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## The Diagnostic and Prognostic Hormones in Common Endocrine Disorders in Adult Males

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#### Authors' contributions

This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.

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#### ABSTRACT

Hormones mediate many body functions, including growth, differentiation, reproduction, and metabolism. There are variations in hormone release, response in different individuals, in relation to age, gender, stress situations, and body mass index. This article outlines the diagnostic and prognostic value of hormones in common endocrine disorders in adult males. Hormones are used in specialty clinics as markers for diagnosis and prognosis of medical disorders, that result from disturbances of pattern of release, deficiency, or resistance. Hormones are measured in specialized laboratories for these purposes, taking into consideration, age, gender and timing variations in hormone release. Lab findings, coupled to the clinical history of patients, help to complete the picture for diagnosis and prognosis of hormone disorders.

Keywords: Hormone disorder; endocrine disorder; pathological condition.

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#### **1. INTRODUCTION**

A hormone disorder, also referred to as an endocrine disorder, is a pathological condition characterized by an imbalance in the synthesis, secretion, or action of hormones. These hormones, produced by the glands of the endocrine system, are vital chemical messengers that regulate a multitude of physiological processes. metabolism, growth, including reproduction, and mood regulation. Hormone disorders may arise from either an overproduction or underproduction of hormones, dysfunction of endocrine glands, or impaired hormonal response by the body. Common causes include genetic factors, autoimmune conditions, neoplasms, infections, and various lifestyle influences [1].

Recent advances in understanding the genetic basis of endocrine disorders have indeed highlighted the significant role that mutations play in the onset and progression of conditions like congenital adrenal hyperplasia (CAH). This genetic disorder is most commonly associated with mutations in the 21-hydroxylase gene, which leads to impaired cortisol synthesis and the subsequent accumulation of androgens, disrupting normal endocrine function and leading to clinical disease [2].

Illustrative hormone cases of disorders diabetes hypothyroidism, include mellitus, hyperthyroidism, Cushing's syndrome, Addison's disease, and polycystic ovary syndrome (PCOS). These disorders affect a wide range of physiological functions and can result in diverse clinical manifestations, such as fatique, weight fluctuations, mood alterations, and changes in appetite. Diagnosing these disorders typically involves blood tests, genetic assessments, and in studies. Advances diagnostic imaging including technologies, next-generation sequencing and advanced imaging techniques, have improved the accuracy and speed of diagnosis in endocrine disorders [3].

Therapeutic strategies for hormone disorders, such as Cushing's syndrome, continue to evolve with a combination of traditional pharmacological approaches and surgical procedures. Pharmacological interventions, including drugs like metyrapone and ketoconazole, are used to lower cortisol levels, especially when surgery is not feasible. However, adrenalectomy remains the definitive treatment for conditions like adrenocortical adenomas, providing long-term relief from hypercortisolism and its complications [4].

In addition to these traditional methods, CRISPR-Cas9 technology is being explored for its potential to treat genetic forms of endocrine disorders. This gene-editing tool allows for precise modification of mutations that lead to hormonal imbalances, offering hope for more targeted and effective interventions [5].

#### 2. COMMON ENDOCRINE GLANDS

The endocrine system consists of several glands, each producing specific hormones that target different organs and tissues. The primary endocrine glands include:

The pituitary gland, often referred to as the "master gland," is a small but pivotal endocrine gland situated at the base of the brain, within the bony structure known as the Sella turcica. It is connected to the hypothalamus, a region of the brain that plays a central role in regulating autonomic functions, by a narrow stalk called the infundibulum. This connection is not merely physical but also functional, as the hypothalamus controls the pituitary gland through a complex network of hormonal signals.

**The Pituitary Gland is Divided into Two Main Lobes**: The anterior (front) lobe and the posterior (back) lobe, each with distinct roles and hormonal outputs.

