The CREB-Binding Protein Inhibitor ICG-001 Suppresses Pancreatic Cancer Growth

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Abstract

Pancreatic ductal adenocarcinoma (PDAC) is a highly lethal cancer due in part to a lack of highly robust cytotoxic or molecular-based therapies. Recent studies investigating ligand-mediated Wnt/β-catenin signaling have highlighted its importance in pancreatic cancer initiation and progression, as well as its potential as a therapeutic target in PDAC. The small-molecule ICG-001 binds cAMP-responsive element binding (CREB)-binding protein (CBP) to disrupt its interaction with β-catenin and inhibit CBP function as a coactivator of Wnt/β-catenin-mediated transcription. Given its ability to inhibit Wnt/β-catenin-mediated transcription in vitro and in vivo, as well as its efficacy in preclinical models of colorectal cancer and other Wnt-driven diseases, we examined ICG-001 and its potential role as a therapeutic in PDAC. ICG-001 alone significantly inhibited anchorage-dependent and -independent growth of multiple PDAC lines, and augmented in vitro growth inhibition when used in combination with gemcitabine. ICG-001 had only variable modest effects on PDAC apoptosis and instead mediated PDAC growth inhibition primarily through robust induction of G1 cell-cycle arrest. These effects, however, seemed decoupled from its inhibition of Wnt/β-catenin-mediated transcription. DNA microarrays performed on PDAC cells in the context of ICG-001 treatment revealed ICG-001 altered the expression of several genes with well-established roles in DNA replication and cell-cycle progression, including direct actions on SKP2 and CDKN1A. ICG-001 also significantly prolonged survival in an in vivo orthotopic xenograft model of PDAC, indicating ICG-001 or derived compounds that disrupt CBP activity are potentially useful small-molecule therapeutics for pancreatic cancer. Mol Cancer Ther; 13(10); 2303–14. ©2014 AACR
ICG-001 significantly inhibited malignancies (NCT01606579; ref. 26). Advanced solid tumors (NCT01302405) and myeloid differentiation as a Wnt inhibitor in early-phase clinical trials for fibrosis (24), and salivary tumorigenesis (25). The ICG-20, renal interstitial fibrosis (21), acute lymphoblastic leukemia (22), chronic myeloid leukemia (23), dermal fibrosis (24), and salivary tumorigenesis (25). The ICG-001-derived compound PRI-724 is now under investigation as a Wnt inhibitor in early-phase clinical trials for advanced solid tumors (NCT01302405) and myeloid malignancies (NCT01606579; ref. 26).

Given the importance of Wnt signaling in pancreatic carcinogenesis, we have now explored therapeutic potential and mechanism of action of ICG-001 in PDAC. ICG-001 significantly inhibited in vitro and in vivo PDAC growth by inducing G1 cell-cycle arrest through effects that were largely decoupled from its activity as a Wnt inhibitor. Instead, ICG-001 seemed to more broadly affect CBP function as a cotranscriptional activator, directly and indirectly perturbing the expression of numerous genes with key roles in cell-cycle progression.

Materials and Methods

Cell lines and reagents

All cell lines were cultured as previously described (7). AsPC-1, MiaPaCa-2, and PANC-1 were obtained in 2005 from the ATCC (Rockville, MD). L3.6pl was obtained in 2010 from Hong Wu (UCLA). The cell lines have not been subsequently authenticated since receipt. ICG-001 was purchased from Selleck Chem and gemcitabine was kindly provided by Timothy Donahue (UCLA). Wnt3a and β-catenin (Sigma; C2206) and tubulin (Santa Cruz; sc-5546) antibodies. Western blot analyses to verify knockdown were performed using primers listed in Supplementary Table S1.

Cell growth, proliferation, and apoptosis assays

MTT cell viability assays (ATCC) were carried out per the manufacturer’s instructions at initial plating of 5,000 (AsPC-1 and L3.6pl) or 3,000 (MiaPaCa-2 and PANC-1) cells per well in 96-well plates. Soft agar assays were prepared as previously described (7) in 48-well format. Media and drug were replenished once every 3 to 4 days. For cell-cycle analysis, treated cells were stained using hypotonic DNA staining buffer: 0.1 mg/mL propidium iodide (PI; Calbiochem), 20 μg/mL RNaseA, 1 mg/mL sodium citrate, and 0.3% Triton X-100. Flow cytometry analysis was performed in the UCLA Flow Cytometry Core using a Becton Dickinson FACSCalibur cytometer and using ModFit LT v3.1 software. Annexin-V–PI flow cytometry analysis was performed with Cellquest v3.3 software using an Annexin V FITC Apoptosis Detection Kit (BD Biosciences).

