Chronic GLP-1 Receptor Activation by Exendin-4 Induces Expansion of Pancreatic Duct Glands in Rats and Accelerates Formation of Dysplastic Lesions and Chronic Pancreatitis in the Kras$^{G12D}$ Mouse Model

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Pancreatic duct glands (PDGs) have been hypothesized to give rise to pancreatic intraepithelial neoplasia (PanIN). Treatment with the glucagon-like peptide (GLP)-1 analog, exendin-4, for 12 weeks induced the expansion of PDGs with mucinous metaplasia and columnar cell atypia resembling low-grade PanIN in rats. In the pancreata of Pdx1-Cre; LSL-Kras$^{G12D}$ mice, exendin-4 led to acceleration of the disruption of exocrine architecture and chronic pancreatitis with mucinous metaplasia and increased formation of murine PanIN lesions. PDGs and PanIN lesions in rodent and human pancreata express the GLP-1 receptor. Exendin-4 induced proproliferative signaling pathways in human pancreatic duct cells, cAMP–protein kinase A and mitogen-activated protein kinase phosphorylation of cAMP-responsive element-binding protein, and increased cyclin D1 expression. These GLP-1 effects were more pronounced in the presence of an activating mutation of Kras and were inhibited by metformin. These data reveal that GLP-1 mimetic therapy may induce focal proliferation in the exocrine pancreas, and, in the context of exocrine dysplasia, may accelerate formation of neoplastic PanIN lesions and exacerbate chronic pancreatitis. Diabetes 61:1250–1262, 2012

GLP-1 receptor (GLP-1R) is a G-protein–coupled receptor that is expressed in pancreatic islets and exocrine duct cells (2,3). The increased GLP-1 released after meal ingestion amplifies postprandial nutrient-driven insulin secretion, the so-called incretin effect (4). Based on this property, GLP-1R activation became an attractive therapeutic target for type 2 diabetes mellitus (T2DM). To overcome the short half-life of circulating GLP-1 that is rapidly degraded by dipeptidyl peptidase (DPP)-4 (5), two small molecule inhibitors, such as sitagliptin, prolong the half-life of endogenously secreted GLP-1 (6). Alternatively, GLP-1R peptide agonists given by injection, such as exenatide (7) and liraglutide (8), are resistant to DPP-4 degradation. Pancreatitis emerged as an unexpected side effect of GLP-1–based therapy in case reports (9,10), and in the U.S. Food and Drug Administration adverse-event reports, liraglutide and sitagliptin showed a signal of pancreatitis (11–13), although analysis of insurance claims records have been reported to show no association between GLP-1–based therapy and pancreatitis (14). Because the human pancreas is inaccessible in treated patients, the question as to whether GLP-1 mimetic therapy acts on the exocrine pancreas has been a subject of animal-based studies. Pancreatic duct cell proliferation increased transiently with a GLP-1 infusion in Wistar rats (15). Sprague-Dawley rats treated with exendin-4 for 12 weeks developed low-grade chronic pancreatitis (16). Furthermore, DPP-4 inhibition with sitagliptin for 12 weeks was associated with increased pancreatic duct cell replication and acinar-to-ductal metaplasia and, in 1 of 10 rats, chronic pancreatitis (3). However, GLP-1–based therapy also has been reported to not exacerbate chemically induced pancreatitis in mice (17). Also, exenatide was reported to have no effect on ductal turnover in mice or rats, as well as to have a beneficial action in chemically induced pancreatitis (18).

Pancreatic duct glands (PDGs), under conditions of chronic injury, such as chemically induced pancreatitis, may give rise to lesions resembling pancreatic intraepithelial neoplasia (PanIN) (19). To date, there is no information on the actions of GLP-1–based therapy on PDGs or the development of PanIN in pancreata predisposed to dysplasia. Here, we sought to address the following questions. First, does chronic activation of GLP-1Rs by exendin-4 lead to proliferation of the PDGs? Second, is GLP-1R expression present in PDGs and PanIN-like dysplastic lesions? Third, does chronic activation of GLP-1Rs alter the phenotype of Pdx1-Cre; LSL-Kras$^{G12D}$ (Pdx1-Kras) mice?

