The true significance of urine sediment findings from in-house urinalysis needs to be considered in broader context with other key clinical information, such as patient history, examination results, and findings on diagnostic imaging.

The following cases demonstrate the importance of evaluating the total clinical picture and explore how preparing dry-mount urine sediment cytology can maximize the potential of in-house urinalysis. Dry-mount cytology is a quick and relatively easy technique useful not only for evaluating conditions such as urinary tract infection (UTI) or neoplasia but when seeking a pathologist’s second opinion regarding cells found in urine sediment. Cells prepared for dry-mount urine sediment cytology are preserved indefinitely and, therefore, are not subject to degradation during shipment to an outside laboratory.
A previously healthy castrated German shepherd dog (5 years of age) presented after being hit by a car. As part of the diagnostic workup, urinalysis was performed (Table; Figures 1 and 2).

Diagnosis
Incidental Crystalluria

Interpretation & Discussion
Incidental crystalluria from amorphous phosphate and struvite crystals was present in this patient. Based on cytology alone, amorphous crystals could be either phosphate or urate; further urinalysis information is required for specific identification. Because the crystals are colorless and the patient’s urine is mildly alkaline, these are amorphous phosphate crystals. Amorphous urate crystals are less common, tend to form in acidic urine, and are yellow or yellow–brown. Although amorphous crystals may be seen with various conditions, their presence here is a nonspecific finding and does not typically augment the clinical picture.

Struvite crystalluria may be seen in clinically normal patients, patients with struvite or mixed-content uroliths, or patients with UTI. Struvite crystals can also develop during refrigerated storage of urine samples. In this case, clinical context is essential when determining the significance of the patient’s struvite crystalluria. This dog did not exhibit lower urinary tract signs. Abdominal radiography to detect injuries related to trauma revealed no radiopaque cystic calculi. Leukocytes were not observed in the urine sediment, indicating that lower urinary tract inflammation and, therefore, infection or presence of uroliths were unlikely.

The 1+ heme reaction was secondary to mild hematuria noted in the wet mount and may have been secondary to cystocentesis or blunt trauma. The presence of erythrocytes, hemoglobin,
CASE 1: ACUTE TRAUMA

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
<th>Reference Interval or Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Color</td>
<td>Yellow</td>
<td>Yellow</td>
<td>Brown*</td>
<td>Yellow to amber</td>
</tr>
<tr>
<td>Clarity</td>
<td>Cloudy*</td>
<td>Hazy*</td>
<td>Cloudy*</td>
<td>Clear or slightly hazy</td>
</tr>
<tr>
<td>Specific gravity</td>
<td>1.055</td>
<td>1.019*</td>
<td>1.033</td>
<td>1.020–1.050 (range, 1.001–1.070)</td>
</tr>
<tr>
<td>pH</td>
<td>7.50</td>
<td>6.00</td>
<td>8.50*</td>
<td>5.50–7.50</td>
</tr>
<tr>
<td>Dipstick protein</td>
<td>2+*</td>
<td>Trace*</td>
<td>4+*</td>
<td>Negative to trace, depending on USG</td>
</tr>
<tr>
<td>Glucose</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Ketones</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Bilirubin</td>
<td>1+</td>
<td>Negative</td>
<td>1+</td>
<td>Negative to 1+, depending on USG</td>
</tr>
<tr>
<td>Heme</td>
<td>1+*</td>
<td>2+*</td>
<td>3+*</td>
<td>Negative</td>
</tr>
</tbody>
</table>

* Values outside reference interval or value

USG = urine-specific gravity

Urinalysis via Cystocentesis: Physical and Chemistry Results*

and myoglobin may also contribute to the heme reaction. Given this patient’s history, muscle damage and myoglobinuria could have contributed to the positive heme reaction. Multiple factors likely contributed to the 2+ proteinuria, including urine concentration, the presence of heme protein, and possibly other factors related to recent trauma or occult disease. Also present was 1+ bilirubin, which is a normal finding in concentrated canine urine; this is not a normal finding in some other species.

Struvite crystalluria was incidental and may have been secondary to the highly concentrated urine or possibly refrigerated storage, as handling factors can promote precipitation of crystals in urine.¹

Did You Answer?

