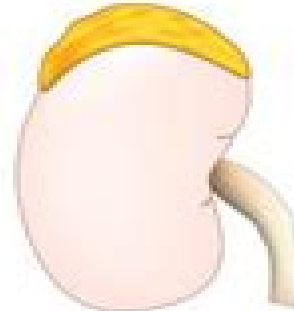
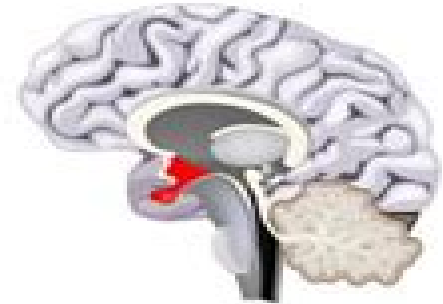


Cushing's disease



Production of large amounts of cortisol by the adrenal glands



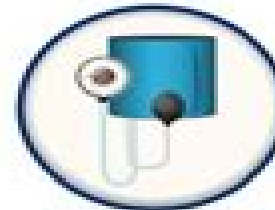
Increased secretion of adrenocorticotropic hormone (ACTH) from the anterior pituitary

Signs and symptoms

Weight gain



High blood pressure



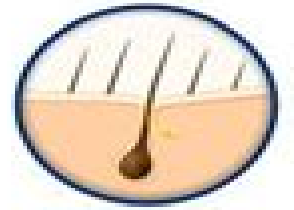
Poor memory



Red stretch marks



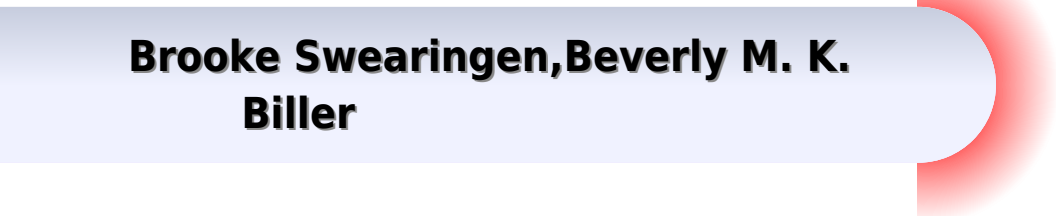
Ruddy face



Excess hair growth

Cushing S Disease Cushing S Disease

**Brooke Swearingen, Beverly M. K.
Biller**



Cushing S Disease Cushing S Disease:

Cushing's Disease Brooke Swearingen, Beverly M. K. Biller, 2011-09-08 In *Cushing s Disease* leading authorities in the field offer a thorough review of the pathogenesis diagnostic algorithm and treatment options for this complex disease Beginning with a fascinating history of Cushing s disease that outlines its historical significance to both endocrinology and neurosurgery the book goes on to cover the full range of important issues including the molecular pathogenesis of Cushing s anatomic pathology the diagnosis of Cushing s syndrome the differential of pseudo Cushing s syndromes hypercortisolemia surgical removal of the corticotroph adenoma post operative management and assessment of remission radiotherapeutic options and the exciting developments in medical therapy In addition the book also addresses Cushing s disease in the pediatric population given that its clinical manifestations and impact on growth can be severe silent corticotroph adenomas as a distinct clinical entity diagnosis and management of Cushing s disease during pregnancy bilateral adrenalectomy and finally the long term psychological manifestations of hypercortisolemia Comprehensive and an invaluable addition the literature *Cushing s Disease* is an essential reference for enhancing diagnosis and treatment of this debilitating disorder

