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Congenital brain tumors, a series of seven patients

Farideh Nejat; Syed Shuja Kazmi; Shahin Behjati Ardakani

Abstract:

Objective: Congenital brain tumors are very rare. We review these tumors in patients younger than 2 months diagnosed in our Department. **Material & Methods:** Seven congenital brain tumors were diagnosed during five years. Clinical and radiological findings and prognosis are analyzed. **Findings:** The study included 5 female and two male infants. Two cases were diagnosed antenatally by means of ultrasonography. All patients presented with intracranial hypertension. The tumor was non-homogenous with cystic and solid components in all neuroimaging, except for the case with choroid plexus papilloma. Hydrocephalus was evident in all of them. Most findings were infra-tentorial lesions. There were three teratomas, one primitive neuro-ectodermal tumor, one ependyoblastoma and one choroid plexus papilloma. Six patients were operated on, with one intra-operative death. Two passed away postoperatively with aspiration pneumonia. One patient died due to complications of chemotherapy and another one due to tumor recurrence one year after surgery. Only the patient with choroid plexus papilloma is alive after 2 years. **Conclusion:** Today, the availability of noninvasive imaging procedures such as computerized tomography scan and magnetic resonance imaging has improved the diagnosis of congenital brain tumors. In spite of development in prenatal diagnosis, appropriate pre and post operative management, the mortality associated with these tumors still remains high. The final prognosis in these patients is still discouraging despite early surgery and operative and anesthetic improvements. Choroid plexus papilloma accompanies the best prognosis, whereas teratoma and primitive neuroectodermal tumors have the worst prognosis.

Keywords:

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