Differential O- and Glycosphingolipid Glycosylation in Human Pancreatic Adenocarcinoma Cells With Opposite Morphology and Metastatic Behavior

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Changes in the glycosylation profile of cancer cells have been strongly associated with cancer progression. To increase our insights into the role of glycosylation in human pancreatic ductal adenocarcinoma (PDAC), we performed a study on O-glycans and glycosphingolipid (GSL) glycans of the PDAC cell lines Pa-Tu-8988T (PaTu-T) and Pa-Tu-8988S (PaTu-S). These cell lines are derived from the same patient, but show an almost opposite phenotype, morphology and capacity to metastasize, and may thus provide an attractive model to study the role of glycosylation in progression of PDAC. Gene-array analysis revealed that 24% of the glycosylation-related genes showed a ≥ 1.5-fold difference in expression level between the two cell lines. Subsequent validation of the data by porous graphitized carbon nano-liquid chromatography coupled to a tandem ion trap mass spectrometry and flow cytometry established major differences in O-glycans and GSL-glycans between the cell lines, including lower levels of T and sialylated Tn (sTn) antigens, neoexpression of globosides (Gb3 and Gb4), and higher levels of gangliosides in the mesenchymal-like PaTu-T cell s compared to the epithelial-like PaTu-S. In addition, PaTu-S cells demonstrated a significantly higher binding of the immune-lectins macrophage galactose-type lectin and galectin-4 compared to PaTu-T.

In summary, our data provide a comprehensive and differential glycan profile of two PDAC cell lines with disparate phenotypes and metastatic behavior. This will allow approaches to modulate and monitor the glycosylation of these PDAC cell lines, which opens up avenues to study the biology and metastatic behavior of PDAC.

Keywords: glycosylation, gene array analysis, glycosyltransferase, O-glycosylation, glycosphingolipid (GSL) glycans, pancreatic ductal adenocarcinoma

INTRODUCTION

Pancreatic adenocarcinoma is an important cause of cancer-related death in the western world. Especially, pancreatic ductal adenocarcinoma (PDAC), accounting for 90% of the pancreatic adenocarcinoma, has a very poor prognosis. The 1-year survival rate is reported to be 20%, and the 5-year rate <5% for all stages of pancreatic cancer combined (1). These low survival rates are
mainly due to a lack of effective diagnostic strategies, the rapid progression of pancreatic cancer at an early stage, and unreliable targets for therapeutic intervention. It is of great importance to develop reliable diagnostic biomarkers and to understand molecular and cellular mechanisms in PDAC metastasis.

Glycosylation is one of the most common and complex post-translational modifications of proteins, and glycans also occur conjugated to lipids (2). Changes in glycosylation are involved in the regulation of many cancer-related processes such as signaling, cell adhesion, tumor proliferation, invasion, metastasis, and angiogenesis (2–5). Importantly, cancer-associated glycans also provide a set of diagnostic biomarkers and potential targets for therapeutic strategies (2, 5–7). For example, CA 19-9 which is an antigen carrying the glycan epitope siaI Lewis A (sLe^A) is a widely used biomarker in clinical diagnosis of PDAC (4, 8). The recently reported non-invasive diagnostic biomarker for early stages of pancreatic cancer glypican-1 is also a cell surface proteoglycan (9), indicating again the importance of glycosylation in PDAC. Although, many glycan changes have been observed in pancreatic cancer [reviewed in (4)], the causes as well as the consequences for cancer progression are still incompletely understood.

Many factors can contribute to changes in cancer glycosylation, including regulation and localization of the enzymes that are involved in their biosynthesis [glycosyltransferases (GTs) and glycosidases], enzymes synthesizing glycan precursors, nucleotide sugar donors and transporters, and alteration of the peptide backbone (2, 10, 11). Among them, GTs catalyze the transfer of a monosaccharide to the initiating and growing glycans, thereby showing a key role in the regulation of glycosylation and aberrant biosynthesis of core-glycans in cancer (12, 13). Importantly, changes in enzymes involved in the modification and termination like fucosyltransferases (FUTs) (14) and sialyltransferases (STs) (15, 16) have been associated with progression of invasive malignancies by increasing apoptosis resistance, affecting tumor growth, metastasis formation, or resistance to therapy.

For pancreatic cancer, several aberrations in N- and O-glycosylation and their regulation have been described Munkley (4), Taniuchi et al. (13), Radhakrishnan et al. (17), Hofmann et al. (18), and Chugh et al. (19). To further understand the expression and regulation of glycosylation in metastasis of pancreatic cancer, we performed a comprehensive glycosylation study, including O- and GSL-glycosylation analysis starting from differential gene expression and activity of GTs, to the structural and relative quantitative analysis of their glycan end-products, in two closely related pancreatic cell lines. These two cell lines, originate from a single liver metastasis of a PDAC (20) and display disparate grades of differentiation and have distinct properties that are relevant for PDAC progression. PaTu-8988S (PaTu-S) cells exhibit an epithelial-like phenotype, with polarity, E-cadherin expression, preserved cell-cell contacts, and low migratory properties. Pa-Tu-8988T (PaTu-T) cells are spindle-like with increased cell projections and minimal cell-cell contacts (21). They have a more mesenchymal phenotype and are highly migratory in vitro. Most remarkable is the much stronger metastatic potential of PaTu-T, compared to PaTu-S cells in a zebrafish model (21–23). Therefore, these sister cell lines are an attractive model to study the role of glycosylation in the cell biology of human PDAC.

