

Equine pituitary pars intermedia dysfunction: current perspectives on diagnosis and management

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Abstract: Equine pituitary pars intermedia dysfunction (PPID) is a neurodegenerative disease of the hypothalamus, resulting in the loss of dopaminergic inhibition of pars intermedia. An oxidative stress injury of unknown etiology has been suggested to initiate the neurodegeneration. While hypertrichosis (formerly known as hirsutism) is considered pathognomic for advanced disease, the antemortem diagnosis of subclinical and early disease has continued to prove difficult. Numerous tests have been used with varying sensitivities and specificities. The overnight dexamethasone suppression test, originally documented to have 100% sensitivity and specificity in horses with advanced disease, has proven to be less valuable in identifying early disease. Basal plasma adrenocorticotropin concentrations have improved sensitivity and specificity when sampled during the autumn months, and α -melanocyte-stimulating hormone, while not yet commercially available, shows promise as a sensitive and specific single sample test. Recent advances in our knowledge include the strong association between laminitis and hyperinsulinemia, both common clinical signs associated with PPID. The pathogenesis of hyperinsulinemia, laminitis, and their association with this disease is a focus of current research. The dopamine agonist pergolide mesylate is still the mainstay of medical management, with studies on oral bioavailability, pharmacokinetics, and long-term survival rates now published.

Keywords: PPID, ACTH, α -MSH, laminitis, pergolide, hypertrichosis, pars pituitary intermedia dysfunction

Introduction

Equine Cushing's disease, now known as pituitary pars intermedia dysfunction (PPID), was first described in 1932.¹ The classic clinical picture is one of advanced disease, the aging horse or pony with a persistent, long, curling hair coat; epaxial muscle wastage; pendulous abdomen; and an increased risk of laminitis.² In recent epidemiological surveys, 21%–22% of aged horses and ponies had PPID based on endocrine testing, an increase in reported prevalence from previous studies.^{3–6}

Considerable research into the pathophysiology of PPID has occurred in recent years. The current pathophysiological theory for PPID is an age-related oxidative stress-induced neurodegeneration of the dopaminergic neurons of the hypothalamus.^{7–10} This in turn causes a loss of dopaminergic inhibition on the pars intermedia resulting in an increased production of pro-opiomelanocortin (POMC)-derived peptides including adrenocorticotropin (ACTH), α -melanocyte-stimulating hormone (α -MSH), β -endorphin, and corticotrophin-like intermediate peptide. Of these hormones, the production and actions of ACTH and α -MSH have been most closely studied and assessed for diagnostic test potential. The cause of the oxidative stress compared

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to age-matched controls is yet to be determined. The very slow progression of this disease makes the elucidation of an initiating cause or insult difficult as clinical signs are only evident after a substantial time lapse (years), when normal hormonal function is lost.¹¹

In the normal pituitary, the pars distalis is responsible for the majority of ACTH production with small amounts of ACTH from melanocyte-derived POMC.⁹ The pars intermedia's primary product is α -MSH, a metabolically active peptide involved in metabolism and obesity, which has potent anti-inflammatory properties. The corticotrophs in the pars distalis are under a glucocorticoid negative feedback loop. The production and secretion of the POMC peptides from the melanotropes in the pars intermedia are under dopamine inhibitory control from the hypothalamus. Hypothalamus and pituitary activities are influenced by season and photoperiod length, with significant up-regulation of α -MSH and ACTH production in the late summer and autumn months (periods of decreasing day length).^{12–14} The increased activity is seen in both unaffected and PPID-affected horses and ponies. This has a significant effect on basal endogenous ACTH and α -MSH levels and therefore on diagnostic testing during this period.

