found in partial or complete androgen insensitivity syndrome. The aim of this study was to determine whether even the most minor forms of isolated hypospadias are associated with AR mutations and thus whether all types of hypospadias warrant molecular analysis of the AR.

**Methods** 292 Caucasian children presenting with isolated hypospadias (no micropenis, no cryptorchidism) and 345 controls were included prospectively. Mutational analysis of the AR through direct sequencing (exons 1–8) was performed.

**Results** Five missense mutations of the AR were identified in 9 patients with glandular or penile anterior (n=5), penile midshaft (n=2) and penile posterior (n=2) hypospadias, i.e., 3%; p.Q58L (c.174 G>T), 4 cases of p.F592S (c.1776 C>T), 2 cases of p.A475V (c.1425 C>T), p.D551H (c.1651 G>C) and p.Q799E (c.2397 C>G). None of these mutations was present in the control group. Four of these novel findings since 1 has never been reported to date (p.D551H) and 3 have never been reported in patients with genital malformation but only in isolated infertility: p.Q58L, p.F592S, and p.A475V. It is notable that microgenit, a cardinal sign of AIS, was not present in any patients.

**Conclusion** AR mutations may play a role in the cause of isolated hypospadias, even in the most minor forms. Identification of this underlying genetic alteration is important for proper diagnosis and may significantly improve the follow-up of these patients during puberty, especially regarding future fertility.

**THE FEASIBILITY OF TISSUE EXPANDERS IN RECONSTRUCTION OF GIANT CONGENITAL MELANOCYTIC NEVI IN CHILDREN**

Background Children with Giant Congenital Melanocytic Nevi (GCMN) carry a great challenge to the pediatric and reconstructive surgeons to cover the widely exposed area after its excision. A variety of treatment options exists for the management of such cases. In this prospective review of selected group of children who had a GCMN of their abdominal walls were managed with implantation of tissue expanders (TE) for staged reconstruction, patients evaluated with respect to complications, general and esthetic patient and parents satisfaction.

Objective Our purpose was to study the feasibility of use tissue expanders in the management of children who had GCMN with special emphasis to the complications and children and their parents satisfaction.

Material and Methods Retrospective data collection of the 12 patient’s charts, operative data of 86 surgical procedures and follow-up visits were calculated and analyzed, with Statistical analysis done using the Student t test, and P<0.05 was considered statistically significant.

Results In a period of 4 years, from 2004 to 2008, the results of using 37 different sizes tissue expanders and 86 operative procedures in a 12 child, 9 boys and 3 girls aged from 2 to 12 years with different types of abdominal wall GCMN will be discussed.

Conclusion Tissue expander is a useful and feasible tool for reconstruction of the abdominal wall in cases of GCMN in children. Parents and children satisfaction and body image are generally acceptable.

**ROLE OF THORACOSCOPY IN THE TREATMENT OF LOCULATED PLEURAL EMPYEMA IN PEDIATRIC PATIENTS: ABOUT 73 CASES**

Objective Evaluate the results of thoracoscopy for the treatment of located pleural empyema.

Methods Retrospective study of 73 patients with located pleural empyema in pediatric surgery department of Monastir during the period between 1997–2010. Located empyema was confirmed in all patients through imaging (ultrasonography or tomography of chest).

Results The age of patients ranged between 2 months-16 years (mean age: 41.5 months) with a sex ration of 1:2.8. Fever was present in 98% of the cases. 25 patients had severe respiratory distress and mechanical ventilation was necessary in 4 cases. Bacteriology of the pleural liquid was positive in 12 cases. All patients had intravenous antibiotics therapy during a mean period of 20 days. Sixteen patients (22%) were treated by thoracic drainage. Intrapleural fibrolysis was indicated in 5 cases (7%). One patient had an open thoracotomy for lung abscess with a located pleural empyema. Thoracoscopy was performed in 51 cases (70%). It was indicated after thoracic drainage failure in 14 cases. It was indicated for first-line before any prior thoracic drainage in 37 cases because of a multiloculated aspect at ultrasonography.

Conclusion Management of pleural empyema is still controversial in children and surgical indication is often delayed. Early first-line thoracoscopy yields a better clinical outcome for pediatric patients with pleural localized empyema, with apparent decreased morbidity, earlier