Loss of PTEN Expression Is Associated with Poor Prognosis in Patients with Intraductal Papillary Mucinous Neoplasms of the Pancreas

Dario Garcia-Carracedo1, Andrew T. Turk2, Stuart A. Fine1, Nathan Akhavan1, Benjamin C. Tweel1, Ramon Parsons1,2,5, John A. Chabot3, John D. Allendorf3, Jeanine M. Genkinger6, Helen E. Remotti2, and Gloria H. Su1,2,4

Abstract

Purpose: Previously, we reported PIK3CA gene mutations in high-grade intraductal papillary mucinous neoplasms (IPMN). However, the contribution of phosphatidylinositol-3 kinase pathway (PI3K) dysregulation to pancreatic carcinogenesis is not fully understood and its prognostic value unknown. We investigated the dysregulation of the PI3K signaling pathway in IPMN and its clinical implication.

Experimental Design: Thirty-six IPMN specimens were examined by novel mutant-enriched sequencing methods for hot-spot mutations in the PIK3CA and AKTI genes. PIK3CA and AKTI gene amplifications and loss of heterozygosity at the PTEN locus were also evaluated. In addition, the expression levels of PDK1, PTEN, and Ki67 were analyzed by immunohistochemistry.

Results: Three cases carrying the E17K mutation in the AKTI gene and one case harboring the H1047R mutation in the PIK3CA gene were detected among the 36 cases. PDK1 was significantly overexpressed in the high-grade IPMN versus low-grade IPMN (P = 0.034) and in pancreatic and intestinal-type of IPMN versus gastric-type of IPMN (P = 0.020). Loss of PTEN expression was strongly associated with presence of invasive carcinoma and poor survival in these IPMN patients (P = 0.014).

Conclusion: This is the first report of AKTI mutations in IPMN. Our data indicate that oncogenic activation of the PI3K pathway can contribute to the progression of IPMN, in particular loss of PTEN expression. This finding suggests the potential employment of PI3K pathway-targeted therapies for IPMN patients. The incorporation of PTEN expression status in making surgical decisions may also benefit IPMN patients and should warrant further investigation.

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Introduction

Pancreatic adenocarcinoma is a malignancy of extremely poor prognosis with high mortality and short survival. Methods for its early detection and effective treatments, which will require an understanding of the underlying mechanisms of its development, are urgently needed.

Pancreatic adenocarcinoma is thought to develop from precancerous lesions such as pancreatic intraepithelial neoplasia (PanIN), intraductal papillary mucinous neoplasms (IPMN), and mucinous cystic neoplasm (MCN; ref. 1). IPMN is the most common cystic precancerous lesion of the pancreas, representing ~20% of surgically resected pancreatic neoplasms (2, 3). The prognosis for noninvasive IPMN is excellent and detection and surgical intervention at the preinvasive stage is curative (4, 5). The prognosis is poor for tumors associated invasive adenocarcinoma, but still more favorable than for conventional pancreatic ductal adenocarcinoma (PDAC) arising from PanIN lesions (6).

By definition, IPMN involve the main pancreatic duct and/or its branches; they are characterized by papillary projections of ductal epithelium and dilatation of the pancreatic duct. Histologically, IPMN are distinguished by the replacement of normal ductal epithelium with a mucinous epithelium showing a broad spectrum of histopathological changes, ranging from IPMN with low-grade dysplasia (adenoma), IPMN with moderate dysplasia (borderline tumor), and IPMN with high-grade dysplasia/carcinoma in situ (IPMC). IPMN may be associated with invasive adenocarcinoma showing stromal invasion (7). Currently, 4 histologic subtypes of IPMN by immunophenotypic characteristics of the lining epithelium are defined: a gastric foveolar-type (MUC1+ , MUC2−, MUC5AC+, MUC6+), an intestinal-
type (MUC1−, MUC2−, MUC5AC+, CDX2+), a pancreatobiliary-type (MUC1+, MUC2−, MUC5AC+, CDX2−), and an oncocytic-type (variable expression of MUC1 and MUC2; ref. 8). The intestinal-type usually affects the main pancreatic duct and if an associated invasive carcinoma is present, it is commonly of the mucinous/colloid type with a more favorable prognosis than the conventional PDAC. The pancreatobiliary-type often shows high-grade dysplasia with or without associated invasive PDAC, with a tubular morphology showing histologic features of the conventional type of PDAC. The gastric foveolar-type often affects branch ducts and is less commonly associated with an invasive tumor. The oncocytic-type is exceptionally rare. Approximately 20% to 45% of resected IPMNs have an invasive adenocarcinoma component (9, 10).

The specific mutations leading to the development of various histologic grades of IPMN have been partially characterized in previous studies (11–13). Reported genetic alterations identified in IPMN include mutations in KRAS (11, 14, 15), GNAS (16, 17), PIK3CA (12), and Braf (11). Other changes include the loss of expression of STK11/LKB1 (18–20) and overexpression of TP53 and ERBB2 proteins in the IPMNs (15), and allelic loss of PTEN in the pancreatic cyst fluid DNA of IPMN patients (21). A genetic analysis of microdissected IPMN of different grades within the same tumor has demonstrated early polyclonal epithelia gradually replaced by monoclonal neoplastic cells gaining tumor. The oncocytic-type is exceptionally rare. Approximately 20% to 45% of resected IPMNs have an invasive adenocarcinoma component (9, 10).