Orthotopic xenograft tumor assay

Animal work was performed with oversight by the UCLA Division of Laboratory Animal Medicine and approval from the UCLA Animal Research Committee. Orthotopic tumors were established by injecting 5 × 10⁵ AsPC-1 cells suspended in 50% Matrigel (BD Biosciences)/RPMI into surgically exposed pancreatic tails of 6-week-old female athymic nude mice (Charles River) as previously described (27). Intraperitoneal drug injections were begun 10 days after tumor cell injections in animals randomized to four treatment arms: vehicle alone (20% PEG300, 5% DMSO/1% DMSO in PBS; n = 12); ICG-001 (5 mg/kg 6 days/week; n = 12); gemcitabine (25 mg/kg 2 days/week; n = 11); or ICG-001 + gemcitabine (n = 12). Treatment was for 8 weeks and then discontinued for the remainder of the study (up to 120 days after surgery). Animals were monitored daily and sacrificed per institutional guidelines when reaching either a Body Conditioning Score of 2 (28) or palpable tumor of 1.5 cm in diameter. Full necropsy was performed at the time of death with snap-frozen and formalin-fixed tissue sections taken of primary and metastatic tumors, as well as normal spleen, lung, liver, small, and large intestine. Histologic analysis was performed with hematoxylin and eosin (H&E)–stained sections by a practicing surgical pathologist (D.W. Dawson) blinded to treatment arms. Ten random high power fields per tumor were analyzed for mitotic counts. A previously described dissemination score awarding points based on tumor infiltration and metastasis (29) was used to quantify tumor spread.

Wnt reporter assays

Dual luciferase assays (Promega) were performed as previously described (7) on cell lines stably transduced with lentiviral constructs containing β-catenin–activated reporter (BAR) driving firefly luciferase expression and separate normalization control construct with constitutive EF1α promoter driving Renilla expression.

Gene knockdown

Lipofectamine 2000 was used per the manufacturer’s instructions to transfect AsPC-1 cells with 20 nmol/L control siRNA (#12935-200) or CTNNB1 #437 (#4390824) siRNA (all purchased from Life Technologies). Western blot analyses to verify knockdown were performed using β-catenin (Sigma; C2206) and tubulin (Santa Cruz; sc-5546) antibodies.

Real-time PCR

RNA extraction, cDNA synthesis, and SYBR Green–based qPCR were performed as previously described (7) using primers listed in Supplementary Table S1.
Western blot analyses and immunoprecipitation

Western blot analyses were performed as previously described (7) using antibodies specific for β-catenin (Sigma; C2206), PARP [Cell Signaling Technology (CST); 9542], cleaved caspase-3 (CST; 9664), cleaved caspase-7 (CST; 9491), SKP2 (CST; 2652), MYC (Santa Cruz; sc-40), p21 (CST; 2947), or tubulin (Santa Cruz; sc-5546). CBP coimmunoprecipitations were performed on nuclear extracts as previously described (30) with anti-CBP (Santa Cruz; sc-369) or control isotype-matched IgG (Abcam; ab46540) antibodies.

Chromatin immunoprecipitation

AsPC-1 cells (70% confluence) were cross-linked with formalin. Chromatin immunoprecipitation (ChiP) was performed using the Pierce Agarose ChIP Kit (Thermo Scientific) on chromatin digested with 2U micrococcal nuclease. Digested chromatin extracts were precleared with 20 μg agarose beads and incubated overnight with beads prebound with either 5 μg IgG control or CBP antibody (Abcam; ab46540 and ab2832, respectively) blocked with 1 μg fish sperm DNA. PCR was performed using primers (Supplementary Table S1) covering gene regulatory regions for CDKN1A (−231 to −117 relative to transcriptional start site, TSS), SKP2 (1772 to 1870 relative to TSS), and CDC25A (−31 to −310 relative to TSS).

