Islet amyloid deposits preferentially in the highly functional and most blood-perfused islets

Sara Ullsten¹, Sara Bohman¹, Marie E Oskarsson¹, K Peter R Nilsson², Gunilla T Westermark¹ and Per-Ola Carlsson¹,³

¹Department of Medical Cell Biology, Uppsala University, Uppsala, Sweden
²Department of Chemistry, Linköping University, Linköping, Sweden
³Department of Medical Sciences, Uppsala University, Uppsala, Sweden

Abstract

Islet amyloid and beta cell death in type 2 diabetes are heterogeneous events, where some islets are affected early in the disease process, whereas others remain visibly unaffected. This study investigated the possibility that inter-islet functional and vascular differences may explain the propensity for amyloid accumulation in certain islets. Highly blood-perfused islets were identified by microspheres in human islet amyloid polypeptide expressing mice fed a high-fat diet for three or 10 months. These highly blood-perfused islets had better glucose-stimulated insulin secretion capacity than other islets and developed more amyloid deposits after 10 months of high-fat diet. Similarly, human islets with a superior release capacity formed more amyloid in high glucose culture than islets with a lower release capacity. The amyloid formation in mouse islets was associated with a higher amount of prohormone convertase 1/3 and with a decreased expression of its inhibitor proSAAS when compared to islets with less amyloid. In contrast, levels of prohormone convertase 2 and expression of its inhibitor neuroendocrine protein 7B2 were unaltered. A misbalance in prohormone convertase levels may interrupt the normal processing of islet amyloid polypeptide and induce amyloid formation. Preferential amyloid load in the most blood-perfused and functional islets may accelerate the progression of type 2 diabetes.

Introduction

Wide heterogeneities among and within pancreatic islets have previously been identified. Beta cells vary in size, granularity, proliferation, functionality and maturation (1, 2, 3). Some subsets of beta cells are more adaptable during pregnancy and metabolic challenges, and beta cells have also shown age-dependent variations (4, 5, 6). On an inter-islet level, we have previously described two subpopulations of islets distinguished by their differences in blood supply; one low blood-perfused, low-oxygenated population with dormant metabolic activity (7), and another population with a high blood supply exhibiting superior functionality and beta cell proliferation indexes (8). Noteworthy, the latter population of high-functioning islets exhibited an increased sensitivity to hypoxia and cytokines leading to reduced survival upon stress (9).

Also, islet lesions have been shown to be of a heterogeneous nature and the beta cell loss in type 2 diabetes (T2D) is not consistent between islets or between different parts of the pancreas (10). Islet amyloid, a clinical finding seen in approximately 90% of T2D patients, certainly affects islets in this irregular pattern; some islets may contain large amounts of amyloid in contrast...
Materials and methods

Experimental animals

Male mice deficient in mouse IAPP (mIAPP), but expressing hIAPP (mIAPP<sup>−/−</sup>, hIAPP<sup>+/−</sup>) (28) had free access to water and standard chow. Throughout the experimental studies, the animals additionally had free access to lard (HFD) (28). Experiments were performed after short-term (3-months) or long-term (10-months) HFD (28). All experimental procedures were approved by the Ethical Committee at Uppsala University.

Detection of highly blood-perfused islets

The animals were anesthetized with i.p. injections of avertin (Sigma-Aldrich) and placed on a heating pad to maintain body temperature. Polyethylene catheters were inserted into the ascending aorta for continuous blood pressure measurement and subsequent microsphere injection and into the left femoral artery for blood sampling. After the surgical procedure, the animals were allowed to stabilize their blood pressure for 10 min before injection of black (10 µm, E-Z Trac Ultraspheres; IMT, Stason Labs., Irvine, CA, USA) or fluorescent microspheres (10 µm, Fluosphere Polystyrene Microspheres; Molecular Probes), into the ascending aorta. A reference blood sample was obtained by free flow from the femoral artery. It was collected for 60 s, with a start 5 s before microsphere injection, to enable quantitative determination of blood flow (8, 9). The pancreas was thereby immediately retrieved for islet isolation or immunohistochemistry.