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IPMN in this study were all positive for MUC5A immunolabeling (data not shown) and included gastric, intestinal, and pancreatobiliary histologic subtypes. Detailed clinicopathological findings of the 36 IPMN cases are presented in Table 1. Patients with pancreatic neoplasms were recruited to our institutional prospective longitudinal outcomes study. Patients being followed for premalignant cystic neoplasms were evaluated with cross-sectional imaging and/or endoscopic ultrasound every 6 to 12 months depending on patient-specific demographics. Patients who have undergone pancreatic resection are evaluated and managed by our multidisciplinary group at regular intervals for the remainder of their lives.

Patients who underwent resection of a benign IPMN lesion were evaluated yearly with magnetic resonance imaging/magnetic resonance cholangiopancreatography (MRI/MRCP) to screen the remnant pancreas for new or recurrent disease. For patients with benign IPMN, no evidence of disease (NED) was defined as the absence of cystic changes within the remnant pancreas on follow-up MRI/MRCP imaging. Patients who underwent resection of a malignant IPMN were offered chemotherapy to reduce the risk of recurrence. Once this was completed, patients were seen every 3 months with repeat surveillance imaging at 6-month intervals. For patients with malignancies, recurrences were detected by imaging studies and serum tumor markers, and then confirmed by tissue biopsy. Patients were considered as NED when there was no elevation in serum tumor markers and no new findings on surveillance imaging. Length of follow-up in this study was calculated from the date of surgery to the last clinic appointment on record at our center.

Preparation of DNA extracts
To enrich the number of neoplastic cells procured from each sample, laser capture microdissection (LCM) was performed on the IPMN. The regions containing the IPMN neoplastic cell populations were microdissected for each case. Paraffin-embedded tumor samples were deparaffinized by incubating the slides in xylene for 2 minutes and rehydrated in 99.9% ethanol for 2 × 10 minutes, in 96% ethanol for 2 × 10 minutes, and in 70% ethanol for 2 × 10 minutes. Slides were stained with hematoxylin and eosin. Microdissection was carried out using a laser microdissection microscope (P.A.L.M., Bernried, Germany). Approximately between 10,000 and 14,000 cells were collected into 50 μL of ATL buffer (Lysis buffer from QIAamp DNA Mini Kit; QIAGEN). Surrounding nonneoplastic tissues were treated, microscopically defined, and dissected the same as the tumors, and served as the corresponding normal control for each sample. DNA extraction was performed according to manufacturer’s instructions.

Mutational analysis of the PIK3CA gene
Mutations in exons 9 and 20 of the PIK3CA gene were analyzed by direct genomic sequencing methods and confirmed by our previously described mutant-enriched sequencing method (44). PCR amplification of genomic DNA (40 ng each) and direct sequencing of the PCR products were performed using the same primers and conditions as previously described (44). All PCR fragments were purified using ExoSAP-IT kit (Affymetrix) and sequencing was performed with ABI Prism 3730xl DNA analyzers by Genewiz, Inc. using the PCR primers (44). Any alteration detected was further verified by sequencing of a second PCR product derived independently from the original DNA template.

Conventional direct genomic sequencing of the AKT1 gene
The point mutation G > A at nucleotide 49 of the AKT1 gene (E17K) was first examined by direct genomic sequenc-

ing. Genomic DNA was amplified with primers designed to amplify exclusively the hot-spot (AKT1-F 5'-ACATCGTGCAC-3' and AKT1-R 5'-GCCACGTCGTGTGCGCTGTT-3'; ref. 40). All PCR fragments were purified using Invitrogen PureLink PCR purification kit (Life Technologies) and sequencing was carried out with ABI 3730xl/DNA analyzers by Genewiz, Inc.

Mutant-enriched sequencing for detecting AKT1 mutation E17K
Based on the same principles that we have used to design the mutant-enriched sequencing method for PIK3CA (44), here we developed a sensitive mutant-enriched sequencing method specific for the E17K hot-spot mutation of the AKT1 gene (Fig. 1A). A mismatched primer was designed to create a unique restriction enzyme site for EcoRI in the AKT1 exon 2 region in the first round of PCR. The mismatched primer AKT1ME-F1 (5'-GCCGCCCGAGCTCTGATGAAAT-3') was used as the reverse primer for both rounds of PCR. The forward primers for the first and second PCR were AKT1ME-F1 (5'-GGCTGTGCGACAGTGCCCACAG-3') and AKT1ME-F2 (5'-ACACAGCTCGGGGTGCCGCTT-3'), respectively. EcoRI digestion was performed at 37°C overnight. The forward PCR primer AKT1ME-F2 was also used as the DNA-sequencing primer (Fig. 1A). An alteration detected was further verified at least twice by repeating the process starting with a second PCR product derived independently from the original DNA template.

The PCR condition for all the PCR reactions is 95°C, 5 minutes; (95°C, 30 seconds; 60°C, 30 seconds; 72°C, 30 seconds) × 25 and 40 cycles for first and second PCR, respectively; 72°C, 7 minutes.