Using glyco-gene microarray analysis we show that a quarter of the ~1,171 glycosylation-related genes displayed differential expression in PaTu-S and PaTu-T. A large variation in the N-glycosylation signature between these two cell lines has been previously reported (24). Here, we report the analysis of their O- and GSL-glycosylation patterns, and demonstrate divergent abundance of glycan structures and binding to lectins between the cell lines using porous graphitized carbon nano-liquid chromatography coupled to a tandem mass spectrometer (PGC nano-LC-ESI-MS/MS) and flow cytometry, respectively. In addition, we show that both PDAC cell lines differentially interact with immune-related lectins, likely as a consequence of the dissimilarity in surface glycosylation, which may result in their differential recognition by innate immune cells.

RESULTS

Gene Expression Profiling
To determine the differential glycosylation potential of the PaTu-S and PaTu-T cell lines, we analyzed the expression profiles of glycosylation-related genes in these cells via Affymetrix microarray analysis, and selected for genes that showed a significant ≥1.5-fold difference in expression levels between the two cell lines (P < 0.001). Remarkably, 281 genes (24% of all genes within the array), met this criterion (Figure 1A). The largest group of these 281 differentially expressed genes consists of GT genes (26%), indicating a major difference in glycan biosynthetic potential between PaTu-T and PaTu-S. A pathway analysis was performed manually by grouping these GTs in known glycosylation pathways and subsequently a Treeview analysis was generated utilizing selected groups of GT genes (Figure 1B).

The observed changes in expression levels of 7 genes involved in core N-glycosylation have been described previously in detail (24). In the GAG family gene, 20 gene transcripts displayed a more than 1.5-fold difference in expression between the two cell lines, and most of them were higher expressed in PaTu-T (Figure 1B). The enzymes encoded by these genes are almost exclusively involved in the synthesis of heparan sulfate (HS) or chondroitin sulfate (CS). The important role of expression of HS and CS and their sulfation on proteoglycans in cancer metastasis has been demonstrated previously (6). Remarkably, whereas two transcripts that encode enzymes (XYLT1 and B3GALT6, Figure 1B) involved in synthesis of the common core for HS and CS were higher expressed in PaTu-S, genes for extension of HS, and in particular sulfation in CS, were mostly higher expressed in PaTu-T.

Of the differentially expressed GT genes, 11 are putatively involved in mucin-type core O-glycosylation. Many of these gene transcripts were higher expressed in PaTu-S compared to PaTu-T cells, including several ppGALNTs, enzymes that catalyze the addition of GalNAc to Ser or Thr residues in glycoproteins. Most results observed in the gene array were confirmed by quantitative real-time PCR (Figure 2) using a set of specific primers (Table 1).
FIGURE 1 | Expression levels of glycosylation-related genes in PaTu-T and PaTu-S cell lines. (A) Glyco-gene transcripts with more than 1.5-fold difference in transcription levels between the two cell lines (281 from a total of 1,171 gene transcripts) clustered according to their putative function. The miscellaneous group (Continued)
FIGURE 1 | Includes genes related to glycosylation such as growth factors, receptors, interleukins, and adhesion molecules. Glycosyltransferases (GTs) comprises 73 of the 281 genes (26%) and are classified according to their assumed role in biosynthesis of target structures like O-glycan core, N-glycan core, GAG (glycosaminoglycan), GSLs (glycosphingolipids), Term (terminal modifications of various types of glycan structures), and Ext (extension of the various types of glycan structures). (B) TreeView of GT gene expression levels in PaTu-S and PaTu-T. Expression levels are depicted from low expressed genes (dark blue), through middle range gene expression (black) to highly expressed genes (yellow).

Remarkably, GALNT3, GALNT5, and GALNT12 are almost exclusively expressed in PaTu-S, whereas GALNT14 was 6 times higher expressed in PaTu-T (Figure 2). Furthermore, C1GALT1, encoding the core 1 enzyme and GCNT3 coding for the core 2/4 enzyme were expressed 5-fold and >1000-fold higher in PaTu-S compared to PaTu-T. No significant difference was observed in expression levels of GALNT2, which showed no differential expression between PaTu-T and -S in the gene array and was included as a control. In summary, our data indicate that the two cell lines differ profoundly in their potential to synthesize mucin-type O-glycans. The array data showed differential expression of 7 GT genes that are involved in GSLs biosynthesis. Interestingly, the UDP-glucose ceramide glucosyltransferase (UGCG) transcript which catalyzes the first glycosylation step in GSL-biosynthesis showed 5 times higher expression in PaTu-T compared to PaTu-S (Figure 2). The gene expression of four key enzymes responsible for the synthesis of globosides, gangliosides, and (neo)lacto-series GSLs (nsGSLs), namely A4GALT, ST3GAL5, B4GALNT1, and B3GNT5, was also analyzed (Figure 2). It revealed a neoexpression of A4GALT, a 9-fold higher level of ST3GAL5 and a 1.5-fold higher expression of B4GALNT1 in PaTu-T cells. However, a 2-fold lower expression of B3GNT5 was observed. These different expression levels may lead to specific expression of globosides, elevated expression of gangliosides, and a decreased level of nsGSLs in PaTu-T cells, respectively.

