<table>
<thead>
<tr>
<th>Title</th>
<th>Primitive small round cell tumour of the adrenal gland presenting with fever of unknown origin and t(12;22) (q13;q12) cytogenetic finding</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Lam, KY; Lo, CY; Shek, TWH; Ma, ESK; Au, WY; Chan, GCF</td>
</tr>
<tr>
<td>Citation</td>
<td>Journal Of Clinical Pathology, 2001, v. 54 n. 12, p. 966-969</td>
</tr>
<tr>
<td>Issued Date</td>
<td>2001</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/10722/42418">http://hdl.handle.net/10722/42418</a></td>
</tr>
<tr>
<td>Rights</td>
<td>Journal of Clinical Pathology. Copyright © B M J Publishing Group.; This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License.</td>
</tr>
</tbody>
</table>
Primitive small round cell tumour of the adrenal gland presenting with fever of unknown origin and t(12;22)(q13;q12) cytogenetic finding

K Y Lam, C Y Lo, T W H Shek, E S K Ma, W Y Au, G C F Chan

Abstract
This report describes a left adrenal tumour in a 16 year old Chinese girl who presented with fever of unknown origin. The histological and ultrastructural features of the adrenal tumour were those of a primitive small round cell tumour with neuroendocrine differentiation. Cytogenetic analysis of cultured tumour cells showed a reciprocal translocation t(12;22)(q13;q12). This is the first example of such a tumour being reported in the adrenal gland. The adrenal tumour was also the cause of the fever, which subsided after the removal of the tumour.

Keywords: adrenal; cytogenetic; primitive tumour; pyrexia of unknown origin

Tumours that arise from the adrenal medulla show either neuroendocrine or neural differentiation. These include phaeochromocytoma, ganglioneuroma, ganglioneuroblastoma, neuroblastoma, neurilemmoma, malignant peripheral nerve sheath tumour, malignant melanoma, primitive neuroectodermal tumour (PNET), and tumours with mixed differentiation. In this report, we describe a primitive adrenal medullary tumour with unique clinicopathological and cytogenetic features.

Case report
PATIENT
A 16 year old Chinese girl, previously in good health, was admitted because of prolonged intermittent fever. Physical examination was unremarkable. Laboratory studies revealed anaemia (haemoglobin concentration, 78 g/litre), mild leucocytosis (white blood cell count, 11.9 x 10^9/litre), and a raised erythrocyte sedimentation rate (> 140 mm/hr). The concentration of C3 was 2700 mg/litre (range, 600–1300) and the C reactive protein concentration was 260 mg/litre (normal, < 10 mg/litre). However, immunoglobulin concentrations were within the normal range. Renal and liver functions were normal except for increased concentrations of alkaline phosphatase (674 U/litre; normal range, 36 – 117) and γ glutamyl transferase (215 U/litre; normal range, 11–49). Screening for hepatitis virus was negative. Bone marrow aspiration and repeated culturing of blood, urine, and bronchoalveolar lavage showed no evidence of bacterial growth. Screening of blood for malaria, rickettsia, and brucella was negative. There was no clinical or laboratory evidence of collagen vascular disease. During her hospitalisation, the patient suffered from a high swinging temperature; a maximum temperature of 41°C was recorded. Subsequent abdominal ultrasonography revealed a left suprarenal heterogeneous lesion that was confirmed by computed tomography (CT) and magnetic resonance imaging to be a heterogeneous left adrenal mass with a maximum dimension of 6.2 cm (fig 1). Hormonal investigations (24 hour urinary catecholamine values, serum cortisol, and overnight dexamethasone suppression test) were within the normal range. Fine needle aspiration of the adrenal mass was performed under CT scan, and revealed features consistent with a malignant tumour. Total adrenalectomy was performed. There was no definite evidence of invasion into the adjacent soft tissue, enlarged lymph nodes, or distant metastases.