In 1994, Dybdal et al described the overnight dexamethasone suppression test (ODST) as a 100% sensitive and specific diagnostic test based on a population of horses with clinically advanced PPID.² The challenge since then has been the diagnosis of PPID in subclinical and early clinical horses and ponies. Plasma POMC peptide levels are secreted along a concentration continuum. Horses and ponies will have normal to mildly elevated levels early in the disease process. There is a transition period or “gray zone” within this continuum, where hormone levels may be normal in clinically suggestive cases or elevated in clinically normal horses and ponies.¹⁵ Clinically overt cases will have significantly increased plasma POMC concentrations than either clinically normal or early clinical cases. Cutoff levels for the current diagnostic tests are based on statistical methodology to maximize sensitivity and specificity. To minimize the number of false negative and positive results, a number of recommendations by various equine endocrinology groups have been made including the use of seasonally adjusted reference ranges and further dynamic testing.^{16–18}

Diagnostics

Diagnosis of PPID has included the presence of overt clinical signs, endogenous hormone levels including ACTH, insulin and cortisol levels, dynamic endocrine testing such as the

ODST, thyrotropin-releasing hormone (TRH) stimulation test measuring cortisol or ACTH, combined TRH–dexamethasone suppression test, domperidone stimulation test, and histology.^{2,19–23} Most recently, α -MSH has also been evaluated as a possible diagnostic option.²⁴ Of these tests, the current recommendations are the use of endogenous plasma ACTH with seasonally adjusted reference ranges in combination with suggestive clinical signs. Horses and ponies with ambiguous results (suggestive clinical signs with normal ACTH results) or in the gray zone (clinically normal horses with mildly elevated ACTH levels) should either be resampled during autumn or undergo a TRH stimulation test measuring ACTH to improve sensitivity.

Clinical signs

A long, wavy, or curling hair coat, known as hirsutism, has long been considered pathognomonic for advanced PPID disease. The long hair coat in horses with advanced PPID has recently been determined to be due to the persistence of hair follicles in the anagen stage and is therefore more correctly termed “hypertrichosis”.²⁵ Follicles arrested in the anagen stage fail to shed, as occurring in PPID-affected horses and ponies. Hirsutism, the term commonly used to describe the hair coat changes in PPID-affected horses and ponies, has been borrowed from the medical field and refers to the excessive androgen-dependent growth of hair in women with Cushing's disease. The term “hirsutism”, therefore, has been incorrectly used to describe the hair coat abnormalities seen in PPID.

Laminitis is commonly associated with PPID.^{1,9,22,26} In all, 13% of PPID confirmed horses had a history of or concurrent laminitis in a recent survey.³ Historically, it was thought that the loss of the diurnal cortisol fluctuations with exposure to increased daily cortisol levels was responsible for the PPID-associated laminitis.^{27,28} Recent work on the pathogenesis of endocrinopathic laminitis, however, has demonstrated a strong association with a concurrent hyperinsulinemia.^{29–31} The exact mechanisms of this association and the association between hyperinsulinemia and PPID are currently unknown and are the subject of ongoing research.

Other common clinical signs include bulging supraorbital fat pads, lethargy or docility, epaxial muscle wastage, a pendulous abdomen, sweating abnormalities, polyuria/polydipsia, poor wound healing, chronic infections or increased susceptibility to infection, endoparasitism, and very rarely, seizure-like activity.^{1,9,22,32–34} The presence of three or more common clinical signs (hypertrichosis, wasted epaxial muscles, pendulous abdomen, bulging supraorbital fat pads,

and divergent laminar rings) in combination with a seasonally adjusted endogenous ACTH value has recently been documented to be the “clinically diagnostic gold standard”.³

Plasma ACTH

In the hypothalamus–pituitary–adrenal axis, ACTH secretion from the pituitary stimulates the adrenal glands to increase the secretion of cortisol. In PPID-affected horses and ponies, only 20% have adrenomegaly, supporting the theory that the excessive ACTH produced in PPID-affected horses and ponies is biologically inert, resulting in ACTH–cortisol dissociation.^{9,35} ACTH secretion is affected by numerous external factors including stress, transport, illness, marked pain, and debilitation and exercise.^{36–39} ACTH secretion is also affected by day length, with increased secretion associated with decreasing photoperiods during the late summer and autumn (Table 1).⁴⁰ This effect does not appear to be universal though, with varying degrees of seasonal fluctuations documented at different latitudes.^{15,41–44} PPID-affected horses and ponies still show the greatest elevations. Increased plasma ACTH secretion in the autumn months in both normal and PPID-affected horses and ponies initially resulted in reluctance to measure endogenous plasma ACTH due to a presumed decrease in specificity during these months. However, recent work and determination of seasonally adjusted ACTH reference ranges has significantly increased the sensitivity and specificity of endogenous plasma ACTH levels during this period. The seasonally adjusted values are latitude dependent. Large, multicenter collaborations to establish regional seasonally adjusted values are needed. In lieu of these data, using a local laboratory with its own established reference ranges is recommended.