Islet isolation

Mouse islets were isolated by collagenase digestion, hand-picked and incubated in groups of 100 overnight at 37°C (air/CO<sub>2</sub>, 95:5) in 5 mL RPMI-1640 medium (Sigma-Aldrich) supplemented with 2 mmol/L l-glutamine, 11.1 mmol/L glucose, 10% (v/v) fetal calf serum (FCS) and 0.1 mg/mL streptomycin (Roche Diagnostics Scandinavia) (29). The islets were thereafter dichotomously sorted in a fluorescence microscope. Microsphere-containing islets were size-matched to control islets without microspheres from the same mouse (8, 9).

Amyloid evaluation in mouse islets

Islets of animals fed 10-month HFD were following their isolation fixed in 10% (v/v) buffered formalin and...
embedded in paraffin. Consecutive sections (5 µm) were stained for amyloid with Congo red (Sigma-Aldrich) (30). Sections of size-matched islets with and without microspheres were photographed in a fluorescence microscope and the amyloid area was calculated as percentage of the total area of the islet. Image analysis was performed with ImageJ (NIH).

Location of microsphere-containing islets

From 3-month HFD microsphere-injected animals, the pancreas was oriented with a suture in the tail part, immobilized and consecutively sectioned (10 µm) perpendicularly to the longitudinal axis of the pancreas. Microsphere-containing islets were identified in a light microscope after hematoxylin–eosin staining. Whole section scanning of whole pancreas was performed by Aperio Scanscope AT (Leica Biosystems, Nussloch, Germany). The distance from the center of each microsphere- or non-microsphere-containing islet to the pancreatic center, defined as the origo of an image fitting the whole pancreatic section, was calculated with Aperio ImageScope software (Leica Microsystems, Wetzlar, Germany).

Mouse islet glucose-stimulated hormone release

Islets isolated from hiAPP animals fed 3-month HFD were sorted based on their microsphere content or not and incubated at low (1.67 mmol/L) and high glucose (16.7 mmol/L) concentrations for 60 min, respectively (31). The supernatants from both incubations were collected and analyzed for mouse insulin and human IAPP with ELISA (Mercodia, Uppsala, Sweden and EZHAR1-5,2K, Merck Millipore, respectively). Stimulation index was calculated accordingly (insulin release in high glucose)/(insulin release in low glucose). The islets were harvested to measure total hormone content and to ensure equal islet size in the groups by DNA measurement (Quant-iT PicoGreen dsDNA assay kit, Molecular Probes).

Mouse islet gene expression

RNA was extracted from isolated islets of animals fed 3-month HFD by the use of RNeasy Plus Mini Kit (Qiagen) followed by first-strand cDNA synthesis by SuperScript III First-Strand Synthesis SuperMix (Invitrogen). qPCR was performed using a Light Cycler 480 (Roche Diagnostics) and Light Cycler FastStart DNA Master PLUS SYBR Green I kit (Roche Diagnostics). Gene expression was normalized to the reference gene Tata-box-binding protein (TBP). For primer information, see Supplementary Table 1 (see section on supplementary data given at the end of this article).

Prohormone convertase (PC) protein levels in mouse islets

Pancreas sections obtained from animals fed 3-month HFD were incubated with monoclonal mouse PC1/3 and PC2 antibodies (22) together with an insulin antibody (Fitzgerald, Acton, MA, USA). Heat-induced antigen retrieval (20 min) in 0.01 mol/L sodium citrate was performed for PC2 staining only. Fluorescent anti-mouse 488 (1:200, Jackson ImmunoResearch Laboratories) and anti-guinea pig 555 (1:1000 Molecular Probes) were used for primary antibody detection. To quantify PC protein levels on sections, images were acquired using a confocal laser scanner microscope (Zeiss LSM780) and analyzed with ImageJ (NIH).

