A detailed diagnostic work-up for renal calculi is beyond the scope of this presentation. Therefore, this presentation is related mainly to calcium oxalate (CaOx) stone formation.

**Diagnosis of nephrolithiasis**

Very briefly, conditions such as genetically acquired diseases (e.g. cystinuria, Dent's disease, renal tubular acidosis (RTA), primary hyperparathyroidism, granulomatous hypercalcemia or, especially, hypercalcuria, autosomal dominant polycystic kidney disease (ADPKD) associated with kidney stones should be referred to the relevant discipline (e.g. urologist, endocrinologist, etc.).

The first principle is to know when a patient with CaOx stone disease needs a detailed work-up. Therefore, exclusion and inclusion criteria have been formulated.

**Exclusion criteria**

A single (first) attack of CaOx stone disease. Patients with uric acid calculi are also excluded, as are those with calculi associated with infection and/or obstruction.

**Inclusion criteria**

The first clinically symptomatic attack of kidney stone disease is included in the metabolic work-up if the radiological investigation depicts multiple stone disease. Patients <19 years of age, pilots, fire-workers or armed forces personnel who present with a first stone all require a full metabolic work-up. The same applies to persons with a single kidney, and a first stone in obese patients with diabetes and metabolic syndrome.

The management of renal stones in pregnancy requires specific knowledge of how to handle the situation, and is well dealt with in the review by Meher et al.[1] on renal stones in pregnancy. As the urine volume and mineral content vary vastly during pregnancy, a pathological urinary work-up is contraindicated. In fact, very little work has been done on this subject, and our personal advice is to recommend that the work-up be delayed until 6 months post partum. Of note, 70% of calcium stones formed during pregnancy consist of calcium phosphate (CaP). These are non-infective in origin, and mandate a full stone work-up post partum. Idiopathic stones consisting of calcium and phosphorus are much more likely to affect kidney function and occasionally result in chronic kidney disease.[2] General management and kidney stones in pregnancy is shown in Table 1.

Point 8 in the table recommends that pregnant patients take an appropriate dose of K-Cit. Although citrate inhibits the crystallisation of CaP, it also alkalinises the urine, which could do more harm than good by resulting in CaP stone formation, and is therefore not recommended. Thus the main therapy in pregnancy is a very large fluid intake, which will aid spontaneous stone passage. About 70% of all calcium stones are passed spontaneously during pregnancy owing to the presence of marked urinary tract dilatation.[3]

**Work-up in secondary causes of CaOx nephrolithiasis**

Although not in the scope of this article, secondary causes are briefly mentioned for the sake of clarity. They include the exclusion of primary hyperparathyroidism, the granulomatous hypercalcemias, drug-induced RTA, recurrent CaOx stone formation in patients with ADPKD[4] and stones associated with inflammatory bowel disorders.

**Stone work-up protocols**

- Routine blood tests (Table 2)
- Routine diagnostic radiology (ultrasound or computed tomography scan)
- Patient information – what to discuss with the patient – see
Table 1. Nephrolithiasis during pregnancy: Eight essential guideline points

1. As far as is possible, pregnancy should be planned in patients with RCaOx stone formation.
2. These patients should, if required by their attending urologist, be stone-free because of the seriousness of possible stone passage during pregnancy.
3. If patients are on either maintenance therapy on thiazides or indapamide, these should be avoided until the postpartum period.
4. If on allopurinol, this agent should be discontinued as some animal studies have indicated fetal toxicity.[3,4]
5. Urinary tract infection is common in pregnancy in patients with recurrent nephrolithiasis and must be carefully monitored and managed.
6. The care team should include, in addition to an informed obstetrician, both the attending urologist and kidney stone nephrologist, if possible.
7. Antenatal visits are guided by the clinical situation but must be more frequent in patients with stones, especially if they have the metabolic syndrome, or are diabetic or have secondary causes of recurrent CaOx stone formation.
8. Of paramount importance is the necessity to always remain well hydrated, aiming at 2.5 L. (at least) of water per day. Although not studied, it would seem desirable that pregnant women should either commence with or remain on a three times per day appropriate dose of K-Cit with careful monitoring of the urinary pH.

RCaOx = recurring calcium oxalate; CaOx = calcium oxalate.
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Table 2. Routine blood tests*

<table>
<thead>
<tr>
<th>Test</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Full blood count</td>
<td>Platelets</td>
</tr>
<tr>
<td>Erythrocyte sedimentation rate</td>
<td>C-reactive protein</td>
</tr>
<tr>
<td>Albumin</td>
<td>Urea</td>
</tr>
<tr>
<td>Serum creatinine</td>
<td>Electrolytes</td>
</tr>
<tr>
<td>Estimated glomerular filtration rate</td>
<td>Calcium</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>Parathyroid hormone</td>
</tr>
<tr>
<td>Aspartate aminotransferase</td>
<td>Gamma glutamyl transferase</td>
</tr>
<tr>
<td>Fasting lipid profile</td>
<td>Fasting blood sugar (glucose)</td>
</tr>
<tr>
<td>Thyroid-stimulating hormone</td>
<td>Vitamin D (25(OH)D (25-hydroxy vitamin D))</td>
</tr>
<tr>
<td>Uric acid</td>
<td>Serum magnesium (not commonly done in South Africa)</td>
</tr>
<tr>
<td>Arterial blood gas (only if required)</td>
<td>Other tests (to be specified on request)</td>
</tr>
</tbody>
</table>