To develop the method, a 583 bp product containing the E17K hot-spot mutation was amplified using a known mutant DNA sample as a template, using primers AKT1-G49AFwd 5'-ACATCGTGCAC-3' and AKT1-G49ARvs 5'-GCCACGTCGTGTGCGCTGTT-3'. The PCR product was subcloned into a pcDNA3.3-TOPO expression vector using the pcDNA3.3-TOPO cloning kit (Life Technologies). Mutant and wild-type colonies were sequenced and selected. To determine the approximate
#### Table 1. Summarized report of the 36 patient samples and genetic alterations identified in the PI3K/AKT/PTEN pathway

| Patient no. | Gender (age) | Cyst size (cm) | Location within pancreas | Main duct or mixed (MD) versus branch duct (BD) | IPMN lesion analyzed histologic subtype gastric = G, intestinal = I, pancreatobiliary = P, nuclear grade (1–3) | Resection specimen (highest grade of invasive cancer) | Outcome | PIK3CA mutant | AKT1 mutant | PIK3CA amp | AKT1 amp | PTEN LOH |
|-------------|--------------|----------------|--------------------------|-----------------------------------------------|-------------------------------------------------------------------------------------------------|---------------------------------------------------|---------|--------------|-------------|-------------|------------|----------|------------|
| 1           | F (70)       | 3.5            | Head                     | Main                                          | IPMN-2                                                                                          | Recurrence                                       | NED     | No           | No          | No          | Yes        | No       |
| 2           | F (79)       | 2.2            | Body                     | Main                                          | IPMN-3                                                                                          | DOD                                               | No      | No           | Yes         | No          | No         | Yes      |
| 3           | M (64)       | 2.0            | Head                     | Main                                          | IPMN-3                                                                                          | Recurrence                                       | No      | No           | No          | Yes         | No         | No       |
| 4           | M (64)       | 4.1            | Head                     | Main                                          | IPMN-1                                                                                          | Recurrence                                       | No      | No           | No          | No          | No         | No       |
| 5           | F (79)       | 4.5            | Head                     | Main                                          | IPMN-2                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 6           | M (78)       | 4.5            | Head                     | Main                                          | IPMN-3                                                                                          | DOD                                               | No      | No           | No          | No          | No         | No       |
| 7           | M (85)       | 5.5            | Head                     | N/A                                           | IPMN-3                                                                                          | pT3N1 DOD                                        | Recurrence | Yes         | No          | No          | No         | Yes      |
| 8           | F (44)       | 1.0            | Head                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | Yes         | No          | No         | No       |
| 9           | F (70)       | 3.0            | Head                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 10          | M (75)       | 3.0            | Head                     | Branch                                        | IPMN-3                                                                                          | Recur/DOD                                        | No      | No           | No          | No          | No         | No       |
| 11          | F (79)       | 2.0            | Head                     | Main                                          | IPMN-1                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 12          | F (75)       | 2.0            | Tail                     | Branch                                        | IPMN-2                                                                                          | DOD                                               | No      | No           | No          | No          | No         | No       |
| 13          | F (85)       | 4.3            | Body                     | Main                                          | IPMN-2                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 14          | F (36)       | 4.8            | Head                     | Main                                          | IPMN-2                                                                                          | NED                                               | Yes     | No           | No          | No          | No         | No       |
| 15          | F (66)       | 2.1            | Head                     | Main                                          | IPMN-1                                                                                          | Recur/DOD                                        | No      | No           | No          | No          | Yes        | Yes      |
| 16          | F (68)       | 1.7            | Tail                     | Main                                          | IPMN-3                                                                                          | H1047R                                            | No      | No           | No          | No          | No         | No       |
| 17          | M (59)       | 5.0            | Head                     | Branch                                        | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 18          | M (70)       | 1.6            | Body                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 19          | M (70)       | 1.5            | Head/neck                | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 20          | M (73)       | 1.3            | Body                     | Main                                          | IPMN-3                                                                                          | NED                                               | Yes     | No           | Yes         | No          | No         | No       |
| 21          | F (58)       | 1.2            | Body                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 22          | M (66)       | 1.5            | Head                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 23          | F (58)       | 4.0            | Head                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 24          | M (58)       | 5.1            | Body                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 25          | M (73)       | 0.5            | Body/neck                | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | Yes         | No          | No         | No       |
| 26          | F (78)       | 4.5            | Distal                   | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 27          | F (67)       | 2.6            | Body                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 28          | M (66)       | 1.3            | Distal                   | Branch                                        | IPMN-3                                                                                          | NED                                               | Yes     | No           | No          | No          | No         | No       |
| 29          | F (55)       | 1.2            | Distal                   | Head/neck                                     | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 30          | F (58)       | 2.0            | Distal                   | Body/neck                                     | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 31          | F (82)       | 2.0            | Head                     | Main                                          | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |
| 32          | M (72)       | 1.1            | Head/Body                | Branch                                        | IPMN-3                                                                                          | NED                                               | No      | No           | No          | No          | No         | No       |

IPMN-1 (low-grade dysplasia) = IPMN, adenoma; IPMN-2 (moderate dysplasia) = IPMN, borderline tumor; IPMN-3 (high-grade dysplasia) = IPMC, carcinoma in situ; NED, no evidence of disease; DOD, death of disease; NA, not informative.
sensitivity of our assay, we made serial dilutions of the mutant plasmid with the wild-type plasmid. When the ratio of the mutant to wild-type DNA copies reached 1:100, the mutant-enriched sequencing result still contained a recognizable mutant peak. However, in conventional direct genomic sequencing, the mutant peak disappeared when the ratio of mutant and wild-type DNA was lower than 1:10. This indicated that the mutant-enriched sequencing method was at least 10 times more sensitive than the conventional sequencing method.

Figure 1. The detection of AKT1 E17K mutation with the mutant-enriched sequencing method. A, the schematic for the mutant-enriched sequencing method for AKT1 E17K. Briefly, a unique restriction enzyme site EcoRI was introduced by mismatch PCR to the wild-type DNA strand but not the mutant sequence in the first round of PCR. The mismatch primer (AKT1ME-R) has 2 nucleotide substitutions (G → A and A → T). Subsequent EcoRI digestion of the wild-type DNA strands would allow preferential amplification of the mutant templates in the second round of PCR. B, detection of AKT1 E17K mutation by mutant-enriched method. AKT1 E17K mutation was not detectable by the conventional direct sequencing method in cases #7, #20, and #35, but was enriched and visible with the mutant-enriched sequencing method.
AKT1 E17K Taqman mutation detection assay

AKT1 E17K mutation was further confirmed using a Competitive Allele-Specific Taqman PCR (castPCR; Life Technologies). Fresh DNA from each of the potential positive cases was extracted as follows: five to ten 10-μm serial sections were microdissected for each case using a clean sterile scalpel for each of the 3 samples. After paraffin-containing tissue was scraped, DNA was extracted using QIAamp DNA FFPE Mini Kit (QIAGEN) following manufacturer’s instructions.

