The Interplay Between Pituitary Health and Diabetes Mellitus – The Need for ‘Hypophyseo-vigilance’

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The anterior and posterior hypophyseal hormones alter glucose metabolism in health and disease. Secondary diabetes may occur due to the hypersecretion of anterior pituitary hormones like adrenocorticotropic hormone in Cushing’s disease and growth hormone in acromegaly. Other hormones like prolactin, gonadotropins, oxytocin and vasopressin, though not overtly associated with causation of diabetes, have an important physiological role in maintaining glucose homeostasis. Hypoglycaemia is not an unusual occurrence in hypopituitarism. Many of the medications that are used for the treatment of hypophyseal diseases after glucose metabolism. Agents like pasireotide should be used with caution in the setting of diabetes, whereas pegvisomant should be given preference. Diabetes mellitus itself, on the other hand, can alter the functioning of the hypothalamic pituitary axis; this is documented in both type 1 and type 2 diabetes. This review focuses on the clinically relevant interplay of hypophyseal hormones and glucose homeostasis. The authors define ‘hypophyseo-vigilance’ as an approach which keeps the bidirectional, multifaceted interactions between the pituitary and glucose metabolism in mind while managing diabetes and pituitary disease.

Keywords
Diabetes mellitus, pituitary, hypophysitis, acromegaly, Cushing’s disease, hypogonadism

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Glucose metabolism in anterior pituitary disease

Anterior pituitary dysfunction can be due to either hypersecretion or hyposecretion of pituitary hormones. The impact of anterior pituitary dysfunction on glucose metabolism has been described in Table 1.

Hyperfunction

This encompasses four recognised conditions of hormone hypersecretion: growth hormone (GH)-secreting pituitary adenoma, or acromegaly; adrenocorticotropic hormone (ACTH)-secreting pituitary adenoma, or Cushing’s disease; prolactin-secreting pituitary adenoma; and very rarely, thyroid-stimulating hormone (TSH)-secreting pituitary adenoma. Gonadotrophin-secreting adenomas are usually clinically non-functioning, but secretion of intact follicle-stimulating hormone (FSH) in women of reproductive age has been associated with ovarian hyperstimulation.

Growth hormone hypersecretion

In a database of 3,173 patients recently diagnosed with acromegaly, prevalence of DM was 27%. In the same study in subjects without diabetes, the glucose levels correlated with insulin-like growth factor-1 (IGF-1) levels, but not with GH levels. GH enhances hepatic glucose production by opposing
The action of insulin on the liver and stimulates gluconeogenesis. GH also increases hepatic and peripheral insulin resistance. One of the mechanisms behind peripheral insulin resistance is substrate competition with glucose due to higher free fatty acid availability from GH-induced lipolysis. On the other hand, IGF-1 improves peripheral insulin sensitivity but is not enough to override the diabetogenic action of chronic GH excess. Studies have demonstrated the association of IGF-1 levels and DM as predictors of mortality in acromegaly.

### Glucose metabolism in anterior pituitary dysfunction

<table>
<thead>
<tr>
<th>Disease/condition</th>
<th>Effect on glucose metabolism</th>
</tr>
</thead>
<tbody>
<tr>
<td>GH hypersecretion</td>
<td>Insulin resistance</td>
</tr>
<tr>
<td></td>
<td>Diabetes mellitus</td>
</tr>
<tr>
<td>ACTH hypersecretion</td>
<td>Impaired glucose tolerance</td>
</tr>
<tr>
<td></td>
<td>Diabetes mellitus</td>
</tr>
<tr>
<td>Prolactin hypersecretion</td>
<td>Prolactinoma – higher FPG in small studies</td>
</tr>
<tr>
<td>TSH hypersecretion</td>
<td>Insulin resistance, worsening of diabetes</td>
</tr>
</tbody>
</table>

### Adrenocorticotrophic hormone hypersecretion

Cushing’s disease, or ACTH-secreting pituitary adenoma, is another recognized cause of DM. The prevalence of DM in Cushing’s disease is 40–45%, and an additional 10–30% have impaired glucose tolerance. A state of chronic cortisol excess stimulates hepatic gluconeogenesis partly resulting from increased mobilisation of gluconeogenic substrate from peripheral tissues. Cortisol also has a permissive effect on glucagon and epinephrine in inducing gluconeogenesis and glycogenolysis. Insulin secretory defects coupled with increased peripheral insulin resistance have an additional role in the development of hyperglycaemia.