*Tests included are routine blood tests for urolithiasis management (both diagnostic and follow-up).
Reproduced with permission from Meyers.[3,4]


- The work-up in a patient with any form of RTA should be done by a nephrologist
- Routine 24-hour urine tests – see supplementary material: block 3 on patient instructions (http://samj.org.za/public/sup/15992.pdf) as well as vital instructions for all chemical pathology laboratories.

There is one more pivotal factor to stress. Neither the doctor nor the dietician should give the patient any stone management advice until the total work-up is complete and the patient has returned for his or her follow-up. Although the reasons are obvious, it is worthwhile to stress them further. If, at the first visit, advice is given to increase fluid intake or to start dietary manipulation (especially on salt intake), the result of the tests will be falsely skewed, making accurate assessment impossible. Important information to be discussed with the patient is included in the supplementary material.

Management and prevention of stones
Extracorporeal shock wave lithotripsy (ESWL), lithotripsy and surgical interventions are beyond the scope of this article. Other rarer conditions such as management of cystinuria or primary hyperoxaluria (i.e. genetic) and other rare non-genetic disorders of stone formation will be briefly mentioned.

Lifestyle factors contributing to stone formation
- Fluid intake: this must be aimed at 2.2 - 2.5 L per day. Apart from water, lemon juice, coffee and herbal or rooibos tea are fluids of choice. No Ceylon tea is allowed. The most valuable study ever published on stone prevention addressed this simple remedy as recently as 1996.[9] The frequency of non-adherence and reinforcement of adequate hydration must continue indefinitely.
- Dietary assessment and advice: this is presented in more detail in the appendix. Apart from the assessment of a skilled dietician, the doctor should also become acquainted with the patient's food and eating habits.
- Of importance is to train the 'stone dieticians' and attending medical staff never to give advice to the patient until all the tests and work-up are complete. This also dictates that the doctor will have seen the results of the tests, which will then be discussed with the patient. Failure to carry out these instructions will be the cause of many errors. An important example of this is the assessment of fluid intake. If the patient is instructed by the doctor or dietician to increase fluid intake or to avoid certain foods, the assessment will be false. The initial two to three 24-hour urine work-up assessments yield vital information of the true fluid balance between intake and output.
- In obese patients, weight loss and exercise must be repeatedly advised by the dietician, making follow-up assessments essential.

Medications
- The use of K-Cit: in practice, it is common to come across stone patients who are advised by their physicians to go onto citrate therapy without justification. Although hypocitraturia is common, about 50% of patients will not have this condition, and therefore urinary alkalisation with all its attending hazards could well occur.[2,3] Another common problem seen in practice is that other urinary alkalisers have potential problems. The commercial products Uroline U, Citro-Soda and sodium bicarbonate all contain significant quantities of sodium.[7] In stone formers, this increased quantity of sodium will result in increased sodium excretion, increasing the theoretical chance of treatment failure with new stone formation. Simple K-Cit is the treatment of choice.[8,9] and is inexpensive. The downside of K-Cit ingestion is the bad taste, which can be eliminated by diluting.
Table 3. Urinary oxalate excretion (μmol/24 hour) after 'standard' diets

<table>
<thead>
<tr>
<th>Variable</th>
<th>Black</th>
<th>White</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Female</td>
<td>Male (study 1)*</td>
</tr>
<tr>
<td>n</td>
<td>19</td>
<td>12</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>244 (127)</td>
<td>239 (135)</td>
</tr>
</tbody>
</table>

SD = standard deviation.
*Columns represent results from two separate studies.
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Table 4. Clinical points to remember

<table>
<thead>
<tr>
<th>Clinical aspect</th>
<th>Points to remember/interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>RCaOx stone presentation</td>
<td>Citrate is not the panacea for all CaOx or CaP nephrolithiasis treatment.</td>
</tr>
<tr>
<td>Total body of Mg deficiency states</td>
<td>Subclinical magnesium deficiency states are dangerous and found more in children. Also found in patients on chronic loop diuretics.</td>
</tr>
<tr>
<td>Use of allopurinol</td>
<td>This is safe and even important in CaOx stone formation in normouricosuria patients.</td>
</tr>
<tr>
<td>CT scan diagnosis and endo-uroscopic stone removal procedures</td>
<td>Remember that Randall's plaques are not stones. Patient safety is of paramount importance.</td>
</tr>
<tr>
<td>Remember the host of secondary and the large list of drug causes of nephrolithiasis</td>
<td>Varied pathogenesis and treatment. Recognition of and treatment of cause and/or discontinuation of offending drug.</td>
</tr>
<tr>
<td>High salt intake</td>
<td>Of paramount importance in causing RCaOxSF and can result in serious osteoporosis. Can only be diagnosed with 24-hour urine examination.</td>
</tr>
<tr>
<td>The most important dietary advice to all stone formers (and everyone else)</td>
<td>Eat a little of everything and a lot of nothing. Note: When on a weight-reducing diet, this does not apply and only kicks in as maintenance when the target weight has been reached.</td>
</tr>
<tr>
<td>General scientific advice behind the overall dietary pattern</td>
<td>Diet imposes patterns of metabolism upon the organism. 57</td>
</tr>
</tbody>
</table>

