Ellis-van Creveld syndrome is a rare congenital genetic disorder having autosomal recessive inheritance. It is a syndrome affecting the Amish population of Pennsylvania in USA with prevalence rate of 1/5,000 live at birth. In non-Amish population, the birth prevalence is 7/100000. The syndrome is characterized by bilateral postaxial polydactyly of the hands, chondrodysplasia of long bones resulting in acromesomelic dwarfism, ectodermal dysplasia affecting nails as well as teeth and congenital heart malformation. There were very rare reports (1,2,3) of this syndrome in dentistry. 16 year old girl, 4th in birth order with a history of polydactyly and neonatal death in one of her siblings, born of non-consangious marriage presented to our clinic for evaluation of short stature. On examination she had a height of 134 cms; upper segment 74 cms, lower segment 60 cms, with an arm span of 76 cms and waist circumference of 89 cms. She had hemidystrophy of the face on left side ,large tongue, high arched palate; lips attached to the gums by 2 frenular bands, microdontia, congenitally missing teeth, peg like dysplastic teeth with caries ,discolored, dystrophic and hypoplastic nails of fingers and toes, 6 digits in both hands (Fig:1 & 3) and left foot (Fig-2) with syndactyly in left toe. Thus, our case had all these limb anomalies. The effective management of this syndrome will require a team which includes pedodontist, oral and maxillofacial surgeon, prosthodontist, clinical geneticist, cardiologist, pulmonologist, orthopaedician, urologist, psychologist, pediatrician, and pediatric neurologist.

References