**Anterior Pituitary:** This lobe produces several key hormones that regulate other endocrine glands. The hormones produced by the anterior pituitary include:

**Growth Hormone (GH):** Essential for growth and metabolism.

**Thyroid-stimulating** hormone (TSH): Regulates the thyroid gland, which in turn controls metabolism.

Adrenocorticotropic Hormone (ACTH): Controls the adrenal glands, influencing stress response and the production of cortisol.

Luteinizing Hormone (LH) and Follicle-Stimulating Hormone (FSH): Both are involved in regulating reproductive functions, including the menstrual cycle in females and sperm production in males.

**Prolactin:** Stimulates milk production after childbirth.

**Posterior Pituitary:** Unlike the anterior lobe, the posterior pituitary does not produce hormones but stores and releases two hormones produced by the hypothalamus:

Antidiuretic Hormone (ADH), also known as vasopressin: Regulates water balance by controlling reabsorption of water by the kidneys.

**Oxytocin:** Plays a role in childbirth (labor and delivery) and lactation, and is also involved in social bonding and behavior [6].

The thyroid gland, situated within the neck, is an integral component of the endocrine system, responsible for the synthesis and secretion of hormones that play a pivotal role in modulating metabolic processes, energy generation, and growth. The principal hormones produced by the thyroid gland include:

**Thyroxine (T4):** This hormone is instrumental in regulating metabolic rate and growth.

**Triiodothyronine (T3):** Representing a more active form of thyroid hormone, T3 also plays a significant role in metabolic regulation.

**Calcitonin:** This hormone contributes to the regulation of calcium levels within the bloodstream, thereby assisting in the maintenance of skeletal integrity and systemic calcium homeostasis [7].

The parathyroid glands, located behind the thyroid gland, produce parathyroid hormone (PTH), which regulates calcium and phosphate levels in the blood [8].

# The adrenal glands, located on top of the kidneys, produce several important hormones:

**Cortisol:** Manages metabolism by regulating blood sugar levels, influences immune responses, and helps the body respond to stress by controlling inflammation.

**Aldosterone:** Maintains sodium and potassium balance, thereby regulating blood pressure and fluid balance in the body, essential for normal cardiovascular function.

Adrenaline (epinephrine) and noradrenaline (norepinephrine) are stress-response hormones that increase heart rate, energy, and blood pressure, and prepare the body for "fight-orflight." **Androgens:** Promote the development of male secondary sexual characteristics, including increased muscle mass, and body hair growth [9].

The pineal gland produces melatonin, a hormone that regulates sleep-wake cycles [10].

The organs that are part of the endocrine system include the hypothalamus, which links the endocrine and nervous systems, producing hormones stored in the pituitary gland. The pancreas regulates blood sugar with insulin and glucagon. Adipose tissue secretes hormones like leptin. Ovaries control reproduction and menstruation, while testicles produce sperm and testosterone [11].

## 3. COMMON HORMONE DISORDERS IN ADULT MALES

Hormone disorders in adult males can be broadly categorized into non-malignant and malignant conditions. Each disorder has distinct clinical features, diagnostic criteria, and treatment approaches.

#### 3.1 Non-malignant Disorders

• **Hypogonadism** is characterized by low testosterone levels, which can result from primary testicular failure (primary hypogonadism) or hypothalamic pituitary dysfunction (secondary hypogonadism). Symptoms include reduced libido, erectile dysfunction, fatigue, decreased muscle mass, and osteoporosis [12].

• Hyperthyroidism is the overproduction of thyroid hormones, leading to an accelerated metabolism. Common causes include Graves' disease (an autoimmune disorder) and toxic multinodular goiter. Symptoms include weight loss, heat intolerance, anxiety, palpitations, and tremors [13].

• **Hypothyroidism** is the underproduction of thyroid hormones, leading to a slowed metabolism. Common causes include Hashimoto's thyroiditis (an autoimmune disorder) and iodine deficiency. Symptoms include weight gain, cold intolerance, depression, fatigue, and constipation [14].

• **Cushing's syndrome** results from prolonged exposure to high levels of cortisol. It can be caused by endogenous factors (e.g., pituitary adenomas producing ACTH) or exogenous factors (e.g., prolonged use of corticosteroid medications). Symptoms include central obesity, moon face, hypertension, muscle weakness, and diabetes [15].

