Lung cancer as a predominant feature in a patient with Peutz–Jeghers syndrome: Case report

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Abstract
Peutz–Jeghers syndrome (PJS) is characterized by mucocutaneous pigmentation and gastrointestinal hamartomatous polyposis, which can lead to intussusception. PJS patients face high lifetime risks for various cancer types, with the majority of patients being diagnosed with tumors along the gastrointestinal tract. Herein, we present the case of a 34-year-old man who carried a germline STK11 pathogenic variant, while lacking the cardinal features of PJS syndrome. Interestingly, he was diagnosed with lung adenocarcinoma despite being a never-smoker. Tumor testing revealed clinically relevant molecular alterations, including the known germline pathogenic variant STK11, a KRAS somatic pathogenic variant, and FGFR3 gene amplification. Treatment with standard chemotherapy and immunotherapy did not have a clinical benefit. Due to clinical deterioration, the patient deceased 18 months after his initial diagnosis prior to having the chance for targeted therapy. Identification of rare hereditary cancer syndromes and the respective presence of tumor biomarkers can provide important alternatives to targeted treatments, including immunotherapy in patients with tumors unresponsive to conventional treatment protocols. This case highlights that although only a small proportion of lung cancer diagnoses will be due to hereditary predisposition, STK11 germline carriers should be under close surveillance for early detection of lung cancer.

KEYWORDS
case report, immunotherapy, lung cancer, Peutz–Jeghers syndrome, STK11

INTRODUCTION
Peutz–Jeghers syndrome (PJS) is a rare hereditary syndrome characterized by mucocutaneous pigmentation along with gastrointestinal hamartomatous polyposis and increased risk for multiple types of cancer. Malignancies of the gastrointestinal tract, i.e. gastric, small bowel, colon, rectum, and pancreas, followed by breast, ovary, and testicular cancers are the most frequently diagnosed malignancies among patients with PJS. Due to their elevated lifetime cancer risks, estimated to be around 85%, individuals with PJS should be offered multidisciplinary care and follow specialized screening protocols. Germline pathogenic STK11/LKB1 variants account for most individuals with a PJS clinical diagnosis.

Lung cancer, although described in the PJS tumor spectrum, is a relatively rare entity, with men and women with PJS facing a 13% and 1% risk, respectively, for diagnosis by the age of 60 years. Overall, lung cancer is a tumor type mostly associated with environmental factors, prominently tobacco smoking, with hereditary predisposition identified in a very small subset of cases.

Herein, we present an unusual case of a young, never-smoker, PJS patient, who lacked the characteristic PJS...
features, with lung adenocarcinoma being his prominent diagnosis.

CASE PRESENTATION

In February 2017, a 34-year-old man with a medical history of schizophrenia presented with anemia and underwent diagnostic gastroscopy. One and two nondysplastic hyperplastic polyps were identified in his stomach and duodenum, respectively (Figure 1). In June 2017, colonoscopy revealed a tubular adenoma with low-grade dysplasia in the sigmoid, while a follow-up colonoscopy revealed two additional polyps.

In December 2017, the patient was hospitalized with small bowel obstruction due to intussusception and underwent surgical resection, followed by end-to-end anastomosis. Histology report described two hamartomatous polyps with a central core of branching smooth muscle, typical of PJS (Figure 2). At this point, genetic testing for STK11 was ordered, based on recent diagnosis, and his family history. His father was diagnosed with ampulla of Vater cancer at the age of 51 years, while at subsequent endoscopies hamartomatous polyps were identified his small intestine and jejunum. He died from metastatic pancreatic cancer at
the age of 56 years. A detailed pedigree is illustrated in Figure 3.

Germline testing led to the identification of a STK11 pathogenic variant, c.169G>T, p.(Glu57Ter). The variant has an entry on ClinVar database and has been previously reported in the literature, but has not been reported in patients of Greek descent. Archival blood sample was used to verify the presence of the variant in the patient’s father.

On January 2018, the patient underwent a positron emission computed tomography (CT) scan which demonstrated an enhanced lobulated lesion of the middle lobe of the right lung (Figure 4a). A subsequent CT-guided biopsy was diagnostic for lung adenocarcinoma. The patient, who reported no history of smoking or alcohol consumption, underwent right pneumonectomy and mediastinal lymphadenectomy in June 2018. Histologic examination revealed a T2bN2 non-small-cell lung (NSCLC) adenocarcinoma (Figure 5). Following diagnosis, adjuvant chemotherapy with four cycles of carboplatin and paclitaxel was administered.

