

2.2 Cytogenetics

Nonrandom structural aberrations, most often involving chromosomes 1 and 12 have been described in germ cell tumors [21]. The chromosomal patterns of malignant ovarian and testicular germ cell tumors are alike, and thus, these tumors are similar and may have the same origin and pathogenesis. The cytogenetic anomalies described are a hyperdiploid/hypertriploid chromosomal number, presence of t(12p), and overrepresentation of chromosomes [3, 21–23]. Investigations in embryonal and testicular germ cell tumors show that they arise somatically from either a somatic or a germ cell [22, 23].

Transcription factors GATA-4 and GATA-6 are expressed in pediatric yolk sac tumors and in teratomas [22, 23]. They play critical roles in mammalian yolk sac differentiation and function. GATA-6 is expressed not only in many yolk sac tumors but also in nonmalignant tissues including gut and respiratory epithelium, effusion cells, and neuroepithelium in mature and immature teratomas. GATA-6 can be used to identify yolk sac components in pediatric germ cell tumors. In addition, it is expressed in specific tissues in teratomas [12, 23].

2.3 Teratoma

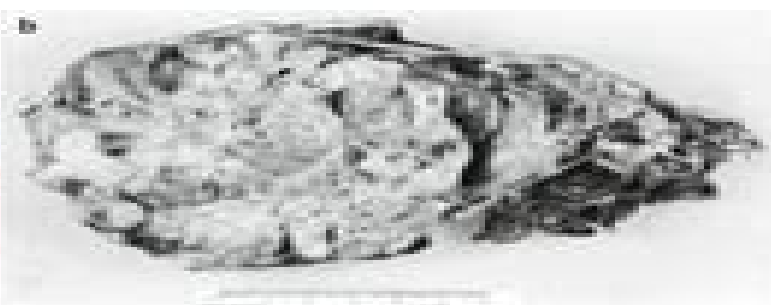
Teratoma is defined as a true tumor composed of multiple tissues foreign to, and capable of growth in areas of, those characteristic of the part from which it is derived [10].

However, it is sometimes difficult to make a distinction between teratomas and structures that result from abortive attempts at twinning. An even progression can be traced from normal twins to conjoined twins, parasitic twins, fetus-in-fetu, and teratomas [10]. Careful studies reveal a break in the progression from an oriented, longitudinal, partially symmetrical structure of a twin to the jumbled, disoriented, irregular growth of a teratoma in which one, two, or three tissues predominate [10]. Despite the apparent progression from twins to fetus-in-fetu to teratomas, some investigators feel that this relationship does not exist. This distinction is based primarily on the fact that teratomas are capable of independent growth, whereas structures included under malformations are limited in their potentiality for growth to a rate similar to the part of the body they resemble [10].

Teratomas are observed in several locations at birth, but the most common ones are the sacrococcygeal, cervical, and retroperitoneal areas [1–4, 7, 12, 14, 15, 17] (Table 2.7) (Figs. 2.2, 2.3, 2.4, and 2.10). Other locations are the brain, anterior mediastinum, stomach, heart and pleura, pharynx, base of the skull, upper jaw, gonads, pelvis, liver, and subcutaneous tissue [3, 4, 7, 10, 15, 24–28] (Figs. 2.5, 2.6, 2.8, 2.9, 2.10, 2.11, and 2.12). Intracranial teratomas are not uncommon in the fetus and newborn [14, 27]. The types of tissues found in fetal and infant teratomas are practically the same regardless of the site of origin [10, 15].



Fig. 2.2 Solid sacrococcygeal teratoma. Sonography in 30 weeks gestation revealed a large abnormality in the caudal region. Shortly before delivery, fetal distress was noted and a cesarean section was performed. (a) The 4.8 kg female was born at 37 weeks gestation with a large sacrococcygeal tumor. (b) The tumor, removed at 1 day of age, weighed 1.5 kg, 13.7% of the baby's birth weight, and measured 13 × 13 × 10 cm.



The dissected tumor with a variegated solid and cystic appearance. Mass partially covers the periphery of the specimen. Microscopically, the tumor consists of immature neuroepithelial elements in addition to a variety of mature tissues typically found in these teratomas (see Figs. 2.13, 2.14, 2.15, 2.16, 2.17, 2.18, 2.19, 2.20, 2.21, and 2.22) (Reprinted from Huser [29], with kind permission of G. Springer-Verlag, 2002).

Tumors Of The Fetus And Infant Tumors Of The Fetus And Infant

Hart Jr. Isaacs



Tumors Of The Fetus And Infant Tumors Of The Fetus And Infant:

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