Targeted Overexpression of CKI-Insensitive Cyclin-Dependent Kinase 4 Increases Functional β-Cell Number Through Enhanced Self-Replication in Zebrafish

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Abstract

β-Cells of the islet of Langerhans produce insulin to maintain glucose homeostasis. Self-replication of β-cells is the predominant mode of postnatal β-cell production in mammals, with about 20% of rodent β-cells dividing in a 24-hour period. However, replicating β-cells are rare in adults. Induction of self-replication of existing β-cells is a potential treatment for diabetes. In zebrafish larvae, β-cells rarely self-replicate, even under conditions that favor β-cell genesis such overnutrition and β-cell ablation. It is not clear why larval β-cells are refractory to replication. In this study, we tested the hypothesis that insufficient activity of cyclin-dependent kinase 4 may be responsible for the low replication rate by ectopically expressing in β-cells a mutant CDK4 (CDK4R24C) that is insensitive to inhibition by cyclin-dependent kinase inhibitors. Our data show that expression of CDK4R24C in β-cells enhanced β-cell replication. CDK4R24C also dampened compensatory β-cell neogenesis in larvae and improved glucose tolerance in adult zebrafish. Our data indicate that CDK4 inhibition contributes to the limited β-cell replication in larval zebrafish. To our knowledge, this is the first example of genetically induced β-cell replication in zebrafish.

Introduction

Controlled genesis of β-cells in situ is a potential treatment for diabetes, as β-cell deficit underlies both type 1 and type 2 diabetes.1–4 Physiologically, β-cells are generated in three ways, neogenesis from precursor cells, self-replication, and transdifferentiation from other cell types. Understanding the molecular mechanisms that control β-cell genesis is essential to harness the capacity for cell-based therapy.5–7 In early stage mammalian embryos, β-cells are generated primarily through neogenesis, and self-replication is rare due to high levels of expression of inhibitors of cyclin-dependent kinases.8 In contrast, in perinatal and postnatal rodents, β-cell replication is the major contributor to β-cell mass expansion. β-Cell replication is highest during the perinatal period, occurring in about 20% of β-cells per day in rodents.9,10 In prenatal human embryos, 3.4% of β-cells were found to be positive for Ki67,11 a proliferation marker for all phases of cell cycle. The fraction of dividing β-cells per day may be much higher since the cell cycle length is likely shorter than 24 hours. Replicating β-cells decline precipitously to about 0.07%–3% in adults.9,10,12,13 The decline may result from the increased expression of cell cycle inhibitors and epigenetic modifications of key genes involved in cell cycle regulation.14,15 Under metabolic stress such as obesity and insulin resistance, some β-cells can re-enter the cell cycle to compensate for the increased insulin demand.16,17 However, the molecular underpinnings regulating self-replication of β-cells are not well understood.

Several groups have started to use zebrafish to study β-cell neogenesis, transdifferentiation, and replication.18–23 The zebrafish pancreatic islet shares morphological and physiological similarities with that of mammals.24 Many of the signaling pathways and transcription factors regulating β-cell development in zebrafish are homologous to mammals.24–26 However, unlike rodents, β-cell replication is rare in newly hatched embryos and free feeding larvae. Several groups reported that replicating β-cells are very rare in larvae at 3 dpf,27 4–6 dpf,22 and 6–8 dpf18 using BrdU incorporation as a measure of proliferation. Using PCNA expression as a measurement, Moro and associates found only 3 of 57 larvae at 7 dpf have a single PCNA-positive β-cell.20 Even under metabolic stress that promotes β-cell expansion, such as overnutrition or after β-cell ablation, replicating β-cells are still rarely observed.18,22
Nevertheless, certain adenosine receptor agonists, such as NECA, can stimulate β-cell replication after ablation, suggesting that larval β-cells have the capacity to replicate. A possible reason for the inability of larval β-cells to self-replicate is insufficient activity of cyclin-dependent kinases (CDKs), master regulators of proliferation. In addition to cyclins that activate CDKs, CDK inhibitors (CDI) that inactivate CDKs are also critical for regulation of CDK activity. There are two families of CKIs, the CIP family (p21, p27, and p57) and INK4 family (p15, p16, p18, and p19).

A first step in profloration stimulated by extracellular mitogens is induction of D-type cyclins which in turn activates CDK4. Active CDK4 phosphorylates and inactivates retinoblastoma (Rb), initiating G1-S progression. A mutant CDK4 found in human cancers, CDK4R24C, promotes tumorigenesis because it is insensitive to inhibition by CKIs, contributing to the lack of cell replication in zebrafish larvae.