In December 2019, disease progression was confirmed via CT scan, with the presence of new pulmonary nodules and bone lytic lesions (Figure 4b). At this point, tumor molecular profiling was ordered to identify potential therapeutic targets, which revealed the pathogenic STK11 variant with Variant Allele Frequency (VAF) of 74%, suggesting loss of heterozygosity at the specified position, the KRAS c.35G>C pathogenic variant (VAF-19%), and FGFR3 gene amplification (copy number gain-5.29 copies). No other targetable molecular alterations (EGFR mutation or ALK fusions) were identified, while programmed death-ligand 1 (PD-L1) expression was negative.

In January 2020, first-line treatment with carboplatin, pemetrexed, and pembrolizumab was initiated. After four cycles of treatment, CT scan documented disease progression with the development of new lung and liver metastatic lesions. Gemcitabine was administered as second-line treatment, resulting in significant clinical improvement, with pain reduction and improved mobility. The patient completed three cycles of chemotherapy and was scheduled for restaging examinations; however, in June 2020, he presented with acute respiratory distress and despite resuscitation efforts he died due to cardiorespiratory arrest. No autopsy was performed (Figure 5).

DISCUSSION

We herein present an unusual case of a 34-year-old man with PJS who lacked the characteristic pigmentation, had only a small number of the typical hamartomatous polyps, while his prominent diagnosis was an advanced stage lung adenocarcinoma. A year after his initial diagnosis, disease recurrence was observed. The patient received standard treatment based on current guidelines, but despite all therapeutic efforts he died 18 months after his initial diagnosis.

Tumor molecular profiling revealed clinically relevant molecular alterations, including the known germline pathogenic variant STK11, a KRAS pathogenic variant, and FGFR3 gene amplification. Even though no targeted treatment was at that time available, these alterations warrant further investigation as prognostic or predictive biomarkers of response or resistance to immunotherapy and/or targeted treatments.

While somatic STK11 pathogenic variants are frequent in lung adenocarcinomas, germline pathogenic variants are rarely identified in these patients. Inactivating STK11 molecular alterations have been associated with decreased PD-L1 expression, cytokine release, T-cell tumor infiltration, and antigenicity, resulting in a “cold” tumor immune response.
microenvironment. Lung cancer patients with co-occurring mutations in KRAS and STK11 have a trend towards shorter overall survival, while STK11 mutational status does not seem to be a standalone prognostic factor.

Notably, patients with STK11/KRAS co-mutated tumors have significantly shorter progression-free and overall survival, along with decreased overall response rates, after treatment with anti-PD-1 agents when compared to patients having KRAS mutations only. Interestingly, the presence of STK11 pathogenic variants has been associated with worse clinical outcomes, in terms of survival and response rates, in patients with NSCLC who received treatment with immunotherapy combined with doublet chemotherapy.

Hereditary lung cancer is rare, with germline STK11 pathogenic variants being a significant genetic predisposing factor. We experienced the case of a patient who did not have the cardinal PJS features and was diagnosed with NSCLC bearing both germline and tumor STK11, along with tumor KRAS pathogenic variants, and who did not respond to combinational treatment with chemotherapy and immunotherapy. The identification of rare hereditary cancer syndromes, along with tumor biomarkers, can lead the way to select individualized treatments in the future.

CONFLICT OF INTEREST
E.F.: Advisory role: LEO Pharma. Speaker fees: Roche, Pfizer, AstraZeneca. Stock ownership: GENPREX INC, Deciphera Pharmaceuticals, Inc. Travel grant: Merck, Pfizer, and K.A.M Oncology/Hematology and DEMO. G.F.: Advisory Board: Pfizer, Novartis and Roche. Honoraria: Astra Zeneca. Stock ownership: ARIAD, GENPREX, Daiichi Sankyo, RFL Holdings, FORMYCON. G.M.: Advisory and consultation fees: AstraZeneca, BMS, MSD, Roche, Takeda, Novartis, AMGEN. Travel and accommodation fees: AstraZeneca, BMS, MSD, Roche, Takeda, Novartis. PI in sponsored clinical trials: Novartis, Roche, MSD, AstraZeneca, Merck, BMS, AMGEN. The other authors have no conflict of interest to declare.

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