### Prolactin hypersecretion

Unlike acromegaly and Cushing’s disease, hyperprolactinaemia is not a recognised cause of DM. Small studies have demonstrated that prolactinoma is associated with higher fasting plasma glucose (FPG) that can be corrected with treatment with cabergoline. However, the relationship between DM and prolactin is complex. In a population-based study of 2,377 Chinese men and postmenopausal women, elevated serum prolactin levels were associated with lower prevalence of DM. In another population-based study of 370 Japanese men without diabetes, elevated prolactin in the physiological range showed an association with insulin resistance. Experimental studies also show conflicting results. On the one hand, prolactin has been shown to induce beta-cell growth in the pancreas, while on the other hand, prolactin is also hypothesised to increase insulin resistance by blocking adiponectin and interleukin (IL)-6 production in fat cells.

### Others

Thyrotoxicosis can cause insulin resistance and predispose to diabetes. TSH secreting pituitary adenoma by producing thyrotoxicosis can worsen glucose intolerance like in any other thyrotoxic state, but the condition is very rare and to our knowledge, diabetes arising from this condition has not been reported. Gonadotropin-secreting tumours are non-functional; only in exceptional circumstances they can produce manifestations related to gonadotropin excess, which is not usually linked to glucose homeostasis.

### Hypofunction

Pituitary hypofunction or hypopituitarism implies the deficiency of one or more anterior and posterior pituitary hormones. The common causes of hypopituitarism are pituitary adenoma or other sellar and suprasellar lesions, surgery or radiation to the pituitary gland, traumatic brain injury, and Sheehan’s syndrome. In most of these situations, multiple hormones are deficient; however, in lymphocytic hypophysitis, isolated deficiency of a single hormone can occur. Hypoglycaemia can be a presenting feature of hypopituitarism. It results primarily from hypocortisolism, secondary to defect in ACTH secretion coupled with deficiency of GH.

GH deficiency has a different impact on glucose homeostasis at various stages of life. Children with GH deficiency have a tendency to be insulin sensitive and can develop spontaneous fasting hypoglycaemia due to reduced hepatic glucose production. GH-deficient adults who are not receiving GH substitution, however, are insulin resistant.

Besides overt hormonal deficiency, functional hypogonadism, presumed to be due to a non-organic defect in the hypothalamic-pituitary-testicular axis, is a common cause of low plasma testosterone in adult males with comorbidities like diabetes and obesity. Low plasma total testosterone levels have been linked with insulin resistance, impaired glucose tolerance, higher visceral adipose tissue and adverse cardiovascular outcomes. Whether metabolic syndrome is the cause behind hypogonadism or its effect remains an area of controversy.

### Glucose metabolism in posterior pituitary disease

The posterior pituitary secretes two major hormones – oxytocin and vasopressin.

#### Oxytocin

An imbalance in oxytocin secretion related to a disease process is not known to cause any specific defect in glucose metabolism; however, over the last few years the physiological role of oxytocin in glucose and lipid metabolism has been recognised. Peripherally, oxytocin maintains electrolyte balance, glucose metabolism and adipogenesis, and centrally it is involved in food selection, reward and satiety. Oxytocin preferentially suppresses intake of sweet-tasting carbohydrates. Animal studies indicate that defects in oxytocin signalling and receptor expression can lead to weight gain despite normal food intake. Oxytocin levels were found to be lower in individuals with newly diagnosed T2DM or who were obese. A review analysing the weight-lowering and glycaemic benefits of intranasal oxytocin in a few small studies showed a positive trend.

#### Vasopressin

Two conditions, diabetes insipidus, a condition caused by vasopressin or antidiuretic hormone (ADH) deficiency, and syndrome of inappropriate ADH (SIADH), whereby ADH is secreted in excess, do not have any direct effect on glucose metabolism. However, other than its role in fluid balance, there is emerging evidence that vasopressin influences
glucose metabolism. Murine and in vitro studies have demonstrated that the vasopressin receptor V1a is present in the liver where its activation causes glycogenolysis and gluconeogenesis. The pancreas also expresses V1b receptors on both alpha and beta cells. At higher glucose concentration, vasopressin predominantly causes insulin secretion from beta cells, whereas at lower glucose concentration it primarily releases glucagon from alpha cells.

