Aarskog-Scott Syndrome: Clinico-Radiological Illustration of a Rare Case

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Aarskog-Scott Syndrome is a rare syndrome and is estimated to occur in 1 in 1 million individuals in general population. It is transmitted in an X-linked recessive manner and occurs due to FGD1 gene mutation. It has facial, genital and digital hands symptoms, musculoskeletal anomalies and affected growth.

A Rare Cause of Oligohydramnios: A Case Report

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Anamnios is due to extrinsic and intrinsic conditions. Intrinsic causes include maternal and fetal abnormalities mostly due to cystic renal changes or absence of kidney. Tubular dysgenesis characterized by a lack of proximal tubules should be considered.

Hemlock Poisoning in Emergent Patients

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Background: Although poisoning from the ingestion of toxic plants is rarely encountered in Emergency Departments, it can cause serious complications and even death. It is difficult to recognize, diagnose, and differentiate if the patient is
Dilemma in Managing Thyroglossal Duct Cyst Carcinoma

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The thyroglossal duct cyst (TGDC) is a well-known developmental abnormality encountered in the neck. It represents over 75% of childhood midline neck masses and 7% of the adult population. It typically presents as a mobile, painless mass in the anterior midline of the neck, usually in close junction to the hyoid bone.