INDICATORS OF POLYURIA AND POLYDIPSIA

- Horses rarely drink more than 5% of their bodyweight daily (25 litres per 500 kg)
- Horses rarely urinate more than 3% of their bodyweight daily (15 litres per 500 kg)
- The only common causes of PUPD are psychogenic polydipsia and PPID

Several logical steps below should lead to a firm diagnosis:

Polydipsia (PD) in adult horses can be defined as water intake >100 ml/kg daily (>10% BWT) although under UK management and environmental conditions it is probable when intake is > 70 ml/kg daily (>7% BWT). Typical water intake for horses is 40 - 60 ml/kg daily (4-6% BWT) although it can be as low as 10-15 ml/kg daily (1-1½% BWT) in grazing horses or as high as 80-90 ml/kg daily (8-9% BWT) in lactating mares, horses in hard work and in hot environmental conditions. Smaller breeds tend to drink relatively more per kg BWT than larger breeds due to the effects of metabolic body size. Polyuria (PU) is usually defined as urine production > 50 ml/kg daily (5% BWT) though in practice it is far harder to measure than polydipsia. Normal urine production is typically between 15 and 30 ml/kg daily (1½ - 3% BWT) and faeces represent the major route of water loss in normal horses.

Before investigating PUPD cases it is important to rule out physiologic explanations such as hot weather, hard work, lactation, excessive dietary protein, excessive salt consumption, administration of glucocorticoids or diuretics

CAUSES OF PU & PD:

- PSYCHOGENIC POLYDIPSIA
- PPID (Cushing’s Disease)
- Chronic Renal Failure
- Hepatic Insufficiency
- Diabetes Mellitus
- Diabetes Insipidus

1. QUANTIFY AND CONFIRM THE PRESENCE OF PD (AND PU?)

Pathophysiologic causes will invariably lead to both PU and PD even though only one and not the other may be recognised and reported by the owner. Therefore, it is both easier and diagnostically acceptable to verify and quantify water intake over a full 24 hour period with the horse stabled.

If water intake is >100 ml/kg/day (>10% BWT) then PD is confirmed (and PU is almost inevitable).

If water intake is 70-100 ml/kg/day (7-10% BWT) then PD may be suspected if there are no apparent physiologic causes (see above).

If water intake is <70 ml/kg/day (<7% BWT) then PD is not confirmed.
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2. INITIAL SCREENING BLOOD AND URINE TESTS

A panel of initial screening tests is worthwhile when investigating PUPD cases comprising the following:

**Blood tests**

**Haematology**

- Anaemia is a common finding with chronic renal failure. Polycythaemia may arise through dehydration suggesting that PU is the primary problem rather than PD (e.g. diabetes insipidus).
- Neutrophilia may indicate a glucocorticoid response or inflammatory disease.

**Basal plasma ACTH or TRH stimulation test** – for PPID (see Endocrine section)

**Creatinine and Urea**

- Very high in chronic renal failure cases: urea is usually >15mmol/L and creatinine >300μmol/L.
- A bit high (e.g urea 8-12mmol/L, creatinine 180-250μmol/L) more commonly indicates dehydration or acute kidney injury but could also suggest early/mild chronic renal failure. Worth comparing with urine creatinine – see below.
- Low values (urea < 4mmol/L and creatinine <75 μmol) may occur in hepatic insufficiency or in cases of primary (psychogenic) polydipsia with washout.

**Glucose**: persistent hyperglycaemia indicates diabetes mellitus – mostly as a consequence of PPID. Short term hyperglycaemia will follow alpha-2 agonist sedatives and also pain/stress.

Hypercalcaemia (total Ca > 3.5mmol/L, ionised Ca > 1.7mmol/L) is often seen in chronic renal failure cases. The major other differential for hypercalcaemia is paraneoplastic disease.

**GGT and AST** can be used to rule-out liver disease.

**Urinalysis**

**Specific gravity (always measure with refractometer and not a dipstick)**

- Low (<1.008, hyposthenuria) suggests that the kidney is actively excreting water and is typical of primary (psychogenic) PD and diabetes insipidus.
- Medium (1.008-1.012, isosthenuria) suggests that the kidney is neither actively concentrating nor diluting the filtrate and is consistent with (but not diagnostic for) chronic renal failure (check serum urea and creatinine).
- High (>1.020, hypersthenuria) is not really compatible with persistent PUPD and confirms renal concentrating ability.
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Glycosuria

- diabetes mellitus (usually caused by PPID)
- Acute stress or transport
- alpha-2 agonists

Comparing urine creatinine to serum creatinine can be useful in cases with mild increases in serum creatinine (e.g. 180-250μmol/L). If increased serum creatinine is due to dehydration then this should be matched by high urine creatinine concentrations (>50 x serum concentration) whereas in chronic renal failure cases will have lower urine creatinine (e.g. <40 x serum creatinine).