Even with the determination of seasonal variation and adjusted cutoff values, there still exists a transitional or “gray” zone.⁴⁵ Horses and ponies that have endogenous plasma ACTH value of between 29 and 40 pg/mL (northern hemisphere non-autumn reference range) during the winter, spring, and summer months with no or few subtle clinical signs present a clinical dilemma and should be retested during the autumn when the test has greater sensitivity and specificity, decreasing the likelihood of false positive results.⁴⁶ Alternatively, a TRH stimulation test measuring ACTH, instead of cortisol, can be performed (see TRH stimulation test measuring ACTH).

Ultradian and circadian rhythms and effects of feeding have been investigated to optimize sample timing. No ultradian rhythms were found in neither unaffected or PPID-affected horses and ponies.⁴⁵ Further to this, paired sampling was not found to increase sensitivity and specificity.⁴⁷ Rendle et al identified a circadian rhythm in unaffected horses and ponies with plasma ACTH values highest at 8 am and decreasing throughout the day, whereas Diez de Castro et al did not demonstrate a circadian rhythm in healthy horses.^{45,48} A significant postprandial increase in endogenous ACTH has been documented, suggesting that the feeding status of the animal may also be a cofounder for both endogenous and dynamic ACTH testing.⁴⁸

Sample handling and storage of plasma for endogenous ACTH measurement has also been investigated. Contrary to previous recommendations, while there is a decrease in endogenous ACTH levels over time, clinically relevant results can be obtained from chilled, non-centrifuged plasma processed within 48 hours of collection.^{9,49} Samples held for greater than 48 hours prior to processing should be centrifuged, the plasma separated and frozen while awaiting

Table 1 ACTH upper reference ranges, sensitivity, and specificity data from different locations and seasons

Location/latitude	Subtropics			Temperate Northern hemisphere ^{a,44,46}
	Southern hemisphere (–27° 24' to –24° 54') ¹⁵			
	Sn (%)	Sp (%)	Cutoff	Cutoff
ACTH				
Autumn	100	95	77.4 pg/mL	
Winter–summer	80	82	29.7 pg/mL	
Late summer to mid-autumn				47 pg/mL
Late autumn to mid-summer				29 pg/mL
α-MSH				
Autumn	100	97	165.4 pmol/L	
Winter–summer	59	93	52 pmol/L	
Late summer to mid-autumn				No precise value documented
Late autumn to mid-summer				35 pmol/L

Note: ^aNorthern hemisphere reference ranges have been calculated on normal distributions and 95% confidence intervals, with no sensitivity or specificity data to date.

Abbreviations: ACTH, adrenocorticotropin hormone; cutoff, upper limit of normal reference range; α-MSH, α-melanocyte-stimulating hormone; Sn, sensitivity; Sp, specificity.

transport to the laboratory. The inclusion of the preservative *N*-phenylmaleimide had no effect on plasma ACTH levels over time.⁴⁹

TRH stimulation test measuring ACTH

The TRH stimulation test originally measured cortisol levels 10 and 30 minutes post-TRH administration and had a sensitivity of 92%. Specificity, positive and negative predicative values were improved when combined with the dexamethasone suppression test.¹⁹ In horses and ponies with PPID though, cortisol levels are rarely elevated in the face of significant elevations in ACTH, suggesting that the melanotropic-derived ACTH is immunologically active but biologically inert and, therefore, dissociated from serum cortisol levels.^{9,35} This observation has led to endogenous ACTH being measured 10 and 30 minutes post-TRH administration (Table 2), resulting in sensitivity of 94% and specificity of 78% at 30 minutes using a cutoff of 36 pg/mL.³⁵ In this study, plasma ACTH was shown to be superior than serum cortisol for the diagnosis of PPID.