Heparan sulfate quantification in mouse islets

In pancreases obtained from 3-month HFD animals, heparan sulfate (HS) glycosaminoglycans were identified with Alcian blue CEC staining (0.65 mol/L MgCl₂, pH 5.8, Sigma-Aldrich) (32). Reciprocal staining intensity was analyzed with ImageJ (NIH).

Human islets

Human islets from brain-dead, heart-beating donors, provided by the Nordic Network for Clinical Islet Transplantation (22–42 islets/donor, for donor data, Table 1), were individually cultured in CMRL 1066 supplemented with 10% (v/v) FCS, 2 mmol/L L-glutamine, 0.1 mg/mL streptomycin (Roche Diagnostics Scandinavia) and 5.5 or 20 mmol/L glucose. Experiments involving human islets were approved by the regional ethical board in Uppsala, Sweden.

<table>
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<tr>
<th>Donor</th>
<th>Sex</th>
<th>Age</th>
<th>BMI (kg/m²)</th>
<th>HbA1C % (mmol/mol)</th>
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</tr>
<tr>
<td>Donor 2</td>
<td>F</td>
<td>59</td>
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<td>5.8 (40)</td>
</tr>
<tr>
<td>Donor 3</td>
<td>F</td>
<td>73</td>
<td>22.3</td>
<td>5.8 (40)</td>
</tr>
<tr>
<td>Donor 4</td>
<td>F</td>
<td>69</td>
<td>23.9</td>
<td>6.5 (47)</td>
</tr>
</tbody>
</table>

Data received from the Nordic Network for Clinical Islet Transplantation. BMI, body mass index; HbA1C, glycated hemoglobin.
Human islet insulin release

Prior to high glucose culture (day 0), insulin release experiments were performed on individual islets to evaluate their functionality. Islets were pre-incubated in 1.67 mmol/L glucose for 30 min. Glucose-stimulated insulin release of single islets was then determined by incubation of them in 1.67 and 16.7 mmol/L glucose for 60 min, respectively. The supernatants were analyzed by an Ultrasensitive Insulin ELISA (Mercodia). Stimulation index was calculated accordingly: (insulin release in high glucose)/(insulin release in low glucose).

Islet amyloid quantification in human islets

Human islets were cultured individually (~30 islets/donor) in 20 mmol/L glucose for 18 days to stimulate IAPP secretion and subsequent amyloid formation (27). To visualize islet amyloid, the live islets were incubated with the amyloid-specific oligothiophene p-FTAA (30 µmol/L) (33, 34) for 60 min and thereafter imaged in a fluorescence microscope at 488 nm excitation. Image analysis was performed with ImageJ (NIH).

Statistical analysis

All values are given as mean±S.E.M. Parametrical data were analyzed by unpaired or paired Student's t-test, and the distribution of amyloid between islets were calculated by a chi-square test. Gene expression analysis was performed on the ΔCT values for each gene. *P<0.05 was considered significant.

Results

Animal characteristics and blood flow

hIAPP+/+mIAPP−/− mice increased in weight during HFD treatment and weighed 33±1 g (n=23) and 44±1 g (n=7) after 3 and 10 months on HFD, respectively. The animals were all normoglycemic (6.7±0.2 mmol/L; n=30). In the experiments, each microsphere corresponded to a blood flow of ≥0.21±0.02 µL/min (n=30).

Amyloid in mouse islets

After 10-month HFD, amyloid deposits (≥0.1% islet area) were more frequent in islets with microspheres than in those without microspheres (n=7, Fig. 1A, B and C).

The amyloid deposits were also twice as extensive (percentage of islet area) in the highly blood-perfused islets when compared to size-matched islets without microspheres (n=7, Fig. 1D).

Location of highly blood-perfused islets

Islets with microspheres were found throughout the pancreas. Islets with and without microspheres were similarly distributed in the central and peripheral parts of the gland (1.39±0.18 vs 1.34±0.17 mm to center, respectively, n=4).