Materials and Methods

Zebrafish strains, maintenance, and nutrient treatment

Adult zebrafish were raised in Aquatic Habitats systems on a 14/10-h light/dark cycle. Embryos were obtained from natural crossing. Fertilized eggs were reared in 0.3x Danieau solution at 28.5°C in an incubator with lights on a 14/10-h light/dark cycle, and staged following standard methods. For egg yolk feeding, chicken eggs were obtained from local grocery stores, the yolk separated and diluted to 5% by volume with 0.3x Danieau solution. Larvae were cultured in the egg yolk solution according to Maddison et al. The transgenic lines used for this study were Tg(−1.2ins:H2BmCherry)18. All procedures have been approved by the Vanderbilt University IACUC Committee.

Generation of transgenic zebrafish

We generated a miniTo12-based transgene consisting of human CDK4R24C cDNA under the control of the zebrafish insulin gene promoter, using a 1.2 kb fragment from upstream of the initiation ATG and tagRFP under the control of 0.9 kb promoter of zebrafish zα crystalline gene (Fig. 1A). The transgene, Tg(−1.2ins:CDK4R24C, −0.9cryaatz:tagRFP)10, was in vitro transcribed and injected into zebrafish embryos at the one-cell stage using standard protocol as described. Embryos with tagRFP expression in lens were selected at 4 dpf after injection, and grown to adulthood. Individual F0 founders were outcrossed to the wild-type AB line, and their F1 progeny were screened for tagRFP expression in the lens and confirmed by PCR using primers zINSSTF:5′-CCAC- CATTCTCGCCTCTGCCT-3′ and hCDK4NR: 5′- CAGATATGTCCTTAGTGCCTGTC-3′. Initial analyses were performed in two independent lines and similar results were obtained. All results reported here were from F2 or F3 fish of one line that has stronger red fluorescence in the lens, Tg(−1.2ins:CDK4R24C, LR)18.

Counting of β-cells

After fixation in 4% paraformaldehyde (PFA) overnight in 4°C, larvae were washed with 1x PBS plus 0.1% Tween-20 (PBST) and flat mounted in Aqua-Mount (Richard-Allan Scientific) with their right side facing the coverslip. The larvae were flattened sufficiently to disrupt the islet slightly to allow better resolution of individual nuclei. The β-cell number was counted according to the nuclear mCherry signal using a Zeiss AxiosImager under a 40x lens or using confocal projections taken by Zeiss LSM510 under a 40x lens (Carl Zeiss).

Whole mount immunofluorescence

The larval zebrafish or dissected pancreata from Tg(+/ins:CDK4R24C, LR)18 or wild-type zebrafish were fixed in 4% PFA overnight at 4°C and then transferred to 100% ethanol and stored at −20°C. Larvae were rehydrated, perméabilized in aceton for 30 min at −20°C, washed in PBST, and nonspecific binding blocked with PBST plus 1.0% DMSO, 1% BSA, and 5% normal goat serum. Subsequently, antibodies for Cdk4 (1:50, Santa Cruz sc-260) or insulin (1:1000, DakoCytomation A0564) antibodies were diluted in blocking buffer and incubated overnight at 4°C. Antibodies were detected using the appropriate Alexa Fluor 488-conjugated

**FIG. 3.** Increased proliferation in Tg(−1.2ins:CDK4R24C, LR) fish. Twenty-four-hour cell proliferation analysis using EdU labeling in 6-dpf unfed Tg(−1.2ins:H2BmCherry)µ506 (A), Tg(−1.2ins:CDK4R24C, LR)µ506 (B), Tg(−1.2ins:H2BmCherry)µ506 (C), and Tg(−1.2ins:H2BmCherry)µ506 (D) fish. In Tg(−1.2ins:H2BmCherry)µ506 larvae β-cells (red) rarely incorporated EdU, while Tg(−1.2ins:CDK4R24C, LR)µ506, Tg(−1.2ins:CDK4R24C, LR)µ506, Tg(−1.2ins:H2BmCherry)µ506 have multiple EdU positive β-cells. The images are confocal projections and scale bars indicate 10 µm. (E) Quantification of the β-cell number in 6-dpf Tg(−1.2ins:H2BmCherry) and Tg(−1.2ins:CDK4R24C, LR)µ506, Tg(−1.2ins:H2BmCherry)µ506, in unfed and egg yolk fed conditions. (F) Quantification of 24 h EdU labeled β-cells, in each group, n = 8–9. (G and H) EdU was incubated from 2-dpf to 6-dpf in unfed Tg(−1.2ins:H2BmCherry)µ506 (G) and Tg(−1.2ins:CDK4R24C, LR)µ506, Tg(−1.2ins:H2BmCherry)µ506 (H), the images are confocal slices and scale bars indicate 10 µm. (I) Quantification of 2-dpf to 6-dpf EdU labeled β-cells, in both group, n = 10. All the values shown are means±S.E., **p<0.001, *p<0.05.**
secondary antibodies (1:2000, Invitrogen). Slides were imaged on a LSM 510 confocal microscope (Carl Zeiss).