3. FURTHER LABORATORY TESTS

On the basis of the above results it should be possible to confirm/rule out most differential diagnoses including PPID, chronic renal failure, diabetes mellitus and liver disease. If these have been ruled out then 2 main differentials remain: primary (psychogenic) polydipsia and diabetes insipidus. Further tests are needed to differentiate these two conditions.

Although a full water deprivation test is possible, it is generally more practical, safer and just as diagnostically useful to perform a partial water restriction test.

**Water Restriction Test**

The purpose of this test is to help differentiate primary (psychogenic) polydipsia from diabetes insipidus, having already ruled out further differentials with the previously described tests. The important diagnostic principle is that psychogenic polydipsia cases CAN concentrate their urine whereas diabetes insipidus cases CANNOT.

Suitable subjects for this test will be systemically well, producing hyposthenuric urine (urine SG<1.008) and have normal serum creatinine and urea. This test MUST NOT be performed on azotaemic horses suspected to have renal compromise. There is no point in performing this test if the urine is already concentrated (SG >1.020).

### Performing a water restriction test

1. Weigh horse (if possible) or make an informed estimate (weigh tape etc)
2. Check serum urea and creatinine are normal (if not, don’t proceed)
3. Take baseline urine sample and measure SG (if > 1.008, don’t proceed)
4. Keep horse stabled and provide water at a rate of 1% bodyweight every 6 hours (i.e. 4% bodyweight per 24h)
5. Check for signs of dehydration, serum urea and creatinine and urinary SG at least every 6 hours (and reweigh if possible)

### The test is terminated when one of the following occurs:

- 72 hours has been reached
- 5% reduction in bodyweight (25 kg per 500 kg)
- clinical signs of dehydration
- creatinine > 200μmol/L or urea >12mmol/L
- urinary SG > 1.020
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Interpretation

- If SG rises above 1.020 this confirms renal concentrating ability is present and therefore rules-out diabetes insipidus. This indicates psychogenic polydipsia.
- If the horse fails to concentrate the urine and urine SG stays <1.020, or the horse becomes dehydrated or loses 5% bodyweight, this suggests diabetes insipidus (often happens by 12 hours).

Further Tests to verify and classify diabetes insipidus

Measure serum vasopressin at end of water restriction period

- Vasopressin > 5 pmol/L – indicates normal vasopressin secretion
- Vasopressin < 5 pmol/L – indicates central diabetes insipidus

Vasopressin response test.

- Inject 0.05 micrograms/kg desmopressin acetate iv and measure urine SG over 24 hours.
  - Urine SG > 1.020 – indicates normal response to vasopressin
  - Urine SG < 1.020 – indicates nephrogenic diabetes insipidus

Diagram:

- Confirm water consumption >10% bwt/d and/or urine production >5% BWT/d
  - Urine SG <1.008
  - No azotaemia

- Urine SG <1.020 after water restriction
  - Diabetes insipidus

- Urine SG >1.020 after water restriction
  - Diabetes insipidus

- Old, hairy laminitis etc. Hyperglycaemia Glycosuria High Plasma ACTH

- CRF

- PPID

- Psychogenic Polydipsia
When catheterised urine samples are obtained, consideration of the effects of sedation is important. Alpha-2 agonists have both acute diuretic and hyperglycaemic effects. The diuretic effect is unlikely to alter the urine sample if collected promptly as a few mLs of most recently produced diuresed urine should not significantly affect the litres of previously accumulated urine in the bladder. However, even a small amount of glycosuria might confuse diagnostic interpretation. Hence acepromazine (5-10 mg/100kg IV) perhaps supplemented by a nose twitch or a small dose of detomidine (0.5 mg/100 kg IV) should be used as necessary in geldings. In mares the use of stocks or a stable door may avoid the need for sedation.