Each specific AKT1 E17K mutant allele assay (Assay ID: Hs0000986_mu) contains: an allele-specific primer that detects the mutant allele, a MGB oligonucleotide blocker that suppresses the wild-type allele, and a locus-specific Taqman FAM dye-labeled MGB probe. A gene reference assay (assay ID: Hs0001010_rf) designed to amplify a mutation-free and polymorphism-free region of the target gene was used in parallel. Each assay contains: a locus-specific pair of forward and reverse primers and a locus-specific Taqman FAM dye-labeled MGB probe.

AKT1 E17K mutation detection experiments were performed in ABI 7500 Sequence Detector (Applied Biosystems) following the manufacturer’s instructions. In brief, after amplification, the Ct values were determined by the Applied Biosystems real-time PCR instrument software. Using 4 different normal (wild-type) gDNA samples in triplicates, a mutation detection ΔCt cutoff value was determined [(Detection ΔCt cutoff = Average ΔCt − (3 × the standard deviation)]. Data files containing the samples Ct values were imported into Life Technologies Mutation Detector Software. In the analysis calculations, the difference between the Ct value of the mutant allele and the Ct value of the gene reference assay was calculated; this ΔCt value represents the quantity of the specific mutation allele detected within each sample. Values less than the Detection ΔCt cutoff value (ΔCt = 7.17) were considered positive for the mutation.

PIK3CA and AKT1 gene amplification

Gene amplification was evaluated by quantitative real-time PCR (Q-PCR), performed in ABI Prism 7500 Sequence Detector (Applied Biosystems) using Power SyBr Green PCR Master Mix and the following oligonucleotides: for the PIK3CA gene (Chr. 3q26.3), Fwd (5'-ATCTTGTCTTCTAATGATGGTGTTTCGAATG-3'), Rvs (5'-CTAGGGTGTTCAGGAAATGTTATG-3'); COL7A1 (collagen, type VII, α1; Chr. 3p21.1) as the reference gene, Fwd (5'-ACCAGTACCAGCATCA-TTGTG-3'), Rvs (5'-TCAGGGTTCAATCTCAGTGTG-3'). For the AKT1 gene (Chr. 14q32.3), Fwd (5'-ACGGGCA-CATAAGATACACA-A-3'), Rvs (5'-TGCCCAGAAAAGGTGTCATT-CATG-3') and DHSR4 (dehydrogenase/reductase (SDR family) member 4; Chr. 14q11.2) as the reference gene, Fwd (5'-GGTAGCTAGGAGCCAGCTCA-3'), Rvs (5'-ATGATTGGGCCCAGAAGGG-3'). Optimal primer concentrations were determined using optimization protocols from Applied Biosystems SYBR Green PCR master mix manual.

Dissociation curve analysis of all the PCR products showed a simple peak and the correct size of each product was confirmed by agarose gel electrophoresis. The relative copy number for PIK3CA and AKT1 was calculated using the 2−ΔΔCt method. ΔΔCt represents the difference between the pair of tissue samples (ΔCt tumor – ΔCt of matched normal), with ΔCt being the average Ct for the target gene (PIK3CA and AKT1) minus the average Ct for the reference gene (COL7A1 or DHSR4, respectively). Values greater than 2.0 were considered positive for gene amplification.

SNP-PCR-RFLP PTEN LOH analysis

LOH analyses of the PTEN locus were assessed by single nucleotide polymorphism-PCR-restriction fragment length polymorphism (SNP-PCR-RFLP) analysis using 7 SNP markers on the tumor and matched non-neoplastic DNA from 36 IPMN samples. All SNPs were intragenic to the PTEN gene. To select SNP markers, genotypic and allelic frequency of 680 SNPs present in the PTEN gene was obtained from the Ensembl database. SNPs reported to have high heterozygosity indices were chosen. Twenty-seven SNPs with high heterozygosity indices were checked for the presence of restriction endonuclease site using the SNP cutter program (available at http://bioapp-psych.uic.edu/SNP_cutter.htm.) as described previously (45). We identified 8 SNPs; each harbors a unique restriction site. Primers were designed for each SNP to generate a PCR product of 200 to 350 bp. Of the 8 SNPs, 7 yielded bands of distinguishable unequal sizes upon restriction digestion that were detectable on an agarose gel. The selected 7 SNPs were then used for the LOH analyses. Digestion of the PCR products was carried out with appropriate restriction enzymes (New England Biolabs Inc.) and resolved on 1% agarose gel. The relative copy number for PIK3CA and AKT1 was calculated using the 2−ΔΔCt method. ΔΔCt represents the difference between the pair of tissue samples (ΔCt tumor – ΔCt of matched normal), with ΔCt being the average Ct for the target gene (PIK3CA and AKT1) minus the average Ct for the reference gene (COL7A1 or DHSR4, respectively). Values greater than 2.0 were considered positive for gene amplification.
Table 2. The list of primers and restriction enzymes for the SNP-PCR-RFLP analyses at the PTEN locus