Gene expression microarray and data analysis

RNA was extracted from three biologic replicates of AsPC-1 cells 6 and 24 hours after treatment with 10 μmol/L ICG-001 or vehicle control (DMSO) or in separate experiments from two biologic replicates of AsPC1 cells transfected for 48 hours with either 20 nmol/L control or CTNNB1 siRNA. Affymetrix U133 plus 2.0 oligonucleotide arrays were processed in the UCLA Clinical Microarray Core. Arrays are available in Gene Expression Omnibus database repository (accession code GSE57728). Data analysis was performed with dChip Analysis software package (31) using invariant set normalization and signal intensities summarized using the model-based expression index algorithm with mismatch probe option for background subtraction. Initial gene filtering criteria were set at >10% present call and variance across samples of 0.4 < standard deviation/mean < 1,000. ICG-001-regulated genes were determined by comparative analysis of ICG-001 versus vehicle control (selection criteria >1.67-fold change and P < 0.01 at 6 hours or >2-fold change and P < 0.01 at 24 hours). β-catenin–regulated genes were determined by comparative analysis of CTNNB1 versus control siRNA transfection (selection criteria >1.33-fold change and >50 absolute change). Functional enrichment was examined using default parameters in the web-based Database for Annotation, Visualization and Integrated Discovery (DAVID, v6.7; http://david.abcc.ncifcrf.gov; ref. 32). Analysis was limited to the gene ontology (GO) biologic process FAT (GOTERM_BP_FAT) category terms and ranked on P values after removal of categories with less than 6 annotated genes.

Statistical analysis

Student t tests were used to compare continuous variables. The Cox proportional hazard model was used to estimate HRs between treatment groups. The assumption of proportionality was assessed both by visual inspection of Kaplan–Meier survival curves and formal analysis of Schoenfeld residuals (P = 0.13). Kaplan–Meier survival curves and log-rank tests were performed using GraphPad Prism.

Results

ICG-001 inhibits in vitro PDAC cell line growth

ICG-001 (Fig. 1A) affects on in vitro anchorage-dependent PDAC cell growth were addressed by MTT cell viability assays. ICG-001 showed dose-dependent cell growth inhibition of AsPC-1, L3.6pl, PANC-1, and MiaPaCa-2 cell lines at IC50 values of 5.48 μmol/L, 14.07 μmol/L, 3.43 μmol/L, and 3.31 μmol/L, respectively (Fig. 1B and Supplementary Fig. S1A and S1B). ICG-001 also significantly inhibited anchorage-independent growth in soft agar assays (Fig. 1C). Addressing the potential for combined use of ICG-001 with cytotoxic chemotherapy, concurrent treatment with 5 μmol/L ICG-001 and 5 nmol/L gemcitabine further increased growth inhibition relative to individual treatments (Fig. 1D).

ICG-001 inhibits in vivo PDAC tumor growth

ICG-001 was next addressed in an in vivo orthotopic xenograft model of PDAC. AsPC-1 tumor cells were injected into the distal pancreas of nude mice 10 days before the start of treatment. Tumor-bearing mice were then randomized into four different treatment arms, including vehicle (N = 12), ICG-001 (N = 12), gemcitabine (N = 11), or ICG-001 + gemcitabine (N = 12). Treatment was continued for a total of 8 weeks, after which all surviving animals were followed until the end of the study (120 days). ICG-001 significantly improved survival compared with vehicle alone as determined by Cox proportional hazard analysis [HR, 0.207; 95% confidence interval (CI), 0.079–0.543; P = 0.001]. Gemcitabine also significantly improved survival compared with vehicle (HR, 0.123; 95% CI, 0.045–0.334; P < 0.001). There was no survival difference between gemcitabine alone versus ICG-001 alone (HR, 0.592; 95% CI, 0.250–1.399; P = 0.23). In line with in vitro results, the greatest survival benefit relative to vehicle was seen with combined gemcitabine + ICG-001 (HR, 0.074; 95% CI, 0.026–0.212; P < 0.001). Although this combination improved survival relative to ICG-001 alone (HR, 0.360; 95% CI, 0.152–0.848; P = 0.02), there was no significant improvement relative to gemcitabine alone (HR, 0.608; 95% CI, 0.254–1.455; P = 0.66). Kaplan–Meier survival curves (Fig. 2A) also demonstrated improved survival relative to vehicle control (mean survival, 46 ± 3.1 days) for ICG-001 (mean survival, 64 ± 7.1 days, log-rank P = 0.01), gemcitabine (mean survival, 83 ± 5.1 days,
Figure 1. ICG-001 inhibits in vitro anchorage-dependent and -independent growth of pancreatic cancer lines. A, chemical structure of ICG-001. MTT cell viability (B) and soft agar assays (C) were performed at the indicated concentrations of ICG-001. D, MTT assay on AsPC-1 cells treated for 6 days with vehicle (V), 5 μmol/L ICG-001 (I), 5 nmol/L gemcitabine (G), or 5 μmol/L ICG-001 plus 5 nmol/L gemcitabine (I + G). Representative experiment from at least three replicates is presented for each cell line. *, P < 0.05; **, P < 0.01; ***, P < 0.001; and ns, not significant.
log-rank \( P < 0.00001 \), and combined ICG-001 + gemcitabine (mean survival, 95 ± 6.0 days, log-rank \( P < 0.00001 \)) treatments.