As with endogenous plasma ACTH, the TRH stimulation test is affected by season and seasonally adjusted reference ranges are needed to improve specificity. Factors that confound the evaluation of endogenous ACTH results will impact on the TRH stimulation test as well. TRH has been both expensive and difficult to obtain and is therefore a test usually performed at referral and university institutions.

A compounded TRH product stored at room temperature has been assessed and found to give comparable test results. The availability and cost of this product outside North America is unknown at this time.⁵⁰

Overnight dexamethasone suppression test

The ODST has, for many decades, been considered the “gold standard” antemortem diagnostic test. This was based on a sensitivity and specificity of 100% achieved in a population of horses and ponies with advanced PPID.² The sensitivity and specificity in early and subclinical disease are 65% and 76%, respectively.¹⁹ The potential induction or exacerbation of existing laminitis post-ODST and the requirement of either two visits or overnight hospitalization of the patient have limited the practicality of this test.¹ As a downstream product of the hypothalamus–pituitary–adrenal axis, cortisol secretion is also seasonally affected. To date, there have been no seasonally adjusted reference ranges documented for the ODST.

α -Melanocyte-stimulating hormone

α -MSH is primarily derived from the pars intermedia and undergoes increased secretion occurring in the later summer and autumn months (Table 1).¹⁴ Its role in metabolism and increased secretion during periods of decreasing day length is a proposed survival mechanism to increase fat

Table 2 PPID test protocols

Test	Protocol	Sample type required
ACTH	<ul style="list-style-type: none"> Blood collected in plastic EDTA vacutainer If ≤ 48 hours to laboratory processing <ul style="list-style-type: none"> No further processing required If ≥ 48 hours until laboratory processing <ul style="list-style-type: none"> Centrifuge sample Separate plasma and place in a plain (red top) plastic vacutainer, marked EDTA Plasma Freeze plasma and send on ice to laboratory 	Plasma (EDTA)
TRH stimulation test measuring ACTH	<ul style="list-style-type: none"> Collect blood in plastic EDTA (purple top) tube Administer 1.0 mg (total dose) TRH intravenously Collect blood in plasma EDTA (purple top) tube 30 minutes later Submit samples for ACTH levels (Sample handling for ACTH is outlined above) 	Plasma (EDTA)
ODST	5 pm <ul style="list-style-type: none"> Collect blood in a plain (red top) vacutainer for basal cortisol level Administer 40 μg/kg of dexamethasone intramuscularly 8 am–12 pm hours the next day (7 pm–12 am later) <ul style="list-style-type: none"> Collect second blood sample in a plain (red top) vacutainer for second cortisol level 	Serum $\times 2$
α -MSH	<ul style="list-style-type: none"> Blood collected in EDTA vacutainer 	Plasma (EDTA)
Insulin	<ul style="list-style-type: none"> Collect blood in a plain (red top) vacutainer, preferably after a short overnight fast 	Serum
Oral glucose challenge	<ul style="list-style-type: none"> Give 0.15 mL/kg of Karo Lyte corn syrup orally Collect blood in a plain (red top) vacutainer for an insulin level 60–90 minutes later 	Serum

Abbreviations: ACTH, adrenocorticotropic hormone; EDTA, ethylenediaminetetraacetic acid; α -MSH, α -melanocyte-stimulating hormone; ODST, overnight dexamethasone suppression test; PPID, pituitary pars intermedia dysfunction; TRH, thyrotropin-releasing hormone.

stores in preparation for anticipated feed restriction in harsh winters. This same mechanism may be responsible for the associated obesity and regional fat deposition seen in PPID cases. PPID-affected horses and ponies have a greater magnitude in plasma α -MSH levels than concurrent ACTH levels, suggesting that α -MSH may rise early in the disease process and, therefore, be a more sensitive diagnostic test.¹⁴ Plasma α -MSH is stable in chilled, whole blood for up to 8 hours, but does require centrifugation, plasma separation, and freezing prior to transportation to the laboratory, unlike ACTH which, when chilled, is stable in whole blood for up to 48 hours.^{14,49} α -MSH is not effected by stress, illness, exercise, or transportation and as such has less confounders on result interpretation than ACTH.¹⁴ In a recent epidemiological study, α -MSH had a higher sensitivity and specificity than ACTH when tested in the autumn months.¹⁵ Currently, α -MSH assays are not commercially available and only used in research facilities.