<table>
<thead>
<tr>
<th>Ref. SNP#</th>
<th>Restriction enzyme</th>
<th>Fwd primer (5'-3')</th>
<th>Rvs primer (5'-3')</th>
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<tbody>
<tr>
<td>rs1903858</td>
<td>HindIII</td>
<td>TTCTGCGAGGAAATCCCATAGC</td>
<td>TAGCCAGCTTAAATGCTGACTTC</td>
</tr>
<tr>
<td>rs1234225</td>
<td>PvuII</td>
<td>AACAATGTGAAAGGTGTCAAA</td>
<td>TGATGGAATCCACAGGTCTCT</td>
</tr>
<tr>
<td>rs10490920</td>
<td>Ncol</td>
<td>TCAAGAGCTCCAGAGGACTT</td>
<td>AGACAAGACAGGCCACCTAA</td>
</tr>
<tr>
<td>rs2735343</td>
<td>HhaI</td>
<td>AGTGGAGACAGACTGACCTG</td>
<td>CTGTAAGATCATATCGTTG</td>
</tr>
<tr>
<td>rs701848</td>
<td>HaeIII</td>
<td>TCTTACATGTGCTTTATTGATTTGC</td>
<td>TTTGAAGACACAAATTTCTGGA</td>
</tr>
<tr>
<td>rs1234224</td>
<td>AsI/SstI</td>
<td>GCAAGTGTGGCAGAAGGTAAACCC</td>
<td>CCACGTGTCCTCTATTCCACCC</td>
</tr>
<tr>
<td>rs34421660</td>
<td>No enzyme</td>
<td>AGAAAGTGACCTGATATTACCTAA</td>
<td>ATTCGCTCTTGGAAACCT</td>
</tr>
</tbody>
</table>

Note that SNP rs34421660 is an insertion/deletion of 32-bp and it does not require restriction enzyme digestion to differentiate the 2 alleles.

1:50 dilution; Cell Signaling Technology). PKB Kinase (E-3; 1:2000; dilution; Santa Cruz Biotechnology), and monoclonal Ki67 (1:100 dilution; Dako). Counterstaining with hematoxylin for 1 minute was the final step. Level of protein expression refers to the intensity of mucinous epithelium only, and was analyzed as follows: PTEN IHC was scored as normal (2), decreased (1), or negative (0). Benign pancreatic tissues sampled in the TMA served as positive and negative control. Pancreatic ducts and centroacinar cells display normal PTEN expression (2) and served as positive internal controls, whereas acinar cells do not express PTEN and served as negative control (0). PKB kinase IHC was scored as weak (1) or strong (2) based on the cytoplasmic intensity of the neoplastic epithelium. Non-neoplastic pancreatic ducts served as positive control for weak expression (1), pancreatic acinar cells served as positive control for strong expression (2). Human normal alveolar lung tissue served as negative control. Ki67 IHC was scored as percentage of neoplastic epithelial cells showing nuclear expression. All the TMAs included a negative control. The pathologists who performed the scoring (H.E.R. and A.T.T.) were blinded with regard to clinical data and genetic data until the completion of our study.

Statistical analysis

For statistical purposes, clinicopathological features were dichotomized as: IPMN lesion: IPMN (IPMN-1/2) and IPMC (IPMN-3). Nuclear grade: low (1–2) and high (3). PTEN expression level was dichotomized as low/absent (≤1) or normal (>1). The expression of PDK1 was adopted as >1-cm. The χ² test (Fisher exact test for small cell counts) was used for comparison between categorical variables and Student t test for parametric continuous variables. Multivariable logistic regression models were conducted to calculate the odds ratio and 95% confidence interval (CI) for the association between PTEN expression and the presence of invasive carcinoma; the model adjusted for PDK1 expression. Survival curves were calculated using Kaplan–Meier product–limit estimate. Differences between survival times were analyzed by the log-rank method.

Multivariable Cox proportional hazards models were used to calculate HRs and 95% CI of the association between risk factors and survival. We included variables in our multivariable model that were statistically significant in the univariate analysis; these variables include: PTEN expression, presence of invasive carcinoma, and age). All tests were 2-sided. P values ≤ 0.05 were considered statistically significant. All statistical analyses were performed using SPSS statistical software version 17.0 (SPSS Inc.).

Results

Mutations in PIK3CA and AKT1

IPMN were examined for the presence of the hot-spot mutations in the AKT1 and PIK3CA genes. We used both direct genomic sequencing and mutant-enriched sequencing methods on the DNA samples extracted from 36 micro-dissected IPMN specimens. One patient with borderline IPMN (moderate dysplasia) harbored the H1047R (A3140G) hot-spot mutation of the PIK3CA gene in the IPMN (data not shown). The E17K (A49G) AKT1 gene mutation was detected in 3 IPMN; 2 of the 3 were intestinal-type and 1 was gastric-type IPMN. Although the PIK3CA mutation was detected by direct genomic sequencing, the 3 AKT1 mutations were identified only when the newly developed mutant-enriched sequencing method was used (Fig. 1). Using Taqman Mutation Detection Assay for AKT1 and PIK3CA gene amplification was dissected IPMN specimens. One patient with borderline IPMN (moderate dysplasia) harbored the H1047R (A3140G) hot-spot mutation of the PIK3CA gene in the IPMN (data not shown). The E17K (A49G) AKT1 gene mutation was detected in 3 IPMN; 2 of the 3 were intestinal-type and 1 was gastric-type IPMN. Although the PIK3CA mutation was detected by direct genomic sequencing, the 3 AKT1 mutations were identified only when the newly developed mutant-enriched sequencing method was used (Fig. 1). Using Taqman Mutation Detection Assay for AKT1 and PIK3CA gene amplification was used to calculate HRs and 95% CI of the association between risk factors and survival. We included variables in our multivariable model that were statistically significant in the univariate analysis; these variables include: PTEN expression, presence of invasive carcinoma, and age). All tests were 2-sided. P values ≤ 0.05 were considered statistically significant. All statistical analyses were performed using SPSS statistical software version 17.0 (SPSS Inc.).

