evaluation in stone disease again, only 10% accepted to continue their routine follow ups. This point out the importance of patient compliance. In our opinion, the compliance of the patient is the most essential factor and it might be less than 80% if the patient had to collect more than one 24 hours urine sample. Also, dietary restriction would help to reduce the participation.

In a recent study it is stated that CE is superior to SE (a single 24 hour urine collection) neither in cost nor in efficacy [15]. We did not investigate the cost effectiveness, but Lotan et al. [15] estimated that only small differences between empiric and conservative therapy were demonstrated in recurrent stone formers in countries such as Turkey and Switzerland. This suggestion is very important when informing the patients about the cost effectivity. It is stated that with appropriate evaluation, a cause for stone formation can be found in 97% of the patients [16]. We determined abnormality in almost 90% of the patients by SE. This is close to the rates determined by CE. When it is thought that 215 patients were out of follow up in the present study, it may be suggested that this rate might be much more than it has been estimated. Stone analysis is warranted in all patients and may provide guidance for the therapeutic treatment. Kourambas et al. [17] stated that non calcareous stone forming patients reflected specific metabolic disorders. The authors suggested that compositional stone analysis should be an integral part of the metabolic evaluation of patients with nephrolithiasis and stone analysis alone may provide guidance for therapeutic treatment and obviate a formal metabolic evaluation. In our study the patients whose stone analysis could not be detected, were excluded.

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Metabolic evaluation and preventive medical therapy are supplementary steps in the treatment of urolithiasis and the patient’s compliance is an essential part of the process. Even in the first visit, SE may enlighten the pathology as high as 87%. However, the participation may be as low as 25% during the follow up. Why do the patients get out of the follow up? The main reason might be the opinion to have undergone an evaluation that is expensive, time-consuming, bothersome and useless. Before treatment, all of the patients must be informed about their disease, risks and the necessity of the evaluation. We conclude that, although SE seems to be safe, inexpensive and easy to perform in the evaluation of PHR, the patients’ compliance may be the restrictive parameter for the success. However, SE may be very useful for the clinician to determine the pathology even in the first visit.

References

1. Lotan Y, Cadeddu JA, Pearle MS (2005) International comparison of cost effectiveness of medical management for nephrolithiasis. Urol Res 33(3): 223-230.
2. Cohen TD, Ehreth J, King LR, Preminger GM (1996) Pediatric urolithiasis: Medical and surgical management. Urology 47(3): 292-305.
3. Sutherland J, Parks JH, Coe F (1985) Recurrence after a single renal stone in a community practice. Miner Electrolyte Metab 11(4): 267-269.
4. Trinchieri A, Ostini F, Nespoli R, Rovera F, Montanari E, et al. (1999) A prospective study of recurrence rate and risk factors for recurrence after a first renal stone. J Urol 162(1): 27-30.
5. Heilberg IP, Schor N (2006) Renal stone disease: Causes, evaluation and medical treatment. Arq Bras Endocrinol Metabol 50(4): 823-831.
6. Pak CY, Britton F, Peterson R, Ward D, Northcutt C, et al. (1980) Ambulatory evaluation of nephrolithiasis. Classification, clinical presentation and diagnostic criteria. Am J Med 69(1): 19-30.
7. Debevecio FC, Preminger GM (2003) Medical management of stone disease. Curr Opin Urol 13(3): 229-233.
8. Heilberg IP, Schor N (2006) Renal Stone Disease: Causes, Evaluation and Medical Treatment. Arq Bras Endocrinol Metabol 50(4): 823-831.
9. Chandhoke PS (2002) When is medical prophylaxis cost-effective for recurrent calcium stones? J Endourol 16(3): 937-940.
10. Pearle MS, Roehrborn CG, Pak CY (1999) Meta-analysis of randomized trials for medical prevention of calcium oxalate nephrolithiasis. J Endourol 13(9): 679-685.
11. Parks JH, Goldfischer E, Asplin JR, Coe FL (2002) A single 24-hour urine collection is inadequate for the medical evaluation of nephrolithiasis. J Urol 167(4): 1607-1612.
12. Yagisawa T, Chandhoke PS, Fan J (1999) Comparison of comprehensive and limited metabolic evaluations in the treatment of patients with recurrent calcium urolithiasis. J Urol 161(5): 1449-1452.
13. Pak CY, Peterson R, Poindexter JR (2001) Adequacy of a single stone risk analysis in the medical evaluation of urolithiasis. J Urol 165(2): 378-381.
14. Ferraz RR, Baxmann AG, Ferreira LG, Nishiura JL, Siliano PR, et al. (2006) Preservation of urine samples for metabolic evaluation of stone-forming patients. Urol Res 34(5): 329-337.
15. Lotan Y, Cadeddu JA, Roehrborn CG, Pak CY, Pearle MS (2004) Cost-effectiveness of medical management strategies for nephrolithiasis. J Urol 172(6 Pt 1): 2275-2281.
16. Pak CY (1997) Southwestern Internal Medicine Conference: medical management of nephrolithiasis: A new simplified approach for general practice. Am J Med Sci 313(4): 215-219.
17. Kourambas J, Aslan P, Teh CI, Mathias BJ, Preminger GM (2001) Role of stone analysis in metabolic evaluation and medical treatment of nephrolithiasis. J Endourol 15(2): 181-186.