656 ENHANCEMENT OF LINEAR GROWTH AND WEIGHT VELOCITY BY CYPROHEPTADINE IN CHILDREN WITH IDIOPATHIC GROWTH HORMONE DEFICIENCY RECEIVING THIS HORMONE
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Background The current study examined the hypothesis that Cyproheptadine can enhance linear growth and increase weight velocity in children with idiopathic GH deficiency.

Method 10 children with idiopathic GH deficiency received Cyproheptadine 0.5mg/kg three times per week plus GH1.6U/kg/day for six days a week for1-year period, alternatively (GH plus Cyproheptadine for first and third trimesters, and GH plus placebo for second and fourth trimesters). Weight velocity and linear growth were assessed at baseline and end of every trimester.

Result The repeated measure ANOVA test showed significant differences in weight velocity across the study trimesters so that the increase in weight velocity within first and third trimesters were significantly higher than other time intervals (ΔWW, first trimester 1.51±0.61kg; second trimester 1.13±0.46 kg, third trimester 1.87±0.65kg; and fourth trimester 0.74±0.34 kg, p=0.026). Similar trend was also observed regarding changes in linear growth that the increase in children height was significantly higher in first and third trimesters as compared with other trimesters (Δheight, GH first trimester 3.1±0.7 ’y’; second trimester 2.07±0.25 ’y’; third trimester 1.75±0.2 ’y’ for RU, C and S bones, respectively, in control (p<0.001), and S bones by a mean 3.5±0.9 ’y’ on hGH and 3.15±0.7 ’y’ in control (p<0.05)). In GHD over 3 y of GH treatment, RU advanced by a mean 3.1±0.4 ’y’ (p<0.10). C advanced by a mean 4.3±0.2 ’y’ (p<0.001), and S bones 2.9±1.0(p<0.06).

These results suggest that GH strongly regulates and GHD interferes with bone maturation by inhibiting chondral osteogenesis and less so through delayed enchondroplasia, observed by RU and S maturation. These profiles help in the diagnosis of GHD.

658 CONGENITAL ANOMALIES IN INFANTS WITH CONGENITAL HYPOTHYROIDISM IN QAZVIN, IRAN
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Background and Aims In the last decade a high frequency of extrathyroidal congenital anomalies has been reported in infants with congenital hypothyroidism detected by neonatal screening. Approximately 10% of infants with congenital hypothyroidism have associated congenital anomalies. Cardiac anomalies are most common, but anomalies of the nervous system and eye have also been reported. The main objective of this study was identification of associated disorders in patients with congenital hypothyroidism.

Methods This descriptive study was performed on newborns with congenital hypothyroidism that had been diagnosed by screening in Qazvin province of Iran from 1385 to 1390. Variables were analyzed by Chi-square test.

Results In the screening 287 newborns with congenital hypothyroidism were diagnosed –122 female (42.2%), 86 (12.5%) of 287 patients had congenital anomalies. 10 (3.5%) Down syndrome, 10 (3.5%) cardiovascular anomalies, 9 (3.1%) hearing disorder, 2 (0.7%) eye disorder, 2 (0.7%) metabolic disease, 2 (0.7%) disorder of face and ear, 2 (0.7%) CNS disorder, one (0.3%) CAH, one (0.3%) panhypopituitism and one (0.3%) pseudohypoparathyroidism were diagnosed.

Conclusions With respect to the association of congenital anomalies with congenital hypothyroidism, every newborn with congenital hypothyroidism should be carefully examined for cardiovascular disorders and other related anomalies.

659 HYPOPARATHYROIDISM AND PSEUDOHYPOPARATHYROIDISM: ABOUT 9 CASES
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Introduction Hypoparathyroidism (HP) is an uncommon disorder of calcium metabolism characterized by hypocalcemia and hyperphosphatemia due to impaired parathyroid hormone (PTH) secretion. The pseudo-hypoparathyroidism (PHP) is characterized by a high level of PTH due to peripheral PTH resistance.

Objective Describe clinical, biochemical, radiological profile, treatment and outcome in 9 patients with HP and PHP seen over a period of 24 years.

Methods This study was performed in the pediatric department of Sfax during 24 years (from January 1988 to April 2012). The diagnosis of HP and PHP was based on demonstration of hypocalcemia, hyperphosphatemia with low or elevated PTH, respectively.

Results During the period of study, 9 children having HP or PHP were admitted. There were 5 girls and 4 boys. The mean age at presentation was 44 months (15 days– 10 years). The most common presenting manifestation was seizures (8cases) followed by carpopedal spasm in one case. The mean serum calcium and inorganic phosphate concentrations were 1, 51 mmol/l and 3 mmol/l,