Insulin

Almost 90% of laminitis diagnosed is classed as endocrinopathic in origin.⁵¹ The initial theorized pathogenesis of PPID-associated laminitis was glucocorticoid induced, due to exposure of the lamellar tissue to increased total daily cortisol as a result of the cortisol diurnal secretory rhythm loss; this has not been clearly demonstrated in studies though.^{27,28,52} Hyperinsulinemia has been repeatedly documented to cause laminitis in healthy non-laminitic horses and may be the major mechanism associated with pasture-associated laminitis.^{29,53} PPID-affected horses and ponies with concurrent hyperinsulinemia experience laminitic episodes more frequently than normoinsulinemic PPID horses and ponies.³¹ The presence of PPID alone though does not impair insulin sensitivity.⁵⁴ Insulin sensitivity decreases with increasing age in equids.^{55,56} The mechanism by which hyperinsulinemia induces laminitis and whether it is the same mechanism/s associated with PPID or a concurrent disease is yet to be elucidated.

Basal serum insulin concentrations or dynamic insulin testing should be undertaken in horses and ponies with PPID and laminitis. The presence of a concurrent hyperinsulinemia will influence the management and feeding practices implemented. To standardize testing, samples for basal serum insulin should be taken in the morning after an overnight fast. A small biscuit of good quality given and consumed before 10 pm is allowed. Insulin levels greater than 20 IU/L are considered hyperinsulinemic.⁵⁷ Dynamic testing has the added benefit of assessing postprandial insulin levels (the pancreatic β -cells response to a glucose challenge).

Dynamic testing involves the oral administration of 0.15 mL/kg of corn syrup (ACH Food Companies, Karo Lyte, Cordova, TN, USA) orally after a short overnight fast (ie, a small biscuit of hay given, consumed, and any remainder removed before 10pm the night before), and serum insulin samples taken 60 and 90 minutes later. Serum insulin of less than 45 IU/L at both time points is found in healthy horses. Serum insulin levels between 45 and 60 IU/L are equivocal and greater than 60 IU/L is evidence of postprandial hyperinsulinemia and insulin dysregulation.⁵⁸ Alternatively, 1 g/kg of dextrose powder can be fed in a small meal of chaff or low-sugar chopped feed and mixed in water after a short overnight fast. Serum insulin levels are measured 60 minutes later. Insulin levels greater than 85 IU/L are considered diagnostic for hyperinsulinemia.⁵⁹ Other tests include the combined insulin and glucose test and the insulin tolerance test. Neither of these tests assesses both the pancreatic secretion and clearance of insulin. They also require more time and sampling with more confounders when compared with the oral glucose tests.⁶⁰

Histopathology

For many other diseases, confirmation of a previously described histopathological appearance is considered the gold standard to which all antemortem tests are compared. Confirmation of the disease on histopathology is therefore used to calculate the sensitivity and specificity of a given antemortem test. This test standard has been unreliable in PPID, with documented inconsistencies between pathologists on classification of histological changes and therefore final diagnosis.²³ The pituitary architecture and size is also subject to seasonal variation.⁶¹ The current accepted description is a hyperplastic pars intermedia with a singular large or multiple small adenomas.⁹

Treatment

There is currently no cure for PPID. Pergolide mesylate, an ergot-derived D2-dopamine agonist, is still the drug of choice for the medical management of PPID.⁹ The replacement of dopamine and reestablishment of dopamine inhibition on the POMC-derived peptides results in normalization of pituitary hormone levels and resolution of clinical signs.

