Recapitulation of embryonic neuroendocrine differentiation in adult human pancreatic duct cells expressing neurogenin 3

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Regulatory proteins have been identified in embryonic development of the endocrine pancreas. It is unknown whether these factors can also play a role in the formation of pancreatic endocrine cells from postnatal nonendocrine cells. The present study demonstrates that adult human pancreatic duct cells can be converted into insulin-expressing cells after ectopic, adenovirus-mediated expression of the class B basic helix-loop-helix factor neurogenin 3 (ngn3), which is a critical factor in embryogenesis of the mouse endocrine pancreas. Infection with adenovirus ngn3 (Adngn3) induced gene and/or protein expression of NeuroDβ2, Pax4, Nkx2.2, Pax6, and Nkx6.1, all known to be essential for β-cell differentiation in mouse embryos. Expression of ngn3 in adult human duct cells induced Notch ligands Dll1 and Dll4 and neuroendocrine-β-cell–specific markers: it increased the percentage of synaptophysin- and insulin-positive cells 15-fold in ngn3-infected versus control cells. Infection with NeuroDβ2 (a downstream target of ngn3) induced similar effects. These data indicate that the Delta-Notch pathway, which controls embryonic development of the mouse endocrine pancreas, can also operate in adult human duct cells driving them to a neuroendocrine phenotype with the formation of insulin-expressing cells.

Introduction

Several studies have suggested that adult β-cells might originate from duct or duct-associated cells (Slack, 1995; Bouwens and Pipeleers, 1998; Edlund, 1999; Bonner-Weir et al., 2000). Evidence for this concept is largely indirect, and the underlying mechanisms are unknown. It is also conceivable that acinar cells can become a source of new β-cells in view of their plasticity, allowing them to transdifferentiate into hepatocytes (Shen et al., 2000) and into duct cells (Rooman et al., 2000). If postnatal acinar and/or duct cells could form new β-cells, they would become a particularly useful target for therapies that aim β-cell replacement in diabetic patients (Keymeulen et al., 1998; Shapiro et al., 2000), since both cell types are abundantly available in the pancreas of these patients and in donor organs. To assess such potential, we examined whether expression of key embryonic transcription factors in adult human duct cells could induce their differentiation into insulin-expressing cells.

Experiments with transgenic mice have indicated key factors in the embryonic development of their endocrine pancreas. Analysis of null mutants for Pdx1/Ipf1, ngn3, NeuroDβ2, Pax4, Nkx2.2, Nkx6.1, or Pax6 has identified a hierarchy of transcription factors that control embryonic formation of pancreatic islets (for review see Sander and German, 1997; Jensen et al., 2000a; Edlund, 2001). It is unknown whether these factors play a role in the postnatal growth of the pancreatic β-cell mass and whether they can be used to induce formation of human β-cells from postnatal nonendocrine cells. In this perspective, we examined the endocrinogenic potential of the class B basic helix-loop-helix (bHLH)* transcription factor neurogenin 3 (ngn3), which seems to function...
as a major and timely switch in the rodent embryonic pancreas (Gradwohl et al., 2000). When expression of ngn3 is directed ectopically into the embryonic epithelium, pancreas precursor cells develop prematurely and exclusively into glucagon-producing cells (Apelqvist et al., 1999; Schwitzgebel et al., 2000). Similarly, ngn3 induced premature differentiation into glucagon- and somatostatin-producing cells when introduced into early chicken endoderm (Grapin-Botton et al., 2001). The failure to induce insulin-producing cells might indicate that these immature cells lack the competence to drive β-cell differentiation. The differentiating activity of ngn3 is under control of Notch signaling. Indeed, null mutant mice for the Notch ligand Dll1, for an intracellular mediator of Notch signaling RBP-Jκ (Apelqvist et al., 1999), or for a downstream bHLH repressor HES1 (Jensen et al., 2000b), which possibly controls ngn3 (Tanabe and Jessell, 1996), all show premature endocrine differentiation. Therefore, we ectopically expressed ngn3 in adult duct cells to assess its role as a switch activating the expression of other developmental transcription factors and Delta-Notch proteins and consequently resulting in the appearance of endocrine differentiation markers, in particular insulin.

