The postnatal growth of the β-cell mass in pigs

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Abstract

Studies of the postnatal growth of the β-cell mass in rats have revealed some unexpected and apparently paradoxical results, the most prominent being a β-cell mass plateau in the early phase of life. We have studied the postnatal growth of the β-cell mass in the domestic pig to investigate its development in a larger mammal. The pancreas from a total of 86 male pigs from 5 to 100 days of age were studied. The β-cell mass increased linearly from day 5 to day 40, reached a plateau from day 40 to day 60, and then increased further into adulthood. The relative β-cell mass (β-cell mass per body mass) was increased in the early postnatal period but reached a constant level from day 60, after which there was a linear relationship between the β-cell mass and the body mass. There were high rates of both β-cell apoptosis and mitosis at 50 and 60 days of age, while the volume-weighted mean islet volume increased from birth and reached a plateau at approximately 60 days of age. A β-cell mass plateau early in life accompanied by a wave of β-cell apoptosis coinciding with the relative β-cell mass decreasing to reach a constant level, and a linear relationship between the β-cell mass and the body mass in later life is exactly what has previously been reported in rats. The coincidence of these events in both rats and pigs, although occurring at different ages in the two species, suggests a causal relationship as previously suggested in a proposed explanatory model for postnatal β-cell growth. 


Introduction

The β-cell mass in rats, following a slow and apparently constant increase from postnatal day 4 to postnatal day 16, reaches a plateau phase from day 16 until day 24 followed by a further increase into adulthood (Finegood et al. 1995, Scaglia et al. 1997, Svenstrup et al. 2002). This plateau phase seems to be paradoxical when the pronounced increase in whole body and organ weight in this time-interval is considered. We have previously reported that in rats the β-cell mass plateau phase coincides in time with the decrease in the relative β-cell mass (β-cell mass per body mass), and that the β-cell mass plateau phase ends when the relative β-cell mass reaches a constant level that is maintained into and during adulthood (Svenstrup et al. 2002). We have suggested that these two events are causally related, and that the β-cell mass plateau is part of an adaptation from the β-cells to the high relative β-cell mass, implying an increased tendency for apoptosis. In the present study, we have investigated the postnatal growth of the β-cell mass in the domestic pig to explore whether the characteristics observed in rats could be reproduced in a larger mammal.

Materials and Methods

Pigs

All pigs were purchased from Sealand III (Roskilde, Denmark), which is a research plant owned and managed by The Danish Slaughteries, Copenhagen, Denmark. All breeder females were Danish–Landrace/Danish–Yorkshire while all breeder males were Danish–Duroc, a constellation commonly used in commercial pig breeding. The piglets were fed by their mothers until weaning at postnatal day 28. From then they were fed a conventional diet as used in commercial pig breeding (Roskilde Andel AMBA, Roskilde, Denmark).

Study groups

A total of 86 male piglets at different ages were studied. Ages studied were postnatal day 5 (n=5), 10 (n=5), 15 (n=8), 20 (n=6), 25 (n=6), 30 (n=6), 40 (n=6), 50 (n=9), 60 (n=12), 70 (n=6), 80 (n=6), 90 (n=5) and 100 (n=6). Piglets from one litter were randomized into only one of the study groups, and no sow delivered more than three piglets into one age group. The piglets were killed by either a lethal injection of barbituric acid or by a bolt pistol followed by exsanguination depending on age. All such procedures were performed by authorized staff members according to Danish laws. The total pancreas was immediately removed, weighed and fixed in 10% buffered formalin, pH 7.4.

