

Research Article

Immunohistochemical Localization of Stem Cell-Related Markers (CD133, Nestin and Bmi-1) and mTOR in the Pancreas of Rat Models of Type 1 and Type 2 Diabetes Mellitus

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Abstract

Stem cell-related markers (CD133, nestin and Bmi-1) and mammalian target of rapamycin (mTOR) were detected in the pancreas of rat models of type 1 and type 2 Diabetes Mellitus (DM). As a model of type 1 DM, Komeda Diabetes-Prone (KDP) rat and control rat (KND rat) were used. As a model of type 2 DM, spontaneously diabetic Torii (SDT) rat and control rat (SD rat) were used. In each control rat, pancreatic islets were strongly positive for CD133, nestin, Bmi-1 and mTOR. In the KDP rat, CD133, nestin, Bmi-1 and mTOR positive cells did not show cellular mass like pancreatic islets. These positive cells were associated with invasion by mononuclear cells. Partial acinar cells showed increased immunoreactivity for CD133, Bmi-1 and mTOR, although nestin immunoreactivity was almost negative. In the SDT rat, CD133, nestin, Bmi-1 and mTOR positive cells also did not show cellular mass like pancreatic islets. These positive cells were associated with increased fibrillar element. Strong immunoreactivity for CD133, Bmi-1 and mTOR was observed in the pancreatic ductal epithelial cells and partial acinar cells compared with control rat. Nestin immunoreactivity was almost negative. Although β -cell denaturation in the pancreatic islet is present in the rat models of DM, pancreatic acinar and ductal epithelial cells express immunoreactivity for CD133, Bmi-1 and mTOR. Present study suggests that pancreatic acinar and ductal epithelial cells expressing CD133, Bmi-1 and mTOR are considerable as an alternative source of pancreatic stem/progenitors, even individuals who showed a diabetic condition.

Keywords: CD133; Nestin; Bmi-1; mTOR; Type 1 diabetes; Type 2 diabetes; KDP rat; SDT rat; Pancreas

Introduction

Type 1 and type 2 Diabetes Mellitus (DM) are the two main forms of DM. Both types are characterized by progressive β -cell failure in the pancreatic islet [1]. In type 1 DM, this is typically caused by an autoimmune assault against the β -cells, inducing progressive β -cell death [1]. The pathogenesis of type 2 DM is more variable, comprising different degrees of β -cell failure relative to varying degrees of insulin resistance [1]. Pancreatic islet transplantation is a promising method to restore functional islet β -cell mass for patients with DM [2]. Pancreatic islet transplants hold significant potential advantages over whole-gland transplants [3]. However, pancreatic islet transplantation is associated with several clinically relevant risks [4-6]. Because of the limited supply of human donor islets, it is critical that new strategies are explored as alternative renewable sources of transplantation [7]. Stem cells are characterized by extensive proliferation and multilineage differentiation capacity [8]. Mouse embryonic stem cells differentiate into cells of all three primary germ layers including endodermal cells that produce insulin *in vitro* [9]. Transplanted embryonic stem cells [9-11] and fetal pancreatic progenitor cells [12, 13] have been successfully used to treat DM

in murine models. Multipotent adult stem cells are harvested from bone marrow [14,15], adipose tissue [16] and umbilical cord and placenta [17]. Stem cells, as alternative islet cell sources, may be a valuable source for cell replacement therapy. In the pancreatic islet, CD133 [18,19], nestin [7,20-22] and B cell-specific Moloney murine leukemia virus integration site 1 (Bmi-1) [23] have been reported as molecular markers of pancreatic stem cells. Recently, Yilmaz et al. [24] show that reduction in caloric intake inhibits the nutrient-sensitive kinase mammalian target of rapamycin (mTOR) pathway and amplify intestinal stem cell numbers during caloric restriction.