Furthermore, gene transcripts involved in the extension and termination of the core structures of N-glycans, O-glycans, or GSLs, were differentially expressed between PaTu-S and PaTu-T (Figures 1, 2). Examples are GCNT2, which was 19 times higher in PaTu-T and which is involved in the extension of type 2 glycans and glucosylceramide-based GSLs. By contrast, B3GNT3, involved in polyactosamine synthesis, was higher expressed in PaTu-S. Transcripts involved in terminal reactions of different glycan-types, which were higher in PaTu-S compared to PaTu-T include FUT1 and FUT2. These results are consistent with
our previous findings on the elevated expression of FUT1 and FUT2 regulated by hypoxia inducible factor α (25). In addition, PaTu-S cells displayed higher levels of ST6GAL1 (5.3 FC), which suggests an elevated capacity for the α2,6 sialylation of terminal galactose. Similarly, the expression of ST6GALNAC1, which can sialylate Tn antigens to form the cancer-associated sialylated Tn antigen (sTn) (26), was higher in PaTu-S compared to PaTu-T (Figure 1B). By contrast, PaTu-T cells show an increased potential for α2,3 sialylation encoded by ST genes ST3GAL3 (3.0 FC), ST3GAL4 (8.2 FC), and ST3GAL6 (4.2 FC). Also, expression levels of ST6GALNAC4 (5.1 FC) and ST6GALNAC6 (2.2 FC), involved in sialylation of core 1 and 2 O-glycans, globosides, and gangliosides were more than 2 fold higher in PaTu-T.

In summary, based on the observed GT expression data our findings suggest that PaTu-T and PaTu-S display major differences in cellular glycosylation.

**Biosynthesis of O-Glycans**

Since one of the most remarkable differences observed in GT expression between PaTu-S and PaTu-T was related to initiation of O-glycan synthesis, we studied this aspect in more detail. We determined for two selected mRNA species whether the differences in gene transcript levels between Patu-S and Patu-T were also observed at the protein level by Western blotting for the GT proteins. Substantial amounts of GALNT3 and CGNT3 protein levels were observed in whole cell lysates of PaTu-S, but were hardly detectable in PaTu-T, which correlated with the mRNA expression levels found in these cells (Figures 2, 3A). To define the potential of the cells to catalyze the addition of α-GalNAc to Ser/Thr residues on a peptide, a ppGALNT enzyme assay was performed using two different peptides with multiple Ser/Thr residues, derived from immunoglobin A (IgA) and mucin 2 (MUC2) proteins, respectively. PaTu-S cells showed a much higher ppGALNT activity of T-antigen (Figure 3B), which was associated with elevated levels of surface Tn antigen as detected by using a monoclonal anti-Tn antibody (Figure 3C and Supplementary Figure S1A). Likewise, the activity of β1,3-galactosyltransferase (CIGALT1, T-synthase) converting Tn antigen into T antigen was measured and found to be highest in PaTu-S (Figure 3D). Human Jurkat T cells do not have T-synthase activity due to a mutation in the O-sacm gene (CIGALT1 chaperone) (27), and were therefore used as a negative and baseline control for the assay. In addition, the peanut agglutinin (PNA) showed enhanced binding to PaTu-S compared to the other cell lines, again indicating a relatively higher level of T-antigen (Figure 3E and Supplementary Figure S1B).
conclusion, PaTu-S cells show a higher potential for the synthesis of O-glycans, resulting in the presence of elevated levels of short O-glycans (both Tn and T antigens) compared to PaTu-T.

**O-Glycosylation Analysis in PaTu-S and PaTu-T by PGC Nano-LC-ESI-MS/MS**

To further study the total expression of cellular glycans, we analyzed the O-glycans released from PaTu-S and PaTu-T cells by PGC nano-LC-ESI-MS/MS (Figure 4A). PGC LC-ESI-MS/MS using negative electrospray ionization was employed for structural analysis of glycans, which is a powerful platform enabling discrimination between glycan isomers (28–30). Glycan structures were assigned on the basis of the obtained glycoanalytical information, and general glycobiological knowledge. The powerful chromatographic separation of two O-glycan isomers with m/z 675.30 and five GSL-glycan isomers with m/z 999.30 with characteristic MS/MS spectra are shown in Supplementary Figures S2, S3, respectively.