The in vivo biologic effects of ICG-001 were further addressed by gross and histologic analysis of mice at necropsy. All tumors significantly involved the pancreas.
with direct extension into adjacent spleen. Metastatic tumor deposits were observed in all animals, most frequently involving the lung, small bowel, and diaphragm (Supplementary Fig. S2A–S2F and Supplementary Table S2). Using a previously described scoring system that awards points based on tumor infiltration and metastasis (29), tumor dissemination was quantified at the time of death. No differences were observed between the four treatment arms (Fig. 2B), indicating that animals were reproducibly sacrificed at time points of equal tumor burden. A mitotic index calculated on tumors harvested during drug treatment revealed a significant decrease in mitotic activity in ICG-001 versus vehicle control–treated tumors (Fig. 2C). Real-time qPCR for three well-established Wnt target genes in tumors harvested during drug treatment revealed a significant decrease in MMP7 expression with ICG-001 versus vehicle control treatment, MMP7 treatment revealed a significant decrease in published Wnt target genes in tumors harvested during drug treatment revealed a significant decrease in mitotic activity in ICG-001 versus vehicle control–treated tumors (Fig. 2C). Real-time qPCR for three well-established Wnt target genes in tumors harvested during drug treatment revealed a significant decrease in MMP7 expression with ICG-001 versus vehicle control treatment, but no differences in BIRC5 or AXIN2 expression (Fig. 2D).

Pharmacologic inhibition of Wnt signaling could have potential deleterious effects on normal somatic stem cell maintenance and tissue homeostasis (26). However, no clinical evidence of ICG-001 cytotoxicity was observed on the basis of body weight measurements (Fig. 2E). Furthermore, H&E evaluation of uninvolved liver and intestine revealed no histologic changes to suggest cytotoxicity in animals sacrificed during treatment with ICG-001 or vehicle control. Small bowel from both treatment groups revealed normal villous:crypt ratios and no changes in apoptotic or mitotic activity within crypts, whereas liver showed no evidence of hepatitis, cholestasis, or hepato-cellular injury (i.e., ballooning change or apoptosis; Supplementary Fig. S2G–S2L).

**ICG-001 induces G1 arrest and has variable effects on Wnt/β-catenin signaling**

We next explored mechanisms underpinning ICG-001 pancreatic growth inhibitory effects. As previously shown in colorectal cancer (16), ICG-001 disrupted the interaction between CBP and β-catenin in AsPC1 cells as measured by coimmunoprecipitation (Fig. 3A). To examine ICG-001 effects on Wnt/β-catenin signaling, PDAC cell lines stably transduced with a β-catenin–activated luciferase reporter (7, 33) were treated with increasing concentrations of ICG-001. ICG-001 exhibited strong, dose-dependent inhibition of reporter activity in AsPC-1 (Fig. 3B), a cell line characterized by higher levels of autocrine Wnt/β-catenin activity (7). By comparison, high concentrations of ICG-001 only weakly inhibited reporter activity in PANC-1 and L3.6pl cell lines carrying stably integrated Wnt/β-catenin luciferase reporter (BAR) were treated with indicated concentrations of ICG-001 and measured in dual luciferase reporter assays at 24 hours. Luciferase is normalized to Renilla and shown relative to vehicle. AXIN2 (C) and BIRC5 (D) transcript levels were determined by qPCR in PDAC cell lines treated with indicated concentrations of ICG-001. All values were normalized to ACTB and shown relative to vehicle. One representative experiment from at least two biologic replicates is shown.