Analysis of AKT1 and PIK3CA gene amplification in IPMN

The AKT1 and PIK3CA gene loci were investigated for potential amplification in the 36 IPMN specimens by Q-PCR. Gene amplification of the AKT1 locus was detected in one IPMN specimen (2.8%). PIK3CA gene amplification was detected in 7 (19.4%) of the 36 IPMN. The case with the AKT1 amplification did not show co-amplification with PIK3CA. The amplification of either of these 2 genes did not correlate with any of the clinicopathological parameters.
Analysis of PTEN LOH

Of the 36 cases, 28 (77.8%) showed heterozygosity at the PTEN locus by SNP-PCR-RFLP, and 8 were not informative (as defined in the Material and Methods) for all the 7 SNPs analyzed. Of the 28 heterozygous cases, 10 displayed LOH (10/28, 35.7%; Table 1).

PTEN, PDK1, and Ki67 protein expression in IPMN

The protein expression levels of PTEN, PDK1, and Ki67 were studied by immunohistochemistry. PTEN expression was low or absent in 11 (30.6%) of the 36 cases. PDK1 was overexpressed in 20 (55.6%) of the 36 cases. Ki67 score was high in 18 (50%) of the 36 cases; 16 out of the 18 (88.8%) were IPMC and 2 were IPMN without carcinoma. In this set of IPMN samples, strong immunolabeling of PDK1 was associated with higher Ki67 labeling index (Fisher exact, \( P = 0.018 \)). Representative staining of the IPMN samples expressing different levels of PTEN and PDK1 are shown in Fig. 2.

Associations of PTEN and PDK1 expression with clinicopathological features and patient outcome

Downregulation of PTEN at protein level was associated with the 3 following clinicopathological parameters: nuclear grade, presence of invasive carcinoma, and disease outcome. PDK1 overexpression, however, showed association with IPMN subtypes and progression (Table 3). PTEN protein expression was more frequently downregulated in the IPMN with nuclear grade 3 versus IPMN with nuclear grade 1 and/or 2 (Fisher exact, \( P = 0.027 \)).

PDK1 protein expression was detected in each IPMN subtype but was more frequently overexpressed in intestinal and pancreatic-type IPMN versus gastric-type IPMN (\( \chi^2, P = 0.020 \); Table 3). It was also observed that PDK1 was more commonly overexpressed in IPMC than IPMN adenoma and borderline tumor (Fisher exact, \( P = 0.034 \)), showing an increasing trend with IPMN progression: adenoma (0/2; 0%), borderline (3/9; 33.3%), and in situ (17/25; 68%). All the cases with PDAC associated, (7/7; 100%) showed strong PDK1 expression. Moreover, PDK1 overexpression showed a linear correlation with the maximum cyst diameter detected (\( \chi^2, P = 0.044 \)).

A striking difference in survival was observed between patients with normal expression of PTEN and those with reduced or absent PTEN expression; the latter group experienced shortened disease-specific survival [median survival time: PTEN normal = 60.6 months, PTEN low = 48.9
As expected, IPMN patients with an invasive carcinoma had poorer survival than those without invasive carcinoma; a 12-fold increase risk of mortality was observed in multivariable models adjusting for PTEN expression and presence of invasive carcinoma (HR = 11.9; 95% CI, 1.7–83.5; P = 0.012); addition of age to the model did not affect the result. This result may indicate that the association of PTEN expression with patient survival was only because of the presence of invasive carcinoma (Fisher exact, P = 0.018). When we examined the factors associated with the presence of invasive carcinoma, absent or weak PTEN expression was associated with a 8-fold increase in the odds controlling for PDK1 expression (OR = 8.3; 95% CI, 1.034–67.142; P = 0.046). However, these results are based on a small sample size.

### Discussion

In this study, we conducted a comprehensive evaluation of the PI3K signaling pathway in 36 well-characterized IPMN samples to understand the extent of its dysregulation in IPMN and the potential impact on clinical outcome.

We identified the H1047R mutation of PIK3CA in 1 (2.7%) of the 36 IPMN. We also detected the E17K AKT1 gene mutation in 3 (8.3%) of the 36 IPMN cases. The H1047R mutation of PIK3CA is known to affect the functionally important kinase domain of the protein, conferring an increased lipid kinase activity, which can lead to the activation of PI3K signaling pathway in the absence of growth factors. The mutation has been shown to be sufficient to induce oncogenic cell transformation of chicken embryo fibroblasts (CEF) and NIH3T3 cells (36, 37, 47). It has also been described that p110α mutants induce in vivo angiogenesis and malignant cell growth in chorioallantoic membrane of chick embryo and cause hemangiosarcomas in young chickens (36).