Murine studies demonstrate that vasopressin has a predominant role in inducing insulin resistance and causing hyperglycaemia through activation of the V1a receptor. These animal data have been corroborated through clinical studies that indicate high vasopressin levels can be a risk factor for DM. Coppezin, the C-terminal of the pre/pro-vasopressin peptide, has been used as a surrogate indicator of vasopressin in clinical studies because it is stable and can be assayed easily. A recent review summarises the experimental evidence, as well as clinical studies, that have demonstrated that high copeptin levels are associated with DM, insulin resistance, obesity, hypertension and metabolic syndrome.

### Table 2: Glucose metabolism during treatment with drugs used for pituitary dysfunction

<table>
<thead>
<tr>
<th>Drug/condition</th>
<th>Effect on glucose metabolism</th>
</tr>
</thead>
<tbody>
<tr>
<td>GH-replacement therapy</td>
<td>Children – subtle insulin resistance, new onset DM rare; Adults – association with impaired glucose tolerance and DM in initial years, prolonged therapy – improvement in metabolic parameters and decreased CVD risk</td>
</tr>
<tr>
<td>Oxytocin and lanreotide</td>
<td>Postprandial hyperglycaemia and increase in HbA1c in uncontrolled disease; if disease is under control – neutral or mild improvement in glycaemic indices</td>
</tr>
<tr>
<td>Pasireotide</td>
<td>New onset DM, uncontrolled hyperglycaemia</td>
</tr>
<tr>
<td>Pegvisomant</td>
<td>Improvement in glycaemic status</td>
</tr>
<tr>
<td>Bromocriptine</td>
<td>Quick-release formulation approved for treatment of DM</td>
</tr>
<tr>
<td>Cabergoline</td>
<td>Anecdotal reports of glycaemic and weight benefit</td>
</tr>
<tr>
<td>GnRH analogue</td>
<td>Increased chance of diabetes, more in males on androgen deprivation therapy</td>
</tr>
<tr>
<td>Mifepristone</td>
<td>Approved for treatment of hyperglycaemia in Cushings’s syndrome</td>
</tr>
<tr>
<td>Mecasermin</td>
<td>Adverse effect of hyperglycaemia, must be taken with food</td>
</tr>
</tbody>
</table>

CVD = cardiovascular disease; DM = diabetes mellitus; GH = growth hormone; GnRh = gonadotropin-releasing hormone; HbA1c = glycated haemoglobin.

### Table 3: Practical considerations for the management of glycaemic parameters in pituitary hypersecretory states

<table>
<thead>
<tr>
<th>Drug/condition</th>
<th>Effect on glycaemic parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>GH-replacement therapy</td>
<td>Screen for DM by measuring FPG, OGTT or HbA1c in acromegaly. Cushing’s disease and TSH-secreting pituitary adenoma</td>
</tr>
<tr>
<td>First generation SSA</td>
<td>First generation SSA is the first-line therapy for acromegaly with persistent disease despite surgical resection</td>
</tr>
<tr>
<td>Pegvisomant</td>
<td>Pegvisomant should be the preferred switch-over agent in patients not optimally controlled on first-generation SSAs, in the presence of pre-existing clinically relevant impaired glucose metabolism</td>
</tr>
<tr>
<td>First-generation SSAs</td>
<td>First-generation SSAs increase HbA1c by causing postprandial hyperglycaemia; appropriate adjustment in glucose-lowering treatment is recommended</td>
</tr>
<tr>
<td>Second-generation SSAs, like pasireotide, worsen glycaemic status; treatment with incretin-based or other glucose-lowering therapy to be considered. In case of severe hyperglycaemia, switch over to pegvisomant is recommended</td>
<td></td>
</tr>
<tr>
<td>For Cushing’s disease not cured by surgical treatment, use of pasireotide can worsen glucose metabolism; incretin-based or other glucose-lowering therapy to be considered</td>
<td></td>
</tr>
<tr>
<td>Treatment with mifepristone can improve hyperglycaemia in Cushing’s syndrome</td>
<td></td>
</tr>
</tbody>
</table>

DM = diabetes mellitus; FPG = fasting plasma glucose; HbA1c = glycated haemoglobin; OGTT = oral glucose tolerance test; SSA = somatostatin analogue; TSH = thyroid-stimulating hormone.