**Figure 3.** ICG-001 variably influences Wnt transcriptional activity across pancreatic cancer lines. A, nuclear extracts from AsPC-1 cells treated with vehicle or 30 μmol/L ICG-001 for 6 hours were immunoprecipitated with anti-CBP or control IgG antibodies followed by Western blot analyses for β-catenin and CBP. B, PDAC cell lines carrying stably integrated Wnt/β-catenin luciferase reporter (BAR) were treated with indicated concentrations of ICG-001 and measured in dual luciferase reporter assays at 24 hours. Luciferase is normalized to Renilla and shown relative to vehicle. AXIN2 (C) and BIRC5 (D) transcript levels were determined by qPCR in PDAC cell lines treated with indicated concentrations of ICG-001. All values were normalized to ACTB and shown relative to vehicle. One representative experiment from at least two biologic replicates is shown.

- Western blot for survivin and tubulin (loading control) after 48 hours vehicle or 10 μmol/L ICG-001. One representative experiment of three replicates is shown. *P < 0.05; **P < 0.01; ***P < 0.001; and ns, not significant.
MiaPaCa-2 cells, and not at all in L3.6pl cells (Fig. 3B). Parallel dose-dependent effects of ICG-001 on the expression of the endogenous Wnt/β-catenin target genes AXIN2 were also observed, with L3.6pl cells demonstrating disparate inhibition of AXIN2 expression at high ICG-001 concentration (Fig. 3C). Of note, ICG-001 significantly inhibited the growth of multiple PDAC cell lines at lower concentrations that had little or no effect on Wnt/β-catenin transcriptional activity (compare Fig. 1B and Fig. 3B and C).

ICG-001 has been shown to induce apoptosis via down-regulation of the apoptosis inhibitor BIRC5 (survivin), a known Wnt/β-catenin target gene (16–19). ICG-001 treatment decreased the expression of BIRC5 transcript and survivin protein levels in all four PDAC cell lines (Fig. 3D and E). Despite this significant inhibition of survivin expression, ICG-001 had no significant effects on AsPC1 apoptosis as determined by cleaved PARP, caspase-3, caspase-7, or Annexin-V–PI staining (Fig. 4A and B). PANC-1 and MiaPaCa-2 cells showed mild increases in cleaved PARP, caspase-3, and caspase-7 with no appreciable differences in apoptosis by Annexin-V–PI staining, whereas L3.6pl showed significantly increased apoptosis as measured by Annexin-V–PI flow staining (Fig. 4A and B). These modest changes indicated apoptosis was not the primary mechanism for the observed inhibition of PDAC growth by ICG-001. Therefore, ICG-001 effects on cell proliferation were analyzed by cell-cycle analysis. ICG-001 treatment robustly induced G1 arrest in all PDAC cell lines (Fig. 4C), a finding consistent with the decreased mitotic activity observed in ICG-001-treated orthotopic tumors (Fig. 2C). Significant G1 arrest was observed in all four cell lines at 10 μmol/L ICG-001, a concentration for which only AsPC-1 showed appreciable inhibition of Wnt transcriptional activity.

If ICG-001 exerts antiproliferative effects by inhibiting β-catenin–mediated transcription, this effect should be phenocopied by knockdown of β-catenin expression. However, siRNA-mediated CTNNB1 gene knockdown failed to induce G1 arrest in AsPC-1 cells, whereas parallel ICG-001 treatment did induce significant G1 arrest (Fig. 4D). Furthermore, although exogenous Wnt3A ligand treatment fully rescued ICG-001 inhibition of Wnt reporter activity in the cell lines, it had no effect on ICG-001–mediated G1 cell-cycle arrest (Supplementary Fig. S3). Altogether, these data indicate that ICG-001 mediates growth arrest in PDAC cell lines through mechanisms discrete from its action as an inhibitor of Wnt signaling.

