A possible new syndrome with growth-hormone secreting pituitary adenoma, colonic polyposis, lipomatosis, lentigines and renal carcinoma in association with familial testicular germ cell malignancy: A case report

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Abstract

Background: Germ-cell testicular cancer has not been definitively linked to any known hereditary cancer susceptibility disorder. Familial testicular cancer in the presence of other findings in affected and unaffected family members might indicate a previously-unidentified hereditary cancer syndrome.

Case presentation: The patient was diagnosed with a left testicular seminoma at age 28, and treated with left orchiectomy followed by adjuvant cobalt radiation. His family history is significant for testicular seminoma in his son, bladder cancer in his sister, and lipomatosis in his father. His evaluation as part of an etiologic study of familial testicular cancer revealed multiple colon polyps (adenomatous, hyperplastic, and hamartomatous) first found in his 50 s, multiple lipomas, multiple hyperpigmented skin lesions, left kidney cancer diagnosed at age 64, and a growth-hormone producing pituitary adenoma with associated acromegaly diagnosed at age 64. The patient underwent genetic testing for Cowden syndrome (PTEN gene), Carney complex (PRKAR1A gene), and multiple endocrine neoplasia syndrome type 1 (MEN1 gene); no deleterious mutations were identified.

Discussion: The constellation of benign and malignant neoplasms in the context of this patient’s familial testicular cancer raised the possibility that these might be manifestations of a known hereditary susceptibility cancer syndrome; however, genetic testing for the three syndromes that were most likely to explain these findings did not show any mutation. Alternatively, this family's phenotype might represent a novel neoplasm susceptibility disorder. This possibility cannot be evaluated definitively on the basis of a single case report; additional observations and studies are necessary to investigate this hypothesis further.
Background
Testicular germ cell tumor (TGCT) is the most common malignancy in U.S. adolescents and young adult males ages 15–45. TGCT risk factors include family history, personal history of previously-diagnosed TGCT, cryptorchidism and other disorders of male urogenital differentiation [1].

Approximately 1–3% of individuals with TGCT report having ≥ one affected first-degree relative. The relative risk of TGCT is higher among siblings than among fathers or sons of affected individuals [1], suggesting the possibility of genetic heterogeneity and environmental influences. Segregation analysis suggested that genetic susceptibility is important in familial TGCT, with recent studies indicating that multiple genetic loci of lower penetrance acting in concert contribute to the familial aggregation of this malignancy [2]. Although TGCT has been implicated in a number of genetic disorders, it has not been definitively linked to any known hereditary cancer syndrome [3].

Finally, individuals with TGCT are observed to have increased risks of certain cancers, including pancreas, kidney, urinary bladder, thyroid, lymphatic tissue, leukemia, and connective tissues [4]. Although the excess risk of several of these second malignancies has been attributed to late complications of testicular cancer treatment, the possibility that TGCT is part of a familial syndrome which includes other types of cancer cannot be dismissed.

We established the Multidisciplinary Etiologic Study of Familial Testicular Cancer in 2002 (NCI Protocol 02-C-0178) to identify possible testicular cancer susceptibility genes and to characterize more precisely the clinical phenotype of the familial testicular cancer syndrome. During the course of this study, we encountered a family with seminoma in a father and son, and a provocative constellation of other cancers and benign conditions in the father. The following case report summarizes our attempt to explain these findings on the basis of a known cancer susceptibility disorder.

Case report
The proband was a 63 year-old, Caucasian male who presented to the NCI Clinical Genetics Branch in October 2004 for evaluation of familial testicular cancer. He developed a left testicular pure seminoma at age 28 (per medical records; no pathology was available for review); he underwent a left radical orchiectomy followed by adjuvant cobalt irradiation to the retroperitoneum, and has remained free of testicular cancer.

At his initial NIH visit, the patient reported resection of multiple colonic polyps on successive colonoscopies, beginning in his early 50s. Materials from several excised polyps were reviewed at the NIH Laboratory of Pathology, and revealed a mixed polyposis of 5 adenomatous polyps, 2 hamartomatous polyps (Figure 1a), and multiple hyperplastic polyps.