Many experimental animal models have contributed to basic and clinical research of DM. Immunohistological analysis using these stem cell-related markers and animal models of type 1 and type 2 DM should provide useful information for understanding the molecular mechanism of DM. However, there is almost no information of these stem cell-related markers in the animal models. In the present study, we detected CD133, nestin, Bmi-1 and mTOR in the pancreas of KDP rat, SDT rat and each control rat. Pancreatic islets were strongly positive for CD133, nestin, Bmi-1 and mTOR in each control rat. CD133, nestin, Bmi-1 and mTOR positive cells in the pancreas did not

show cellular mass like pancreatic islets in both rat models of DM. In the KDP rat, partial acinar cells showed increased Immunoreactivity for CD133, Bmi-1 and mTOR. Strong Immunoreactivity for CD133, Bmi-1 and mTOR was shown in the pancreatic ductal epithelial cells and partial acinar cells. Nestin Immunoreactivity in these sites was not observed in both rat models.

Materials and Methods

Animals

Komeda diabetes-prone (KDP) rat: KDP rat developed by Komeda from Long-Evans Tokushima lean (LETL) rat at Tokyo Medical College in 1996. Komeda non-Diabetic (KND) rat was established simultaneously as a control [25]. The KDP rats (male, n=5) and KND rats (male, n=3) were available from Japan SLC (Shizuoka, Japan).

Spontaneously Diabetic Torii (SDT) rat: SDT rats from Japan Tobacco colonies were used in this study. Age-matched Sprague-Dawley (SD) rats were used as control animals. SDT rats (male, n=5) and SD rats (male, n=3) were purchased from CLEA Japan Inc. (Tokyo, Japan).

All animals were given water in drinking water *ad libitum* for 10 weeks, from 6 to 15 weeks of age. All animals were housed in suspended bracket cages and given a standard laboratory diet (EF, Oriental Yeast Co., Ltd, Tokyo, Japan) in a room with controlled temperature ($23 \pm 2^\circ\text{C}$), humidity ($55 \pm 10\%$) and lighting (12:12-h light-dark cycle).

All animal experiments were approved by the Committee on Animal Experimentation, Saitama Medical University (Permission No. 1214, 1396) and carried out in accordance with the "Guidelines for Animal Experimentation at Saitama Medical University".

Diabetes was defined as glycosuria positivity and blood glucose levels ≥ 250 mg/dL under *ad libitum* dietary conditions. Body weight of each rat was measured. Blood glucose levels were measured by a portable glucose meter (Glutest Neo alpha, Sanwa Kagaku Kenkyusho Co, Nagoya, Japan).

Immunohistochemistry

Tissues were fixed in 10% formalin/PBS, and paraffin sections (5 μm thick) were prepared. Streptavidin/peroxidase immunohistochemistry for CD133, nestin and Bmi-1 was performed. The specimens were treated with 0.3% H_2O_2 in methanol to block endogenous peroxidase activity and subsequently incubated with primary antibodies CD133 (1/100 dilution, rabbit polyclonal antibody; Abnova, Taipei, Taiwan), nestin (1/200 dilution, rabbit polyclonal antibody; Santa Cruz Biotechnology, Inc., Santa Cruz, CA), Bmi-1 (1/200 dilution, rabbit polyclonal antibody; Proteintech, Chicago, IL) and mTOR (1/200 dilution, rabbit polyclonal antibody; Abcam, Cambridge, UK) for 18 hours at 4°C . A biotinylated anti-rabbit immunoglobulin was added as a secondary antibody for 1 hour at room temperature. A horseradish peroxidase-labelled streptomycin-avidin complex was then used to detect the secondary antibody. Antibody binding was detected by staining with the chromogens 3,3'-diaminobenzidine (DAB). Cell nuclei were counterstained with hematoxylin. Some paraffin sections were stained with hematoxylin and eosin.