- Amorphous phosphate and struvite (also known as magnesium ammonium phosphate; Figure 2)
- Urine pH is helpful when identifying amorphous crystals; amorphous phosphate crystals occur in alkaline urine, whereas amorphous urate crystals occur in acidic urine.
- Amorphous phosphate crystals typically have little clinical significance. Struvite crystalluria may be seen in clinically normal patients, patients with struvite or mixed-content uroliths, and patients with UTI. Struvite crystals may also form with refrigerated storage of urine samples.
CASE 2: UROLITHIASIS & PROSTATITIS

A castrated English bulldog (4 years of age) presented for pollakiuria. The dog’s history included urolithiasis, cystotomy, and prescrotal urethrostomy 8 months before presentation. Multiple surgical revisions of the poorly healing urethrostomy had been performed, and the patient was receiving cephalixin. Palpation revealed urinary bladder distention and prostatomegaly. Diagnostic evaluation included urinalysis (Table, page 85), imaging, wet-mountain cytology (Figures 3–6), and prostatic fine-needle aspiration (FNA).

Diagnosis
Urolithiasis, Prostatitis, and UTI with Secondary Struvite Crystalluria

Interpretation & Discussion
The urine sediment had increased leukocytes (ie, pyuria) with struvite crystalluria. These findings alone would raise concern about urinary tract inflammation or infection and potential urolithiasis. Given this dog’s urinalysis findings, extensive history of lower urinary tract disease and current pollakiuria, bladder distention, and prostatomegaly, imaging was performed and revealed recurrent urolithiasis. Ultrasound-guided FNA showed chronic prostatitis, which was likely associated with chronic lower urinary tract inflammation and infection secondary to repeated surgical manipulation. Prostatitis is generally uncommon in neutered dogs.

To evaluate the effectiveness of current antibiotic therapy and determine other treatment options, culture and sensitivity testing was performed. In addition, because partial urethral obstruction was a concern based on the history and physical examination, a chemistry panel was performed to evaluate for azotemia and electrolyte derangements. *Pseudomonas aeruginosa* was cultured from the urine sample, and serum chemistry revealed only mild sodium and chloride abnormalities. Antibiotic treatment was modified based on the sensitivity profile, and cystotomy and scrotal urethrostomy were performed.

A low number of leukocytes (Figure 4) can be seen in normal urine; the expected number is influenced by the collection method. Cystocentesis samples should have fewer than 3 leukocytes/40× high-power field, and those collected by catheterization or midstream voiding should have fewer than 8 leukocytes/40× high-power field.
Considerations for pyuria include primary or secondary bacterial infection, urolithiasis, neoplasia, prostatitis, pyometra, or other less common infections. There were multiple causes of this dog’s pyuria: inflammation associated with poorly healed urethrostomy, chronic prostatitis, urolithiasis, and secondary bacterial infection.

The signs and history of chronic lower urinary tract disease along with severe pyuria and lesser hematuria place this case of struvite crystalluria in dramatic contrast with that observed in the patient in case 1. In addition, this case shows variations of struvite crystal morphology. Many of the crystals occurred as 3-dimensional short rectangles or even squares (Figure 5) rather than the more common long rectangles, prisms, or so-called casket covers. X-shaped struvites (Figure 6) were also rarely seen; they are uncommon and typically occur when urine ammonia concentration is high.

The observation of struvite crystalluria may guide initial therapeutic management of urolithiasis; however, full urolith analysis is necessary because many stones are of mixed mineral content, which was of particular concern for this bulldog given the predisposition of bulldogs for urate urolithiasis. In this case, the trace proteinuria may easily be explained by the presence of inflammation.

### Preparing Urine Sediment Dry Mounts for Cytology

- Following wet-mount preparation, recentrifuge the urine as is done for routine urinalysis (at 400–500 × gravity [~1,000–1,500 rpm]) in a conical centrifuge tube for 5 minutes.
- Use a pipette or gently decant to remove as much supernatant as possible without dislodging the pellet.
- Use a pipette to remove the sediment pellet from the bottom of the test tube.
- Place a small drop of aspirated material on one end of a clean microscope slide.
- Use a second clean microscope slide to spread the material in a monolayer, as would be done for FNA cytology.
- Allow the slide to air-dry. Heat fixation is not necessary and will alter cell morphology.
- Stain as for routine cytology using Diff-Quik or a similar stain. Alternatively, store the slide in a covered container at room temperature for transfer to an outside diagnostic laboratory.