Representative O-glycan profile spectra with the main peaks annotated are shown in Figure 4A. Compared to PaTu-S, most of the O-glycans synthesized in PaTu-T, except for two disialylated O-glycans, were detected at lower level or depleted, resulting in less diversity of O-glycan species (Figure 4A) and a lower abundance of estimated total O-glycans (Figure 4B) in PaTu-T. Relative quantification of individual glycans is given in Supplementary Figure S4, revealing the large differences in detail. Glycan traits or ratios reflecting certain biosynthetic steps were calculated to facilitate the biological interpretation of the data and to relate the MS glycomics data to transcriptomics data. This included core 2/4 structures, sTn antigen, T antigen, terminal acetylglucosamine (GlcNAc), terminal N-acetylgalactosamine (GalNAc), α2,3 sialylation of galactose, α2,6 sialylation of galactose, α2,6 sialylation of GalNAc, α1,2 fucosylation of galactose, α1,3/4 fucosylation, and sLe^A_. In general, PaTu-T showed a lower levels of estimated total O-glycans, probably resulting from the lower level of ppGALNTs (Figure 4B). In PaTu-S, core 2 and core 4 O-glycans accounted

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**FIGURE 3 |** Biosynthesis of O-glycan structures. (A) Protein levels of GCNT3 and GALNT3 in PaTu-S and PaTu-T. Total proteins (150 µg) from whole-cell extracts were analyzed by Western blotting using anti-GCNT3 and anti-GALNT3 for detection of the respective proteins and anti-actin as the protein loading control. (B) ppGALNT activity was determined using whole cell lysates from PaTu-S and PaTu-T cells as enzyme source, UDP-[3H]-GalNAc as a donor, and MUC2 and IgA hinge region peptide as acceptor substrates. Radioactivity incorporated in the peptide-products was determined. (C) Binding of anti-Tn mAb to PaTu-S and PaTu-T cells as measured by flow cytometry, and shown as average MFI ± SEM of at least 3 independent experiments. (D) T-synthase activity was measured using whole cell lysates from PaTu-S, PaTu-T, and Jurkat cells (as negative control) as enzyme sources, UDP-Gal as a donor and GalNAc-α-(4-MU) as acceptor. Final product was measured by fluorescence (excitation 360 nm and emission at 460 nm). Results are given as average enzyme activity ± SEM of at least 3 independent experiments. (E) Binding of PNA recognizing terminal Galβ1-3GalNAc (T-antigen) to PaTu-S and PaTu-T cells as measured by flow cytometry, and shown as average MFI ± SEM of at least 3 independent experiments. *P ≤ 0.05 and ***P ≤ 0.001.
for 78.2% of the species in contrast to only 49.4% in PaTu-T (Figure 4C) and in line with the higher expression of GCNT3 in PaTu-S (Figure 2). Importantly, the tumor-associated O-glycans sTn antigen (Figure 4D) and T antigen (Figure 4E) were only detected in PaTu-S cell lines. Figures 4F–G further show that terminal GlcNAc and GalNAc were detected only in PaTu-S cells. For sialylation, an increase of α2,3 sialylation of galactose was observed for PaTu-S compared to PaTu-T (Figure 4H) in accordance with the elevated level of ST3GALs in PaTu-T cells (Figure 2). Interestingly, α2,6 sialylation on galactose was specifically present in PaTu-S cell line, as there was no α2,6-linked sialic acid on galactose in PaTu-T (Figure 4I). Opposite to α2,6 sialylation of galactose, sialylation on the GalNAc with α2,6 linkage was high in PaTu-T with a relative abundance at 42.5%, compared to 18.0% in PaTu-S (Figure 4J), which is in line with the expression patterns of ST6GALNAC1, 4 and 6. Notably, a decrease of α1,2 and α1,3/4 fucosylation was observed in the more mesenchymal-like
PaTu-T cells (Figures 4K, L), in accordance with the decreased level of β-Galactosidase in PaTu-T cells (Figure 1). Remarkably, we also observed many specific glycan structures in PaTu-S cells including sLeA, blood H antigen, blood group A, and Lewis X (Figure 4A).

**Glycosphingolipid-Glycan Analysis in PaTu-S and PaTu-T by PGC Nano-LC-ESI-MS/MS**

Next, GSL-glycans were analyzed with PGC nano-LC-ESI-MS/MS after enzymatic release using endoglycosidase I (EGCase I) on purified GSLs derived from 2 × 10^6 cells per sample. The combined extracted ion chromatograms of GSL-glycans in PaTu-S and PaTu-T cell lines (Figure 5A) and the relative quantification of GSL-glycans (Supplementary Figure S5) are presented. A lower abundance of estimated total GSL-glycans (Figure 5B) in PaTu-T was observed. The data showed vastly different GSL patterns of PaTu-T cells with higher levels of globosides and gangliosides and lower levels of nsGSLs (Figures 5C–E). Importantly, a specific detection of globosides (Gb3 and Gb4; 7.7±1.0%) in PaTu-T was found, indicating a possible important role of globosides in PDAC cells (Figure 5C). Approximately 10-fold higher levels of gangliosides were observed in PaTu-T cells (35.0±1.4%) compared to PaTu-S cells (3.4±0.3%). In contrast, nsGSLs were relatively lower in PaTu-T cells (57.4±0.5%) compared to PaTu-S cells (96.6±0.3%). These results are also in alignment with the gene expression data, which showed elevated expression of A4GALT, ST3GAL5, B4GALNT1, and a slightly decreased B3GNT5 level (Figure 2).