We have previously reported PIK3CA gene mutations in 11% of IPMN, suggesting that the PIK3CA gene and its pathway may have a role in IPMN but not PanIN/PDA tumorigenesis (12). In our previous study, PIK3CA gene mutations were detected in 3 high-grade IPMN, and 1 IPMN with moderate dysplasia, with the caveat that the majority of the cases examined had high-grade dysplasia. One of the 4 mutations detected in the previous study was H1047R in an IPMC with invasion, the other 3 were not hot-spot mutations (12). In this study, more IPMN with low-grade and moderate dysplasia were included and the hot-spot mutation H1047R was detected in an IPMN with moderate dysplasia. The mutation frequency is slightly lower in this

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**Table 3. Associations of reduced PTEN protein expression and PDK1 overexpression with clinicopathological features, type, lesion, nuclear grade, and patient outcome**

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>No.</th>
<th>Weak/absent PTEN expression (%)</th>
<th>P&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Strong PDK1 expression (%)</th>
<th>P&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age, y</td>
<td>68.9</td>
<td>46</td>
<td>0.863&lt;sup&gt;b&lt;/sup&gt;</td>
<td>89</td>
<td>0.667&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Mean cyst size, cm</td>
<td>2.8</td>
<td>2.8</td>
<td>0.979&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.3</td>
<td>0.044&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>IPMN type</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gastric</td>
<td>14</td>
<td>2 (14)</td>
<td>0.060</td>
<td>4 (29)</td>
<td>0.020&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Pancreatic</td>
<td>8</td>
<td>5 (62)</td>
<td></td>
<td>7 (87)</td>
<td></td>
</tr>
<tr>
<td>Intestinal</td>
<td>14</td>
<td>4 (29)</td>
<td></td>
<td>9 (64)</td>
<td></td>
</tr>
<tr>
<td>IPMN lesion</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>IPMN</td>
<td>11</td>
<td>2 (18)</td>
<td>0.439</td>
<td>3 (27)</td>
<td>0.034&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>IPMC</td>
<td>25</td>
<td>9 (36)</td>
<td></td>
<td>17 (68)</td>
<td></td>
</tr>
<tr>
<td>Nuclear grade</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Low (1–2)</td>
<td>18</td>
<td>2 (11)</td>
<td>0.027&lt;sup&gt;c&lt;/sup&gt;</td>
<td>9 (50)</td>
<td>0.738</td>
</tr>
<tr>
<td>High (3)</td>
<td>18</td>
<td>9 (50)</td>
<td></td>
<td>11 (61)</td>
<td></td>
</tr>
<tr>
<td>Disease status</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alive</td>
<td>29</td>
<td>6 (21)</td>
<td>0.018&lt;sup&gt;c&lt;/sup&gt;</td>
<td>15 (52)</td>
<td>0.426</td>
</tr>
<tr>
<td>Deceased</td>
<td>7</td>
<td>5 (71)</td>
<td></td>
<td>5 (71)</td>
<td></td>
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<tr>
<td>Recurrence</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>6</td>
<td>2 (33)</td>
<td>0.871</td>
<td>4 (67)</td>
<td>1.000</td>
</tr>
<tr>
<td>No</td>
<td>30</td>
<td>9 (30)</td>
<td></td>
<td>16 (53)</td>
<td></td>
</tr>
</tbody>
</table>

<sup>a</sup>x<sup>2</sup> test/Fisher exact test.  
<sup>b</sup>Student t test.  
<sup>c</sup>Statistically significant.

The following variables were dichotomized for statistical analysis: nuclear grade was dichotomized as low (nuclear grades 1–2) and high (nuclear grade 3); IPMN lesion was grouped as IPMN (IPMN-1 or IPMN, adenoma and IPMN-2 or IPMN, borderline tumor) and IPMC (IPMN-3 or IPMC, carcinoma in situ); disease status was grouped as alive (no evidence of disease or recurrence) and deceased (death of disease).
study probably because of the facts that more low-grade cases were included and only hot-spot mutation sites were examined. Nevertheless, a number of recent studies encompassing different histologic grades of IPMN are in agreement with our findings, supporting an important role for PIK3CA in IPMN (12, 39, 48). There is one exception in which PIK3CA mutations were not detected in IPMN but were identified in intraductal tubulopapillary neoplasms of the pancreas (ITNP), a rare variant of intraductal neoplasm of the pancreas recently recognized as a new class of pancreatic tumor (13). Consistent expression of MUC5A among our samples definitively confirmed them as IPMN and ruled out ITNP (data not shown; ref. 9).

The mutational status of AKT1 gene has not been examined before IPMN. Here we show that by using a novel mutant-enchanced method, we were able to detect the E17K mutation in 3 (8.3%) of the 36 IPMN cases; 2 of those were IPMC associated with invasive components. The detection of the AKT1 E17K mutation in IPMN but not PDA further signified the unique importance of the PI3K signaling pathway in IPMN progression. Although presence of false positive mutations could be possible because of a low-quantity DNA extracted from the tumors and/or a potential nontemplate extension of the AKT1ME-R primer used in the mutant-enchanced assay, the absence of a mutant peak on any of the paired normal samples examined suggests that this a rather infrequent event. Furthermore, each positive sample was verified from an independent PCR of the original genomic DNA at least 2 more times. To further rule out false positivity, new DNA from the same potential positive cases was extracted, the presence of AKT1 E17K mutations was then analyzed using the AKT1 E17K Taqman Mutation Detection Assay. By using this method, we were able to confirm 2 of the 3 potential positive cases. The apparent discrepancy could be because of a number of reasons: (1) different mutant to wild-type ratio between the different DNA samples from the same patients; (2) different sensitivity of the 2 methods applied; or (3) lack of calibration ΔCt value for the AKT1_33765_mu assay, could lead to a smaller detection ΔCt cutoff calculated with our samples, which could limit the detection of mutant allele.