• Addison's disease is characterized by insufficient production of cortisol and aldosterone due to adrenal gland dysfunction. Common causes include autoimmune destruction of the adrenal cortex and infections (e.g., tuberculosis). Symptoms include fatigue, weight loss, hypotension, hyperpigmentation, and electrolyte imbalances [16].

 Diabetes mellitus is а group of metabolic disorders characterized by chronic hyperglycemia. Type 1 diabetes results from autoimmune destruction of insulin producing beta cells in the pancreas, while Type 2 diabetes is associated with insulin resistance and relative insulin deficiency. Symptoms include polyuria, polydipsia, polyphagia, fatigue, and blurred vision [17].

#### 3.2 Malignant Disorders

Adrenal cortical carcinoma is a rare and aggressive cancer originating from the adrenal cortex. It can produce excess hormones such as cortisol, leading to Cushing's syndrome, or aldosterone, causing primary hyperaldosteronism. Symptoms depend on the secreted, including type of hormone hypertension, weight gain, and metabolic disturbances [18].

Pituitary tumors can be benign (adenomas) or malignant (carcinomas) and can affect hormone production. Prolactinomas are common the most type. leading to excessive prolactin production and symptoms such as galactorrhea and hypogonadism. Other pituitary tumors may cause overproduction of GH (acromegaly) or ACTH (Cushing's disease) [19].

Thyroid cancer can arise from follicular or parafollicular cells of the thyroid gland. The most common types are papillary and follicular thyroid cancers, which have a good prognosis. Medullary thyroid cancer and anaplastic thyroid cancer are less common but more aggressive. Symptoms may include a neck

mass, difficulty swallowing, and voice changes [20].

#### 4. THE ROLE OF HORMONES AS DIAGNOSTIC AND PROGNOSTIC MARKERS

Hypogonadism and Testosterone Levels: Hypogonadism and Testosterone Levels: Testosterone acts both diagnostically and prognostically. Diagnostically, low testosterone levels confirm hypogonadism when combined with symptoms like reduced libido or muscle mass. Prognostically, testosterone replacement therapy is linked to better symptom relief and improved musculoskeletal health, particularly after therapy [21,22].

Thyroid Disorders and Thyroid Hormones: Thvroid hormones play crucial roles both diagnostically and prognostically. Diagnostically, assessing levels of TSH, free T4, and T3 is essential for detecting thyroid dysfunction, such as hyperthyroidism or hypothyroidism. Prognostically, these hormone levels guide the optimization of treatment, especially in therapies like levothyroxine adjusting or medications. antithyroid Monitoring these levels helps ensure effective management of thyroid diseases, particularly in dynamic situations like post-surgery or during pregnancy [23].

Adrenal Disorders and Cortisol<sup>.</sup> Diagnosis of Cushing's syndrome utilizes tests such as 24-hour urine cortisol, late-night salivary cortisol, and dexamethasone suppression to detect hypercortisolism. In Addison's disease, low cortisol levels accompanied by high adrenocorticotropic (ACTH) levels hormone confirm adrenal insufficiency. Prognostically, cortisol levels predict complications like adrenal crisis and inform management during stress, such as surgery or illness [24].

Diabetes Mellitus Insulin/Glucose: and Diagnosis relies on fasting blood glucose, oral glucose tolerance tests, and hemoglobin A1c (HbA1c) levels. C-peptide levels help differentiate between type 1 and type 2 diabetes by assessing endogenous insulin production. Prognostically, HbA1c levels serve as strong predictors of diabetes-related complications such as nephropathy, neuropathy, and retinopathy. Tight glycemic control, reflected in

lower HbA1c levels, correlates with better outcomes [25].