Cushing's Disease Edward R. Laws Jr, Louise Pace, 2016-11-08 *Cushing s Disease An Often Misdiagnosed and Not So Rare Disorder* reviews the epidemiology of Cushing s including statistics on the incidence and prevalence of this disease There are discussions of the signs and symptoms and the most common co morbidities such as diabetes mellitus hypertension osteoporosis amenorrhea and infertility Surgical medical and radiotherapeutic treatments including indications results risks and complications are reviewed Also featured is a chapter on the patient s perspective coping with Cushing s quality of life and psychosomatic issues This book is essential reading for the wide range of physicians who treat patients with Cushing s disease symptoms as well as biomedical researchers who investigate the etiology and mechanisms of rare genetic diseases in particular rare endocrine disorders Reviews the basics of Cushing s disease and its interrelation with hormones the brain and bodily functions Includes chapters on diagnosis surgical medical and radiotherapeutic treatments and variations in presentation including cyclical disease Presents the cognitive and emotional aspects of Cushing s and the long term sequelae Offers an important resource for physicians who are accustomed to treating individual symptoms rather than a disease complex Reviews multidisciplinary management and post treatment management of Cushing s including recommendations for Cushing s Centers of Excellence

Cushing's Syndrome D. T. Krieger, 2012-12-06 The unraveling of our knowledge of the functions of the adrenal gland constitutes one exciting development of modern medicine and biochemistry We owe these advances to the felicitous cooperative efforts of the clinical investigator and the biochemist Three centuries elapsed between the first recorded anatomical description of the adrenals and the demonstration by Dr Addison in the mid nineteenth century of the fatal results of the destruction of these glands by disease It became evident from this observation that the adrenals secreted a factor or factors essential to life It took approximately 90 years to isolate this elusive vital factor cortisone from

beef adrenal cortices independently by both Reichstein and his co workers in Basle and Kendall and his group in the United States and another 10 15 years before it became more generally available for experimental and clinical use It is perhaps difficult to believe that as recently as 35 40 years ago before cortisone and cortisol were clinically available the surgical removal of a benign adrenal cortical tumor in patients with Cushing s syndrome was associated with a prohibitive postoperative mortality rate Within 12 36 h after operation most of such patients developed an intractable state of shock which was not manifested by significant electrolyte abnormalities or hypoglycemia and was unresponsive to the usual treatment for shock plus the generous use of salt retaining hormone **Cushing's Syndrome** Marcello D.

Bronstein,2010-10-01 Cushing s syndrome is a relatively rare clinical disorder that is associated with many co morbidities such as systemic hypertension diabetes osteoporosis impaired immune function and growth impairment in children all of which severely reduce quality of life and life expectancy Cushing s Syndrome Pathophysiology Diagnosis and Treatment reviews the difficulties in distinguishing Cushing s syndrome from these and other common conditions such as central obesity menstrual irregularity and depression It also provides state of the art information on various strategies to establish the diagnosis of Cushing s syndrome and the differential diagnosis among its diverse etiologies as well as therapeutic approaches Additionally a range of conditions that represent challenges for the diagnosis and treatment such as renal failure pediatric age cyclic hypercortisolism and pregnancy are covered in detail A valuable resource not only for endocrinologists but also internal medicine physicians gynecologists pediatricians pituitary surgeons and urologists Cushing s Syndrome Pathophysiology Diagnosis and Treatment provides insights by experts that will help all physicians dealing with Cushing s syndrome to expand their knowledge about the condition and provide targeted comprehensive care **Prolactin** E.

Flückiger,E. DelPozo,K. v. Werder,1981-12-01 Lactogenic hormone activity was first observed in bovine pituitary extracts by Stricker and Griiter in 1928 working in Bouin s laboratory in Strasbourg Since that time prolactin has been shown to exist in anterior pituitary extracts of almost all vertebrate species investigated Although its biology was extensively studied in many mammalian species the existence of prolactin in the human was generally doubted despite the positive evidence produced by such researchers as Pasteels This can partly be explained by the fact that human growth hormone isolated in 1961 is itself a potent lactogen in contrast to nonprimate growth hormones and is present in the normal human pituitary in much greater amounts than prolactin As a result there was a lag of nearly 10 years until prolactin was unanimously accepted as a hormone of the human pituitary separate from human growth hormone In 1970 new bioassay techniques permitted the demonstration of prolactin bioactivity in the serum of postpartum women and galactorrhea patients and chromatographic methods led to the isolation and purification of human prolactin allowing the establishment of a specific radioimmunoassay for this hormone This opened the road to the understanding of prolactin physiology and pathophysiology in the human which has revolutionized clinical neuroendocrinology and reproductive endocrinology Particularly hyperprolactinemia has turned out to