Intriguingly, PIK3CA and AKT1 genes mutations were not detected in PanIN/PDA (35, 41, 49, 50). This suggests that, although IPMN and PanIN share overlapped molecular alterations, such as KRAS, p16, p53, and SMAD4 (9), pancreatic carcinoma associated with IPMN may arise through a different molecular pathway from PanIN/PDA. Although STK11/LKB1 gene is inactivated more frequently in IPMN than in PanIN/PDA (18–20), PIK3CA and AKT1 are the only genes with mutations unique to IPMN and not detectable in PanIN/PDA thus far (12, 35, 41, 48–50). Direct comparisons of IPMN and PanIN progression might be instructive in characterizing the divergent molecular and histologic pathways of pancreatic cancer evolution between the 2 and reveal additional unique alterations in IPMN.

To further examine the status of the PI3K pathway at the genomic level, we investigated the LOH at the PTEN locus and it was found in 10 of the 28 (35.7%) IPMN analyzed. Albeit the small sample number, LOH frequency at PTEN steadily increased as IPMN progressed (0/1 or 0% IPMN, adenoma; 2/7 or 28.6% IPMN, borderline; 8/20 or 40% IPMC, in situ). The same increasing frequency could be observed with the nuclear grade (1/4 or 25% nuclear grade 1; 4/12 or 33.3% nuclear grade 2; 5/12 or 41.7% nuclear grade 3). There was no tumor size, gender, or age bias associated with the LOH status at PTEN. PIK3CA was amplified in 19.4% of the IPMN analyzed, whereas AKT1 was amplified in only one IPMN associated with invasive carcinoma. These events did not correlate with the clinical variables.

Although we found PDK1 to be frequently overexpressed in our IPMN samples (20/36 or 55.6%), this overexpression was not associated with patient survival. However, PDK1 expression was significantly higher in intestinal and pancreatic (8/8 or 100%) and intestinal (13/14 or 92.8%; ref. 51). PDK1 immunolabeling was more frequently upregulated in IPMC versus IPMN (P = 0.034). Both LOH at PTEN and PDK1 overexpression patterns further depict an increasing aberrant PI3K signaling accompanying IPMN progression. Future studies are needed to define the mechanisms leading to PDK1 overexpression on IPMN samples and whether or not this correlates with PDK1 activation; this could help understand the contribution of PDK1 to PI3K signaling in these patients.

Weak or absent PTEN expression was observed in 30.6% of the IPMN analyzed by IHC. This reduced expression was significantly associated with higher nuclear grade.
Furthermore, we found that weak or loss of PTEN expression was associated with reduced survival rate among our IPMN patients ($P = 0.014$). However, multivariate Cox analysis showed that the presence of invasive carcinoma was the only independent factor of poor prognosis. The low number of IPMN samples with associated invasive carcinoma included in the study ($n = 6$) and the strong association between the presence of invasive carcinoma and low/absent PTEN expression in this set of IPMN render it difficult to separate the effect of these 2 variables on survival. Keeping this limitation in mind, our findings suggest that PTEN expression level may have potential diagnostic and prognostic values, and may need to be taken into consideration when determining treatment options for IPMN patients. Future larger studies are needed to further evaluate the clinical values of PTEN as an independent prognostic marker distinct from invasive cancer. In addition, determining if expression levels of PTEN in cells isolated from cyst fluid or biopsy and within IPMN are correlated with the presence of invasive cancer and patient prognosis will be key to preoperative risk assessment and stratification. These studies will be crucial next steps in determining the utility of PTEN expression in making surgical decisions.

Within this manuscript, we examined a large number of associations, and we observed a limited number of significant associations. Thus, because of the issues of multiple comparisons, we are cautious in our interpretations of the results; some of the statistically significant associations we observed may be explained by chance. Using a conservative approach of Bonferroni correction, we would require a $P$-value of $< 0.003$ to achieve statistical significance. Based on this $P$-value, none of our results would be statistically significant, and may all be a result of chance. However, these statistically significant findings are supported by well-defined biologic rationale (e.g., PDK1 overexpression and PTEN loss in IPMN progression are logical; refs. 26 and 31). Thus, these results require replication in larger studies.

Taken together, these data indicate that aberrations of the PI3K pathway are common in IPMN and suggest an important role for this signaling pathway in cystic tumors of the pancreas. This is the first report of AKT1 mutations in IPMN tumors. Although a larger scale study designed to determine if PTEN is an independent variable from invasive carcinoma is needed, nevertheless, these results indicate that dysregulated PI3K signaling pathway is involved in the malignant progression of IPMN. If inhibition of the PI3K signaling pathway can slow progression of IPMN and development of invasive cancer, patients may benefit from novel therapeutic specifically targeting PDK1, mTOR, and PIK3CA as neoadjuvant or adjuvant therapies.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Authors’ Contributions
Conception and design: D. Garcia-Carracedo, H. Remotti, G.H. Su
Development of methodology: D. Garcia-Carracedo, N. Akhavan, B. Tweel, R. Parsons, G.H. Su
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): D. Garcia-Carracedo, A. Turk, B. Tweel, J.A. Chabot, J. Allendorf, H. Remotti, G.H. Su
Analysis and interpretation of data (e.g., statistical analysis, bioinformatics, computational analysis): D. Garcia-Carracedo, A. Turk, S. Fine, N. Akhavan, J. Genkinger, G.H. Su
Writing, review, and/or revision of the manuscript: D. Garcia-Carracedo, A. Turk, J. Allendorf, J. Genkinger, H. Remotti, G.H. Su
Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): D. Garcia-Carracedo, S. Fine, N. Akhavan, J.A. Chabot, H. Remotti, G.H. Su
Study supervision: D. Garcia-Carracedo, G.H. Su

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References

Loss of PTEN Expression Is Associated with Poor Prognosis in Patients with Intraductal Papillary Mucinous Neoplasms of the Pancreas