Pituitarv Disorders and Growth Hormone/Prolactin: Diagnosis of acromegaly involves measuring growth hormone and insulinlike growth factor-1 (IGF-1) levels, while elevated prolactin levels confirm prolactinomas. Prognostically, treatment response, such as somatostatin analogs for acromegalv or dopamine agonists for prolactinomas. is assessed by monitoring growth hormone, IGF-1, or prolactin levels [26].

#### 4.1 Other Hormones and Conditions

**Parathyroid Hormone (PTH):** Used in the diagnosis and management of hyperparathyroidism and monitoring bone health [27].

**Calcitonin:** Elevated levels may indicate medullary thyroid cancer [28].

#### **5. CONCLUSION**

hormones play a pivotal role in the diagnosis and prognosis of endocrine disorders in adult males. From hypogonadism and thyroid disorders to diabetes and adrenal diseases, hormone measurements provide critical information for confirming diagnoses, guiding treatment decisions, and monitoring disease progression or response to therapy. Moreover, in the context of malignancies, hormone levels can serve as valuable biomarkers for early detection, staging, and evaluating the effectiveness of treatment. Overall, the use of hormones as diagnostic and prognostic markers is indispensable in the management of endocrine disorders, ensuring that patients receive timely and appropriate care to improve their quality of life and outcomes.

#### DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc) and text-to-image generators have been used during writing or editing of this manuscript.

#### CONSENT

It is not applicable.

#### ETHICAL APPROVAL

It is not applicable.

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#### **COMPETING INTERESTS**

Authors have declared that no competing interests exist.

#### REFERENCES

- 1. Professional CC. Medical. Hormonal imbalance: Causes, symptoms & treatment. Cleveland Clinic; 2020. Available:https://my.clevelandclinic.org/hea lth/diseases/22673-hormonal-imbalance
- 2. Endocrine News. (2015, September). Heir apparent: A multi-faceted look at one of the most common inherited metabolic disorders, congenital adrenal hyperplasia. EndocrineNews.https://endocrinenews.end ocrine.org/heir-apparent-a-multi-facetedlook-at-one-of-the-most-common-inheritedmetabolic-disorders-congenital-adrenalhyperplasia/<u>.</u>
- National Institute of Diabetes and Digestive and Kidney Diseases. Endocrine diseases. U.S. Department of Health and Human Services.https://www.niddk.nih.gov/healthinformation/endocrine-diseases.
- Di Dalmazi, G., & Reincke, M. (2018). Adrenal surgery for cushing's syndrome: An Update. Endocrinology and metabolism clinics of North America. 2018;47(2):385– 394.

Available:https://doi.org/10.1016/j.ecl.2018. 01.004

5. Zhang B. CRISPR/Cas gene therapy. Journal of Cellular Physiology. 2021;236(4):2459–2481. Available:https://doi.org/10.1002/jcp.30064

 El Sayed SA, Fahmy MW, Schwartz J. Physiology, pituitary gland. In StatPearls. Treasure Island, FL: StatPearls Publishing; 2023.

Available:https://www.ncbi.nlm.nih.gov/books/NBK459247/

7. InformedHealth.org. (2021, June 18). In brief: How does the thyroid gland work? In Institute for Quality and Efficiency in Health Care (IQWiG) (Eds.), InformedHealth.org. Cologne,

Germany.https://www.ncbi.nlm.nih.gov/books/NBK279388/

- 8. Johns Hopkins Medicine. The parathyroid glands. Johns Hopkins Medicine.https://www.hopkinsmedicine.org /health/conditions-and-diseases/theparathyroid-glandsDutt
- 9. M, Wehrle CJ, Jialal I. Physiology, adrenal gland. In StatPearls. StatPearls Publishing; 2023.
- Shochat T, Haimov I, Lavie P. Melatonin-the key to the gate of sleep. Annals of Medicine. 1998;30(1):109–114. Available:https://doi.org/10.3109/07853899 808999392
- 11. Professional CC. Medical. Endocrine system. Cleveland Clinic. Available:https://my.clevelandclinic.org/hea Ith/body/21201endocrine-system (Accessed on:2022b, November).
- 12. Professional CC, Medical. Low testosterone (low T): Causes, symptoms & treatment. Cleveland Clinic; 2022.

