A huge abdomino-pelvis tumour in a prepubertal girl: A case report of dysgerminoma infiltrating both ovaries and uterus in an 11-year-old girl in Maiduguri

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Abstract

Dysgerminoma is one of the rare highly malignant ovarian germ cell tumour. It originate from undifferentiated germ cells that are similar to primordial germ cells, and it is identical to testicular seminoma in males. Despite the tumour being malignant, prognosis is excellent with surgery and adjuvant chemotherapy, with promising results concerning future fertility. In this case, we report an 11-year-old girl with abdominal swelling for 7 years. She had gross abdominal distension, with a huge palpable mass. Abdominopelvic ultrasound scan showed a huge heterogeneous mass measured 9.8 cm x 15.7 cm and computed tomography showed a huge lobulated mass measured 19.2 cm x 19.0 cm x 16.3 cm. Exploratory laparotomy and excision of the mass was done.

Ovarian dysgerminoma are uncommon in prepubertal girls. Prognosis depends on the stage at presentation and histology. Multidisciplinary management with a Gynaecologists and Oncologists with joint decision on the need to maintain the child’s developmental and reproductive potential as much as possible should aimed at.

Introduction

Solid ovarian tumours are very rare in the prepubertal girls’ population, but when occurred, they are a major source of anxiety especially for the family of the patient. Paediatric ovarian masses account for an estimate incidence of 2.6 cases per 100,000 girls per year, and malignant ovarian tumours account for 0.9% of all childhood and adolescent malignancies.1,2 Dysgerminoma is one of the rare highly malignant ovarian germ cell tumour (MOGT) with a peak incidence in young women. Approximately one-third of all dysgerminomas show KIT mutations and these are associated with advanced stage at presentation.3 Dysgerminoma originate from undifferentiated germ cells that are similar to primordial germ cells, and it is identical to testicular seminoma in males. Most cases occur in the 2nd and 3rd decades of life, but 10% of cases occur in the 1st decade of life.4 Most patients with dysgerminoma (75%) are diagnosed with early-stage disease, and in such situations surgery alone is curative, and the prognosis is usually excellent,5 however, there is a significant dilemma with the management of advanced disease in children and adolescents.6 A rare case of ovarian dysgerminoma in an 11-years-old prepubertal school girl who was referred to the Paediatric oncology unit of the University of Maiduguri Teaching Hospital is being reported. The paucity of information concerning the frequency and pattern of malignant ovarian tumours including dysgerminoma in children in most Nigerian literature and the rarity of this malignancy in prepubertal children and the recognized controversies in its management prompted us to report our experience.

Case Report

An 11-years-old girl presented with complaints of abdominal swelling of 7 years duration. Her parents reported a gradually increasing abdominal distention from the age of 4 with no other symptoms until 3 months before presentation when she developed abdominal pain, more in the lower abdomen associated with progressive weight loss, and decreased appetite. The patient also had occasional urine retention, which resolved spontaneously, no dysuria, no passage of stones or fleshy material in the urine. The
patient was yet to attain menarche, no breast enlargement, no abnormal discharge per vagina.

Physical examination showed a chronically ill-looking, wasted child, pale, not jaundiced, acyanosed, and no dysmorphic facie with Tanner stage II.B of breast development. She weighed 27 kg (3rd percentile), and height of 140cm (25th percentile). The patient had a soft, grossly distended, and non-tender abdomen, with a large firm palpable mass at the suprapubic region extending to the epigastrium, with a nodular surface and mobile. No demonstrable ascites. Haemoglobin, alpha-fetoprotein, and human chorionic gonadotropin beta subunit levels were normal.

Computed tomography scan showed a huge lobulated abdominopelvic mass measured 19.2 cm x 19.0 cm x 16.3 cm and extended from the pelvis to the epigastric region and laterally to the abdominal and pelvic side walls. Central specks of calcifications were also present. A small area of hypodensity was also noted within the mass, suggestive of necrotic area. The lesion did not enhance significantly after intravenous contrast injection (Figure 1).

The preoperative diagnosis of this case was a teratoma to rule out fibrolipoma. Subsequently, an exploratory laparotomy was performed under general anesthesia, and excision of a huge rubbery mass from the pelvis was done. Tumour was mobilized bluntly on sides (after ligating both uterine arteries) and also from the rectum. The ureter on the left was dissected off the tumour wall before the tumour was excised in piecemeal with part of the bladder wall. The distal component was excised off the proximal third of the vagina. Grossly, the tumour weighed 1900g (Figure 2). Histologic section of the mass showed tissue displaying malignant neoplasm composed of aggregate strands and island of fairly uniform large cells separated by delicate fibrous septae that are infiltrated by lymphocytes. The neoplastic cells have round to oval nuclei some of which have prominent eosinophilic nucleoli and moderately pale cytoplasm. Areas of necrosis were also present (Figure 3). Based on histopathologic features, a final diagnosis of dysgerminoma was rendered. The patient tolerated the surgical procedure well without complication and was transferred to Paediatric oncology unit for chemotherapy on postoperative day seven. She had two cycles of adjuvant chemotherapy before discharge. The patient was lost to follow up after two cycles of the adjuvant chemotherapy. All efforts at locating the patient were unsuccessful as she was said to have been residing in a remote village which is not readily accessible due to Boko Haram insurgency in North-eastern Nigeria.