be one of the most common endocrine syndromes

Cushing's disease and aggressive pituitary tumours

Daniel Bengtsson, 2021-04-07 This thesis focuses on clinical and epidemiological aspects of aggressive pituitary tumours carcinomas and Cushing's disease Pituitary carcinomas account for only 0.1-0.2% of the tumours originating from the anterior pituitary gland and are defined solely by the event of distant metastases whereas aggressive pituitary tumours are defined by their clinical behaviour of rapid progressive growth despite optimal treatment with surgery radiotherapy and medical agents The prognosis for individuals with aggressive tumours carcinomas has been poor with few treatment options However case reports indicated better outcomes after treatment with the alkylating agent temozolomide In study I and III we investigated 24 patients 16 aggressive tumours and 8 carcinomas given treatment with temozolomide We found an initial response rate tumour regression 30% in 10/21 evaluable patients with complete regression in two carcinomas Favourable response was associated with low tumour expression of the DNA repair protein MGMT in responders median 9% range 5-20% vs non responders median 93% 50-100% Our results also indicated a longer survival in patients with low MGMT Out of 11 patients with MGMT 10% nine died with an estimated median survival of 26 months 95% CI 14-38 whereas only 1/6 patients with lower MGMT died from tumour progression during a follow up of median 83 months range 12-161 One of the patients in study I and III had a corticotroph pituitary carcinoma and in addition Lynch syndrome LS a hereditary cancer predisposing syndrome caused by germline mutations in DNA mismatch repair MMR genes and primarily associated with colon and endometrial carcinomas In study II we investigated the characteristics of the pituitary carcinoma and found loss of MSH2 and MSH6 protein expression consistent with the patient's germline mutation in MSH2 This was the first published case of a pituitary tumour associated with LS In addition we identified all known Swedish patients with LS n=910 and searched for diagnostic codes consistent with a pituitary tumour in the Swedish national patient register We found in total three patients with clinically relevant pituitary tumours the reported prevalence in the background population is around 1/1000 The last two studies in the thesis focused on Cushing's disease CD i.e. an ACTH secreting pituitary tumour resulting in excess levels of cortisol CD is associated with multiple comorbidities and increased mortality The reversibility of comorbidities and mortality risk after remission of cortisol levels have been under debate Study IV examined psychiatric consequences of CD measured by the use of psychotropic drugs 179 patients with CD and a quadrupled matched control group were followed from diagnosis and at 5 and 10 year follow up We found that use of antidepressants remained at around 25% of patients with CD regardless of remission status at diagnosis and follow up whereas drugs for somatic comorbidities decreased Use of antidepressants sleeping pills and anxiolytics was higher in patients with CD compared to controls at diagnosis and 5 year follow up A cross sectional analysis of 76 patients in sustained biochemical remission for median 9.3 years showed that 25% were taking antidepressants a significantly higher use than controls OR 2.0 95% CI 1.1-3.8 In addition patients with CD had a higher use of psychotropic drugs already in the 5 year period before diagnosis Study V investigated mortality and causes of