**Results**

**Absence of regulators of embryonic endocrine differentiation in postnatal human pancreatic duct cells**

Adult human β-cell preparations express a series of transcription factors that are crucial for embryonic development of mouse endocrine pancreas (Fig. 1 A). Transcripts encoding Pdx1/Ipf1, NeuroD/B2, Pax4, Pax6, Nkx2.2, and Nkx6.1 were abundant in adult human islets. The expression of Pax6 in human islets is at variance with its absence in postnatal mouse islets (Smith et al., 1999) or rat-purified β-cells (see Fig. 4 A). A parallel analysis of adult human duct cell transcripts shows the presence of relatively high levels Pdx1/Ipf1 and Nkx6.1 transcripts (Fig. 1 A). Pdx1/Ipf1 is also expressed in adult human duct cells at the protein level (Heimberg et al., 2000), but this is not the case for Nkx6.1 (Fig. 1 B), suggesting it is subject to posttranscriptional regulation. Sections of adult human pancreas with both β-cells and duct cells clearly indicated that Nkx6.1-positive nuclei were associated with insulin-containing cells and not with cells that expressed the ductal cell marker CA19.9 (Bouwens and Pipeleers, 1998) (Fig. 1 B). Ngn3 mRNA level was low in both islets and duct cells. Compared with islet cells, the duct cell levels of transcripts coding for Notch 1, 2, and 3 receptors were higher, those for Jagged 1 and 2 ligands were similar, and those for Dll1 and Dll4 ligands were much lower (see Fig. 3 C).

**Ngn3 induces expression of regulators of embryonic endocrine differentiation in postnatal human pancreatic duct cells**

We ectopically expressed ngn3 and its downstream target NeuroD/B2 in adult human duct cells. Common transfection methods were unsuccessful, but infection with recombi-
nant adenoviruses (Ad) resulted in efficient expression of the transgenes that were under control of a cytomegalovirus (CMV) promoter (Fig. 2). Both Adngn3 and AdNeuroD/β2 coexpressed GFP as a reporter, and AdGFP served as control for nonspecific viral effects. A multiplicity of infection (MOI) of 50 gave a favorable balance between infection efficiency (30–40% GFP expression) (Fig. 2) and cell survival (>85% living cells).

1 d after ngn3 infection, Pax4 gene expression was activated (Fig. 3 A) together withDll1 and Dll4, the latter two to similar levels as in control islet preparations (Fig. 3 C). 2 d later, NeuroD/β2 was induced, followed byNkx2.2 and Pax6 (Fig. 3 A). The Pdx1/Ipf1 gene remained silent or constant in, respectively, monolayer or suspension cultures of Adngn3-infected duct cells (unpublished data). These effects were confirmed by in situ hybridization and immunocytochemistry. 10 d after infection, many cells expressed either Nkx2.2 or ngn3 (Fig. 3 B, f, arrowheads) but rarely both. However, some Nkx2.2-expressing cells still contained GFP (Fig. 3 B, f, arrow). There was no significant increase in the level of Nkx6.1-encoding mRNA, but the protein appeared abundant (Fig. 3 B, j). Nuclear protein extracts from Adngn3-infected duct cells showed gel retardation of E box-1 and E box-3 sequences from the NeuroD/β2 promoter (unpublished data). Compared with ngn3, NeuroD/β2-induced changes in gene expression were similar but appeared with a several days delay (unpublished data). The same recombinant Ad constructs were used to infect the clonal neuroendocrine cell line PC12 and the unrelated HeLa cell line. In PC12 cells, which endogenously express NeuroD/β2 and Nkx2.2, both ngn3 and NeuroD/β2 induced Pax4 but not Nkx6.1 (Fig. 4 A). In HeLa cells, none of these endocrinogenic transcription factors were induced after infection with Adngn3 or AdNeuroD/β2.
Ngn3 induces expression of neuroendocrine markers, in particular insulin, in postnatal human pancreatic duct cells