![Photomicrographs](a) Haematoxylin and Eosin stain (100×) of a hamartomatous polyp from the descending colon showed proliferation of dilated mucinous glands surrounded by smooth muscle fibers. This latter feature is characteristic of hamartomatous polyps observed in patients with Peutz-Jegher syndrome. (b) Photomicrograph of the left kidney tumor was composed of a solid proliferation of spindle cell with large irregular nucleus, abundant cytoplasm, and prominent mitotic figures. The findings are consistent with a high-grade clear cell renal cell carcinoma, sarcomatoid type.
Physical examination revealed a muscular, mesomorphic white male (he is an avid body-builder) with multiple subcutaneous lipomas in the extremities and trunk. He had multiple dark brown to black, irregularly-shaped pigmented macules on the chest, back, arms, and legs, which were clinically suggestive of dysplastic nevi. Biopsy of the clinically most atypical lesion showed a simple lentigo, without melanocytic atypia. Protocol-related blood work, including testosterone, estradiol, chemistry panel, prostate specific antigen, alpha-fetoprotein, follicle stimulating hormone, luteinizing hormone, and beta-HCG, were normal. Ultrasound of the right testis showed early micro-lithiasis. Computed tomography of the chest, abdomen, and pelvis revealed an unexpected left inferomedial renal mass (Figure 2b), which was confirmed by MRI. Further evaluation of the left kidney mass was recommended, but not pursued due to other medical issues.

He returned for a scheduled clinic visit one year later and reported having an elevated growth hormone (GH) level detected during a complementary/alternative health evaluation. The patient also stated that his head and hands had been “getting bigger”. He reported taking various dietary supplements, none of which was found to contain androgens upon review of their ingredients, as well as testosterone gel, which was started at the time of his complementary/alternative health evaluation due to a low serum testosterone level. On examination, he had an acromeg-

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**Figure 2**

**Imaging studies.** (a) MRI of pituitary adenoma – T1 weighted sagittal MRI of the pituitary gland after intravenous contrast reveals an enlarged gland that is centrally hypoenhancing, with a rim of relative hyper-enhancement at the superior margin. The longest dimension of the pituitary is approximately 2 cm. (b) Composite of CT images of the left kidney, first visit and one year later- Serial contrast media-enhanced CTs of the abdomen demonstrate an enlarging left renal mass. The initial scan (October 2004) demonstrates a solid enhancing mass (white arrow) measuring approximately 2.6 cm. Follow-up CT approximately one year later shows slight enlargement (3.0 cm) (white arrow).
genic appearance, including protruding supra-orbital ridges, a prognathic jaw, wide nose, and diffuse soft-tissue thickening of the hands, all of which were judged clin-
cally to be more prominent when compared with the year before.

Laboratory studies demonstrated a normal fasting glu-
cose, elevated levels of insulin-like growth factor-I (IGF-I) [943 ng/ml; normal range 71–290 ng/ml] and IGF bind-
ing protein-3 [8.4 µg/ml; normal range 3.2–6.6 µg/ml]. A pituitary MRI revealed a hypoenhancing, diffusely-
enlarged pituitary measuring approximately 2 cm, consist-
ent with an adenoma (Figure 2a). A follow-up renal CT
showed the left inferomedial kidney mass had grown compared with the study performed one year earlier; the adrenal glands were normal.

On January 23, 2006, the patient underwent trans-sphe-
noidal resection of the pituitary mass, yielding a GH-pro-
ducing adenoma. On February 22, 2006, he underwent a
left nephrectomy for a stage III, poorly-differentiated renal cell carcinoma, with sarcomatoid features (Figure 1b). By November 2006, his growth hormone levels had returned to normal, and he had no evidence of recurrent kidney cancer.