Differences in GSL abundance between the two cell lines with respect to α2,3 and α2,6 sialylation of galactose residues, α1,2 fucosylation of galactose and α1,3/4 fucosylation were very much in line with those found for O-glycosylation (Figures 5G–J). Decreased abundance of α2,6 sialic acid on galactose (74.8±1.2% in PaTu-S, Figure 5H), α1,2 fucosylation (24.2±0.3% in PaTu-S, Figure 5I), and α1,3/4 fucosylation (1.4±0.5% in PaTu-S, Figure 5J), were observed in PaTu-T compared to the PaTu-S. Remarkably, α2,3 sialylation of galactose was higher in the mesenchymal-like PaTu-S cells than in the epithelial-like PaTu-S cells (Figure 5G). No significant difference was found in terminal GaINAc of GSL-glycans between PaTu-S and PaTu-T (Figure 5F). Similar to O-glycans, the Lewis X moiety was found to be specifically detected on GSLs of PaTu-S cells (Figure 5K).

**Differential Abundance of Terminal Glycan Structures in PaTu-S and PaTu-T**

In order to confirm the glycan expression and to understand the effects of the differential GT expression levels described above, the binding of different glycan-binding proteins (GBPs) including lectins and anti-glycan antibodies to the cell lines was analyzed by flow cytometry.

As shown in Figure 6 and Supplementary Figure S6, the lectins Helix pomatia agglutinin (HPA; terminal α GalNAc), soybean agglutinin (SBA; terminal α/βGalNAc and Gal) and Wistera floribunda (WFA; terminal α/βGalNAc) showed enhanced binding to PaTu-S compared to PaTu-T. Together with the observation that several GALNT genes were expressed at higher levels in PaTu-S compared to PaTu-T, these data indicate the presence of relatively higher amounts of terminal GaINAc residues on PaTu-S. In addition, PaTu-S showed substantial binding of the Wheat germ agglutinin (WGA; βGalNAc), suggesting the presence of structures with terminal GlcNAc. These results are in accordance with the higher abundance of terminal GlcNAc in O-glycans (Figure 4F).

By contrast, PaTu-T cells showed higher binding of the α2,3 sialic acid–recognizing lectin *Maackia amurensis* (MAA, α2,3 sialic acids) than PaTu-S, which was in agreement with a higher expression of ST3GAL3, -4 and -6 genes and a higher abundance of α2,3 sialylation of galactose in both O-glycans (Figure 4H), GSL-glycans (Figure 5G), and N-glycans (24) in PaTu-T (Figure 5G). The binding of SNA lectin, specific to α2,6 sialic acids, to PaTu-T cell lines was elevated, indicating higher levels of α2,6 sialylation in PaTu-T. More details on the relevance of transcriptomics and MS glycomics data will be discussed in the discussion.

*Lotus tetragonolobus* lectin (LTA) bound almost exclusively to PaTu-S, indicating the presence of terminal α1,3/4-linked fucose. In addition, binding of UEA-1, which detects terminal α1,2-linked fucose (H-antigen), to the surface of PaTu-S cells was observed, which is in agreement with the higher levels of *FUT1/FUT2* transcripts (25) and the higher fucosylation level in O-glycans (Figures 4K, L) and GSL-glycans in these cells (Figures 5I, J). In addition, considerably higher levels of sLeA structures were found in O-glycans of PaTu-S than those of PaTu-T (Figure 4M), indicating the presence of sLeA antigen on PaTu-S, in line with our previous findings of more anti-sLeA antibody binding to PaTu-S than to PaTu-T (24).