Control duct cell preparations were negative by RT-PCR analysis for the endocrine cell markers synaptophysin, chromogranin A, and prehormone convertases (PC1/3 and PC2), and for the islet cell markers insulin, somatostatin, glucagon, glucose transporter type II, and glucokinase. Transcripts encoding these proteins were clearly present in the human islet cell fraction (Fig. 5 A; not depicted). 10 d after infection with AdNgn3 or AdNeuroD/B2, duct cell preparations exhibited a strong activation of synaptophysin, chromogranin A, and PC1/3 gene expression, and a weaker increase for the glucokinase and insulin genes (Fig. 5 A). No signals were detected for Glut2, somatostatin, or glucagon transcripts. These effects of ectopic ngn3 expression were also reflected at the protein level (Fig. 5 B). The fraction of cells that were immunopositive for insulin and synaptophysin increased from 1% in control preparations to, respectively, 13 and 22% 10 d after Adngn3 infection. The percentage of glucagon- and somatostatin-positive cells remained under 2% (unpublished data). However, although ectopic ngn3 did not affect the number of glucagon-containing cells, it increased the percentage of somatostatin-positive cells 3.5-fold (0.2 ± 0.1% versus 0.7 ± 0.1%, P < 0.05; n = 4). A similar effect was observed after infection with Ad-NeuroD/B2. Interestingly, most cells that expressed these endocrine markers were negative for ngn3 (Fig. 5 B, b, c, and g, arrowheads point to single positive cells). At this time point, the percentage of ngn3-positive cells was markedly lower than at day 3. Furthermore, the rare ngn3-positive cells that coexpressed endocrine markers exhibited a much lower ngn3 fluorescence intensity than those that did not (Fig. 5 B, b and g, arrows).

10 d after Adngn3 infection, many but not all synaptophysin-positive cells expressed chromogranin A (Fig. 5 B, k and l). More than 90% of the synaptophysin-expressing cells were still positive for the duct cell marker cytokeratin (CK19 (Fig. 5 B, i); however, this was the case for only part of the insulin- or somatostatin-positive cells (Fig. 5 B, h and j; not depicted). Electronmicrographs confirmed the presence of a large percentage of ngn3-infected cells with secretory granules (Table I) that were, however, smaller than the characteristic large granules of fully differentiated endocrine islet cells (Fig. 5 C).

Ectopic ngn3 or NeuroD/B2 expression in postnatal human pancreatic duct cells resulted in a threefold increase in the insulin content and insulin release of the preparations (Table II). In view of the 10-fold increase in the percentage of insulin-positive cells, these data suggest that the newly formed insulin-positive cells have a low insulin content.

In PC12 cells, ngn3 and NeuroD/B2 induced insulin expression, both at the transcript and protein level (Fig. 4) but failed to induce glucokinase, Glut2, somatostatin or glucagon. No ngn3-induced gene expression was seen in HeLa cells.
Discussion

The present study demonstrates that adenovirus-mediated delivery of ngn3, a key transcription factor for the generation of endocrine islet cells in mouse embryos, shifts adult human pancreatic duct cells into a neuroendocrine phenotype with expression of insulin in a significant fraction of transdifferentiated cells. It is unlikely that this effect requires the participation of Pdx1/Ipf1. This transcription factor is expressed in adult human pancreatic duct cell suspensions, be it at lower abundance and with a different phosphorylation status and DNA-binding activity compared with mature human islet cells (Heimberg et al., 2000); however, the ngn3-induced (neuro)endocrine differentiation was also achieved in duct cell monolayers, which are negative for Pdx1/Ipf1 (unpublished data). Of the known developmental transcription factors that operate downstream of Pdx1/Ipf1 to specifically control embryonic pancreas formation, ngn3 comes first in sequence. Ngn3 is a major regulator of lateral inhibition that controls endocrinogenesis in the embryonic mouse pancreas (Apelqvist et al., 1999). It has been proposed as a marker for pancreatic islet progenitor cells during embryogenesis and in adult mice (Jensen et al., 2000a; Gu et al., 2002). The amplified signal for ngn3 transcript in duct cells was similar to islets and suggests very low but specific expression in this cell fraction. Adenovirus-mediated overexpression of ngn3 in adult human pancreatic duct cells was found to activate expression of neuroendocrine differentiation markers and of Pax4 and insulin. Despite their independence of Pdx1/Ipf1, the requirement for ngn3 and the induction of Pax4 and somatostatin expression suggest that the ngn3-transdifferentiated (neuro)endocrine cells resemble cells of the second rather than the first wave of pancreatic endocrinogenesis. The described effects are cell type restricted, since they were reproduced in the PC12 neuroendocrine cell line but not in HeLa cells.