Detailed family history review revealed a son with semi-
noma (confirmed by outside pathology report) at age 34
(cancer-free at age 36), and a sister who developed urinary bladder cancer (death certificate confirmation) at age 53. His son had no evidence of lipomatosis or multiple lentig-
ines on examination at the time of study enrollment, and has had a negative screening colonoscopy. The proband’s
father was reported to have lipomatosis, but was deceased. There is no history of other benign or malignant neoplasms reported in the rest of the family.

Based on his personal history of multiple cancers, a pitui-
tary adenoma, multiple pigmented skin lesions, multiple lipomas, and multiple colon polyps with varying histol-
ogy, the patient underwent genetic testing for Cowden
syndrome (the PTEN gene), Carney complex (the PRKAR1A gene), and multiple endocrine neoplasia syn-
drome type 1 (the MEN1 or menin gene). No germline mutations were detected in any of these genes.

Discussion and conclusion

Our patient presented with multiple malignant and benign neoplasms including left testicular seminoma, left kidney renal cell carcinoma, a growth hormone-secreting pituitary adenoma, multiple pigmented skin lesions, lipomas, and colon polyps with varying histology (adenoma-
tous, hyperplastic, and hamartomatous). The proband’s
son had testicular cancer and microlithiasis, but none of
the other neoplasms, and his father is reported to have
had lipomatosis. The clinical phenotype of the familial
TGCT syndrome, as currently understood, is comprised of
a site-specific predisposition to TGCT and does not appear to include an increased risk for other benign or malignant neoplasms [1, 3]. Various aspects of testicular dysgenesis
and genitourinary development have been implicated in
the pathogenesis of familial TGCT, but this hypothesis
remains unproven at present. As part of the protocol
under which the current family was seen, we have evaluated
136 members of 35 multiple-case TGCT families, including 56 men with a history of TGCT; the constella-
tion of abnormalities seen in the proband reported here is
unique in our experience (Korde L, Kramer J, Mueller CM, and Greene MH, personal communication). Therefore, the possibility that the neoplasms observed in this patient represent unrecognized manifestations of the familial
TGCT syndrome seems unlikely.

We considered whether these observations might be
explained on the basis of a known genetic cancer suscep-
tibility disorder and felt that Cowden syndrome, Carney
complex, and multiple endocrine neoplasia type 1 war-
ranted serious consideration in this regard, although none

Cowden syndrome (CS), an autosomal dominant disease
associated with mutations in the PTEN tumor suppressor
gene on chromosome 10q10.22-23, is associated with ele-
vated risks of breast and thyroid cancer, as well as diverse
benign neoplasms including gastrointestinal hamartoma-
tous polyps, lipomas, giant fibroadenomas of the breast,
hemangiomas, and multiple early-onset uterine leiomyo-

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excluded, but this diagnosis is unlikely since his family does not meet the clinical diagnostic criteria.

Carney complex (CNC) is a multiple neoplasia syndrome characterized by skin lesions (e.g., lentigines, compound nevi, and blue nevi), psammomatous melanotic schwannoma, cardiac myxomas, large-cell calcifying Sertoli cell tumors and Leydig cell tumors of the testis, as well as endocrine tumors, including primary pigmented nodular adrenocortical disease and pituitary adenoma, most of which are GH-producing [6,7]. CNC is associated with germline mutations in the \textit{PRKAR1A} gene on chromosome 17q22-24, which are detected in 50–65% of CNC patients [6]. About 30% of CNC families have been linked to a susceptibility locus on chromosome 2p16 (CNC2), but the gene at this locus has not yet been identified. Our patient had a GH-producing pituitary adenoma and testicular cancer. However, CNC does not include colon polyps, lipomas or renal cancer among its recognized manifestations. He underwent testing for germline mutations in the \textit{PRKAR1A} gene; no mutation was identified.

Multiple endocrine neoplasia type 1 (MEN1) involves tumors of the parathyroid, endocrine pancreas, anterior pituitary, adrenal glands, and carcinoid tumors of the GI tract. It is associated with germline mutations in the \textit{MEN1} gene, on chromosome 11q13 [8]. Anterior pituitary adenomas, about 1/4 of which are GH-producing, are found in 10%–60%, and multiple lipomas are found in about 30% of patients with MEN1 [8]. Both the patient and his father had multiple lipomas. The father was not known to have manifested other signs or symptoms suggestive of MEN1. The patient underwent germline mutation testing of the \textit{MEN1} gene, with no mutation detected.