Available:https://my.clevelandclinic.org/hea lth/diseases/15603-low-testosterone-malehypogonadism

- Kravets I. Hyperthyroidism: Diagnosis and treatment. American Family Physician. 2016;93(5): 363–370.
- Almandoz JP, Gharib H. Hypothyroidism: etiology, diagnosis, and management. The Medical clinics of North America. 2012;96(2):203–221.
  Available:https://doi.org/10.1016/i.mena.20

Available:https://doi.org/10.1016/j.mcna.20 12.01.005

- Ferriere A, Tabarin A. Cushing's syndrome: Treatment and new therapeutic approaches. Best practice & research. Clinical Endocrinology & Metabolism. 2020;34(2):101381. Available:https://doi.org/10.1016/j.beem.20 20.101381
- Betterle C, Presotto F, Furmaniak J. Epidemiology, pathogenesis, and diagnosis of Addison's disease in adults. Journal of endocrinological investigation. 2019;42(12):1407–1433. Available:https://doi.org/10.1007/s40618-019-01079-6
- Cloete L. Diabetes mellitus: an overview of the types, symptoms, complications and management. Nursing standard (Royal College of Nursing (Great Britain); 2022; 1987;37(1):61–66. Available:https://doi.org/10.7748/ns.2021.e 11709

- Dackiw AP, Lee JE, Gagel RF, Evans DB. Adrenal cortical carcinoma. World journal of surgery, 2001;25(7):914–926. Available:https://doi.org/10.1007/s00268-001-0030-7
- Melmed S. Pituitary tumors. Endocrinology and metabolism clinics of North America. 2015;44(1):1–9. Available:https://doi.org/10.1016/j.ecl.2014. 11.004
- Mazeh H, Orlev A, Mizrahi I, Gross DJ, Freund HR. Concurrent medullary, papillary, and follicular thyroid carcinomas and simultaneous Cushing's syndrome. European thyroid journal, 2015;4(1): 65–68. Available:https://doi.org/10.1159/00036875 0
- 21. Wang C, Swerdloff RS. Testosterone Replacement therapy in hypogonadal men. Endocrinology and metabolism clinics of North America. 2022;51(1):77– 98.

Available:https://doi.org/10.1016/j.ecl.2021. 11.005

- Bhasin S, Brito JP, Cunningham GR, Hayes FJ, Hodis HN, Matsumoto AM, Snyder PJ, Swerdloff RS, Wu FC, Yialamas MA. Testosterone therapy in men with hypogonadism: An Endocrine Society clinical practice guideline. The Journal of Clinical Endocrinology & Metabolism. 2018;103(5):1715–1744. Available:https://doi.org/10.1210/jc.2018-00229
- 23. Royal Australian college of general practitioners. Thyroid disease: Using diagnostic tools effectively. Australian Journal of General Practice; 2021. Available:https://www1.racgp.org.au/ajgp/2 021/january-february/thyroid-diseaseusing-diagnostic-tools-effectively
- 24. Park J, Didi M, Blair J. The diagnosis and treatment of adrenal insufficiency during childhood and adolescence. Archives of disease in childhood. 2016;101(9):860–865.

Available:https://doi.org/10.1136/archdischi Id-2015-308799

 Burke MD. Diabetes mellitus: test strategies for diagnosis and management. Postgraduate Medicine. 1979;66(5):213– 220.

> Available:https://doi.org/10.1080/00325481 .1979.11715306

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- 26. Tritos NA, Miller KK. Diagnosis and management of pituitary adenomas:A Review. JAMA, 2023;329(16):1386–1398. Available:https://doi.org/10.1001/jama.202 3.5444
- 27. Pokhrel B, Leslie SW, Levine SN. Primary Hyperparathyroidism. In StatPearls. StatPearls Publishing; 2024.
- 28. Okamoto T. Gan to kagaku ryoho. Cancer & Chemotherapy. 2001;28(4):561–565.

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