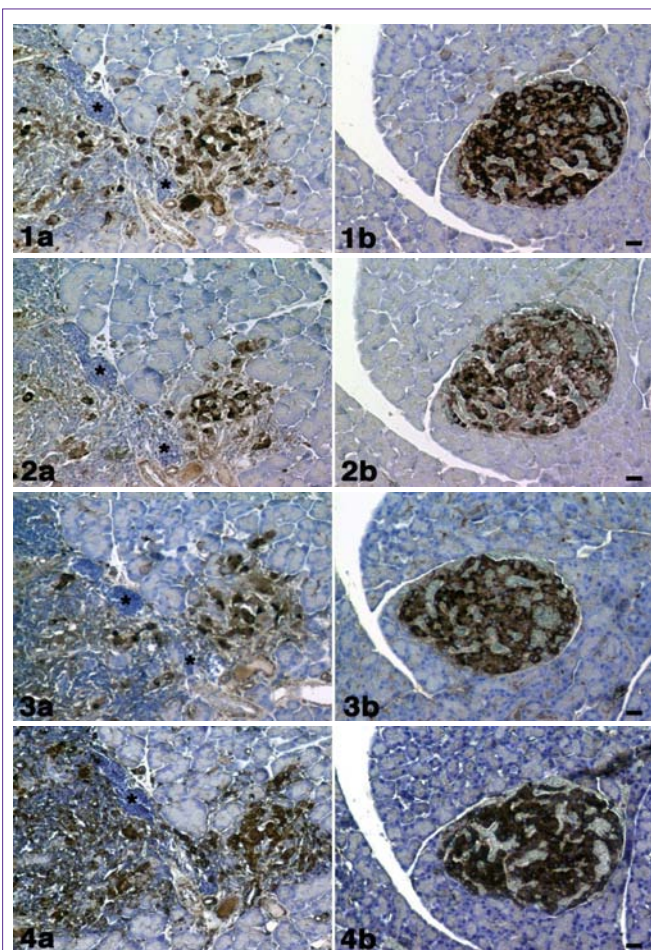
Results

Komeda diabetes-prone (KDP) rat (model of type 1 DM)

Pancreatic islets were positive for CD133, nestin, Bmi-1 and mTOR in the KND rat (control). In the KDP rat, CD133, nestin and Bmi-1 positive cellular mass observed in the control rat was not observed. CD133, nestin, Bmi-1 and mTOR positive cells were associated with invasion by mononuclear cells (Figures 1-4). Figures 1a-4a are the pancreas of a KDP rat (blood glucose level = 577 mg/dL). Figures 1b-4b are the pancreas of a KND rat (blood glucose level = 95 mg/dL). Partial acinar cells showed the increased Immunoreactivity for CD133, Bmi-1 and mTOR, although nestin immunoreactivity was almost negative (Figure 5).

Spontaneously Diabetic Torii (SDT) rat (model of type 2 DM)

Pancreatic islets were positive for CD133, nestin, Bmi-1 and mTOR in the SD rat (control). In the SDT rat, CD133, nestin, Bmi-1



Figures 1a-4a: KDP rat (model of type 1 DM).

1a) CD133, 2a) nestin, 3a) Bmi-1 and 4a) mTOR immunostaining. CD133, nestin, Bmi-1 and mTOR positive cells (dark brown) were observed with invasion of mononuclear cells (*).

Figures 1b-4b: KND rat (control).

1b) CD133, 2b) nestin, 3b) Bmi-1 and 4b) mTOR immunostaining. CD133, nestin, Bmi-1 and mTOR positive cells (dark brown) were observed in the pancreatic islet. There were no invasions of mononuclear cells. Cell nuclei were counterstained with hematoxylin. Scale bar = 20 μm .

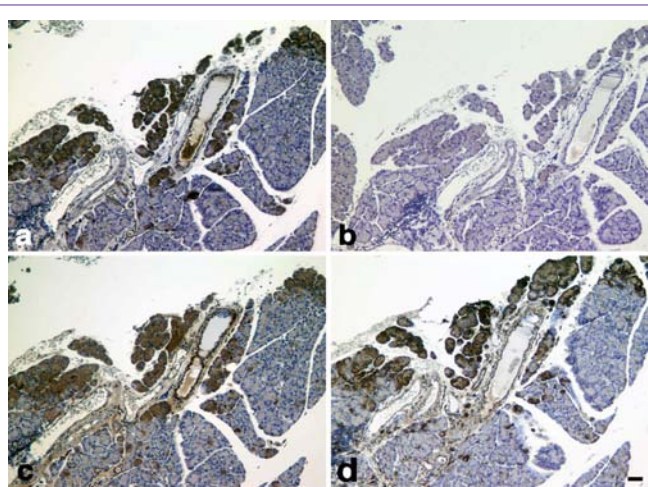


Figure 5: KDP rat (model of type 1 DM).

a) CD133, b) nestin, c) Bmi-1 and d) mTOR immunostaining. Partial acinar cells (dark brown) showed the increased immunoreactivity for CD133, Bmi-1 and mTOR (Figs. 5a, c, d). Cell nuclei were counterstained with hematoxylin. Scale bar = 40 μ m.