### Table 4: Practical considerations for the management of glycaemic parameters in pituitary hyposecretory states

<table>
<thead>
<tr>
<th>Drug/condition</th>
<th>Effect on glycaemic parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>GH-replacement therapy</td>
<td>Hypoglycaemia can be a manifestation of panhypopituitarism, children with GH deficiency have fasting hypoglycaemia. Appropriate screening should be considered in relevant situations</td>
</tr>
<tr>
<td>Hypogonadotropic hypogonadism and adult GH deficiency are associated with insulin resistance. Screen for DM when appropriate</td>
<td></td>
</tr>
<tr>
<td>Glucocorticoid replacement therapy can induce hyperglycaemia, with the characteristic pattern of an afternoon or evening rise in plasma glucose. Consider short-acting insulin prior to lunch</td>
<td></td>
</tr>
<tr>
<td>Hypoglycaemia is a recognised adverse effect of mecaermin. Appropriate precautions are advocated, including administration with food</td>
<td></td>
</tr>
</tbody>
</table>

DM = diabetes mellitus; GH = growth hormone.

In adults with a higher BMI and/or family history of T2DM, GH therapy can be associated with the development of glucose intolerance or T2DM in the initial years of treatment, though the incidence decreases with prolonged treatment. Close monitoring of glycaemic status is advisable for adults with risk factors for T2DM on GH replacement. Treatment of GH deficiency in adults decreases fat mass and improves lean body mass, lipid profile, carotid intima medial thickness, and CV inflammatory markers, and more than offsets for mild derangement in glycaemic parameters.

### First-generation somatostatin analogues

Oxytocin and lanreotide are considered as first-generation somatostatin analogues (SSAs) having a strong affinity to somatostatin receptor 2 (SSR2) and SSR5, and a weaker affinity to SSR3. They are the first-line
medical therapy for acromegaly. First-generation SSAs exert a glucose-lowering effect by decreasing GH secretion from the tumour, which is counterbalanced by its action on the pancreas where somatostatin and its analogues inhibit insulin secretion. The net effect on glucose metabolism in acromegaly is variable. In a recently published meta-analysis, the authors showed that a mild but significant increase in HbA1c occurred with first-generation SSAs, resulting from an increase in post-prandial glucose levels without any alteration in FPG. It was also demonstrated that first-generation SSAs significantly reduced insulin secretion which was not completely counterbalanced by the reduction in GH and IGF-I levels. The authors concluded that the overall effect of first-generation SSAs on glucose metabolism depends on the net balance between these two opposing factors and on the patient’s predisposition. Different trends were observed in a retrospective 1-year study comparing SSAs to surgery. SSAs caused improvement in all metabolic syndrome parameters and significantly lowered FPG and HbA1c when the disease activity was controlled, the main determinant of glycaemic improvement. Higher dose and higher frequency of SSAs do not have an unfavourable glycaemic impact. To summarise, octreotide and lanreotide might have a mild detrimental effect on post-prandial hyperglycaemia and HbA1c, especially in the setting of uncontrolled disease activity. If the disease is under control, first-generation SSAs have either a neutral or even a slightly beneficial effect on glycaemic indices.

**Pasireotide**

This novel SSA has strong affinity for SSR5 and SSR1, moderate affinity for SSR3 and lower affinity for SSR2 in comparison to octreotide. In addition to better efficacy for the treatment of acromegaly as compared with first-generation SSAs, pasireotide, by virtue of its affinity for both SSR2 and SSR5, suppress ACTH hypersecretion in patients with corticotroph tumours. Its predominant adverse effect is hyperglycaemia in both acromegaly (57.3–67.0%) and Cushing’s disease (68.4–73.0%), and uncontrolled hyperglycaemia is a common cause of treatment discontinuation. Insulin release from beta cells of islets of Langerhans is mediated by SSR5 and SSR2, and glucagon release from alpha cells depends on SSR2. Pasireotide has a prominent inhibitory effect on insulin release as a result of its higher affinity for SSR5, whereas the inhibitory effect on glucagon secretion due to SSR2 affinity is only modest. First-generation SSAs are more potent suppressors of glucagon secretion because of stronger binding to SSR2. Apart from this imbalance in insulin and glucagon secretion, pasireotide also reduces incretin response, and incretin-based therapy might be more effective than other forms of therapy for the management of hyperglycaemia in this condition.