Histology

Following fixation, the pancreas was embedded in 4% agar to allow sectioning of the entire pancreas into 4 mm thick...
slices. The first cut was placed at a random position between 0 and 4 mm in each pancreas to ensure systematic uniform random sampling (SURS) (Gundersen & Jensen 1987). Between six and ten of the 4 mm thick slices were sampled according to SURS, each of these slides was embedded into paraffin, and one 3 µm thick section was then cut from the top of each block. The β-cells were visualized by immunohistochemistry using a guinea pig anti-insulin antibody (Dako, Glostrup, Denmark; dilution 1:50) as primary antibody followed by Envision-AP (a polymer backbone to which alkaline phosphatase, goat anti-rabbit immunoglobulin antibody and goat anti-mouse immunoglobulin antibody are attached; Dako; dilution 1:1). Antibody binding was visualized by incubation with Fast Red (Sigma Fast red; Sigma, St Louis, MO, USA). Mayer’s haematoxylin was used as counterstaining.

Stereology

The sections were examined using an Olympus BH-2 microscope equipped with a projecting arm to project the image onto the table (final magnification × 145), and an automated x-y stepper (Lang GMBH, Hüttenberg, Germany) to allow sampling of fields of vision according to SURS. Using a point-counting grid with 150 points (Fig. 1A, one of them encircled (the unit-point), the total number of points that hit β-cells and the number of unit-points that hit the pancreas, and the number of unit-points that hit non-pancreatic tissue such as fat or lymph nodes was counted on all sections from each pancreas. The β-cell mass and the pancreatic mass were then calculated according to:

$$M(\beta) = \frac{M(tis) \times P(\beta)}{150 \times (P_U(\text{pan}) + P_U(\text{non-pan}))}$$

$$M(\text{pan}) = \frac{P_U(\text{pan})}{P_U(\text{pan}) + P_U(\text{non-pan})}$$

where $M(\beta)$ is the total β-cell mass, $M(tis)$ is the weight of the tissue removed from the abdomen of the pig, $P(\beta)$ is the number of points that hit β-cells, $P_U(\text{pan})$ is the number of unit-points that hit pancreatic tissue, $P_U(\text{non-pan})$ is the number of unit-points that hit non-pancreatic tissue, and $M(\text{pan})$ is the total mass of the pancreas.

β-Cell mitosis and apoptosis

From three randomly selected pigs in each of the age groups postnatal day 5, 10, 20, 30, 40, 50, 60, 70, 80, 90 and 100, two randomly selected sections among the sections stained for insulin as described above were used to determine the rate of β-cell mitosis and apoptosis. The sections were investigated systematically according to SURS with a counting frame attached to the table.
At a final magnification of \( \times 583 \) we counted how many \( \beta \)-cell nuclei were sampled by the counting frame in each section and, in addition, how many of these nuclei displayed the morphological characteristics of either mitosis (Fig. 2A–B) or apoptosis (Fig. 2C–D). For each pig an average of 1304 ± 115 (means ± s.e.m.) \( \beta \)-cell nuclei were counted in the counting frame. The data for \( \beta \)-cell mitosis and apoptosis are expressed as the ratio of counted \( \beta \)-cell nuclei that displayed the characteristics of either mitosis or apoptosis.

### Volume-weighted mean islet volume

The volume-weighted mean islet volume was determined as previously described (Skau et al. 2001) by investigating the same sections used for the investigation of \( \beta \)-cell mitosis and apoptosis. Briefly, at a final magnification of \( \times 308 \), each section was investigated systematically according to SURS with a point-grid with direction lines attached to the table (Fig. 1C). Each time a point hits an islet, the length of the intersect of a line through the point and oriented according to the direction lines was measured. These sets of intersect lengths were then corrected for magnification and expressed in \( \mu m \). As previously described (Skau et al. 2001), the volume-weighted mean islet volume was then calculated according to:

\[
v_{V}(isl) = \frac{\pi}{3} \times l_0^3
\]

where \( v_{V}(isl) \) is the volume-weighted mean islet volume and \( l_0 \) is the length of an intersect through a sampling point in the direction given by the direction lines as described above and illustrated in Fig. 1C. The volume-weighted mean islet volume is the mean islet volume when the islets are sampled (weighted) proportional to their volume. Unless all islets are of identical volume, then the volume-weighted mean islet volume is higher than the number-weighted (ordinary) mean islet volume, as shown by:
\[ v_v = v_N \times (1 + CV_N^2) \]

where \( v_v \) is the volume-weighted mean islet volume, \( v_N \) is the number-weighted (ordinary) mean islet volume, and \( CV_N \) is the coefficient of variation between islet volumes in the number-weighted distribution.