ICG-001 inhibits the expression of genes involved in cell cycle

To explore mechanisms underlying ICG-001 inhibition of cell cycle in PDAC, gene expression microarrays were performed on AsPC-1 cells after 6 and 24 hours of treatment with ICG-001. In total, 569 transcripts (419 upregulated and 150 downregulated) showed altered expression

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**Figure 4.** ICG-001 robustly induces G1 cell-cycle arrest. A, cells were incubated with vehicle, 10 μmol/L ICG-001, or 500 nmol/L staurosporine (Stauro, positive control for apoptosis) for 24 hours and analyzed by Western blot for PARP, cleaved-caspase-3, cleaved-caspase-7, and tubulin (loading control). Vehicle or 10 μmol/L ICG-001–treated cells were analyzed by flow cytometry for (B) apoptosis by Annexin-V–PI staining at 48 hours or (C) cell cycle by PI staining at 24 hours. D, cell-cycle analysis was determined for AsPC-1 cells treated simultaneously with indicated combinations of vehicle, 10 μmol/L ICG-001, control siRNA (20 nmol/L), or CTNNB1 siRNA (20 nmol/L) for 48 hours. Western blot confirmed >75% knockdown of β-catenin with CTNNB1 siRNA. One representative experiment of three replicates is shown for all figures. *, P < 0.05; **, P < 0.01; and ns, not significant.
in response to ICG-001 treatment (P < 0.01; 1.67-fold change at 6 hours or 2-fold change at 24 hours; 0.2% median false discovery rate; Fig. 5A; and Supplementary Table S3). Several of these changes were further validated by qPCR in separate ICG-001 treatment experiments (Fig. 5B). To explore biologic processes linked to ICG-001 treatment, we performed enrichment analysis of GO functional annotations. In line with phenotypic observations, the top five GO terms for genes downregulated by ICG-001 were DNA replication, DNA metabolic processes, DNA replication initiation, DNA-dependent DNA replication, and cell cycle (P < 0.00013; Supplementary Fig. S4A). Top GO terms for genes upregulated by ICG-001 included multicellular organismal homeostasis, negative regulation of cell proliferation, cell-cycle arrest, regulation of angiogenesis, and regulation of cell proliferation (P < 0.001; Supplementary Fig. S4B).

Although its inhibition of PDAC cell proliferation did not seem mechanistically linked to its effects on Wnt/β-catenin transcriptional activity, ICG-001 did inhibit several known Wnt transcriptional targets, including MMP7 and LGR5 that were further validated by qPCR (Fig. 5B). To address ICG-001–regulated genes representing actual Wnt transcriptional targets in PDAC, we compared ICG-001 results with a second microarray study comparing gene expression changes in AsPC-1 transfected with either control or CTNNB1 siRNA. In the context of greater than 80% CTNNB1 knockdown, there were 441 altered transcripts (absolute fold change ≥ 1.33; median false discovery rate 10.4%), including 228 upregulated and 213 downregulated genes (Supplementary Table S3). Top GO terms for genes downregulated by CTNNB1 knockdown included cell fate commitment, tubular morphogenesis, tube development, morphogenesis of a branching structure, and angiogenesis (P < 0.0005; Supplementary Fig. S5A). Top GO terms for genes upregulated by CTNNB1 siRNA included response to wounding, cell adhesion, biologic adhesion, response to inorganic substance, and regulation of cell migration (P < 0.0002; Supplementary Fig. S5B). Cross-referencing both ICG-001 and CTNNB1 siRNA microarray results revealed 117 overlapping transcripts (Fig. 5C; Supplementary Table S3). GO functional annotations were only marginally enriched in this overlapping dataset (P < 0.01), but included regulation of cell proliferation, positive regulation of developmental processes, homeostatic process, response to wounding, and regulation of the cell cycle.

Among the most highly regulated genes identified in the ICG-001 arrays were SKP2, CDC25A, and CDKN1A (p21), each regulators of cell-cycle progression. We