**PaTu-S and PaTu-T Differentially Bind to Lectins of the Immune System**

In the immune system, dendritic cells (DCs) are specialized in the control of innate and adaptive immunity by sensing of pathogens as well as cancer cells via receptors present on their cell surface (31). The different glycan structures on PaTu-S and PaTu-T will very likely provoke a differential interaction with C-type lectin receptors present on DCs lectins, as well as with soluble imunolectins like galectins. Such an interaction could directly influence immune responses toward the tumor cells, or mediate immune escape. Here, the interaction of immature DCs with both PDAC cell lines was observed by fluorescence microscopy (Figure 7A) and the level of binding was quantified using a cell adhesion assay (Figure 7B). The DCs showed a similar level of binding to the two PDAC cell lines. This binding was C-type lectin dependent, since it was abrogated in the presence of the chelating agent EGTA (Figure 7B). In addition, the binding of recombinant soluble galectins (Gal-1, Gal-3, and Gal-4) and lectins (DC-SIGN, MGL, DCIR, and Dectin-1), which are normally present on the surface of immature DCs, to both PaTu cell lines was studied. The results showed that whereas the dendritic cell immunoreceptor (DCIR) bound
both cell lines, and no binding of Dectin-1 could be detected. Macrophage galactose-type lectin (MGL) and dendritic cell-specific intercellular adhesion molecule-3-grabbing non-integrin (DC-SIGN) showed a differential binding pattern, with elevated binding to PaTu-S cells, compared to PaTu-T (Figure 7C). In addition, more Gal-1 and Gal-4 bound to PaTu-S compared to PaTu-T, whereas there was no difference in binding of Gal-3 to both cell lines observed (Figure 7D). We have previously shown that the binding of Gal-4 to both cell lines was glycan-dependent (22). Although the binding of both Gal-1 and Gal-3 has not been tested for exclusively be glycan-dependent, it is assumed to be glycan-dependent, without excluding non-glycan dependent interactions. Because there is hardly any binding by Gal-3 detected on both cell lines, this indicates binding specific by Gal-1. These data indicates differences in lectin binding properties between PaTu-S and PaTu-T suggest a disparate effect on the immune response, the exact nature of which is subject for future investigations.
Alterations in glycosylation of cancer cells play an important role in the progression of cancer, however the individual roles that specific glycans play are incompletely understood. A rational design of experimental approaches to selectively detect and functionally analyze the role of specific glycans in the progression of PDAC requires a detailed understanding of the biosynthetic pathways that lead to cancer-associated glycan expression. Recently a direct correlation was reported between the formation of CA 19-9 and the development of rapid and aggressive development in pancreatic tumorogenesis. The work of Engle et al. showed that the expression of FUT3 and B3GALT5 in mouse PDAC cells were essential to the cell surface expression of CA 19-9. In addition, the authors showed that the expression of glycan CA 19-9 together with KrasG12D oncogene was a causal origin of PDAC progression. N-glycosylation of the mesenchymal-like PaTu-T and the epithelial-like PaTu-S cell lines is observed at gene transcript levels of enzymes involved in synthesis of core 1/2 N-glycosylation and GSL-glycosylation, some of which were confirmed to be reflected at the glycan level by Western blotting, binding to plan lectins and interaction with immunolectins. Combined, these results suggest a more dominant role of O-glycosylation and GSL-glycosylation in pancreatic cancer biology, compared to N-glycosylation.

DISCUSSION

By contrast, in this study we demonstrate significant changes in the expression of genes involved in O-glycosylation and GSL-glycosylation, some of which were confirmed to be reflected at the glycan level by Western blotting, binding to plan lectins and interaction with immunolectins. Combined, these results suggest a more dominant role of O-glycosylation and GSL-glycosylation in pancreatic cancer biology, compared to N-glycosylation.

One of the most remarkable differences between the two cell lines is observed at gene transcript levels of enzymes involved in synthesis of core 1/2 O-glycans, which are important determinants of the mucin-layer of epithelial cells. Mucins, especially the highly-glycosylated MUC1, MUC4, MUC5AC, and MUC16, are prominently observed in pancreatic cancer and major carriers of glycans including the CA 19-9 antigen (36, 37). Mucin-type O-glycans usually undergo drastic changes during PDAC progression (13, 18, 36), thereby influencing important properties of the tumor cells, including growth, adhesion, and interactions with the immune system. The enhanced mucin-type O-glycan synthases (ppGLALT5s) and T-synthase activity, as well as enhanced levels of the Tn antigen and T antigen observed in PaTu-S compared to PaTu-T, demonstrate an elevated biosynthesis of short mucin-type O-glycans in PaTu-S. The expression of both antigens is mainly regulated by ppGLALT5s and inefficient conversion of Tn antigen to T antigen. The inefficient conversion to T antigen is regulated
by activity of T synthase and Cosmc, which is a chaperone involved in the proper folding and activity of T-synthase (38). Such inefficient Cosmc expression could be due to lowered expression caused for example by epigenetic silencing (17, 39). In addition, knocking out of C1GALT1 in PDAC cells leads to inefficient conversion of Tn antigen to T antigen, and further increases tumor progression and metastasis (40). In these cells, the presence of Tn antigens most probably is due to a strongly enhanced expression of ppGALNT genes, encoding a family of enzymes that catalyze the first synthetic step in mucin-type O-glycosylation, including GALNT3, -T4, -T5, -T7, and -T12. This may result in an extended O-glycan initiation platform in PaTu-S cells as compared to PaTu-T. Consequently the higher expression of these ppGALNTs could possibly lead to an incomplete conversion of Tn to T antigen by active T-synthase or a relocation of the GalNAc-Ts from the Golgi apparatus to the endoplasmic reticulum (10). Similar to PaTu-S, a high GALNT3 expression was observed in human pancreatic cancer tissues (13). Suppression of its expression significantly decreased cell growth and increased apoptosis (13). Recently, loss of GALNT3 in poorly differentiated PDAC was shown to alter glycosylation of Erythroblastic Leukemia Viral Oncogene Homolog (ErbB) family proteins, and further associated with increased tumor aggressiveness (19). GALNT3 expression was found to be restricted to glandular epithelial cells (41) in lung adenocarcinoma and to be downregulated...
in mesenchymal-like cells in epithelial-mesenchymal transition models (42, 43). GALNT7 upregulation has previously been shown to be involved in growth and invasion in cervical cancer (44) and its downregulation increases metastasis formation in melanoma cells (45). These findings agree with our data, showing a much higher expression of GALNT3 and GALNT7 in the more differentiated epithelial-like PaTu-S, compared to the mesenchymal-like PaTu-T cells. In addition to an elevated level of Tn antigen, also elevated levels of T antigen were found in PaTu-S compared to PaTu-T cells. This indicates that despite the high expression of GCNT3 in PaTu-S cells, the T antigens were not efficiently converted to core 2 structures. In contrast to PaTu-S, the mesenchymal-like PaTu-T expressed lower levels of Tn synthases, which could explain the decrease in total O-glycan expression. The expression of certain mucins was elevated in PaTu-S, however, PaTu-T showed increased expression of other glycoproteins such as mucolipins (data not shown). Therefore, it cannot be excluded that lower expression of glycoproteins with many O-glycan attachment sites such as mucolipins contributes to the effect. In addition, elevated ST6GALNAC4 and 6 terminate β1-6 branching of core GalNac, prohibiting the conversion to core 2/4 structures, which may result in a lower abundance of core 2/4 in PaTu-T.