**Pegvisomant**

Pegvisomant is a genetically engineered potent GH receptor antagonist that acts at the receptor level to decrease IGF-1 levels. Unlike SSAs, it does not have any inhibitory action on insulin secretion on the pancreas, but it does have a favourable effect on glycaemic status. Pegvisomant improves glucose metabolism by improving insulin resistance in acromegaly through the blocking of excess GH action. It has a positive impact on glycaemic control in patients with acromegaly and uncontrolled DM. Patients with acromegaly and DM have a higher requirement for pegvisomant to control the disease and have a lower rate of IGF-1 normalisation compared to those without DM. A possible hypothesis is that hepatic GH receptor expression may be increased due to hyperinsulinism and a higher dose of pegvisomant may be needed to saturate it. The recently published consensus statement by Melmed et al. recommends shifting from first-generation SSAs to pegvisomant in case there is pre-existing clinically relevant impaired glucose metabolism.

**Bromocriptine**

Dopamine and dopaminergic signals in the central nervous system (CNS) decrease liver gluconeogenesis via the sympathetic nervous system. In obese individuals with DM, decreased dopaminergic activity alters signalling from the CNS, resulting in increased hepatic gluconeogenesis and insulin resistance. Antipsychotic medicines that block dopaminergic activity cause weight gain, dyslipidaemia and insulin resistance. Bromocriptine, a D2-dopamine agonist with sympatholytic activity, is primarily indicated for the treatment of prolactinoma. Bromocriptine quick-release tablet ingestion within 2 hours of waking is assumed to enhance low hypothalamic dopamine levels and inhibit excessive sympathetic tone within the CNS. This is the only approved glucose-lowering medication that predominantly acts on the CNS and causes reduction of FPG, HbA1c, triglyceride and free fatty acid levels. In a 1-year prospective study, HbA1c was decreased by 0.6% and composite CV outcome was reduced by 40%. However, the mechanism behind reduction in CV events is not defined. The additional advantages of bromocriptine include absence of hypoglycaemia, no change in weight and lack of major side effects.

**Cabergoline**

Cabergoline, a long-acting D2-dopamine agonist, is better tolerated than bromocriptine and is considered as the first-line treatment of hyperprolactinaemia. It also has demonstrated benefit in DM, obesity and insulin resistance in small studies. In a recently published trial, cabergoline did not exhibit any difference to placebo in maintaining weight loss in obese subjects who had lost at least 5% weight with an 800 kcal/day commercial meal replacement programme. Cabergoline, despite being a stronger D2 receptor agonist, has minimal glycaemic benefit compared with bromocriptine quick-release formulation. The unravelling of the intricacies of dopaminergic and other CNS pathways that control metabolism might resolve these discrepancies in the future.

**Gonadotropin-releasing hormone analogue**

Gonadotropin-releasing hormone (GnRH) analogues, such as leuproide, triptorelin, buserelin, goserelin and histrelin, are used in adult females for the treatment of endometriosis, fibroids, premenopausal ovarian suppression, and for the treatment of metastatic prostate cancer in males. The metabolic changes that occur due to androgen-deprivation therapy with GnRH analogues in males include an increase in adipose tissue, increased low-density lipoprotein cholesterol, and insulin resistance and DM. In an observational study of a population-based cohort of 73,196 people, the unadjusted rates per 1,000 person-years for developing incident diabetes was 29.0 for those receiving GnRH analogue therapy versus 20.9 for those not on treatment, with an adjusted hazard ratio of incident DM of 1.44 with GnRH analogues. GnRH analogue-induced hyperglycaemia has been reported to respond well to pioglitazone. Females also may develop worsening of glycaemic control when treated with these agents.

**Mifepristone**

Mifepristone is a glucocorticoid receptor blocker acting on glucocorticoid receptor-II and the progesterone receptor. It is approved for treatment of hyperglycaemia in the setting of Cushing’s syndrome. In a study of 50 patients with endogenous Cushing’s syndrome (including pituitary, adrenal and ectopic causes of hypercortisolism), 60% of patients with glucose intolerance or DM showed improvement. In patients using insulin, their dose was reduced by half and mean HbA1c came down from 7.43% at baseline to 6.29% at study conclusion.
Mecasermin
Mecasermin or recombinant human IGF-1 is used for the treatment of growth failure resulting from severe primary IGF-1 deficiency, or in cases of GH gene deletion with neutralising antibodies to GH that make GH ineffective. Hypoglycaemia is a recognised side effect of mecasermin, especially in small children due to inconsistent oral intake. Individuals should not undertake high-risk activities (e.g. swimming) within 2–3 hours after dosing, particularly at the initiation of therapy until the dose is tolerated. It is injected subcutaneously twice daily and should be administered within 20 minutes of a meal or a snack.