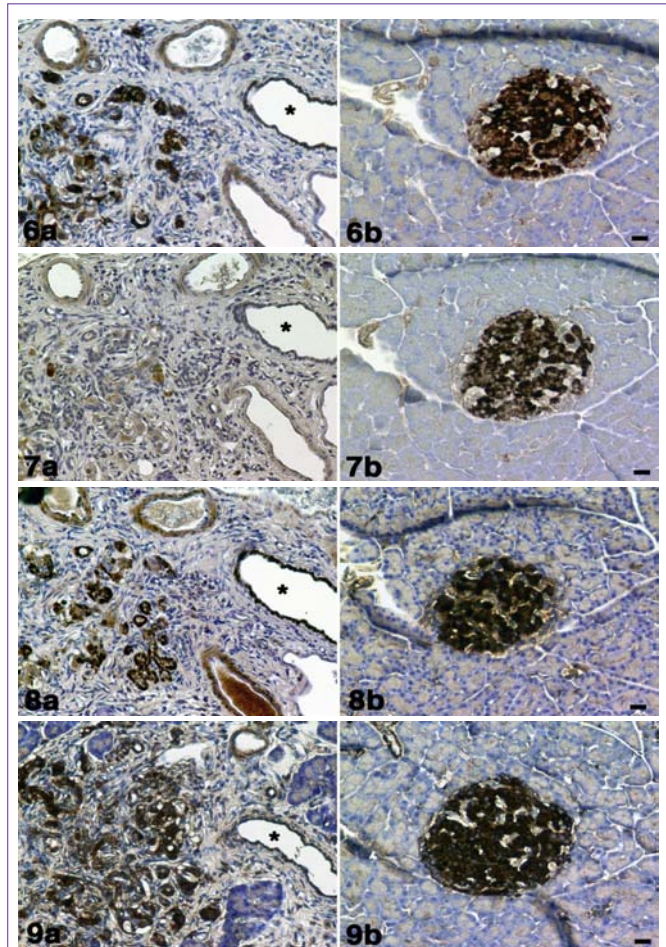
1 and mTOR positive cellular mass observed in the control rat was not observed. CD133, nestin, Bmi-1 and mTOR positive cells were associated with increasing fibrillar element (Figures 6-9). Decreased numbers of nestin positive cells were observed. These pathological changes in the pancreas of SDT rats differ from KDP rat, in which mononuclear cells infiltration into pancreatic islets is observed. Figures 6a-9a are the pancreas of a SDT rat (blood glucose level = 397 mg/dL). Figures 6b-9b are the pancreas of a SD rat (blood glucose level = 126 mg/dL). The strong immunoreactivity for CD133, Bmi-1 and mTOR was observed in the pancreatic ductal epithelial cells (Figures 6a, 8a, 9a) compared with SD rat (Figures 10a, c, d). Nestin immunoreactivity was almost negative (Figures 7b, 10b).

Discussion

There are many rat models of type 1 DM, see review [26] and type 2 DM, see review [27]. In the present study, we used KDP rat as a model of type 1 DM and SDT rat as a model of type 2 DM. The phenotypic features of the KDP rat resemble human type 1 DM because of autoimmune destruction of the pancreatic β -cells, rapid onset of overt DM with no sex difference, and lack of significant T-cell lymphopenia [25,28]. In the human type 2 DM, inflammation and fibrosis in the pancreatic islets have been described [29]. The phenotypic features of the SDT rat resemble human type 2 DM because of inflammation and fibrosis in the pancreatic islets [28,30,31].