GSLs play essential roles in maintenance of membrane stability and regulation of numerous cellular processes, including adhesion and metastasis (46–48). Our result shows higher levels of UGCC, B4GALT5, and B4GALT6 in the mesenchymal-like PaTu-T compared to PaTu-S. Notably, neoeexpression of globosides (Gb3 and Gb4) was observed, matching the increased expression level of A4GALT. This is an interesting observation, since it has been reported that both UGCC and Gb3 are associated with tumor metastasis. Inhibition of UGCC reduced the levels of all cellular GSLs, including lactosylceramide, Gb3 and GM3, thus decreasing lung tumor metastasis (49). Moreover, enhanced ceramide glycosylation by UGCC has been shown to confer multidrug resistance of cancer cells (50) and to selectively maintain pluripotency properties of cancer stem cells (51). Globoside Gb3 has also been reported to be highly expressed in a number of tumors, including metastatic colon cancer (52, 53), breast cancer (54), gastric adenocarcinomas (55), and pancreatic cancer (56). Importantly, the overexpression of Gb3 in cultured human colonic epithelial cells induced cell invasiveness, whereas knocking Gb3 down inhibited cell invasiveness (52). We show in this study that globoside Gb3 and Gb4 are specifically expressed in the mesenchymal-like PaTu-T cells, indicating it may contribute to the metastasis in PDAC. However, further functional studies are necessary to further understand the role of globosides especially Gb3 in cancer progression and metastasis.

A striking difference in terminal glycosylation was observed in the levels of fucosylated structures. Fucose, as a constituent of oligosaccharides, is often associated with cancer and inflammation (57, 58), and can be used as a promising target for cancer diagnosis and therapy (59). The enhanced binding of UEA1 to PaTu-S compared to PaTu-T is in line with the higher levels in PaTu-S of the FUT genes FUT1 and FUT2, which are involved in the synthesis of the H-antigen, the precursor for blood group A/B antigens (BG-A/B). This can be further confirmed by highly expressed blood group H antigen and BG-A in PaTu-S compared to PaTu-T (Supplementary Figures S7A–C). These data are consistent with our previous report (25) where we showed that both the expression of FUT1 and FUT2 genes, together with the binding of anti-BGA and anti-BGB antibodies, using flow cytometry, were much higher in PaTu-S and undetectable in PaTu-T cells. In addition, the α1.3 FUT gene FUT4 is expressed at higher levels in PaTu-S and may be involved in the synthesis of Lewis X (24), and in combination with FUT1 and FUT2 in PaTu-S in the synthesis of Lewis Y. The observation that higher binding of the lectin LTA, which binds to terminal α1,3 fucose, to PaTu-S than to PaTu-T as well as the specific presence of Lewis X on O-glycans (Supplementary Figure S7D) and GSL-glycans in PaTu-S is also in agreement with our previous reports (24, 25) and supports the expression of these known cancer-associated glycan structures (14) on PaTu-S cells.