Discussion

Dysgerminomas are the most common ovarian malignancies in children, constituting 9.5% to 11% of childhood ovarian tumours and 24.5% of paediatric ovarian malignancies.7 There is an increased frequency of dysgerminomas among patients with genetically abnormal gonads.8,9 The rarity of paediatric ovarian malignancies was reported by Ajani et al.10 as only 24 cases of ovarian neoplasms were seen under the age of 15 years over a 22-years period (1991–2003) at University College Hospital, Ibadan. Dysgerminoma was found to be the most common primary malig-

Figure 1. CT images: a) Axial pre-contrast scan showing huge intraabdominal mass with central specks of calcification; b) Axial post-contrast scan showing no significant contrast enhancement; c) Coronal reformatted image demonstrating the extent of the mass from pelvic cavity to epigastrium and laterally to the abdominal and pelvic side walls.

Figure 2. Gross picture showing both ovaries involved with a huge solid capsulated mass.

Figure 3. Photomicrograph section shows nests of fairly uniform polygonal cells with moderate to scanty cytoplasm separated by fibrous septae infiltrated by mature lymphocyte. H and E x 100.
nant paediatric ovarian tumour in Nigerian children.9 However, there were only two reported cases in the Nigerian literature over the past 15 years.6,10 A pure dysgerminoma is endocrinologically inactive. Signs of pronounced hormonal activity indicate the presence of a functioning component, placing the tumor into a mixed germ cell category.11 Ovarian tumours including malignancies can present with diverse signs and symptoms, and most often, the clinical presentation does not permit differentiation between benign and malignant tumour as found in the index case. The rate of bilaterality as found in the index case is reported to be 5% to 20%.11 Abdominal pain is the most common symptom encountered,10 as we found in our patient. A mobile, palpable abdominal mass is the most frequent physical finding,10 as we also observed in our patient. Grossly, dysgerminomas are rubbery, gray, smooth, or bosselated and are surrounded by a dense capsule. The cut surface is soft and homogeneous and has a brainlike consistency.11,12 that is similar to what was found in the index case. Elevated LDH can be an early clinical sign of an ovarian dysgerminoma. However, it is not all dysgerminomas that produce LDH, and elevated LDH is often regarded as a nonspecific finding. Our patient had a normal LDH levels.

The management of childhood ovarian tumour must be balanced with the desire to maintain the child’s reproductive and developmental potential as much as possible. Vicus et al. published data of the occurrence of pure ovarian dysgerminomas with 72.3% at stage I, 4.6% at stage II, and 21.5% at stage III disease. The initial treatment employed was surgery with 72.2% of the patients undergoing unilateral oophorectomy and 21.5% bilateral oophorectomy with or without hysterectomy.13 The indication for bilateral oophorectomy and hysterectomy in our case was intrasurgically made. The reason for that decision was a macroscopically visible infiltration of both ovaries and the uterus which coalesce to constitute the huge tumour mass. Although the main aim is fertility-preserving surgery and is possible in 70% of cases,14,15 the indication for bilateral oophorectomy and hysterectomy in our case was inevitable. We could not see any remote possibility of sparing the infiltrated uterus and both ovaries without putting the child into risk of a further growing of the tumour and probably leading to metastasis into other organs and there by jeopardizing the survival of the child. Similar to our case Nishio et al. did radical surgery and administered chemotherapy in advance stages of ovarian dysgerminoma.16 In contrast to this Vicus et al. showed that fertility-sparing surgery in women with pure ovarian dysgerminoma led to 8 in 65 cases to 12 pregnancies and 12 live births.13 Lymph node metastasis is present in 28% of dysgerminomas and is significantly associated with poor survival.15-18 In order to evaluate the prognosis and find adequate therapeutic options lymphadenectomy is indicated. The result of the lymphadenectomy in our case was negative for tumour cells. Adjuvant chemotherapy in combination with initial surgery shows promising results concerning outcome and fertility. We believe that our report on the challenges of diagnosis and treatment in this case can help clinicians to better understand and manage this kind of malignant tumour.

Conclusions

Ovarian dysgerminoma are uncommon in girls below the age of 12. Pain and abdominal mass are the most common modes of presentation. Dysgerminoma is the most common malignant ovarian tumour. Prognosis depends on the size of the tumor, stage at presentation and histology of the tumour. Conservative surgery should be aimed at during the initial treatment in children and adolescents who are yet to start their reproductive life. Adoption of multidisciplinary management with a Gynaecologists and Oncologists with joint decision on the need to maintain the child’s reproductive and developmental potential as much as possible should be the aim.

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