Effect of diabetes on hypophyseal function
Pituitary hormones may be altered in both T1DM and T2DM.

Diabetes and the growth hormone/insulin-like growth factor-1 axis
Type 1 diabetes mellitus
GH and insulin appear to be metabolically antagonistic to each other, but in reality, insulin plays a permissive role in GH action. Portal insulin up-regulates hepatic GH-receptor expression, and that in turn causes downstream production of IGF-1 and IGF-binding protein-3 (IGFBP3). Low portal insulin levels, decreased IGF-1 levels and elevated proinflammatory cytokines in T1DM lead to IGF-1 resistance. Low IGF-1 in T1DM has multiple deleterious effects like decreased growth, insulin resistance and poor CV outcomes. The imbalance in the GH/IGF-1/IGFBP3 axis, however, does not get corrected with continuous subcutaneous insulin infusion despite optimal glycaemic control, but does improve with continuous portal insulin infusion. The implication of the finding, that even optimal glycaemic control might not restore the GH/IGF-1 axis is not clear. Whether deficient portal insulin in T1DM adversely affects the final height outcome is yet to be proven, and the adult height of individuals with T1DM remains highly variable.

Type 2 diabetes mellitus
The GH/IGF-1 axis plays a possible role in glucose homeostasis and the etiopathogenesis of T2DM and insulin resistance. In T2DM, GH levels are low and free IGF-1 levels are elevated; however, total IGF-1 levels are normal. In addition, IGFBP1 and IGFBP2 levels are low, while IGFBP3 is elevated in T2DM. Epidemiological studies on the connection between IGF proteins and the incidence of T2DM report conflicting results and most fail to show any association. Low levels of IGFBP1, however, have been associated with a future risk of T2DM development. Glucose metabolism and GH, IGF-1 and IGFBPs are interlinked in a complicated manner and their relationship is a subject of ongoing research.

Diabetes and the gonadal axis
It is well established that males with T2DM have a high prevalence of decreased serum testosterone along with inappropriately low luteinising hormone and FSH levels, a condition that has been labelled as functional HH because it is not associated with any structural disease. HH has a close link to insulin resistance and can be found in individuals with T2DM, obesity and metabolic syndrome. C-reactive protein levels are higher in individuals with HH and have an inverse association to testosterone levels. Inflammation secondary to insulin resistance may contribute to its pathogenesis. Testosterone replacement is recommended in symptomatic males with HH, but there is controversy regarding the CV risk associated with testosterone therapy. An interesting observation is that the hypogonadism in these conditions responds well to clomiphene. The prevalence of HH in males with T1DM does not appear to be increased, and risk factors in T1DM for low testosterone levels are similar to that of the general population.

Diabetes and prolactin secretion
An early study reported that prolactin levels are elevated in T2DM. Subsequent findings, however, suggest that high prolactin levels reduce the prevalence of DM and insulin resistance. Additionally, a study reported that those with diabetic retinopathy have lower levels of prolactin. A possible hypothesis behind this is prolactin is broken down to vasoinhibins, with antiangiogenic, vasconstrictive and anti-permeability effects that may exert protective action in diabetic retinopathy. However, a few other clinical studies have not confirmed the benefits of elevated levels of prolactin on retinopathy.

Diabetes and the cortisol axis
Animal studies suggest that the hypothalamus–pituitary–adrenal (HPA) axis may be hyperfunctioning in T2DM and that these changes may be mediated through the endogenous cannabinoid receptor system. The clinical implication of this finding is not yet known.

Diabetes and hypophysis with syndromic connection
Diabetes is a defining feature of numerous syndromes and many of them have associated hypophyseal affection (Table 5).