RESEARCH DESIGN AND METHODS

Rodent studies. All animal studies were approved by the animal use and care committee at the University of California Los Angeles (UCLA). Animals were housed individually in a 12-h light/dark cycle and were weighed weekly to adjust drug doses. Blood glucose and food intake were monitored on a biweekly basis. Sprague-Dawley rats treated with exendin-4. To establish the actions of GLP-1R activation in the exocrine pancreas, we treated 10 male Sprague-Dawley rats (Charles River Laboratories, Wilmington, MA) with daily injections of 10 μg/kg body wt exendin-4 (ChemPept, Miami, FL) administered by subcutaneous injection for 12 weeks starting at 10 weeks of age (20). Animals were fed chow (Teklad; Harlan Laboratories, Madison, WI) ad libitum. A total...
of 15 control rats received daily saline injections. We did not identify PDGs in 5 of 15 controls; therefore, these 5 rats were not included in subsequent analyses. PDGs were identified in all treated rats.

**Pdx1-Kras mice treated with exendin-4.** To investigate the effect of chronic GLP-1 mimetic treatment on pancreatic cancer precursor lesions, the conditional KrasG12D from Hingorani et al. (21) was used. Experimental animals were generated by crossing Pdx1-Cre with LSL-KrasG12D mice on a C57/BL6 background (both gifts of Guido Eibl, UCLA). Mice (6 weeks old) were fed an AIN-76A-based diet (Research Diets, New Brunswick, NJ) ad libitum for 12 weeks, during which either saline (n = 7) or exendin-4 (5 mmol/kg body weight) (as counted in 10 nonoverlapping HPFs) and graded as 0, absent; 1, 1–5%; 2, 6–10%; 3, 11–20%; and 4, >20% fibrosis. Duct profiles were evaluated according to established consensus guidelines for the histologic evaluation of mPanIN (24) and quantified as previously described (25). All duct profiles in one full pancreas cross-section were evaluated to determine the relative proportions of nondysplastic (normal, reactive, and metaplastic) ducts and each category of mPanIN lesion. The proportion of each mPanIN lesion to the overall number of duct profiles was recorded for each animal.

**Duct cell replication frequency.** To determine the frequency of replication of PDG cells and cells in the adjacent main ducts in the head of the pancreas in rats, we quantified the number of Ki-67–positive cells. Thus, the total number of cells in the head of the pancreas evaluated was 57,261 in exendin-4–treated rats and 61,298 in controls. We also evaluated the frequency of duct cell replication in the sections of the tail of the pancreas immunostained for cytokeratin and Ki-67. The total number of duct cells evaluated from the tail was 24,483 in exendin-4–treated rats and 19,706 in controls.

**Duct cell replication in pancreas from Pdx1-Kras mice also was evaluated by Ki-67.** The extensive acinar-to-duodenal metaplasia and frequent dysplastic ductal lesions in GLP-1–treated Pdx1-Kras mice precluded distinguishing replication frequency in the various component of the ductal compartment (PDGs and normal and dysplastic ducts). Slides were analyzed using the Ariol SL-50 automated slide scanner (Leica Microsystems) to quantitate the amount of positive staining for each category of interest containing only ducts and dysplastic ductal tissue. A total number of 121,693 (control group) and 101,830 (exendin-4–treated group) cells were analyzed.

**GLP-1 actions on pancreatic duct cells.** In vitro experiments were carried out to investigate the effects of exendin-4 on human pancreatic duct epithelial (HPDE) cells (26,27). HPDE cells (kindly made available by Dr. Ming-Sound Tsao, University of Toronto) were maintained in keratinocyte serum-free media supplemented with bovine pituitary extract and human epidermal growth factor (In Vitrogen) at 37°C with 5% CO2. HPDE cells transfected with the empty vector (pBabeRho) (HPDE-pBP) or with oncogenic pBP-KrasG12D (HPDE-Kras) also were used to perform the assessment of GLP-1R activation in the presence of an activating Kras mutation.