RCaOx = recurring calcium oxalate; CaOx = calcium oxalate; CaP = calcium phosphate; Mg = magnesium; CT = computed tomography; RCaOxSF = recurring calcium oxalate stone formation.
Reproduced with permission from Meyers. 34

10 -15 mL of the solution in a glass of water, taken three times a day (not less), which will also help ensure adequate hydration.

- Hypomagnesaemia/hypomagnesuria: although magnesium is an important inhibitor of stone formation, there is no clinical or experimental basis to recommend it in the absence of low magnesium levels. Most cases of low magnesium are found in children, or the very occasional adult. The most common magnesium tablet used in South Africa (Slow-Mag) is contraindicated in stone therapy because it consists of magnesium chloride. Magnesium oxide or magnesium hydroxide are the medications of choice for both efficacy and cost-effectiveness. 34

- Hypernatrituria: at the time of writing, the ‘true’ role of excess dietary salt has not yet been adequately assessed. If patients were compliant enough to decrease their dietary salt intake so as to ensure 80 -100 mg of daily sodium loss, most patients with so-called ‘idiopathic hypercalciuria’ might well have completely normal urinary biochemistry and would cease the formation of new CaOx calculi. However, this degree of long-term dietary salt restriction is probably nothing but a pipe dream, because lifestyle changes with salt restriction are so difficult to attain. The treatment of other causes of the hypercalciurias is dealt with below.

- Hypercalciurias: there are three subtypes of hypercalciuria. (i) True (primary) renal hypercalciuria – a very rare disease. Some of these patients may be associated with urinary phosphate excretion changes, resulting in a raised level of 1.25 dihydroxy vitamin D3 and stone formation. (ii) Salt-induced hypercalciuria. (iii) Secondary hypercalciuria associated with hyperparathyroidism, sarcoidosis, other granulomas, RTA and vitamin D excess. Secondary hypercalciuria patients are managed according to the underlying cause. Patients falling into primary and salt-induced hypercalciuria are usually treated with thiazide diuretics. In fact, thiazide diuretics are completely contraindicated, which has been clearly demonstrated in previous publications. 10,11

The doses of thiazide required for the treatment of hypercalciuria (25 - 50 mg per day) are in the toxic range, and their hypocalciuric effect disappears after 6 months to 1 year of therapy. The treatment of choice in these patients is indapamide, and the urinary calcium-lowering effects continue indefinitely. 11,12

- Urinary oxalate excretion: The present laboratory normal 24-hour urinary oxalate excretion range is totally inaccurate. This criticism applies especially to the so-called mild metabolic hyperoxalurias (MMHO). Standard diets are required before MMHO can be verified. In addition, laboratories should report the results, as the excretion of urine oxalate in μmol per 24 hours and not μmol per litre. 13,14

Urinary oxalate excretion has been measured in previous studies. 13,14 In summary, the normal ranges that were found in these studies are shown in Table 3.

There are two more points to be made. Firstly, there was no difference in the excretion of oxalate between males and females, or between black and white patients. Secondly, because oxalate is so ubiquitous and is found in most foodstuffs (particularly red meat), the practical normal excretion on a low oxalate diet has been regarded as 350 μmol per 24 hours. If the dietary standardisation is accurate and the oxalates are above this level, the two methods used in treatment are to increase the dietary calcium ingestion (which binds to oxalate and prevents absorption), or to add pyridoxine, which is only effective in a small minority. 15

However, causes of secondary (i.e. acquired) hyperoxaluria, such as is seen in inflammatory bowel diseases or after barosurgery,
are significantly higher than the normal range (600 - 1 500 μmol per 24 hours). In patients with hereditary oxalosis, the levels are considerably higher than the above, and reach as high as 3 000 μmol per day.

Patients with kidney stone disease are usually worked-up using a ‘best practice available’ basis. They have one 24-hour urine work-up, urine cultures where necessary, relevant blood tests and radiology and, if the stone does not pass spontaneously after fluid load, extracorporeal shockwave lithotripsy or laser therapy are considered. For long-term maintenance, patients are normally referred to a general practitioner. In spite of the skills and good intentions of the attending physicians, the above mode of therapy is not in the best interest of the patient, nor is it cost-effective. However, any long-term and detailed stone management programme is very difficult to conceive without the services of a multidisciplinary team.

In summary, the main aims of this article have been to educate all interested personnel on what could and should be done for patients with recurrent renal calculi, i.e. the ‘office’ approach. The most important clinical points to remember are shown in Table 4, and the most important management aspects are shown in the included appendices.

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