Table 5: Syndromes involving the pituitary and diabetes/the pancreas

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Clinical manifestations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wolfram syndrome</td>
<td>DM, central diabetes insipidus, optic atrophy, sensorineural hearing loss, neurodegeneration, prototype of endoplasmic reticulum disorder</td>
</tr>
<tr>
<td>Prader-Willi syndrome</td>
<td>Hypophagia, obesity, hypotonia, hypopituitarism, DM</td>
</tr>
<tr>
<td>Mitochondrial diseases</td>
<td>Maternal inheritance, DM (mimics T2DM), sensorineural deafness (early onset), HPA axis dysfunction, GH deficiency</td>
</tr>
<tr>
<td>Multiple endocrine neoplasia syndrome 1</td>
<td>Parathyroid adenoma, pituitary tumours, pancreatic endocrine tumour – gastrinoma and insulinoma (small, multiple, early onset)</td>
</tr>
</tbody>
</table>

DM = diabetes mellitus; GH = growth hormone; HPA = hypothalamus–pituitary–adrenal; T2DM = type 2 diabetes mellitus.

Wolfram syndrome
Wolfram syndrome is a rare genetic disease which manifests as childhood-onset DM, central diabetes insipidus, optic nerve atrophy, sensorineural hearing loss and neurodegeneration. It is a prototype of endoplasmic reticulum disorder occurring due to mutation in the WFS1 gene and pancreatic beta cells, and neurons are particularly sensitive to it, most probably due to high rates of protein synthesis.

Prader-Willi syndrome
Prader-Willi syndrome is one of the most common varieties of obesity syndrome. It results from the absence of expression of the paternally active genes on the long arm of chromosome 15. Excessive appetite, obesity and hypotonia are hallmarks of this syndrome. Hypopituitarism is often an accompanying feature with sequential loss of GH, luteinising hormone, FSH, TSH and ACTH secretion.

Mitochondrial disease
Another situation where diabetes and pituitary dysfunction coexist are mitochondrial disorders. The common variants of mitochondrial diseases include mitochondrial encephalomyopathy, lactic acidosis

EUROPEAN ENDOCRINOLOGY
Multiple endocrine neoplasia syndrome 1

Multiple endocrine neoplasia 1 ( MEN1 ) syndrome is an autosomal dominant condition that is characterised by a predisposition to tumour formation in the parathyroid glands, anterior pituitary and islet cells of the pancreas. Menin, a tumour suppressor gene located on the long arm of chromosome 11 (11q13), is inactivated in the majority of affected members.1–10

In a recently published series of patients with MEN1 syndrome, the prevalence of pituitary tumour, followed by GH-secreting tumour, Cushing’s disease, and gonadotroph adenomas and clinically non-functioning tumours.11–111 The prevalence of pituitary tumour was 42%. Prolactinoma was the most common variety followed by GH-secreting tumours, Cushing’s disease, and clinically non-functioning tumours. The other manifestation of MEN1 syndrome is tumour of the pancreas and gastro-intestinal neuroendocrine glands. Zollinger-Ellison syndrome, or gastrinoma, is the most common presentation among this group, followed by insulinoma. Insulinomas in MEN1 syndrome are usually small, multiple and present early in the second to fourth decade.112

Other syndromic associations

There are multiple other syndromes, such as Bardet-Biedl syndrome, Friedrich ataxia, Alström syndrome and Berardinelli-Seip syndrome where diabetes and hypophyseal dysfunction coexist.

Conclusion

Pituitary hormones play an integral role in controlling glucose metabolism. There are diseases like acromegaly and Cushing’s disease which are overtly associated with diabetes. Emerging evidence suggests that prolactin, vasopressin and oxytocin also exert a subtle physiological role in glucose homeostasis.13,15,21 The choice of medication in pituitary diseases such as acromegaly can be governed by the glycaemic impact of these drugs; pegvisomant being preferred in patients with uncontrolled hyperglycaemia, while pasireotide is often administered in those where glycaemic control is optimal. Bromocriptine quick-release formulation, a drug principally used for hyperprolactinaemia, has been used as an antihyperglycaemic agent and might open the door for future medications which target the CNS as a modality to treat DM. Diabetes is a recognised cause of HH in males, and inflammation associated with T2DM has a negative impact on functioning of gonadotropins.12 Whether replacement of testosterone in these individuals is beneficial is again a subject of controversy. Finally, there are two syndromes which link the pituitary and the pancreas, and which throw important light on the mechanism of secretory function of these two major endocrine glands.
The Interplay Between Pituitary Health and Diabetes Mellitus

somatostatin analogue SOM230, compared with octreotide, induces differential effects in several metabolic pathways in patients with acromegaly.