\textit{AIP} (which encodes for the aryl hydrocarbon receptor-interacting protein) has recently been implicated as a low-penetrance, familial pituitary adenoma susceptibility gene [9]. Although there are no other cases of pituitary adenoma in this family, it is not presently known whether other neoplasms occur excessively in pituitary adenoma kindred. We therefore tested the patient for mutations in \textit{AIP}; none were detected (CA Stratakis, personal communication).

Finally, testicular germ cell tumor has been described in acromegalic patients [10]. Renal cancer, colon cancer, and colon polyps have been reported to occur excessively in acromegaly; however, the polyps observed in acromegalic patients are adenomatous or hyperplastic; hamartomatous polyps have not been described in this context [11,12]. Our patient’s seminoma was unlikely to be related to acromegaly, given the temporal sequence of events, and neither lipomatosis nor lentigines are part of the acromegaly phenotype. However, we can not rule out the possibility of an association between acromegaly and either renal cancer and/or adenomatous and hyperplastic colon polyps in this patient.

While our patient had some of the clinical features associated with each of these cancer susceptibility disorders, none can account for his entire clinical phenotype, and germline mutation testing of the three genes implicated in the inherited diseases was negative. Conventional clinical mutation testing may have failed to detect a mutation in one of these genes; however, this likelihood is small since his family does not meet the clinical diagnostic criteria for any of these syndromes. He might represent an atypical presentation of some other known, multiple benign and malignant neoplasm susceptibility disorders (e.g., juvenile polyposis syndrome, Peutz-Jeghers syndrome, hereditary hemorrhagic telangiectasia, or \textit{MYH}-adenomatous polyposis); research evaluation of a wider spectrum of genes is now underway. Or, he may have several co-exist-

| Patient’s Findings  | Acromegaly | Hereditary Syndrome |
|---------------------|------------|---------------------|
|                     |            | Cowden syndrome     |
|                     |            | Carney complex      |
|                     |            | MEN1                |
| Seminoma            | X          | XX                 |
| Pituitary adenoma (GH-producing) | XX | XX | XX |
| Colon adenomatous polyps | XX | | |
| Colon hamartomatous polyps | | XX | |
| Colon hyperplastic polyps | | | |
| Renal cell carcinoma | X | | |
| Lipomas             | XX         | XX                 |
| Lentigines          | XX         | | |

\text{XX}: Classical syndrome-related findings
\text{X}: Findings are less clearly established as part of the syndrome

* Testicular cancers associated with Carney complex are typically large-cell calcifying Sertoli cell tumors or Leydig cells tumors of the testis

Table 1: Summary of patient’s clinical findings relative to acromegaly and the three hereditary cancer susceptibility syndromes considered
ing but etiologically-unrelated conditions, e.g., familial testicular cancer, acromegaly with renal cancer and colon polyps; and idiopathic lipomatosis.

The final possibility is that this family's phenotype represents a novel neoplasm susceptibility disorder, of which TGCT is an important component. Of course, this hypothesis cannot be tested on the basis of a single case report. We await with interest future observations from other clinical investigators, in response to the report we have provided herein.

Competing interests
The author(s) declare that they have no competing interests.

Authors' contributions
PM had primary responsibility for drafting the manuscript. LK, JK, and MHG were responsible for the design and operation of the Familial Testicular Cancer Study, contributed to this patient's evaluation, and contributed to writing the manuscript. JP, CM, and SP were involved in this patient's evaluation, genetic counseling, and data collection and medical record abstraction. CS evaluated the patient for acromegaly and performed genetic testing for Carney complex, MEN 1, and familial pituitary adenoma. PP, WML, GB, MM, and PC were involved in the patient's clinical assessment and treatment. All authors read and approved the final manuscript.

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