Proliferation analysis

Proliferation was analyzed using the Click-iT EdU Alexa Fluor 488 Imaging Kit (C10337; Invitrogen). For 24 hours labeling, 5 dpf zebrafish larvae were injected pericardially with 3 nL of 100 mmol/L 5-ethynyl-2-deoxyuridine (EdU) and harvested after 24 hours. For long-term incubation, 2 dpf embryos was incubated with 100 μmol/L EdU, and harvested at 6 dpf. EdU was detected according published protocols. All images were collected using a Zeiss LSM510 or Zeiss LSM710 (Carl Zeiss).

**FIG. 1.** Schematic representation of the Tg(1.2ins:CDK4^{R24C}; LR) transgenic line and the expression pattern. (A) Expression of a human CDK4 mutant (CDK4^{R24C}) was controlled by the zebrafish insulin promoter with α-crystallin driven tagRFP used as an indication of transgene carriers. (B, C, D) Confocal projections of CDK4^{R24C}-expressing cells in double-transgenic Tg(1.2insCDK4^{R24C}; LR)^{V1208}, Tg(1.2ins:H2BmCherry)^{V1208} fish, with CDK4^{R24C}-expressing cells labeled by an antibody against human specific CDK4 protein (B). β-cells are indicated by nuclear mCherry protein (C).

**FIG. 2.** β-Cell specific overexpression of CDK4^{R24C} increases β-cell number. Images of β-cells Tg(1.2ins:H2BmCherry)^{V1208} (A, C) and double-transgenic Tg(1.2ins:CDK4^{R24C}; LR)^{V1208}, Tg(1.2ins:H2BmCherry)^{V1208} (B, D) fish at 6 dpf (A, B) and 23 dpf (C, D). Scale bars, 10 μm. (E) β-Cell number in Tg(1.2ins:H2BmCherry)^{V1208} and Tg(1.2ins:CDK4^{R24C}; LR)^{V1208}, Tg(1.2ins:H2BmCherry)^{V1208} at 3, 4, 5, 6, 14, and 23 dpf. Expression of CDK4^{R24C} resulted in an increase in the number of β-cells throughout this developmental period. In each group, n = 7–18. The values shown are means ± S.E., **p < 0.01, *p < 0.05.**
Glucose tolerance testing

Glucose tolerance was determined in adult fish based on published protocols with the following modifications. The glucose solution for injection was prepared with Hanks Buffered Salt Solution (Cellgro). Fish were anesthetized using ice water, injected intraperitoneally with 0.5 mg glucose/g fish weight and allowed to recover for the 30, 90, or 180 min after injection. For blood collection, fish were fully anesthetized in ice water, dried with a Kimwipe to remove as much water as possible. An incision was made just posterior to the gill operculum, resulting in puncture of the heart. Blood was collected into heparinized hematocrit tubes (IRIS Sample Processing) and subsequently transferred into microcentrifuge tubes and frozen at −80°C. Glucose concentration in the samples was determined using Amplex Red Glucose Assay (Life Technologies). The glucose concentration was then normalized to hemoglobin content, which was determined using Drabkin’s reagent (Sigma). In our hands, this assay is more sensitive and less variable than the OneTouch Ultra Glucose meter (LifeScan).

Beta-cell area

Adult fish were euthanized and immediately injected with 4% PFA into the peritoneal cavity to ensure rapid fixation of internal organs. The fish were then placed in 4% PFA for 2 days at 4°C. The gastrointestinal tract was removed from each fish and rinsed in PBS + 0.1% Tween-20, dehydrated in methanol and stored at −20°C. After insulin immunofluorescence using the protocol outlined above, the pancreas was carefully dissected away from the rest of the GI tract, placed on slides, and flat mounted avoiding multiple layers of tissue.

To determine relative β-cell area, 6–10 islet-containing images for each fish were analyzed. The images were obtained under a 20× DIC lens on a Zeiss AxioVert 200 microscope equipped with a CCD camera. The insulin-positive area and the area of the exocrine pancreas for each image was determined using ImageJ software.

