Thymoma and thymic carcinoma in children and adolescents: A report from the European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT)

Teresa Stachowicz-Stencel, Daniel Orbach, Ines Brecht, Dominik Schneider, Ewa Bien, Anna Synakiewicz, Julien Rod, Andrea Ferrari, Giovanni Cecchetto, Gianni Bisogno

- 16 children with thymoma (median age 11 years) and 20 with thymic carcinoma (median age 14 years)

- At diagnosis complete primary resection was possible in 11 patients with thymoma and 1 with thymic carcinoma

- Chemotherapy with various regimens was given to 22 children (17 as neoadjuvant); 8 patients with carcinoma had radiotherapy

- 17 children died (15 thymic carcinoma, 2 thymoma)

- 5-year overall survival for patients with thymic carcinoma is 21%

Conclusions

Thymoma generally has good prognosis; radical surgery is the treatment of choice

Thymic carcinoma is associated with paediatric patients to give a very poor prognosis independently of intensive multimodal management: surgery is the most important issue, eventually associated to radiotherapy. The role of chemotherapy is unknown (it seems that it is worth using chemotherapy regimens with cisplatin, etoposide and ifosfamide)

Multidisciplinary, multicenter approach and collaboration with adults’ physician are necessary in order to propose homogenous guidelines.