The postoperative recovery was uneventful and the swinging fever that had persisted on and off for more than two months duration subsided immediately after surgery. Standard staging of the disease, including a CT scan of the thorax and bone scintigraphy, did not reveal metastatic disease. Adjuvant local radiotherapy and systemic chemotherapy were administered. However, the patient had lung metastases two years after surgery, which was confirmed by biopsy.

Methods
Fresh tissue samples (1 mm^3) of the resected adrenal gland were taken. They were (1) fixed in 2.5% buffered glutaraldehyde and then processed for ultrastructural examination under a transmission electron microscope and (2) put into culture medium for cytogenetic studies. Cytogenetic studies were performed...
on short term overnight synchronised and non-synchronised cultures of marrow cells. These cells were supplemented by a direct harvest and analysed in accordance with standard protocol.

Fresh frozen tumour tissues were also analysed for the chimaeric transcripts that encode fusion products found commonly in PNET and clear cell sarcoma. These gene fusion products were identified by the reverse transcription polymerase chain reaction (RT-PCR) using cDNA that was derived from tumour tissue RNA as a template. PCR products were identified by gel electrophoresis using ethidium bromide stained gels. The sequence of the primer used and the experimental conditions of the RT-PCR were as described previously. The presence of amplifiable RNA was confirmed by RT-PCR using primers specific to β actin (a ubiquitously expressed gene in human tissues). Primers specific to the EWS and ATFI genes were used to test for the presence of the gene fusion transcript commonly found in clear cell sarcoma. The DTC1-IA clear cell sarcoma cell line, which produced approximately 950 bp of PCR product, was used as control. Primers specific to EWS, FLI1, or ERG were used to detect the presence of gene fusion transcripts commonly noted in PNET. Two PNET control cell lines known to have a t(11;22) translocation were used as positive controls in these studies. The PCR products were Southern blotted on to a nylon membrane and probed with an internal end labelled oligonucleotide specific to EWS, FLI1, and ERG.

Immunohistochemistry was conducted on 5 μm thick sections from representative paraffin wax embedded blocks using the avidin-biotin–peroxidase complex method. Antibodies to the following antigens were used: vimentin, neuron specific enolase (NSE), synaptophysin, chromogranin, desmin, smooth muscle actin, myoglobin, myosin, leucocyte common antigen (LCA), MAK-6, CAM 5.2, cytokeratin 7, cytokeratin 20, HMB-45, CD34, and CD99.

**Results**

**Gross Pathology**

The resected left adrenal gland weighed 180 g. The contour of the adrenal gland was distorted by a medullary tumour measuring

**Histological Features**

The adrenal tumour was composed of sheets of uniform small and medium oval cells with scanty cytoplasm and irregular nuclei with fine chromatin (fig 2). Mitotic counts were approximately 1/10 high power fields. In some foci, multinucleated tumour giant cells were noted. Rosettes or rosette-like structures were not present. The tumour cells were embedded in a vascular stroma in which haemorrhage and haemosiderin deposits were frequently found. Extensive areas of necrosis with perivascular preservation of viable tumour cells forming a peritheliomatous pattern were also seen.

**Immunohistochemical Findings**

Immunohistochemical studies revealed that the primary tumour cells were positive for vimentin but negative for CD99, neuroendocrine markers (NSE, chromogranin, and synaptophysin), cytokeratins (Mak-6, CAM 5.2, cytokeratin 7, and cytokeratin 20), S-100, HMB-45, muscle markers (actin, desmin, myosin, and myoglobin), CD31, CD34, and LCA. The tumour cells from the metastatic lesion in the lung were positive for CD99.

**Electron Microscopy**

Ultrastructural examination showed closely opposed cells that were uniform and primitive
Cytogenetics
Chromosomal analysis of cultured tumour cells revealed translocation between the long arms of chromosome 12 and 22; t(12;22)(q13;q12) (fig 4). On RT-PCR analysis, the tumour was negative for fusion transcripts for PNET, clear cell sarcoma. The absence of the transcripts was confirmed on labelling of Southern blotted PCR products with a cDNA probe.