Glucose-stimulated hormone release

No difference in islet size was measured between the investigated islets with and without microspheres (81.2±13.5 vs 70.2±12.1 ng DNA/islet, respectively, n=6 animals). There was neither any difference in insulin or IAPP content between the two groups of islets (6218±1136 vs 6891±1817 fmol insulin/islet and 377±80 vs 375±84 fmol IAPP/islet in islets with- and without microspheres, respectively, n=6). There was no difference between the two groups of islets in insulin release at a low glucose concentration (1.67 mmol/L, but in response to a high glucose concentration (16.7 mmol/L), the microsphere-containing islets secreted ~30% more insulin than the size-matched islets without microspheres (Fig. 1E). When calculating the glucose stimulation index (ratio of insulin release in high and low glucose concentration), islets with microspheres had a 44% higher ratio (n=6, Fig. 1F). No difference was however seen between the groups in IAPP release at either a low or a high glucose concentration (Fig. 1G). Islets with microspheres therefore had a ratio of released IAPP/insulin in response to glucose that was ~30% lower than other islets (Fig. 1H).

Amyloid in human islets

Little or no amyloid (<1% of islet area) was found at day 0 in the islets obtained from four different donors. After functional test of single islets, the islets were cultured individually for 18 days in high glucose to allow amyloid formation. At day 18, the islets had developed amyloid covering ~4% (range 0–30%) of the islet area. The islets were thereafter divided into two groups; islets with little or no (<1% of islet area) amyloid (n=58) and islets with amyloid (>1% of islet area) (n=59). The group of islets that formed amyloid after 18 days of high glucose culture was
found to have an ~50% better glucose-stimulated insulin release when measured prior to the high glucose culture (Fig. 2A). This was observed in islets from all four donors (Fig. 2B). In contrast, there was no visible amyloid in any of the islets before high glucose culture (day 0) (Fig. 2C, D, E, F, G, H, I and J).

**ProlIAPP processing enzymes**

No difference in gene expression of the two prohormone convertases PC1/3 and PC2 (n=6, Fig. 3A and B) and the processing enzyme Carboxypeptidase E (CPE) (fold change 0.73 ± 0.20 in islets with microspheres, n=5) could be observed between islets with or without microspheres. However, the expression of the neuroendocrine peptide precursor proSAAS was ~50% lower in the microsphere-containing islets (n=5, Fig. 3C). 7B2, another neuroendocrine protein, did not differ between the two groups (n=6, Fig. 3D). PC1/3 immunofluorescence co-localized with insulin staining and was ~17% higher in islets with microspheres (n=4, Fig. 3E, G, H, I, J, K and L). In contrast, the intensity of PC2 staining in insulin-positive cells did not differ between the islets with and without microspheres (n=4, Fig. 3F, M, N, O, P, Q and R).

**Highly sulfated glycosaminoglycans**

The intensity of HS staining did not differ between the islets with or without microspheres (n=4, reciprocal intensity 136.6 ± 3.1 vs 131.2 ± 6.2 AU, respectively).

**IAPP-degrading enzymes**

A higher gene expression of the two matrix metalloproteinases MMP2 and MMP9 was observed in the islets with microspheres (n=5–6, Fig. 4A and B). No differences could be detected in the expression of the inhibitors tissue inhibitor of matrix metalloproteinases (TIMP) 1–3 between islet with or without microspheres (n=6, Fig. 4C, D and E). There was neither any difference in the expression of insulin-degrading enzyme (IDE) (n=6, fold change 1.54 ± 0.51 in islets with microspheres) and neprilysin (n=4, fold change 1.22 ± 0.67 in islets with microspheres).

**ER stress induced gene expression**

Gene expression of C/EBP-homologous protein (CHOP) (n=5, fold change 1.21 ± 0.37 in islets with microspheres)
and immunoglobulin-binding protein (BiP) \((n=5, \text{ fold change } 1.21 \pm 0.62 \text{ in islets with microspheres})\) did not differ between the islets with or without microspheres.