Management

PPID-affected horses and ponies require excellent husbandry and management practices. The presence of insulin dysregulation and postprandial hyperinsulinemia requires careful dietary management.⁶² Horses and ponies with a

history of laminitis or documented hyperinsulinemia require restricted access to diets high in soluble nonstructural carbohydrates, including lush or fast-growing pastures and sweet feeds to decrease the frequency of laminitic episodes.⁶³ The use of a grazing muzzle aids in dietary restriction without restricting exercise. Grazing should be restricted from mid-morning to late afternoon, as wilting increases the sugar content in the pasture. Likewise, grazing should be restricted after a harsh frost as this also increases the sugar content of pasture. Soaking hay for 60 minutes in cold water will also decrease the sugar content. All PPID-affected horses and ponies benefit from the feeding of good quality senior diets to provide adequate protein intake to minimize muscle catabolism.

PPID-affected horses and ponies are more likely to suffer from moderate to severe endoparasitism compared with age-matched controls.³³ PPID horses and ponies tend to be the “high egg shedders” in the herd and have a shorter egg reemergence period post-worming than their herd mates. A deworming program which incorporates regular fecal egg counts and fecal egg reduction test is optimal.

Regular dental checks and vaccinations are recommended. PPID horses and ponies, while having an increased susceptibility to infections, continue to have adequate immune responses to vaccination.^{64,65} The degree of response and duration of immunity may vary depending on severity of disease compared with age-matched controls. Further research is required to further quantify these changes and assess whether a change in vaccination dosage frequency should be recommended. Early and aggressive medical management of infections is also needed.

Pergolide mesylate

Pergolide mesylate is available in Australia and New Zealand as a liquid (Ranvet Pty Ltd, Ranvet's Pergolide, East Botany NSW, Australia) and as a tablet (Boehringer Ingelheim, Prascend, Ingelheim am Rhein, Rhineland-Palatinate, Germany). Only the tablet form is commercially available in the UK, Europe, and North America. The 200 mg/mL liquid formulation allows for very accurate dosing. Pergolide, however, is unstable and light sensitive in the liquid form, with a short 3-month shelf life, potentially resulting in drug wastage and extra cost to the client. The tablet formulation is more stable with a shelf life of up to 24 months. The 1 mg tablets are half-scored allowing for one division and a dosage accuracy of 0.5 mg. Compounded pergolide mesylate has shown itself to be unreliable in both drug distribution and stability and is not recommended for use.⁶⁶

The current recommended dose rate is 0.002 mg/kg once daily orally.⁹ Lower doses have been successfully used.^{34,67} Once clinical and biochemical resolution has been achieved, the dose should be titrated down to the lowest effective dose to prevent side effects such as anorexia, sweating, colic, diarrhea, and neurological signs.⁶⁸ These side effects are most commonly seen at higher dose rates (0.01 mg/kg) and can be avoided by starting at a lower dose rate and gradually increasing the dose if needed over a period of weeks. Dosage efficacy is based on the demonstration of normalized plasma ACTH levels and the resolution of clinical signs. It should be noted that hair coat shedding will not be seen until the spring, regardless of appropriate dosage administration, due to the day length effect on α -MSH.

The serotonin antagonist cyproheptadine is no longer considered useful either alone or in conjunction with pergolide due to the lack of efficacy achieved above that attributed to either management or pergolide.⁶⁸

Conclusion

The last 3 decades has seen considerable expansion in our knowledge of PPID. There is still much to learn though. A “gold-standard” antemortem diagnostic test continues to be elusive. Currently, basal plasma ACTH is the most sensitive and specific test commercially available for early diagnosis. The sensitivity and specificity increase if sampling occurs during the autumn months. Dynamic testing using the TRH stimulation test measuring ACTH is also useful in detecting early clinical disease. α -MSH shows promise as a single sample diagnostic test, but is not yet commercially available. Pergolide is still the drug of choice for medical management of clinical signs and, when used in conjunction with improved husbandry and management practices, can significantly improve the quality and duration of life of PPID horses and ponies for many years.

Disclosure

The author reports no conflicts of interest in this work.

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