Diagnosis
The diagnosis of a primitive small round cell tumour of the adrenal gland with neuroendocrine differentiation was corroborated by morphological, immunohistochemical, ultrastructural, and cytogenetic findings. The features were in keeping with PNET.

Discussion
The fever pattern of our patient fulfils the criteria for paraneoplastic fever. Paraneoplastic fever has been defined as an unexplained fever that coincides with tumour growth and disappears promptly on tumour removal or control. Lymphomas (especially Hodgkin’s disease) and renal cell carcinomas are the prototypes of tumours frequently associated with paraneoplastic fever. However, descriptions in the literature of adrenal tumours presenting with prolonged fever as the principle symptom are rare, although there have been some patients with adrenal tumours presenting with neoplasic fever. These included phaeochromocytomas, adrenal cortical carcinomas, and one patient with aldosteronoma. In our study, an adrenal small round cell tumour was described in which fever was the primary manifestation. The extensive tumour necrosis may have triggered the release of circulating cytokines that mediated the fever. On the basis of this report, it seems logical that adrenal tumour should be included in the differential diagnosis in patients with fever of undetermined origin.

Abe et al described a small round cell variant of malignant peripheral nerve sheath tumours in five patients with tumours located within the soft tissue of the trunks and limbs. The tumours consisted mainly of uniform, small, round tumour cells with scanty cytoplasm. They stained negative for CD99 and tumour rosettes were seen. These features are similar to those noted in our patient. However, there were histological and ultrastructural features that distinguished small round cell variant of malignant peripheral nerve sheath tumour from that of our present patient. In the cases described by Abe et al, foci of uniform, short, spindle shaped tumour cells arranged in a whorl-like pattern were seen. Ultrastructural examination also revealed a poorly developed basal lamina that was fragmentous and discontinuous along the cell bodies. Dense core granules were absent in all except one case.

The cytogenetic abnormality found in our patient was t(12;22)(q13;q12). This translocation has never been reported in PNET. t(12;22)(q13;q12) has been reported to involve the fusion of the EWS and ATFI genes. EWS-ATFI fusion has often been found in clear cell sarcoma but it is not present in conventional cutaneous melanoma. The clinical presentation and pathological, immunohistochemical, and ultrastructural characteristics of the adrenal tumour reported here (such as the absence of reactivity to HMB-45 or melanomas) excluded the diagnosis of clear cell sarcoma. In addition, RT-PCR studies showed that t(12;22) translocation in the adrenal tumour did not involve EWS–ATFI fusion, as seen in clear cell sarcoma. Thus, the adrenal tumour had unique pathological and cytogenetic features and was neither characteristic of PNET nor clear cell sarcoma on cytogenetic investigation.

The authors thank Dr P Sorenson, Department of Pathology, British Columbia’s Children Hospital, Vancouver, BC, Canada for his help with the RT-PCR analysis.


www.jclinpath.com

Reproduced with permission of the copyright owner. Further reproduction prohibited without permission.
7th European Forum on Quality Improvement in Health Care

21–23 March 2002
Edinburgh, Scotland

We are delighted to announce this forthcoming conference in Edinburgh. Authors are invited to submit papers (call for papers closes on Friday 5 October 2001) and delegate enquiries are welcome.

The themes of the Forum are:
- Leadership, culture change, and change management
- Achieving radical improvement by redesigning care
- Health policy for lasting improvement in health care systems
- Patient safety
- Measurement for improvement, learning, and accountability
- Partnership with patients
- Professional quality: the foundation for improvement
- Continuous improvement in education and training
- People and improvement.

Presented to you by the BMJ Publishing Group (London, UK) and Institute for Healthcare Improvement (Boston, USA). For more information contact: quality@bma.org.uk or look at the website www.quality.bmjgroup.com. Tel: +44 (0)20 7383 6409; fax: +44 (0)20 7373 6869.