To assess the effect of exendin-4 (10 mmol/L) on the phosphorylation of cAMP-responsive element-binding (CREB) protein and the mitogen-activated protein kinases (MAPKs) extracellular signal–related kinase (ERK) 1/2, we studied the phosphorylation of CREB and ERK1/2 with endogenous, oncogenic, and exogenous forms of KrasG12D, and control Pdx1-Kras mice treated with exendin-4.
FIG. 1. The extent and frequency of PDGs surrounding the main pancreatic duct are increased by exendin-4 treatment in rats. Sections from the head of the pancreas from an untreated control rat (A) and after 12 weeks of daily exendin-4 injections (E), in which PDG clusters were identified surrounding the main pancreatic duct. PDGs were confined to the mesenchyme surrounding the main duct in controls but, after exendin-4, expanded to the extent that they projected into the lumen of the pancreatic duct as complex villous-like structures. A and E, insets: PDG cells were columnar in comparison with the cuboidal ductal cells and included goblet-like cells (arrowheads). B and C: PDGs contained mucin confirmed by Alcian blue and PAS staining. D: In contrast to duct cells, PDG cells also expressed Pdx-1 (red; combined staining with the duct cell marker cytokeratin [CK] in green). E: PDGs were more common in exendin-4-treated rats (Table 1). F–H: In addition, the epithelium often showed pseudostratification and pseudopapillary features, which are features characteristic for PanIN-like lesions. Scale bars = 200 μm (A and E) and 100 μm (B–D), and magnification ×20 (F–H). (A high-quality digital representation of this figure is available in the online issue.)
RESULTS

Metabolic actions of exendin-4 in rats. Twelve weeks of daily exendin-4 injections had the anticipated effects of decreasing weight gain (66 ± 8 vs. 164 ± 5 g; \( P < 0.001 \) exendin-4 vs. control) and blood glucose levels (99 ± 2 vs. 108 ± 4 mg/dL; \( P < 0.01 \) exendin-4 vs. control). As expected, exendin-4 decreased daily food intake (153 ± 5 vs. 204 ± 5 mg/day; \( P < 0.001 \) exendin-4 vs. control), but the treated animals did not seem to be in any apparent pain or distress (Supplementary Fig. 1).

Effects of exendin-4 on exocrine pancreas in rats. Pancreas weight was comparable in the treated versus control group (2.3 ± 0.1 vs. 2.3 ± 0.1 g; exendin-4 vs. control). However, relative to body weight, pancreatic weight in exendin-4-treated animals was increased (0.53 ± 0.02 vs. 0.43 ± 0.02; \( P < 0.01 \) exendin-4 vs. control) (Supplementary Fig. 1D).

There was no histological evidence of pancreatitis in either the exendin-4 or control group. Consistent with this, lipase activity was not changed by exendin-4 (330 ± 19 vs. 299 ± 11 units/L; exendin-4 vs. control) (Supplementary Fig. 1E). However, exendin-4 did induce a marked expansion of the PDG compartment (Fig. 1 and Supplementary Fig. 2). PDGs were identified, as previously described, as blind outpouchings from large pancreatic ducts present in the mesenchyme surrounding the ducts. PDG cells were further distinguished from main duct cells by frequently being columnar rather than cuboidal (Fig. 1A and E, insets) and mucin positive (Alcian blue and PAS stains). PDGs also expressed Pdx-1 (Figs. 1 and 3) and 1R expression was readily detected in pancreatic tissue (data not shown).