**Statistics**

Error bars represent the s.e.m. The Pearson product-moment method was used for analysis of correlation. Linear regression analysis was based on the least squares method. The rates of β-cell mitosis and β-cell apoptosis over time was evaluated by one-way ANOVA.

**Results**

Figure 3 shows body weight as a function of age. This curve is in good agreement with prospective growth curves obtained in commercial pig breeding. The β-cell mass (Fig. 4A) showed a constant increase from day 5 until day 40, followed by a plateau until day 60 where it continued to increase into adulthood. The relative β-cell mass (Fig. 4B) was high in the younger age groups and decreased, most steeply from day 40 until day 60, to reach a constant (adult) level at day 60 that was maintained for the rest of the period studied. The pancreatic mass...
(Fig. 4C) increased during the entire age span investigated, while the relative pancreatic mass (Fig. 4D) showed an increase around day 30 followed by a slow decline.

Figure 5 shows the results of the β-cell mass as a function of age in pigs 60 days of age or older (the period during which the relative β-cell mass remained at a constant level). There was a significant correlation between the two variables, $r = 0.92$, $P < 10^{-14}$. The linear regression line shown in Fig. 5 has the equation:

$$M(\beta) = 21.0 \text{ mg} + (9.9 \text{ mg/kg BM(kg)})$$

where $M(\beta)$ is the β-cell mass and BM is the body mass. The S.E.M. for the constant was 22·0 mg and 0·71 mg/kg for the coefficient.

The rates of β-cell mitosis and apoptosis in the different age groups are shown in Fig. 6. As can be seen, there was a postnatal wave of β-cell apoptosis (Fig. 6A) that was most pronounced around days 50–60. But, interestingly, Fig. 6B shows that the rate of β-cell mitosis was also relatively high in the period 50–60 days after birth. The difference between the mean values at different ages were, for both parameters, greater than those that would be expected by chance, $P = 0.026$ for the rate of apoptosis and $P = 0.021$ for the rate of mitosis.

Figure 7 shows typical patterns of the histological appearance of the pancreas in pigs aged 5 and 80 days. The volume-weighted mean islet volume (Fig. 8) increased from a low value at postnatal day 5 to reach a level at approximately day 50 that seemed to be maintained up to day 100 which was the oldest age investigated.

Discussion

This study revealed striking similarities in the postnatal β-cell growth in domestic pigs and rats. In both species, the β-cell mass increases constantly after birth, reaches a plateau and then increases further into adulthood. The age for the plateau phase differs between the two species since it occurs between day 16 and day 24 (Finegood et al. 1995, Scaglia et al. 1997, Svenstrup et al. 2002) in rats and between day 40 and day 60 in pigs. Even though there is a difference in the time at which the plateau phase occurs, the obvious similarities in the appearance of the β-cell mass versus age curves in the two species points to a common etiology.