It is unknown presently whether this forced transdifferentiation is restricted to a subpopulation of duct cells. A major proportion of ngn3-infected cells expressed the (neuro)en-
endocrine markers, but only a fraction was insulin positive. The sequential activation of a comprehensive set of (neuro)endocrine-specific genes rather than the existence of (sub)populations of cells expressing only few individual markers is characteristic for the induction of a coordinated differentiation program. Based on its rapid nature and independence of cell proliferation (unpublished data), this process likely represents immediate transdifferentiation of the duct cells without the need for an intermediate cellular state (Shen et al., 2000; Slack and Tosh, 2001). Transdifferentiated cells are characterized by coexpression of the duct cell marker CK19 and the neuroendocrine marker synaptophysin. Such double positive cells are virtually absent in noninfected duct cell preparations. The expression levels of both markers changed gradually and reciprocally with time after infection. Only little cytokeratin positivity was left in the cells that finally expressed insulin. A combination of the transient nature of the adenoviral expression system and the elimination of fluorescence by GFP after extended fixation allowed simple tracing of cell fate. A direct relation between the ngn3-infected duct cells and the cells that became positive for endocrine markers was observed. Rare cells coexpressed endocrine marker proteins and traces of ngn3, suggesting a causal relationship between expression of ngn3 and the endocrine protein. Furthermore, several endocrine cells still contained the stable green fluorescent protein that remains present for days after its transcript, and consequently the mRNA encoding the less stable ngn3 protein, have disappeared (Corish and Tyler-Smith, 1999).

The mechanism whereby ngn3 induces duct cell differentiation into endocrine β-cells seems to involve activation of the Pax4 and the NeuroD/B2 promoter. Ngn3 is known to activate NeuroD/B2 expression in chicken embryos (Grapin-Botton et al., 2001), Xenopus embryos, and (neuro)endocrine cell lines (Huang et al., 2000). This is also the case in HeLa cells that had been transfected with E47, a class A bHLH heterodimerization partner of ngn3 (Huang et al., 2000). Adult human duct cells contain high endogenous levels of E47 (unpublished data), allowing E-box binding of the ectopically expressed ngn3 and activation of the NeuroD/B2 promoter. The delayed induction of NeuroD/B2 by ngn3 suggests the existence of intermediate transcription factors. Moreover, the delay in NeuroD/B2-induced Pax4 activation compared with ngn3 uncovers that both Pax4 and NeuroD/B2 are ngn3 targets, instead of Pax4 being downstream of NeuroD/B2. The present study thus supplements the hierarchy model of transcription factors involved in the formation of embryonic β-cells (Schwitzgebel et al., 2000). It also demonstrates that the embryological program in mice can be recapitulated in postnatal human duct cells, leading to formation of insulin, and to a minor extent somatostatin-expressing cells. In experimental terms, adult duct cells infected with adenoviruses expressing recombinant transcription factors are a simple in vitro model for studying the molecular biology of endocrine transdifferentiation.

Ectopic expression of ngn3 or NeuroD/B2 in isolated adult duct cells activates several (neuro)endocrine-specific genes, such as insulin and somatostatin, but not glucagon. In mouse or chicken embryonic endoderm cells in vivo, ectopic ngn3 induces glucagon but not insulin (Grapin-Botton et al., 2001). Activation of β-cell–specific Pax4 in duct cells might be responsible for the absence of glucagon expression (Smith et al., 1999; Petersen et al., 2000). The role of Nkx2.2 and Nkx6.1 in this transdifferentiation process is unclear: Nkx2.2 was stimulated by ngn3 at the transcriptional level, and Nkx6.1 was stimulated at the posttranscriptional level. Both transcription factors appear essential during embryonic development of β-cells, with Nkx2.2 acting upstream of Nkx6.1 (Sander et al., 2000). Adngn3-infected duct cells failed to generate glucose-induced insulin release within the limits of the present study, i.e., 10 d after infection (unpublished data). Our study can thus not be taken as evidence for the ability to produce functionally mature β-cells from pancreatic duct cells. Nevertheless, the data are indicative for a selective ability of adult duct cells to differentiate toward (neuro)endocrine and islet cells. Ngn3 induced moderate to high expression of synaptophysin, chromogranin A, PC1/3, and glucokinase, but the degree of insulin gene activation is low and so is the cellular insulin content. Transdifferentiated pancreatic duct cells exhibit the ultrastructural characteristics of immature endocrine cells, which is consistent with the absence of a glucose-regulated secretory activity. The absence of Glut2 induction is compatible with the earlier report that Glut2 is poorly expressed in human β-cells compared with human liver cells or rodent β-cells (De Vos et al., 1995).