To evaluate the extent of the PDG compartment in treated and control animals, we analyzed the PDG compartment in sections from the head of the pancreas from 10 animals in each group (illustrated in Supplementary Fig. 2). The number of PDGs per millimeter of main duct (first row) and the average size of a PDG (second row) revealed a marked expansion of the PDG compartment after exendin-4 treatment. Furthermore, the main duct appears to be dilated because the ratio of main duct lining to length was increased in the treated group (third row), \( *P < 0.05 \), \( \dagger P < 0.001 \).

Actions of GLP-1 mimetic treatment on the exocrine pancreas in the Pdx1-Kras mutant mouse. In Pdx1-Kras mice, 12 weeks of exendin-4 treatment had no impact on body weight (23.2 ± 1.2 vs. 25.8 ± 1.7 g), food intake (18.1 ± 0.7 vs. 19.7 ± 0.5 g per week), or blood glucose levels (83.0 ± 3.4 vs. 75.4 ± 3.7 mg/dL) when compared with littermate control mice. However, GLP-1 mimetic treatment increased pancreatic weight (1.1 ± 0.1 vs. 0.7 ± 0.1 g exendin-4 vs. control) (Supplementary Fig. 3).

While overall lobular architecture was preserved in both animal groups, the exendin-4–treated animals demonstrated more extensive chronic pancreatitis with greater loss of acini with replacement by reactive or metaplastic duct profiles (Fig. 3). The percentage of pancreas composed of acinar tissue was decreased by 61% by exendin-4 treatment (13.0 ± 13.5% vs. 33.6 ± 14.6%; \( P < 0.05 \) exendin-4 vs. control). These changes were accompanied by increased inflammation, more extensive stromal fibrosis, and widespread reactive and metaplastic changes, as determined by pancreatitis score (10.0 ± 1.2 vs. 8.6 ± 0.8; \( P < 0.05 \) exendin-4 vs. control). The plasma lipase activity also was increased with exendin-4 (1,020 ± 164 vs. 678 ± 34 units/L; \( P < 0.05 \) exendin-4 vs. control) (Supplementary Fig. 3). In comparison to control animals, treated animals showed more extensive acinar-to-ductal metaplasia with replacement of acini by ductules lined by mucin-producing cells primarily with small, round basally oriented nuclei without papillary features (mPanIN1). A minority of the duct profiles demonstrated increased nuclear hyperchromasia and pleomorphism with stratification and micropapillary changes (mPanIN2 and mPanIN3) (Fig. 3). Moreover, GLP-1 mimetic treatment induced increased duct cell proliferation (\( P < 0.05 \) in Pdx1-Kras mice when compared with control animals (Fig. 4)).