Statistics

Differences among groups were analyzed by Student’s t-test or by one-way ANOVA, followed by Fisher post-hoc test (SPSS, Chicago, IL). Data are shown as means ± standard error (SE). Statistical significance was accepted when p < 0.05.

Results

Generation of transgenic zebrafish with constitutive expression of CDK4R24C in β-cells

We obtained two independent Tg(−1.2ins:CDK4R24C, LR)VU508 lines. Although the intensity of lens tagRFP differs in the two lines, their β-cell phenotypes are indistinguishable. The body length of Tg(ins:CDK4R24C, LR)VU508 fish was not different from wild-type siblings at all stages measured (data not shown), indicating that the transgene does not impact growth. To ascertain that CDK4R24C was properly expressed, we crossed Tg(−1.2ins:CDK4R24C, LR)VU508 to Tg(−1.2ins: H2BmCherry)VU506 fish that express a nuclear localized mCherry protein only in β-cells18 and performed whole-mount immunostaining. Robust expression of CDK4R24C protein was detected in β-cells of transgenic fish at 6 dpf, but not in wild-type controls (Fig. 1B–D and data not shown).

![Fig. 4](image_url)

**Fig. 4.** Adult Tg(−1.2ins:CDK4R24C)VU508 fish have increased β-cell mass and improved glucose tolerance. (A, B) Images of insulin positive islets in adult Tg(−1.2ins:CDK4R24C, LR)VU508 (B) and nontransgenic siblings (A), scale bars indicate 20 μm. (C) β-Cell area relative to total pancreas area, 10 control and 8 Tg(−1.2ins:CDK4R24C; LR)VU508 fish were analyzed, **p < 0.001. (D) glucose clearance in Tg(−1.2ins:CDK4R24C)VU508 and nontransgenic siblings. Glucose was injected intraperitoneally at 0.5 mg glucose/g body weight and blood glucose levels determined 30 and 180 min after injection. 10–14 fish per time point for each group.
CDK4<sup>R24C</sup> expression increases β-cell number in larvae

To determine the consequence of CDK4<sup>R24C</sup> expression on β-cell number, the number of β-cells was determined in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub>, Tg(1.2ins:H2BmCherry)<sub>V1506</sub> fish, and Tg(1.2ins:H2BmCherry)<sub>V1506</sub> siblings at various stages. Similar to other reports, 18,19 we found 33.2 ± 1.0 (mean ± SE) β-cells in the principal islet of the control group at 6 dpf (Fig. 2A and E). Whereas in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub>, Tg(1.2ins:H2BmCherry)<sub>V1506</sub> fish, the number of β-cells increased to 41.3 ± 0.9 (Fig. 2B and E). No significant difference in pancreatic x-cells was found between the two groups (data not shown). An increased number of β-cells was found in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub>, Tg(1.2ins:H2BmCherry)<sub>V1506</sub> fish as early as 3 dpf (32.4 ± 1.1 vs. 28.2 ± 0.7, p < 0.05). In each of the subsequent 11 days, roughly one extra β-cell was added in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> larvae compared to control siblings. This results in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub>, Tg(1.2ins:H2BmCherry)<sub>V1506</sub> fish having 10 more β-cells than Tg(1.2ins:H2BmCherry)<sub>V1506</sub> fish (66.4 ± 3.2 vs. 56.3 ± 1.7, p < 0.01) at 14 dpf. At some point after 14 dpf, substantially more β-cells were produced in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish. As a result, at 23 dpf there are 40% more β-cells in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish than control (192.6 ± 18.8 vs. 141.1 ± 9.6, p < 0.05) (Fig. 2C–E). Taken together, these data indicate that expression of CDK4<sup>R24C</sup> causes supernumerary β-cells in zebrafish.

CDK4<sup>R24C</sup> expression enhances β-cell replication in larvae

To determine whether the supernumerary β-cells in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish is due to replication, we labeled replicating cells for 24 h by pericardial EdU injection in 5 dpf larvae. In the control group, no EdU labeled β-cells were observed, consistent with previous data 18,20,22 (Fig. 3A). However, we found 2.0 ± 0.5 EdU positive β-cells in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish (Fig. 3B and F). These cells presented predominantly as doublets, consistent with β-cell replication. With long-term EdU labeling from 2 dpf to 6dpf, Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> larvae had 11.2 ± 0.8 EdU-positive β-cells, while control siblings had only 1.9 ± 0.6 (Fig. 3G–I), indicating that all the extra β-cells in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish arose through replication.

