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Extract with Winrar.Introduction {#sec1-1} ======== Situs

1/1,000,000 live births and the majority of cases are sporadic \[[@ref1]\]. The clinical implications of SIT include defects in organ placement at the outset of embryonic development. Despite the multiple cardiac anomalies, SIT does not frequently manifest as a primary clinical problem in the neonatal period, though the onset of cardiopulmonary disease in infancy is more common \[[@ref2]\]. We present a case of SIT, where the patient had features of obstructive sleep apnoea syndrome (OSAS) and severe scoliosis requiring multiple surgical procedures during growth. Case report {#sec1-2} ======= A 7-year-old male patient presented to the outpatient clinic with complaints of excessive sleepiness, nasal discharge, snoring and recent intermittent laryngospasm. His past medical and family histories were noncontributory. He was the first-born child of non-consanguineous parents and his growth and development were normal. On general physical examination, he was found to have bulky neck with a length of 14 cm, and a head circumference of 44 cm. He had axillar hypertonia, exophthalmos, palatal arch, high-arched palate, over-riding nose, bilateral optic nerve atrophy and right pes anserinus atrophy. He had gastroesophageal reflux and hypertrichosis. His blood pressure was 110/70 mmHg, and heart rate was 80 beats/min. The respiratory and cardiovascular systems were normal. The patient was diagnosed as SIT and was advised to undergo periodic review, but unfortunately, he was lost to follow-up. Two years later, he returned with a progressive scoliosis and at presentation, he had a body mass index of 26.8 kg/m². The full-length spine radiography showed a sacral shift, gross barrel-shaped chest, thoracic inlet angle of 120°, mild severe right convex scoliosis and mild left convex scoliosis. Spirometry showed a forced vital capacity of 0.91L (46% of normal predicted), forced expiratory volume in the first second of 1.32L (54% of normal

inversus totalis (SIT) is a very rare congenital abnormality. It is noted in

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