Based on the data from rats reported previously, one could speculate that the etiology of the plateau phase was related to weaning since laboratory rats are usually weaned at postnatal day 21. But since the pigs investigated were weaned at day 28, this theory is not supported by the results in this study. We previously reported that the β-cell mass plateau phase coincides in time with the relative β-cell mass reaching the constant ‘adult’ level in rats (Svenstrup et al. 2002), and this observation was reproduced in pigs in this study, supporting the view that these
two events are causally related. Since the most pronounced decrease in the relative β-cell mass occurs during the β-cell mass plateau phase, and especially since the plateau phase ends as the relative β-cell mass reaches the constant level, the plateau could be interpreted as an adaptive reaction by the β-cells to the high relative β-cell mass that ceases when the relative β-cell mass reaches the constant level as previously suggested (Svenstrup et al. 2002). But this and other theories are limited by the fact that the mechanisms that regulate the β-cell mass and the relative β-cell mass are still not understood in detail. It seems evident, however, that the metabolic demand on the β-cells is important, as shown by the increased β-cell mass under conditions where there is an increased demand for insulin secretion such as pregnancy (Blondeau et al. 1999), continuous glucose infusion (Bonner-Weir et al. 1989) and obesity in ob/ob mice (Edvell & Lindstrom 1995), and the decreasing β-cell mass in the presence of an implanted insulinoma (Blume et al. 1995). But whether the key player is the mean daily insulin secretion, hereditary factors, a complex system of the two, or other factors such as glucagon-like peptide 1 and islet neogenesis-associated protein, is still not known.

The β-cell mass plateau has been shown to be accompanied by a wave of β-cell apoptosis in rats (Scaglia et al. 1997). This phenomenon could also be confirmed in this study in pigs, as seen from the high values of β-cell apoptosis in pigs 50–60 days of age (Fig. 6A). It has previously been shown that two animal models of spontaneous type-1 diabetes, BioBreeding rats and non-obese...
diabetic mice, display a reduced clearing of apoptotic bodies by macrophages, and this defect may play a role in the β-cell targeted autoimmunity that leads to diabetes in these models (Trudeau et al. 2000, O’Brien et al. 2002a,b). In this way, the β-cell mass plateau and the simultaneous increased rate of β-cell apoptosis may be involved in the pathogenesis of diabetes. However, in contrast to the findings in rats (Scaglia et al. 1997), we also found a wave of β-cell mitosis at the end of the β-cell mass plateau phase in pigs. In addition, the volume-weighted mean islet volume increased in the postnatal period but apparently also reached a plateau after approximately 60 days of age in pigs. Together, these findings point to the period around the β-cell mass plateau phase as representing a period with remodelling of the endocrine pancreas, somewhat in parallel to that which has previously been shown in rats (Scaglia et al. 1997). Which mechanisms drive this remodelling are still not clear, but we suggest that the calibration of the relative β-cell mass to the level that is maintained into adulthood is a key player, since only this parameter has an obvious direct impact on the overall glucose homeostasis.

The finding of a linear relation between β-cell mass and body weight in pigs after the end of the plateau phase is also in good agreement with that which has previously been reported in rats (Montanya et al. 2000). The value for the adult steady-state level for the relative β-cell mass seems to be higher in rats than in pigs, approximately 15 mg/kg in rats versus 10.5 mg/kg in pigs. The sudden increase in the relative mass of pancreas around day 30 is probably a consequence of the change in eating behaviour due to weaning at day 28.

In summary, these data describe the postnatal development of β-cells in pigs. We have found pronounced similarities between the postnatal growth of the β-cell mass in pigs and rats such as a plateau phase in the β-cell mass accompanied by a wave of β-cell apoptosis, coinciding with the relative β-cell mass reaching the constant ‘adult’ level, and the linear correlation between β-cell mass and body weight thereafter. Given the data from rats and pigs, it would be highly interesting to investigate whether these results could also be reproduced in humans. However, even with access to the pancreases of young humans who have died suddenly, the number of cases needed to give an adequate description of the development in an organ displaying considerable biological variation and, furthermore, prone to fast postmortem degradation, makes such an investigation unlikely. However, non-invasive techniques to measure the β-cell mass are under development (Moore et al. 2001, Larsen et al. 2003), and these techniques may enable an indirect investigation of human β-cell mass development in the future. The identical patterns found in two such poorly related mammals as rats and domestic pigs suggests that these patterns are highly conserved and may also occur in the human pancreas.

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