GLP-1R expression in PDGs and PanIN lesions. GLP-1R expression was readily detected in pancreatic β-cells in rat and human pancreas, serving as a positive control (data not shown). GLP-1R expression also was present in PDG cells in both rodent and in human pancreas (Fig. 5). GLP-1R expression was not detected in pancreatic acinar cells. GLP-1R expression also was abundantly present in mPanIN lesions.
FIG. 2. PDG cell replication is increased by exendin-4 treatment in rats. The frequency of replication ascertained by Ki-67 immunostaining (red; colabeled with cytokeratin [CK] in green) was increased in PDGs compared with adjacent duct cells (*lumen of the large duct) in both control (A) and exendin-4–treated (B) rats. Replication frequency showed variation within the PDGs in control (C) as well as exendin-4–treated (D) animals. E: However, both the abundance of PDGs and the frequency of replication were increased by exendin-4 treatment. Exendin-4 also increased replication in main duct cells but not in the duct cells in the tail of the pancreas. □, control (Ctrl); ■, exendin-4 (Ex). *P < 0.05, scale bars = 100 μm. (A high-quality digital representation of this figure is available in the online issue.)
FIG. 3. Exendin-4 treatment increased chronic pancreatitis and the frequency of mPanIN lesions in Pdx1-Kras mice. Pancreata from Pdx1-Kras mice treated for 12 weeks with either vehicle (A) or exendin-4 (B) (20× objective). The pancreas from the exendin-4–treated animal demonstrates only scant residual intact acini (white arrow) with more extensive inflammation and fibrosis (stars) and more frequent mPanIN (black arrows). C and D: Low-grade mPanIN1a and mPanIN1b lesions with abundant apical mucin and basally oriented nuclei without significant nuclear pleomorphism or mitotic activity. E and F: Higher-grade mPanIN2 and mPanIN3 lesions with increased nuclear pleomorphism and focal loss of polarity. G: Quantitative analysis of mPanINs showing the percentage of pancreatic ducts with no dysplasia (□, normal [nl]); light-grey box, mPanIN1 (1); medium-grey box, mPanIN2 (2); or ■, mPanIN3 (3) lesions in control (Ctrl) and exendin-4 (Ex)-treated mice. H: Combined amylase (red) and cytokeratin (CK; green) immunofluorescent staining of the pancreas of a control Pdx1-Kras mouse. I: Intact acinar tissue (red) is replaced by cytokeratin-positive (green) ducts, and amylase-positive cells are rarely found in exendin-4–treated animals. Alcian blue staining (blue; counterstained with Nuclear Fast red) reveals mucin-containing lesions in control mice (J) and a higher frequency in treated mice (K). *P < 0.05; **P < 0.01 vs. control. (A high-quality digital representation of this figure is available in the online issue.)
FIG. 4. Duct cell replication frequency is increased in the pancreas of exendin-4–treated Pdx1-Kras mice. Immunohistochemical labeling of Ki-67–positive cells (brown; counterstained with hematoxylin) in benign ducts in areas of intact acinar tissue in control mice (A) and exendin-4–treated mice (B). An area of ductal proliferation embedded in fibrotic tissue shows an increase in Ki-67–positive cells in the exendin-4–treated group (D) compared with controls (C). Note the presence of proliferative ducts and mPanIN1a lesion in the exendin-4–treated animal. E: Analysis of duct cell proliferation by Ki-67 reveals an increase in the replication frequency in Pdx1-Kras mice treated with exendin-4 (Ex; ■) compared with vehicle control (Ctrl; □). *P < 0.05. (A high-quality digital representation of this figure is available in the online issue.)
in the pancreas of Pdx1-Kras mice and humans (Fig. 6). GLP-1R was also detected in areas of acinar-to-ductal metaplasia as well as mPanIN lesions in Pdx1-Kras mice (Fig. 6A and B). In humans, cells with a columnar phenotype had prominent GLP-1R expression. For example, immunoreactivity was present in PanIN1 lesions but only was minimally detected in adjacent cells with normal cuboidal pancreatic duct morphology in the same duct (Fig. 6C). In 6 of 10 human pancreata, GLP-1R expression was detected in a variety of ductal lesions (PanIN1a to PanIN3) (Fig. 6D and E).

**Actions of exendin-4 treatment in human pancreatic duct cells.** GLP-1 activation of G-protein–coupled receptors has been reported to activate multiple signaling pathways in pancreatic β-cells, such as the cAMP–protein kinase A and the MAPK pathways leading to phosphorylation of CREB with increased cyclin levels and β-cell replication in pancreatic β-cells (30–32).

To investigate the mechanism of GLP-1–induced duct cell proliferation, we treated HPDE cells with exendin-4 (Fig. 7). CREB phosphorylation increased after 10 min of exendin-4 exposure, reaching a plateau at ~30 min (1.8 ± 0.2-fold vs. control; \( P < 0.05; n = 3 \)) (Fig. 7A). Exendin-4 induced a time-dependent phosphorylation of the mitogen-activated kinases ERK1 (4.8 ± 0.6-fold) and ERK2 (2.7 ± 0.1-fold, respectively, vs. control at 10 min; \( P < 0.01; n = 3 \)) (Fig. 7B). Consequently, cyclin D1 protein was induced to a maximum at ~6 h (1.5 ± 0.2-fold vs. control; \( P < 0.05; n = 3 \)) (Fig. 7C). However, no changes were observed in cyclin