death in 371 patients with CD compared to a quadrupled matched control group Follow up was median 10.6 years IQR 5.7-18.2 after time of diagnosis Overall mortality was increased in patients with CD HR 2.1 95% CI 1.5-2.8 and remained elevated for patients in remission at last follow up n 303 HR 1.5 1.02-2.2 For patients not in remission n 31 HR was 5.6 2.7-11.6 Cardiovascular diseases 32/66 and infections 12/66 were overrepresented causes of death in patients with CD Main conclusions of the thesis Temozolomide improves outcome in patients with aggressive pituitary tumours carcinomas and a low MGMT expression in the tumour predicts a favourable outcome As additional therapies evolve MGMT may help to tailor the treatment Germline mutations in MMR genes may contribute to the development and clinical course of pituitary tumours and may be a novel cause of hereditary pituitary tumours Patients with Cushing's disease have a high use of psychotropic drugs that remains elevated despite achievement of biochemical remission suggesting persisting negative effects on mental health and highlighting the need for long term monitoring of psychiatric symptoms In addition psychiatric symptoms may be early and important signs of CD Efforts to achieve biochemical remission are crucial to reduce mortality in CD However patients in remission still have an increased mortality compared to controls This underscores the need for life long monitoring and treatment of associated comorbidities in patients with CD *Cushing's Syndrome* Lewis S.

Blevins, 2012-12-06 Cushing's Syndrome provides the reader with an update on the clinical presentation diagnosis and treatment of patients with Cushing's syndrome Molecular mechanisms of pituitary and adrenal causes of Cushing's syndrome are reviewed in detail Successful diagnostic and treatment strategies that have been employed by readers in the field are recommended and discussed Numerous advances in the pathophysiology and diagnosis of Cushing's syndrome speak to the timeliness of this volume that has been penned by experts in the field Handbook of Nurse Anesthesia John J.

Nagelhout, Karen L. Plaus, 2010-01-01 A practical reference for the clinical setting Handbook of Anesthesia 5th Edition puts key information at your fingertips It's an ideal companion to Nurse Anesthesia providing easy access coverage of the subjects you look up most frequently such as common diseases common procedures drugs and protocols It also includes the most current information related to perioperative management and anesthetic care considerations for a wide variety of surgical procedures Written by leading CRNA experts John Nagelhout and Karen Plaus this compact handbook provides the up to date clinical and drug information you'll use in daily practice This book is packed with information is easy to read and easy to navigate The clinical information is authentic and reads like the work of authors with clinical experience and wisdom Reviewed by British Journal of Anaesthesia Date Nov 2014 Convenient quick reference format is organized into three parts common diseases common procedures and drugs Disease monographs include definition incidence and prevalence pathophysiology laboratory results clinical manifestations treatment anesthetic considerations and prognosis Monographs of surgical procedures include a brief description preoperative assessments including health history physical assessment and patient preparation anesthetic technique perioperative management and postoperative implications Drug monographs

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Equine Medicine and Management Cody W. Faerber,2001 **Cushing's Syndrome, an Issue of Endocrinology and Metabolism Clinics of North America** Adriana G. Ioachimescu,2018-06-28 This issue of Endocrinology and Metabolism Clinics edited by Adriana G Ioachimescu will focus on Cushing s Syndrome Topics include but are not limited to 1 Diagnosis of Cushing s syndrome in the modern era Gene mutations in patients with Cushing s syndrome Morbidity in Cushing s syndrome and impact of treatment Localization of ACTH dependent Cushing s syndrome Prognostic factors of long term remission after surgical treatment of Cushing s disease Outcomes of pituitary radiation for Cushing s disease New molecular targets in Cushing s disease Recent advances in subclinical Cushing s syndrome Adrenal surgery for Cushing s syndrome Adrenocortical carcinoma with hypercortisolism Ectopic Cushing s syndrome Medical therapy for Cushing s syndrome in the 21st century Pregnancy in patients with Cushing s syndrome Mortality in patients with Cushing s syndrome and Cushing s syndrome in children and adolescents

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Cushing S Disease Cushing S Disease Introduction

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he also requested arjuna and krishna to take benedictions as he was very much pleased by their valour arjuna asked indra for many divine weapons and mantras to control them indra granted them all he asked krishna to take some benediction krishna asked the following beautiful benediction and it is worth meditating for all aspiring devotees

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