**Discussion**

The present study describes increased susceptibility for islet amyloid formation in a subpopulation of highly blood-perfused and functionally active islets of hIAPP mice. In addition, a subgroup of human islets with a better insulin release capacity developed more amyloid during high glucose exposure. Increased amyloid in the highly blood-perfused functionally active islets of hIAPP mice was associated with an imbalance in PC protein levels. This imbalance with the following incapacity of processing proIAPP to mature IAPP has previously been described to be of great importance for the development of amyloid deposits \((22, 35)\).

Identification of highly blood-perfused, highly functional islets has previously been performed by the microsphere technique \((8, 9, 36, 37)\). In order to be able to study the tendency for amyloid formation in these islets, we used mice expressing hIAPP due to the non-amyloidogenic nature of rodent IAPP \((28)\). These mice lack mIAPP to avoid interference with amyloid fibril formation. Since amyloid formation does not develop spontaneously in the hIAPP mouse model, a trigger of e.g. HFD is needed \((28)\). In order to investigate visible amyloid deposits, we used mice fed a diet high in fat for 10 months to allow sufficient time for the development of the deposits. Also, we tried to identify mechanisms preceding the fibrillation process, and for these experiments, we used animals that had been fed a HFD for three months.

After 10 months of HFD, amyloid was detected in islets from all animals. However, when comparing islets, we observed preferential amyloid development in the islets with a high blood perfusion as identified by...
microspheres. The amount of amyloid was also twice as extensive in these islets when compared to the group with lower blood perfusion.

Consistent with our previous study in rat (8), islets with high blood flow as identified by microspheres had an approximately 30% higher insulin release than other islets in response to high glucose. However, when measuring IAPP secretion from the same islets, no difference could be seen between the two groups in either high or low glucose concentrations.

Since the glucose stimulation index, a measure of insulin secretory capacity, was particularly augmented in the highly blood-perfused mouse islets marked by microspheres, we decided to use this index for studies on human islets as the microsphere technique is restricted to experimental animal models and cannot be used to identify highly perfused human islets. The glucose stimulation index of each single human islet was evaluated at day 0 and the islets were subsequently cultured in high glucose medium for 18 days to induce IAPP secretion and amyloid formation (27). After 18 days of high glucose culture, extensive amyloid was seen in numerous islets (amyloid deposits range 0–30% of islet area). When the islets were divided into no or low (<1%) or high (>1%) amyloid content, islets that developed amyloid were found to have an approximately 50% higher glucose stimulation index prior to the incubation irrespective of the donor.

To further investigate the mechanisms of amyloid formation in the highly blood-perfused, highly functional mouse islets, we studied the expression of the ER stress markers CHOP and BiP as ER stress has been described to be associated with amyloid formation in some (38), but not all studies (39). In this study, we did not see any indications of ER stress in the highly functional islets prone to develop amyloid. Another study has described a preferential location of human islets containing...
We did, in our studies in mice, however, not find any regional differences of the position of microsphere-containing islets.

Processing deficiencies of proIAPP has been suggested to contribute to islet amyloid development in T2D (41, 42, 43, 44). An increased proinsulin-to-insulin ratio with a major proportion of partially processed des-31,32 proinsulin, a proinsulin intermediate partially processed by PC1/3 but not PC2, has also been identified in such patients (45). To investigate if proIAPP processing was altered in the highly blood-perfused islets, we performed gene expression analysis of PC1/3 and PC2, the two major processing enzymes of both IAPP and insulin precursor molecules. Since the regulation of the two prohormone convertases also has been shown to occur on a post-transcriptional level (46, 47), the levels of insulin, PC1/3 and PC2 were evaluated by immunohistochemistry as well. Although the intensity of insulin and PC2 did not differ, the PC1/3 intensity was increased with approximately 17% in islets containing microspheres. Although we were unable to investigate the activity of either PC1/3 or PC2, we also observed a decrease in the gene expression of the endogenous inhibitor of PC1/3, proSAAS, which couples to increased activity of PC1/3 (48). The corresponding inhibitor of PC2, 7B2, was unchanged.