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**FIG. 5.** GLP-1R expression is present in PDGs in rats and humans. A: In the PDGs (shown here for an exendin-4–treated rat), GLP-1R expression (red) was detected by immunofluorescence with combined labeling for the duct cell marker cytokeratin (CK) in green and DAPI to mark the nuclei in blue. B: Colocalization of GLP-1R and cytokeratin is indicated in the merged images by the color orange. GLP-1R expression was similarly apparent in PDGs in duct cells in the human pancreas. Scale bars = 100 μm. (A high-quality digital representation of this figure is available in the online issue.)
FIG. 6. GLP-1R expression is present in PanIN lesions in Pdx1-Kras mice and humans. GLP-1R (red; shown with combined cytokeratin [CK] labeling in green) was detected in areas of acinar-to-ductal metaplasia (ADM) (A) and mPanIN lesion (B) in the pancreas of Pdx1-Kras mice. Colocalization of GLP-1R and cytokeratin is indicated in the merged images by the color orange. C: In human pancreas, GLP-1R expression was more apparent in the columnar cells (arrowheads) in regular ducts compared with adjacent normal cuboidal duct cells shown away from the arrowhead. D: Where duct cells adopt the columnar phenotype (PanIN1a lesion shown), GLP-1R expression becomes more apparent. E: In more advanced PanIN3 lesions, GLP-1R immunoreactivity also was clearly present. Scale bars = 100 µm. (A high-quality digital representation of this figure is available in the online issue.)
A levels (Fig. 7D). We also investigated the actions of exendin-4 with or without metformin in the presence of the activating Kras mutation in HPDE cells. Exendin-4 induced CREB phosphorylation in control (pBP) cells (1.4 ± 0.1-fold pCREB/CREB vs. control; *P < 0.01; n = 3), an effect that was more pronounced in the presence of mutant Kras (1.7 ± 0.1-fold vs. control; **P < 0.001; n = 3), and this effect was abrogated by metformin pretreatment (1.0 ± 0.1-fold vs. control; *P < 0.01 vs. exendin-4 treatment alone) (Fig. 8).

DISCUSSION
The possibility that GLP-1 mimetic therapy might induce sustained proliferative changes in the exocrine pancreas is of concern because therapy for T2DM may be administered for decades (33,34). An increased reported adverse event rate in the U.S. Food and Drug Administration adverse-event reporting system for pancreatitis and pancreatic cancer in patients treated with GLP-1–based therapy underscores this concern (35). Because T2DM with obesity is a risk factor for pancreatitis and pancreatic
cancer (36, 37), administration of a drug that may further amplify those risks requires closer investigation. In contrast, also unexpectedly, the diabetes medication metformin may decrease the risk of pancreatitis and pancreatic cancer (38, 39). Given the recent appreciation that PDGs can give rise to PanIN-like lesions in the context of chronic pancreatitis (19), we first sought to establish the effects of GLP-1R activation on this compartment.

Exendin-4 treatment for 12 weeks induced a marked expansion of the PDG compartment in nondiabetic lean Sprague-Dawley rats. If the pancreas had been sectioned exclusively through the body or tail, no striking abnormalities would have been observed, including no increase in the frequency of replication of duct cells. The normal histology in the most accessible portion of the pancreas and the absence of tumors or overt pancreatitis in lean nondiabetic animals treated with exendin-4 may explain normal exocrine pancreas toxicology screens (40) and some animal studies (17, 18). Therefore, to observe the GLP-1–induced changes in PDGs reported in rats here, methodical analysis of the entire pancreas, to include longitudinal sections through the main pancreatic duct, is necessary.