A different microscopic field shows a large struvite crystal surrounded by cells. The crystal is 3-dimensional, 6-sided, and almost square with central intersecting lines. The two irregularly shaped areas of refraction at either end of the crystal are artifacts created by the focal plane. A degraded remnant of a struvite crystal is also evident (right). (unstained wet mount; 100× objective)

A different microscopic field shows another uncommon struvite morphology. An X-shaped struvite is present with more typical 3-dimensional rectangular struvite crystals in different focal planes. (unstained wet mount; 100× objective)
A spayed crossbreed dog (13 years of age) presented for pollakiuria and dysuria of 2 days’ duration. The wet-mount sediment contained many struvites (not shown), leukocytes, erythrocytes, and abundant amorphous debris (Figure 7). It was difficult to confidently identify bacteria in the active sediment.

Dry-mount urine sediment cytology (Figures 8 and 9) was performed to assist with detection of bacteria and accurate description of their morphology (cocci vs bacilli). Urinalysis via cystocentesis also was evaluated (Table, page 85).

**Diagnosis**
UTI with Bacilli

**Interpretation & Discussion**
Struvite crystalluria in this patient with lower urinary tract signs was secondary to a bacterial infection. Infection was suggested by urinalysis findings of alkaline urine, proteinuria, pyuria, and hematuria and was supported by urine sediment dry-mount cytology. Urease-producing bacteria (eg, *Proteus* spp or *Staphylococcus* spp), produce ammonia that increases urine pH. Alkaline pH and high ammonia concentration favor struvite crystal formation.

The urine sediment wet mount was difficult to interpret because of the density of cells and amount of debris. The presence of pyuria elevated concern that the debris-filled background of the wet mount potentially masked bacteria. Dry-mount cytology was performed to better detect bacteriuria (Figures 8 and 9). This technique has been shown to improve the sensitivity and accuracy of sediment evaluation for the presence of bacterial UTI and can be performed easily in general practice.² ⁴ In this case, dry-mount cytology clearly revealed the presence of a monomorphic population of bacilli and numerous neutrophils with a background of lysed nuclear material. Although angular crystals could be seen in the dry-mount preparation, they were better identified and quantified in the wet-mount preparation, where fine detail could be appreciated. Urine sediment dry-mount cytology can be useful for identification of microorganisms and cells but not for identification and enumeration of crystals or renal tubular casts.

In addition to better detection and morphologic description of bacteria, dry-mount cytology can aid in identification and preservation of atypical nucleated cells and unusual microorganisms (eg, yeast, fungi). In any fluid sample, cells can degrade after collection and during transport to an outside laboratory; this is particularly the case for urine samples, which are naturally inhospitable to cells. The dry-mount technique helps avoid...
the challenge of discerning important cytomorphologic details on in-house wet-mount cytology and the frustration of a nondiagnostic result that may be associated with sending a liquid urine sample to an outside laboratory.

Even if a wet mount has already been prepared and stained, the same sample can be converted to dry-mount cytology using the urine sediment that remains in the test tube (see Preparing Urine Sediment Dry Mounts for Cytology, page 87). Urine sediment wet mounts are prepared by suspending the urine sediment pellet in a volume of urine supernatant that is equal to 10% of the sample volume that was centrifuged to make the sediment. A drop of this sediment suspension is used to prepare the wet mount, and the remaining volume is reserved in the test tube. This preparation will be indefinitely stable and can be either stained and evaluated in-house or sent to an outside laboratory for a pathologist’s opinion. With dry-mount cytology, there is no deterioration during transport.

**Did You Answer?**

- Several degenerate neutrophils with monomorphic bacilli (rods) are in the background and are phagocytosed by the neutrophils. These observations indicate bacterial infection.
- The lattice-like purple material in the background is free nuclear material and proteins (ie, nucleoproteinaceous material) from lysed cells.

**Closing Comments**

All three patients presented here had struvite crystalluria, although each had a different clinical presentation. These cases highlight the need to integrate other clinical information with urinalysis data to make optimal decisions about further workup and treatment. The accuracy of urine sediment evaluation can be improved by using dry-mount cytology. ■ cb

See Aids & Resources, back page, for references & suggested reading.