To determine whether the transgenic fish with extra β-cells have an altered compensatory response to increased insulin demand, we challenged Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> larvae in an overnutrition paradigm that stimulates compensatory β-cell expansion. 18 Interestingly, although culturing in 5% egg yolk increased β-cell number in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish, (43.6 ± 0.9 vs. 40.6 ± 1.2, p = 0.015) (Fig. 3E), the response is dampened compared to WT larvae, with only 3 additional β-cells compared to the 7 β-cells added in WT larvae. The increase is likely through neogenesis, since there was not a significant increase of EdU-positive β-cells (2.6 ± 0.6 vs. 2.0 ± 0.5, p = 0.43) (Fig. 3C, D, and F).

CDK4<sup>R24C</sup> expression improves glucose tolerance

To determine if functional β-cells are continuously produced in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish, we determined β-cell content and glucose tolerance in 1-year-old fish. To determine β-cell content, the pancreas of Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish and wild-type siblings was dissected and stained with an insulin antibody. The insulin-positive areas in Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> pancreas was significantly greater than in the control siblings (Fig. 4A and B). The ratio of islet area to exocrine pancreas was ~2 times greater in the Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish than in their wild-type siblings (Fig. 4C). To investigate glucose clearance, blood glucose concentrations were examined at 0, 30, and 180 min after IP glucose injection of 0.5 mg glucose/g body weight. As shown in Figure 4D, glucose levels peaked at 30 min and declined to nearly pre-glucose administration levels at 180 min. When compared to control siblings, Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish had lower blood glucose following glucose injection, although there was no difference in fasting blood glucose. These data suggest that adult Tg(1.2ins:CDK4<sup>R24C</sup>; LR)<sub>V1508</sub> fish have an increased β-cell mass, and this contributes to improved glucose tolerance.

Discussion

Postnatal plasticity of β-cell mass in mammals results mainly from self-replication. 18 Yet the molecular mechanisms underlying β-cell replication is not fully understood. In contrast to perinatal mammals, larval zebrafish β-cells rarely replicate, even under conditions that promote β-cell mass expansion. Understanding the mechanism of this replication refractory period should also shed light on regulation of β-cell replication.

CDK4 plays a key role in cell cycle re-entry by integrating extracellular mitogenic and antimitogenic signals. CDK4 is normally inhibited by the INK4 family of CDK inhibitors. This inhibition is relieved by D-type cyclins that can be induced by mitogenic signals. 28 To test if insufficient CDK4 activity underlies the replication resistance of larval β-cells, we generated transgenic fish to express CDK4<sup>R24C</sup>, a mutant that is insensitive to the INK4 family of CDK inhibitors, 32,35,45 ectopically in the β-cells. As in mice, targeted expression of CDK4<sup>R24C</sup> result in increased β-cell content and improved glucose tolerance in adults (Fig. 4). 35 Nevertheless, the 2-fold expansion of β-cells compared to wild-type siblings is not as large as seen in the CDK4<sup>R24C</sup> transgenic mice, possibly due to intrinsic differences in the two animal models. In larvae, targeted expression of CDK4<sup>R24C</sup> resulted in one dividing β-cells (or 2%-4%) each day in the first 2 weeks of life. Much more β-cell replication occurred between 14 to 23 dpf. To our knowledge, this is the first report of genetically induced β-cell replication in zebrafish.

Since CDK4<sup>R24C</sup> still requires cyclin Ds for activity, the increased β-cell self-replication in the transgenic fish suggests sufficient cyclin D is present in β-cells to allow for replication. High levels of INK4 inhibitors relative to D-type cyclins may prevent these cells from self-replicating. More direct analyses will be required to dissect the precise mechanisms and contributions of cyclins and CKIs.

The impetus for generating the transgenic zebrafish model was to understand the lack of β-cell replication in larval zebrafish. Expression of CDK4<sup>R24C</sup> in zebrafish does lead to replication in larval β-cells (Figs. 2 and 3). Furthermore, these β-cells seem to be functional, since a less robust expansion was induced by overnutrition in the CDK4<sup>R24C</sup> fish than control.
siblings and adult transgenic fish are more glucose tolerant (Figs. 3E and 4). However, the fraction of replicating \( \beta \)-cells (2%–4%) is still very small compared to perinatal rodents. In addition, no significant increase of replication induced by overnutrition in the CDK4R24C fish. The data suggest that insufficient CDK4 activity plays a minor role in limiting self-replication of larval \( \beta \)-cells and other unidentified factors are more predominant players. The adenosine pathway is unlikely to be involved, since its activation only increases \( \beta \)-cell replication after ablation.\(^\text{22}\) Identification of the factors that control \( \beta \)-cell replication in zebrafish larvae may shed light on the diminished self-replication of \( \beta \)-cells in adult mammals.

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Disclosure Statement

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