**SPECIFIC GRAVITY**

- SG <1.008 (hyposthenuria) suggests that the kidney is actively excreting water and is typical of primary (psychogenic) polydipsia or diabetes insipidus.
- SG between 1.008-1.014 (isosthenuria) suggests that the kidney is neither actively concentrating nor diluting the filtrate and is consistent with (but not diagnostic for) chronic renal failure (CRF). Although isosthenuria is typical of CRF, other causes of PU/PD might coincidentally happen to fall in the isosthenuric range. Therefore, when isosthenuric samples are obtained, serum urea and creatinine should be checked to rule in or rule out possible CRF.
- SG >1.014 (hypersthenuria) indicates that the kidney is actively concentrating urine although SG >1.020 is usually regarded as more convincing of good concentrating ability (and therefore absence of chronic renal failure or diabetes insipidus – both of which lack concentrating ability).

**GLYCOUSRIA** – if persistent then this indicates diabetes mellitus and usually PPID although primary diabetes mellitus is a rare possibility.

**URINE CREATININE : SERUM CREATININE RATIO** is useful in cases with equivocal mild increases in serum creatinine concentration (e.g. 180-250 mmol/L). Mild to moderate increases in serum creatinine due to dehydration will be expected to be matched by high urine creatinine concentrations (>50 x serum concentration) whereas CRF cases will have lower urine creatinine (e.g. <40 x serum creatinine).

**ENZYMURIA** – urinary GGT, AP and LDH may all be raised in renal failure due to tubular damage (more so in acute renal failure). Urinary GGT is the most commonly assayed urinary enzyme and is usually similar in concentration to serum levels (i.e. <40 iu/L urine). Clearly adjustments should be made for urine concentration and this is done by comparison with urine creatinine concentration (expressed in mmol/L, not μmol/L):

<table>
<thead>
<tr>
<th>URINARY GGT (iu/L)</th>
<th>&lt; 5.0 iu/mmol in normal horses</th>
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</thead>
<tbody>
<tr>
<td>URINARY CREATININE (mmol/L)</td>
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Assessment of the overall body electrolyte status is difficult due to the fact that some electrolytes are primarily intracellular (and serum concentrations only reflect extracellular concentrations) and also that there are several strong homeostatic processes that maintain normal serum concentrations whatever the total body status is. Urinary fractional excretion ratios make use of the fact that many homeostatic mechanisms work by altering renal excretion/reabsorption and this can be detected by comparing serum and urine concentrations.

Calculation of electrolyte fractional excretion rates require simultaneously collected serum and urine samples. It is vital that serum is separated from the cellular fraction promptly (within a few hours) as leakage of potassium and phosphate from dying cells will significantly alter serum electrolyte concentrations. For this test it is assumed (slightly incorrectly!) that the sole route of serum creatinine loss is the urine and that it is neither secreted nor absorbed by the kidney. Thus each electrolyte is compared with creatinine, that is supposedly neither secreted nor absorbed, to give an impression of the renal handling of each electrolyte.

Hence, the clearance of any substance (e.g. an electrolyte, “E”) can be compared to creatinine as a fractional excretion rate (or creatinine clearance ratio) thus:

\[
\text{Fractional electrolyte excretion rate} = \frac{\text{urine output} \times [U_{ELEC}]}{[S_{ELEC}]} \div \frac{\text{urine output} \times [U_{CREAT}]}{[S_{CREAT}]}
\]

\[
= \frac{[U_{ELEC}] 	imes [S_{CREAT}]}{[U_{CREAT}] 	imes [S_{ELEC}]}
\]

A value of 1 (or 100%) suggests similar handling to creatinine – i.e. neither secreted nor absorbed. Values <1 (or <100%) signify active reabsorption of the electrolyte whereas values >1 (or >100%) signify active excretion. Typical normal values comprise:

- **Na**: 0.03 – 0.5% (i.e. 99.5 - 99.97% filtered Na is reabsorbed)
- **K**: 15 - 70% (i.e. 30 - 85% filtered K is reabsorbed)
- **Cl**: 0.2 – 1.7% (i.e. 99.8 - 98.3% filtered Cl is reabsorbed)
- **Ca**: < 7% (i.e.>93% filtered Ca is reabsorbed)
- **PO4**: <0.5% (i.e.>99.5% filtered PO4 is reabsorbed)
- **Mg**: < 15% (i.e.>85% filtered Mg is reabsorbed)

High values (signifying decreased reabsorption) across several electrolytes are consistent with renal tubular failure (especially Na and Cl) although excessive electrolyte consumption should also be considered.

Low calcium values (signifying reduced excretion) and high phosphate values (increased excretion) is a typical parathyroid hormone effect indicating calcium deficiency.

Low sodium values (signifying reduced excretion) and high potassium values (increased excretion) is a typical aldosterone hormone effect indicating sodium deficiency.