



## Risk Factor for Incident Renal Stone: A Tertiary Centre Experience

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### Abstract

*Renal stone continues to be a leading cause of chronic kidney disease. Renal stone recurs at high variable rate. There are certain predisposing factors identified for formation of kidney stones. In this retrospective study we identify the risk factor for incident renal stone formation in our local population. Hypocitraturia was the most common predisposing factor and was seen in 95 percent of the patient. The hypocitraturia did not normalize with the standard dosage of citrate supplementation in majority of the patient.*

### Introduction

Renal stone and its complication is a significant cause of morbidity and mortality in the general population. Once a renal stone is formed they tend to recur in a sizable proportion of patient. It is important to identify the risk factor for stone formation so that the risk of recurrence is reduced. In this study we attempt to find the predisposing risk factor for renal stone in our local population.

### Material and Methods

This is a retrospective study with data compiled from a large tertiary government hospital in Imphal.

**Inclusion Criteria:** Adult patients with newly diagnosed renal stone were included in this study. Patients were recruited from Nephrology department of Jawaharlal Nehru Institute of Medical sciences which serve as a referral centre. The study period was from February 2014 till February 2019. The patients were subjected to the following test as a work up for renal stone:

Urine examination to look for pH. Urine microscopic examination. 24 hour urine estimation of urine citrate /uric acid/calcium/oxalate. Kidney function test, Complete haemogram, intact parathyroid hormone, serum bicarbonate were done in all the patients. Nuclear scan of parathyroid gland was performed in suspected case of primary hyperparathyroidism. Stone analysis was done whenever the stone was retrieved.

Hypercalciuria is defined as a 24 hour urine calcium values exceeding more than 300 mg in males and 250 mg in females.

Hyperoxaluria is defined as a 24 hr urine oxalate values exceeding more than 40 mg.

Hyperuricosuria is defined as a 24 hour urine uric acid more than 800 mg in male and 750 mg in females.

Hypocitraturia is defined as a 24 hour urine citrate amount less than 320 mg.

**Exclusion Criteria:** Patients with deranged renal function (serum creatinine more than 1.5 mg/dl)

were excluded. Patients with polycystic kidney disease or those with significant congenital urinary tract anomalies and inborn errors of metabolism were excluded.

### Statistical Analysis

Continuous variables are expressed as means. SPSS 16 statistical software was used in the data analysis.

### Results

102 patients with newly diagnosed renal stone disease were included during the study period .24

**Table 2:** Baseline characteristic of the risk factor

	Number of patients	Mean value
Intact parathyroid hormone level		22.2+/-28.6
Number of patients who had parathyroid level greater than upper limit	4	
Confirmed hyperparathyroidism on nuclear scan	1	
Distal renal tubular acidosis	None	
Hyperoxaluria (no. of patients)	10 /80	27.5 +/- 15.3
Hypercalciuria	5 /80	135.05 +/- 80.4
Hypocitraturia	76/80	47.6 +/-32.11
Hyperuricosuria	2/80	381.4 +/- 141.8
Stone analysis :12 patients	Calcium oxalate 10 Mixed stone 1 Uric acid 1	

### Discussion

Renal stone and its related complication is the second most common cause of chronic kidney disease in our local population. Renal stone can cause obstructive nephropathy with renal failure. It contributes to urosepsis and can also cause chronic tubulointerstitial nephritis .Renal stone have also been associated with cardiovascular disease and bone disease<sup>(1,2)</sup>.

Renal stones are known to recur at a high rate<sup>(3)</sup>. The cause of recurrence is because the primary predisposing factor remained uncorrected .As such it is important to identify the risk factor in order to reduce the risk of recurrence .Once the risk factors are identified then proper corrective measures can be adopted.

In our study population, the most common risk factor for stone formation was hypocitraturia which was noted in 95 percent of (76 /80) the patient. The mean urinary citrate level was 47.6 +/- 32.11. Urinary citrate complexes with calcium

hour urine biochemistry results were available in 80 patients. Stone analysis results were available in 12 patients .The findings are as shown in table 2.

**Table 1** Baseline characteristics of the patient

Age	45.4+/-16.33
Sex(M/F)	74/28
Dietary pattern	
Non vegetarian diet	88
Vegetarian diet	14
Serum calcium (corrected for albumin)	9.13+/-0.13
Serum creatinine	1.12 +/- 0.17

and prevent calcium super saturation as well as nucleation of calcium oxalate and calcium phosphate complexes. Approximately 65-90% of filtered load of citrate is reabsorbed<sup>(4,5)</sup>. Citrate are absorbed through the apical membrane from proximal renal tubules which is believed to be pH dependent. The tubular citrate exists mostly as citrate -3 ion although it is reabsorbed as citrate -2 ion through the sodium citrate co transporter. With decreasing tubular pH, citrate reabsorption increases<sup>(6)</sup>.

The most common cause for hypocitraturia is idiopathic. Other factors include distal renal tubular acidosis<sup>(7,8)</sup>, chronic diarrhoea, gastrointestinal disorders<sup>(9,10)</sup> and high animal protein intake<sup>(11,12)</sup>. Certain medications are also known to affect urinary citrate excretion<sup>(13,14,15)</sup>. In our population we could not find any case of obvious distal renal tubular acidosis. Patients were not subjected to evaluation for incomplete distal renal tubular acidosis .In the absence of any other

obvious predisposing factor they are presumed to be idiopathic.

The patients received potassium citrate and potassium magnesium citrate supplementation after diagnosis of hypocitraturia. Even after 3 months of citrate supplementation (around 40-80 meq) along with dietary restrictions, it was difficult to correct the hypocitraturia. Only 7 of the 71 patients who had undergone a repeat urinary citrate level could achieve the normal desired range.

Hyperoxaluria was noted in 10 patients .4 of these patients also had concomitant hypocitraturia .In the absence of any renal failure , history of prior bowel rejection and other congenital diseases, Hyperoxaluria is believed to be secondary to increased oxalate consumptions.

Hypercalciuria was noted in 5 patients. However primary hyperparathyroidism was diagnosed in only one patient.

The most common stone was oxalate stone which comprised more than 83% in our study population. Uric acid stone is rare in our population and seen in only one patient.

Our study is limited by its retrospective nature. Secondly we do not have detailed information about the etiology of the hypocitraturia /hyperoxaluria. Detailed work up into the etiology is warranted.

Thus to summarize, in this study we noted that hypocitraturia is the most common predisposing factor for patients with incident renal stone. The hypocitraturia is mostly idiopathic in nature and does not improve even with standard treatment suggesting that the standard citrate supplementation may not be able to correct the biochemical abnormalities. Isolated hyperoxaluria and hypercalciuria are uncommon predisposing factors in our study population.

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