Present study showed that pancreatic islets were CD133, nestin and Bmi-1 positive in both control rat. In the KDP rat, mononuclear cells infiltration into pancreatic islets is observed. Pancreatic β -cells are the target of an autoimmune assault in type 1 DM, with invasion of the islets by mononuclear cells in an inflammatory reaction termed "insulinitis," leading to loss of most β -cells after prolonged periods of disease [32]. On the other hand, SDT rat characterized by inflammation and fibrosis in and around the islets [28,30,31].

Present study also showed that increased immunoreactivity for CD133 and Bmi-1 was observed in the pancreatic ductal epithelial cells and acinar cells. Bmi-1 is essential for maintenance of adult stem cell self-renewal primarily due to its ability to repress the Ink4a/Arf



Figures 6a-9a: SDT rat (model of type 2 DM).

6a) CD133, 7a) nestin, 8a) Bmi-1 and 9a) mTOR immunostaining. CD133, nestin, Bmi-1 and mTOR positive cells (dark brown) were observed in the increasing fibrillar element. Decreased numbers of nestin positive cells were observed. Pancreatic ducts (*). Immunoreactivity for CD133 was moderately positive in the epithelial cells of pancreatic ducts. Immunoreactivity for nestin was almost negative. Immunoreactivity for Bmi-1 was strongly positive in the epithelial cells of pancreatic ducts. Immunoreactivity for mTOR was moderately positive in the epithelial cells of pancreatic ducts.

Figures 6b-9b: SD rat (control).

6b) CD133, 7b) nestin, 8b) Bmi-1 and 9b) mTOR immunostaining. CD133, nestin, Bmi-1 and mTOR positive cells (dark brown) were observed in the pancreatic islet. There was no increasing fibrillar element. Cell nuclei were counterstained with hematoxylin. Scale bar = 20 μ m.

locus [33-36]. Dhawan et al. [23] reported that Bmi-1-dependent regulation of the Ink4a/Arf locus plays a critical role in regulating pancreatic β -cell proliferation during aging and regeneration. On the other hand, the biological function of CD133 is not well understood. Recently, we reported the co-localization of CD133 with F-actin. It is strongly suggested that CD133 is involved in cell migration [37]. We also reported that CD133 was mainly localized to plasma membrane in the late stages of cultivation. After treatment with CD133 antibody, cell-cell contact was decreased, and apoptosis was increased. It is suggested that CD133-mediated regulation may be required for cell survival [38]. Increased immunoreactivity for CD133 and Bmi-1 may be attributed by the denaturation of β -cells. Insulin and Insulin-Like Growth Factors (IGFs) are prominent inducers of mTOR activity [39,40]. In the diabetic rats used in this study, reduced insulin secretion

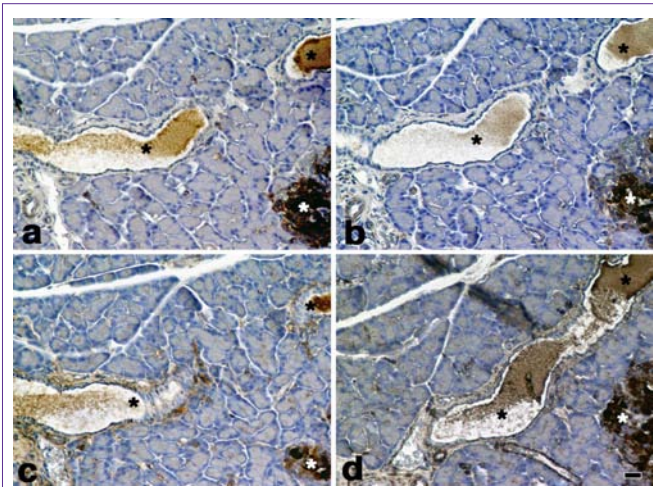


Figure 10: SD rat (control). a)CD133,b) nestin, c)Bmi-1 and d) mTOR immunostaining. Pancreatic ducts (*). Immunoreactivity for CD133 was negative or slightly positive in some epithelial cells of pancreatic ducts. Immunoreactivity for nestin was almost negative. Immunoreactivity for Bmi-1 was weekly positive in the epithelial cells of pancreatic ducts. Immunoreactivity for mTOR was slightly positive in the epithelial cells of pancreatic ducts. Pancreatic islets (white *).Cell nuclei were counterstained with hematoxylin. Scale bar= 20 μ m.