PC1/3 appears to be more important for proinsulin processing than PC2, but both enzymes can process both the COOH- and NH2-terminal of proinsulin, although not as efficient as the two enzymes combined (49, 50). In contrast, proIAPP requires PC2 for processing at the NH2-terminus (41, 49). Interestingly, the proIAPP intermediate N-IAPP with an unprocessed NH2-terminal has been suggested to be more amyloidogenic than mature IAPP (42) and has been identified in human amyloid deposits (43, 44). We were not able to measure N-terminal intermediate proIAPP, but in diabetic mice transplanted with mouse islets expressing human proIAPP, lack of PC2 in the islet transplants caused rapid graft failure with recurrence of hyperglycemia (51). The authors concluded that an increased production of prefibrillar proIAPP aggregates is plausible to have contributed.

N-IAPP has been described to have an increased affinity to HS, which is located at the cellular surface. This is suggested to cause a locally increased concentration of the amyloidogenic processing intermediate, which may seed amyloid formation (52). As HS is important for islet function (24), we hypothesized that highly functional islets may have an increased amount of HS facilitating the observed increase in amyloid formation. We did, however, not find any differences in HS intensity between the two groups of islets. Nevertheless, an increased amount of high-affinity N-IAPP in the microsphere-containing islets may by itself seed the process.

Amyloid peptide Aβ has been described to increase transcription of vascular MMP9 (53). This could be of importance for amyloid formation, since both MMP2 and MMP9 have shown the ability to degrade Aβ to soluble and less fibrillogenic fragments (54). MMP9, but not MMP2, has also been described to degrade IAPP (55). We could detect an increase in both MMP2 and MMP9 expression in the microsphere-containing islets, a feature which might be a response to an increased fibrillation process in these islets. However, the effects of islet amyloid on MMP2 and MMP9 expression or the degradation capacity of the two matrix metalloproteinases have not been studied. An increased
expression may simply reflect the specific phenotype of the highly vascularized islets (56). In conclusion, in the present study, we have found that the most functional islets of hIAPP mice and humans are more prone to develop amyloid than other islets. This may be explained by the observed imbalance between the two prohormone convertases PC1/3 and PC2 in these islets. As the subpopulation of islets that was affected by amyloid was the most functional one, the amyloid formation may result in unfavorable effects on long-term glucose homeostasis, which per se increases the metabolic demand and may speed up and worsen disease development.

Supplementary data
This is linked to the online version of the paper at http://dx.doi.org/10.1530/EC-17-0148.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

Funding
The Swedish Research Council (55X-15043), the Swedish Child Diabetes Foundation, the Swedish Diabetes Foundation, Diabetes Wellness Sweden, the Novo Nordisk Foundation, and the strategic grant Excellence of Diabetes Research in Sweden (EXODIAB).

Author contribution statement
S U, S B, M O, P N, G T W and P O C designed the study and interpreted data. S U, S B and M O acquired and analyzed data. S U and P O C wrote the manuscript and S B, M O, P N and G T W revised the manuscript critically for intellectual content. All authors approved the final version of the paper.

Acknowledgements
Human islets were generously provided through the Nordic Network for Clinical Islet Transplantation. The skilled technical assistance of My Quach, Petra Franzen, Astrid Nordin, Birgitta Bodin and Lisbeth Sagulin is gratefully acknowledged.

References


41 Marzban L, Radke CJ, Steiner DF, Haataja L, Halban PA & Verchere CB. Impaired NH2-terminal processing of human proislet amyloid polypeptide by the prohormone convertase PC2 leads to amyloid formation and cell death. Diabetes 2006 55 2192–2201. (doi:10.2337/db05-1566)


Received in final form 13 July 2017
Accepted 8 August 2017
Accepted Preprint published online 8 August 2017