Figure 5. ICG-001 alters the expression of cell-cycle genes. A, gene expression was determined with Affymetrix Human U133 plus 2.0 microarrays using RNA collected from AsPC1 following 6 or 24 hour treatment with vehicle or 10 μM ICG-001. Biologic replicates of significantly regulated genes are shown in the heatmap along with a partial list of genes curated based on known roles in tumorigenesis, cell-cycle control, cancer diagnosis/prognosis, or Wnt/β-catenin transcriptional targets. B, qPCR validation was performed on selected genes in AsPC-1 cells separately treated with vehicle or 10 μM ICG-001 for 24 hours. C, Venn diagram comparing AsPC-1 gene expression microarray changes observed with either 10 μM ICG-001 treatment or CTNNB1 siRNA transfection.
therefore explored whether these genes were direct transcriptional targets of CBP that are altered in response to ICG-001 treatment. After 1-hour ICG-001 treatment, ChIP revealed reduced CBP occupancy at SKP2 and increased CBP occupancy at CDKN1A (Fig. 6A), corresponding, respectively, to aforementioned decreased SKP2 and increased CDKN1A gene expression results (Fig. 5). No change in CBP occupancy was observed at CDC25A (Fig. 6A), suggesting that it is transcriptionally downregulated indirectly in response to ICG-001. Further detailed time-course analyses of gene and protein expression revealed SKP2 transcript levels were reduced within 4 hours, whereas CDKN1A transcript levels were increased within 2 hours of ICG-001 treatment (Fig. 6B and C). Corresponding decrease in SKP2 protein expression and increase in p21 protein expression were observed within 4 to 6 hours of ICG-001 treatment (Fig. 6D). MYC protein levels were also decreased after 6 hours, although MYC transcript levels were unchanged (Fig. 6D and E), indicating that a posttranscriptional mechanism controls MYC protein levels following ICG-001 treatment.

Discussion

At least subsets of PDAC are dependent upon Wnt/β-catenin signaling for initiation and/or progression (6-10, 34), warranting exploration of ICG-001 and other Wnt inhibitors as potential therapeutics. PRI-724, an ICG-001 derivative, is presently under investigation in combination with gemcitabine in a phase I clinical trial for patients with metastatic pancreatic cancer (NCT01764477). We have demonstrated that ICG-001 significantly inhibits in vitro PDAC growth through induction of G1 cell-cycle arrest and improves survival in an in vivo orthotopic xenograft model of PDAC. Despite augmented PDAC growth inhibition in vitro, combined ICG-001 and gemcitabine did not further improve survival over gemcitabine alone in the xenograft model. It is possible that robust induction of G1 arrest with ICG-001 compromises the action of cytotoxic chemotherapy such as gemcitabine, which primarily kills cancer cells undergoing DNA synthesis in S phase. An analogous situation has been observed with CDK4 inhibitors that by inducing G1 arrest can reduce the efficacy of cytotoxic chemotherapy in certain circumstances (35). Further optimization of the dosing and delivery of ICG-001 (or PRI-724) is needed to resolve whether ICG-001 can be used effectively in combination with gemcitabine. Intriguingly, ICG-001 reduced expression of RRMI in AsPC-1 microarrays. RRMI expression directly correlates with gemcitabine resistance (36) and worse overall survival in PDAC (37, 38). Furthermore, ICG-001 was shown to act synergistically with
conventional chemotherapy to prolong survival in NOD/SCID mice engrafted with primary acute lymphoblastic leukemia (22). Thus, optimized combinations of cytotoxic regimens with ICG-001 may still offer additional survival benefit in PDAC or other tumors.

ICG-001 treatment and β-catenin knockdown regulated an overlapping set of genes in AsPC-1, including known targets of canonical Wnt signaling (i.e., MMP7, EDN1, LGR5, RN43, and PORCN). Originally identified in a small-molecule library screen of compounds that inhibited β-catenin/T-cell factor (TCF)–dependent reporter activity, ICG-001 inhibits Wnt transcriptional activity by binding to the N-terminal domain of CBP and preventing its interaction with β-catenin (16). This may only have the potential to disrupt CBP-dependent β-catenin–TCF transcription, but also may increase the pool of β-catenin available to interact with p300. Like CBP, p300 functions either as a transcriptional coactivator or corepressor in a context-dependent manner (26, 39, 40). Kahn and colleagues have proposed a model of CBP/p300 differential coactivator usage whereby TCF–β-catenin–CBP-mediated transcription drives a nondifferentiated proliferative state with ICG-001 inducing a switch to TCF–β-catenin–p300-mediated transcription favoring cellular differentiation (39, 41). In keeping with this model, ICG-001 strongly induced G1 cell-cycle arrest in PDAC cell lines and inhibited the expression of certain genes linked to pluripotency, including LGR5 that marks progenitor cells in intestinal crypts and DCLK1 which was recently shown to mark a unique tumor initiating cell population in PDAC (42).