by the denaturation of β -cells can cause the decreased activity of mTORC1. In SDT rats, blood insulin concentration tended to be lower than in normal SD rats even before the onset of diabetes, and marked hypoinsulinemia developed after the onset of hyperglycemia [31,41,42]. Matsubara et al. [43] noted that the percentage of CD133+ cells in xenografts from pancreatic cells showed the tendency to increase after rapamycin (mTOR inhibitor) treatment, which was not shown *in vitro*. They noted some differences between *in vitro* and *in vivo*. In the present study, increased immunoreactivity of mTOR was observed in the partial acinar and pancreatic ductal epithelial cells. Increased immunoreactivity of mTOR is not attributed insulin. Influence of hyperglycemia must also be considered. mTOR pathway is involved in sensing of nutrient availability and modulation of insulin action *in vivo* [44]. Um et al. [45] and Khamzina et al. [46] indicate that the activity of the mTOR pathway is increased in rodent models of obesity and that mice deficient in S6 kinase (S6K), a downstream effector of mTOR, are protected against diet-induced insulin resistance. Further study is required to elucidate the issue. Nestin expression was significantly decreased in the heart of diabetic rats [47]. Liu et al. [48] reported that nestin expression in the glomeruli was gradually decreased with the development of diabetic nephropathy. On the other hand, the expression of cyclin-dependent kinase 5 (Cdk5) was increased. They suggest that Cdk5 may relate degradation of nestin.

Koblas [19] used human pancreatic acinar tissue that remains after islet isolation and is discarded. They found that population of human CD133-positive pancreatic cells contains endocrine progenitors expressing neurogenin-3 and cells expressing human telomerase, ABCG2, Oct-3/4, Nanog, and Rex-1, markers of pluripotent stem cells. These cells were able to differentiate into insulin-producing cells *in vitro* and secreted C-peptide in a glucose-dependent manner [19]. Ohshima et al. [49] isolated mouse pancreatic ductal progenitor cells expressing CD133 by flow cytometric cell sorting. Bonner-Weir

et al. [50] showed that human pancreatic ductal epithelial cells could be expanded and differentiated into glucose responsive islet tissue *in vitro* given insulin. Ramiya et al. [51] isolated murine pancreatic ductal epithelial cells into culture, and induced them into functional islets containing alpha, beta and delta cells. These resulting islets showed temporal changes in mRNA transcripts for islet cell associated differentiation markers, responded to glucose challenge *in vitro* and reversed insulin-dependent diabetes after being implanted into non-obese diabetic mice. Multipotent stem/progenitors are present in peribiliary glands of extrahepatic biliary trees from humans of all ages and in high numbers in hepato-pancreatic common duct, cystic duct, and hilum [52]. It was shown that extrahepatic bile ducts in mice have β -cells, with secretory granules that are immunoreactive for insulin and that exhibit glucose-stimulated insulin secretion [53]. Their histological studies indicate that the β -cells form directly from the bile duct epithelium in late embryogenesis.

Pancreatic ductal epithelial cells and acinar cells expressing CD133, Bmi-1 and mTOR are considerable as an alternative source of stem/progenitors for targeted therapies and regenerative medicine for diabetes.

Conclusion

Stem cell-related markers (CD133, nestin and Bmi-1) and mTOR were detected in the pancreas of rat models of type 1 and type 2 DM. Although β -cell denaturation in the pancreatic islet was present in both rat models of DM, pancreatic acinar and ductal epithelial cells express CD133, Bmi-1 and mTOR. Nestin immunoreactivity in these sites was not observed in both rat models. Pancreatic acinar and ductal epithelial cells expressing CD133, Bmi-1 and mTOR are considerable as an alternative source of stem/progenitors, even individuals who showed a diabetic condition.

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