BACKGROUND Primary immature mediastinal teratomas pose a diagnostic challenge, as they are extremely rare whereas can be cured if promptly diagnosed. The authors report a case of immature mediastinal teratoma in a dead newborn.

METHOD AND RESULTS

A 36 years old pregnant woman was admitted at 32-weeks gestation and had a cesarean section with an outcome of a dead newborn. A previous ultrasound examination showed a living fetus with generalized edema. A necropsy was performed on the newborn according to the Perinatal Autopsy Manual (AFIP - 1983). Received was a dead fetus of 2780,0 g of weight, measuring 43,0 cm of length. The inspection of the thoracic cavity showed a 88,0 g, 7,5 x 6,0 x 3,5 cm, bosselated, pink - tan tumor mass in thymic topography, displacing the thoracic organs. All tissues were immersion fixed in formalin, routinely processed, embedded in paraffin and were sectioned and stained with hematoxylin and eosin. Histological samples showed mature epithelial and mesenchymal tissues and also mesenchymal and immature neuroepithelium. A diagnosis of immature teratoma was made.

CONCLUSION

Primary immature mediastinal teratomas are very rare, comprising 1% of all mediastinal teratomas. Immediately after the delivery, these tumors may cause life threatening respiratory obstruction and must be promptly diagnosed and treated, as its prognosis is excellent, without recurrence on the follow-up.
Primary germ cell tumors of the mediastinum in newborns are unusual neoplasms with histopathologic features that are similar to those of germ cell tumors in the gonads. In this location, all types of germ cell tumors were described: seminomatous, non-seminomatous, teratomas, nonteratomatous germ cell tumors (yolk sac tumors, embryonal carcinomas, and choriocarcinomas) and combined germ cell tumors without teratomatous components (1). From those types, the most common were mediastinal teratomas.

Mediastinal teratomas are uncommon in infants and children, constituting 7% to 10% of all teratomas in this age group (2,3). The overwhelming majority of patients are male (1). In newborns, immature teratomas are rare and constitute less than 1% of all mediastinal teratomas. Other uncommon sites where immature teratomas are reported in early infancy include brain (4), neck (5), pharynx (6), stomach (7) and mesentery (8).

Review of literature revealed that respiratory distress is the most common presentation of immature teratoma of mediastinum in newborn (3,4,9). Chest X-ray often demonstrated a mediastinal mass, some of which are calcified, but a definite diagnosis of teratoma is established only after microscopic examination (3).

Im mature teratoma is characterized by the presence of elements that resemble embryonic tissues, including neurogial or neuro-epithelial components that may coexist along with mature tissues. In most instances, immature teratomas occurring in the fetus and newborn are associated with a favorable
prognosis (2,9,10). The behavior of immature teratoma in adolescents and adults is less predictable and may be associated with poor clinical outcome (9). Dehner cites a 15% overall malignancy rate for mediastinal teratomas diagnosed in the pediatric age group (2). However, other workers concluded that the mediastinal teratomas, both mature and immature, occurring in newborns and infants behave in a benign fashion, if resectable, and are associated with a favorable prognosis as compared to those occurring in adolescents and adults (3,9,11). In our case, the prenatal diagnosis was not made and the fetus died in utero.

Conclusiones

Primary immature mediastinal teratomas are very rare, comprising 1% of all mediastinal teratomas. Immediately after the delivery, these tumors may cause life threatening respiratory obstruction and must be promptly diagnosed and treated, as its prognosis is excellent, without recurrence on the follow-up.

Bibliografía