ICG-001 also regulated many genes unaltered by β-catenin knockdown, including several linked to DNA replication and cell-cycle regulation. Beyond β-catenin, ICG-001 could more generally alter gene expression by disrupting CBP interaction with any of a number of other known protein binding partners that includes many transcription factors (43). As an example, ICG-001 was recently shown to block CBP binding to γ-catenin, resulting in increased p300/γ-catenin interaction and decreased BIRC5 expression (22, 44).

ICG-001 can induce caspase-3/7–mediated apoptosis in colon cancer cells via downregulation of BIRC5 (16). Although ICG-001 treatment inhibited survivin protein levels, there were only variable and modest effects on apoptosis in PDAC cell lines. Rather, ICG-001 induced robust G1 cell-cycle arrest, which is perhaps not unexpected given CBP/p300 histone acetyltransferase activity is essential for cell proliferation and orderly G1–S transition (45). Several groups found that Wnt/β-catenin signaling promotes cellular proliferation in PDAC (4, 9) and while Wnt/β-catenin transcriptional targets MYC or CCND1 are commonly linked to its regulation of the cell cycle, transcript levels of these genes were not altered in PDAC cell lines in response to ICG-001 treatment. Moreover, growth inhibition and G1 cell-cycle arrest were observed in multiple PDAC cell lines at ICG-001 concentrations with little or no measurable effect on Wnt reporter activity or endogenous AXIN2 expression, indicating ICG-001 inhibition of PDAC cell proliferation was decoupled from its function as a Wnt inhibitor.

ICG-001 treatment altered the expression of numerous genes linked to DNA replication and cell cycle in AsPC-1 microarray analysis, both downregulating genes that promote (i.e., SKP2, CDC25A, CDC7, CCNE, and MCM2-MCM 6) and upregulating genes that inhibit (i.e., CDKN1A, CDKN1C, CDKN2B, MXD1, MXII, and BTG1) cell-cycle progression. ICG-001 induced rapid changes in CBP occupancy on SKP2 and CDKN1A regulatory regions with corresponding changes in mRNA and protein levels, implicating SKP2 and CDKN1A as direct targets of ICG-001–mediated cell-cycle arrest. SKP2 is a substrate component of the SCF E3 ubiquitin-protein ligase complex that promotes cell-cycle progression through ubiquitination and subsequent degradation of several proteins linked to cell-cycle inhibition, including p21, p27, p57, p130, and RASSF1A among others (46). ICG-001 increased CBP occupancy at CDKN1A promoter site in AsPC-1 cells, perhaps by increasing the availability of CBP to different subsets of transcription factors mediating CDKN1A expression. CDKN1A encodes the cyclin-dependent kinase inhibitor p21 that opposes the function of multiple cyclin–CDK complexes that mediate G1 and S cell-cycle progression. Increased p21 levels with ICG-001 could reflect decreased SKP2-mediated degradation of p21 and/or increased CDKN1A expression via CBP cotranscriptional activation. Although MYC transcript levels were unchanged, ICG-001 reduced MYC protein levels within 6 hours along with corresponding increases in several known targets of MYC-mediated transcriptional repression (i.e., CDKN1A, CDKN1C, and CDKN2B). It is intriguing to speculate how ICG-001 might indirectly antagonize MYC through additional posttranslational mechanisms linked to its protein stability or function. MXD1 and MXII, which were both upregulated in response to ICG-001 treatment, can inhibit the transcriptional activity of MYC by competing for its important heterodimerization partner MAX (47). CBP/p300-mediated acetylation has been shown to stabilize MYC protein and function (48), which could also be perturbed by ICG-001. Finally, ICG-001 could indirectly disrupt MYC activity through its aforementioned downregulation of SKP2, which has been shown to ubiquitylate MYC to license its function as a transcriptional activator (49).

In conclusion, ICG-001 seems to robustly inhibit PDAC growth through both direct and indirect mechanisms that mutually reinforce the induction of G1 cell-cycle arrest. Although further studies are needed to expand upon precise mechanisms underpinning its transcriptional and phenotypic activities, ICG-001 seems to more broadly disrupt CBP function beyond its effects as cotranscriptional activator of Wnt/β-catenin signaling. These future studies should not only further examine the therapeutic potential of ICG-001 and its derivatives in PDAC or other cancers, but also better establish effective drug combinations and patient selection.
Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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Conception and design: M.D. Arensman, T.R. Donahue, D.W. Dawson
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