Efficacies of Eruca Sativa and Raphanus Sativus Seeds’ Oils in Streptozotocin-Induced Diabetic Rats

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Objective: The study was planned to investigate the effect of Eruca sativa seeds’ oil (ESSO) and Raphanus sativus seeds’ oil (RSSO) on impaired glucose tolerance, lipid profile and oxidative stress in streptozotocin-induced diabetic albino rats.
**Agent or Vitamin D and Hypocalcemia or Hypophosphatemia?**

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Author(s): Abdelwahab T H Elidrissy*

Anemia due to iron deficiency is commonly associated with rickets, but rarely myelofibrosis was seen in infants with rickets in the hypocalcaemic phase. The aim of this review is to elucidate the mechanism of development of myelofibrosis in rickets. We reviewed the literature in PubMed with keywords myelofibrosis, hypocalcemia and anemia. The cases diagnosed as mye ...
60-Year-Old Man with Pheochromocytoma and Clinical Picture of Depression

Author(s): Urszula Ambroziak*

Introduction: Pheochromocytoma is an adrenal gland tumour, which usually produces catecholamines. The classical triad of clinical symptoms consists of palpitations, headaches and profuse sweating. Other symptoms include: hypertension, anxiety, pallor, nausea, weakness. However, it can be asymptomatic. Because of unspecific symptoms the diagnosis of this rare neuroendo ...

Kocher-Debre-Semelaigne Syndrome: Response to Thyroxine Replacement Therapy

Author(s): Vishal V Tewari*, Ritu Mehta, Kunal Tewari

Introduction: Congenital hypothyroidism with muscular pseudohypertrophy or Kocher-Debre-Semelaigne syndrome is the result of long standing untreated moderate to severe hypothyroidism. The pathogenesis of this muscular pseudohypertrophy is unknown and it is usually noted in the muscles of the extremities, limb girdle, trunk, hand and feet but is most evident in the mus ...

Congenital Generalized Lipodystrophy: A Multisystemic Metabolic Disorder

Author(s): Vishal V Tewari*, Ritu Mehta, Kunal Tewari
Introduction: Congenital generalized lipodystrophy or Berardinelli-Seip syndrome is a rare autosomal recessive multisystem disorder characterized by the near absence of subcutaneous and visceral adipose tissue from birth or early infancy with severe insulin resistance. It is caused by mutations in the gene for AGPAT-2 on chromosome 9 or BSCL-2/Seipin on chromosome 11 ...