The incomplete differentiation of this particular phenotype might be caused by, or related to, a variety of factors. (a) Although Pax4 is subject to autoregulation in mouse islet cell lines (Smith et al., 2000), ngn3–induced activation of Pax4 in human duct cells was not. This allows the relatively high levels of Pax4 to exert their inhibitory action on insulin gene expression (Qiu et al., 1998; Smith et al., 1999). Low levels of Pax4 also occurred in normal adult human islet cells in contrast to its restriction to the embryonic mouse pancreas (Sosa-Pineda et al., 1997; Smith et al., 1999, 2000). It is likely that in mature islet β-cells a yet unknown factor overrules the Pax4 repressor activity. Given the transient presence of ngn3 and Pax4 in the embryonic mouse pancreas, it needs to be investigated whether a closer simulation of the embryonic situation by conditional expression of ngn3 in adult human duct cells would augment the insulin levels. (b) The state of the Delta-Notch pathway probably also has its influence on the degree of (neuro)endocrinogenesis. In uninfected duct cells, high levels of Notch1, 2, and 3 but not of Dll1, 3, and 4 were found. Ngn3 infection activated transcription of Dll1 and Dll4 and thus increased Notch signaling in neighboring pairs, which is expected to limit overall endocrine differentiation. It may thus be useful to add antimorphic forms of Delta that antagonize Notch signaling (Jen et al., 1997) or to introduce ngn3 targets that bypass stimulation of Delta genes in order to specifically drive the formation of endocrine cells. (c) The transdifferentiated cells likely are dependent on extracellular factors to promote their maturation. (d) Finally, repression of specific proneural regulators induced by ngn3 may be necessary to allow full endocrine differentiation. Although the transdifferentiated duct cells did not express genes that label differentiated neuronal cells (N-Cam, neurofilament, peripherin, and class III β-tubulin [unpublished data]), major overlaps exist between the
expression profile of endocrine pancreas and neurons (Atouf et al., 1997; unpublished data). These similarities were key to design conditions that drive embryonic stem cells to insulin production (Lumelsky et al., 2001). Moreover, insulin-producing cells are present in primary cell cultures from mammalian fetal brain (Clarke et al., 1986), and insulin-producing neurons in *Drosophila* brain show remarkable similarities with β-cells in mammalian islets of Langerhans (Rullivan et al., 2002). All these data are highly suggestive for a common ancestral insulin-producing cell of neural origin, which complicates the identification of specific pronuclear factors that are nonessential for islet cell differentiation.

In more general terms, the present findings extend the role of lateral inhibition in cell differentiation from embryonic, poorly differentiated tissues (Artavanis-Tsakonas et al., 1999) to postnatal differentiated cell populations. Adult human duct cells were shown to transdifferentiate into neuroendocrine and into insulin-producing cells after ectopic expression of ngn3. Pancreatic duct and islet cells have a common embryonic progenitor, but according to a recent lineage tracing study ngn3 is never expressed in duct cells or their progenitors (Gu et al., 2002). However, our current study shows that ngn3 might serve as a master switch that drives transdifferentiation to a (neuro)endocrine phenotype when misexpressed in adult duct cells. It is still unknown whether ngn3 can be activated by environmental factors in normal duct cells of regenerating postnatal pancreas and thus result in the formation of new β-cells. Current efforts focus on finding ways to differentiate embryonic stem cells into insulin-producing cells with the purpose of producing β-cells that can be activated by environmental factors in the body.
Electron microscopy

Cell preparations were fixed in 2.5% glutaraldehyde in 0.1 M sodium cacodylate buffer (pH 7.4), postfixed in 1% osmium tetroxide, stained with 2% uranyl acetate, embedded in Spurr’s resin, and ultra thin plastic sections were examined with a Zeiss EM 952 electron microscope.

Data analysis

Results obtained from infected cells were compared with uninfected and/or control virus-infected cells and statistically analyzed using the paired Student’s t-test.

References

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