Because PDGs have properties of an adult stem cell compartment (19), it is not surprising that short-term activation of the PDGs by GLP-1 therapy coincident with induced pancreatic injury facilitates recovery from that injury, presumably by fostering regeneration and providing increased protective mucin secretion (17, 18). The clinically more relevant question concerns the implication of longer-term stimulation of the PDG compartment and its derivatives.

A total of 12 weeks of exendin-4 therapy in young healthy rats generated mucinous metaplasia and cysticologic atypia resembling low-grade PanIN-like lesions in the PDG compartment, features reminiscent of the response to induced chronic pancreatitis in mice and spontaneous chronic pancreatitis in humans (19) (Fig. 1 and Supplementary Fig. 2). However, we also report that GLP-1R expression is present in PDGs and PanIN lesions in rodents and humans, raising the question, does GLP-1 mimetic therapy stimulate the growth of PanIN lesions? Low-grade PanIN lesions are present in 16–80% of normal adult pancreata, the frequency increasing with age (41). PanIN lesions in humans are considered neoplasms and potential precursors for invasive pancreatic cancer based on both pathological findings in humans and longitudinal studies in mice in which mutant Kras is introduced into the pancreas (42). The activating point mutation in the KRAS gene is the most frequent mutation present in human PanIN lesions and is considered to be the first step in the progression toward pancreatic cancer (42).

To better appreciate the actions of GLP-1–based therapy in a progression model of PanIN to pancreatic cancer, we treated Pdx1-Kras mice for 12 weeks with exendin-4. Exendin-4 treatment increased duct cell replication, increased the formation of dysplastic mPanIN lesions, and accelerated the development of chronic pancreatitis. These data are consistent with the hypothesis that PanIN lesions contribute to the development of pancreatitis by the obstruction of ductal outflow, with the resulting chronic pancreatitis fostering further development of PanINs (42). The dose of exendin-4 used here, although comparable with that used previously to show the benefit in rodents, exceeds the dose (per kilogram) used in humans (20). A lower dose was used in a recent study to evaluate the effects of exendin-4 on the rodent exocrine pancreas in which no adverse actions were reported (18). However, no data were provided in that report as to whether the dosage of exendin-4 achieved the clinically desired metabolic actions of exendin-4. Moreover, the PDG compartment apparently was not evaluated in those studies, and the animals were not predisposed to dysplasia. It is unknown to date whether a dose of GLP-1 mimetic therapy might be identified that has the intended beneficial actions of enhanced glucose-mediated insulin secretion but no proproliferative effects on the exocrine pancreas.
Evaluation of the proliferative actions of GLP-1 in the exocrine pancreas in humans is not technically feasible. Therefore, we examined the actions of exendin-4 on human pancreatic ductal epithelial cells in vitro. These in vitro studies on the actions of GLP-1R activation in pancreatic duct cells revealed a proliferative action mediated through the activation of MAPK pathways and phosphorylation of CREB, which was even more apparent in the setting of an activating Kras mutation and inhibited by the actions of metformin. This provides a mechanistic basis for the association of metformin treatment with decreased risk for pancreatitis and pancreatic cancer in individuals with T2DM (38,39). It is also consistent with a previous rodent study in which metformin attenuated the proliferative actions of the DPP-4 inhibitor sitagliptin on the pancreatic ductal tree (3).

In conclusion, we report that treatment of rats for 12 weeks with exendin-4 induced a marked expansion of PDGs through the mechanism of enhanced PDG cell replication. Moreover, we report that the PDGs in rats and humans express GLP-1Rs and that these also are abundantly expressed in PanIN lesions in human pancreas. GLP-1 treatment advances the rate of formation of dysplastic mPanIN lesions and chronic pancreatitis in a mouse model prone to the development of pancreatic ductal adenocarcinoma. Finally, we report that treatment of human pancreatic duct cells with the GLP-1 analog exendin-4 induces proproliferative signaling pathways, an effect that is inhibited by metformin. Collectively, these studies imply that GLP-1–induced proliferation within the exocrine pancreas is focal and may accelerate the development of dysplastic lesions when present.

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