Altered sialylation has long been associated with metastatic behavior of tumor cells including invasion and enhanced cell survival [reviewed in (60)]. Overexpression of ST3GAL3 in pancreatic cell lines induced increased expression of sLeX together with decreased α2,6 sialylation on α2β1 integrin and E-cadherin, which further regulate adhesion and invasion in pancreatic cancer (61–63). The mesenchymal-like PaTu-T shows enhanced binding of the α2,3-sialic acid-recognizing lectin MAA compared to the epithelial-like PaTu-S, which is in line with the higher expression levels of ST3GALs in PaTu-T for all these three types of glycans. For the α2,6-linked sialic acid, we observed higher SNA lectin binding to PaTu-T cells indicating a slightly higher expression level of α2,6 sialic acid. However, we found a reduced relative abundance of α2,6 sialylation of galactose in all three types of glycans [O-, GSL-, or N-glycans (24)] together with decreased expression of ST6GAL1 in PaTu-T when compared to PaTu-S. This conflicting result could be explained by the higher expression level of α2,6-linked sialic acid on GalNAc in O-glycans (Supplementary Table S1). This type of α2,6 sialylation of GalNAc is highly expressed in the mesenchymal-like PaTu-T cells, in accordance with the increased gene expression of ST6GALNAC 4 and 6. Similar to PaTu-T, ST6GALNAC4 was shown to be upregulated in other mesenchymal type PDAC cells (42).

The profound differences in glycosylation between PaTu-S and PaTu-T may indicate that these cells will differentially interact with immune cells. The preferential binding of DC-SIGN to PaTu-S may be due to the higher abundance of fucose-containing Lewis antigens (64), whereas the higher MGL binding to PaTu-S can be explained by the higher levels of terminal GalNAc, including the Tn antigen. Binding of DCs to cancer cells via MGL and DC-SIGN was previously shown and has been proposed to lead to an immune escape of the tumor cells (65–67). Since MGL binds to terminal α- and β-linked GalNAc residues, including Tn-antigen (65, 68), the differential binding of MGL and DC-SIGN to the cell lines corresponds well with their respective glycosylation profiles, especially in O-glycans. Expression of the Tn antigen on cancer cells is associated with poor prognosis in several types of cancer and instructs DCs to drive Th2 mediated responses, which contribute to tumor development (31).
Also the galectins 1 and 4 show differential binding to PaTu-T and PaTu-S. Galectins belong to a family of soluble immunomodulatory proteins that may act extracellularly by crosslinking their ligands thereby modulating signaling pathways (31). Remarkably, the soluble lectin Gal-4 shows a much higher binding to PaTu-S than to PaTu-T (8-fold), which may be due to the presence of blood group H and A ligands for this lectin on PaTu-S (25, 69). We demonstrated previously that PaTu-S, but not PaTu-T produces significant levels of intracellular and secreted Gal-4 (22). Whereas cytoplasmic Gal-4 can suppress metastatic activity in PDAC cells (22, 23), the role of extracellular Gal-4 in PDAC is unknown, but may include adhesion of the tumor cells to each other, or to extracellular matrix structures.

CONCLUSION

In this work an overview of the difference in cellular glycosylation between two cell lines was obtained using a variety of approaches. Supplementary Table S1 provides an overview on the significant alterations identified in this work and our previous studies (24, 25). The observed differences in expression levels of GTs are in good accordance with the glycan structures highlighted in the significance of current approaches. The differences in expression levels of GTs are in good accordance with the glycan structures identified by mass spectrometry and flow cytometry. This is remarkable, since cancer cells are often reported to display a disorganized ER/Golgi compartment, which would be expected to influence glycosylation patterns (10). Moreover, by using this combined approach it was possible to demonstrate the association of specific glycan structures with their carrying glycan types (Supplementary Table S1). For example, we showed detailed changes in α2,6 sialylation as well as that Lewis X is predominantly detected on GSLs and O-glycans while sLeA was mainly found on O-glycans. In addition, the higher binding ability of HPA, SBA, and WFA lectins to terminal GalNAc in PaTu-S was in accordance with the increased level of terminal GalNAc in O-glycosylation, rather than present on GSL-glycans, highlighting the importance of studying cellular glycosylation in a comprehensive and glycan-specific manner for characterization and functional studies of glycosylation.

The epithelial-like PaTu-S cells and the more mesenchymal PaTu-T cells can be regarded as a differentiation model of cells that undergo the early and later stages of metastasis formation. Here we show pronounced differences in O- and GSL-glycosylation between PaTu-T and PaTu-S cells, which is in line with the observed gene expression and associated with differential interaction with immune cell lectins. The results from this study provide leads for future research to define the role of specific glycan structures in PDAC progression, such as the role of the globoside Gb3 in metastasis, which may reveal targets for future prognostic and/or therapeutic approaches. Recent work has opened new possibilities for pancreatic cancer prevention and possible pancreaticitis treatment through direct targeting of CA 19-9 (3). Aberrant glycosylation leads changes in key factors that lead to cancer development, poor prognosis, and treatment resistance (70). The possibilities and the necessity of current approaches in personalized cancer therapy by tackling these changes in glycosylation are emerging and becoming more prominent.

MATERIALS AND METHODS

Materials, Chemicals, Antibodies, Reagents, and Buffers

Ammonium bicarbonate, cation exchange resin beads (AG50W-X8), trifluoroacetic acid (TFA), potassium hydroxide, ammonium bicarbonate, and sodium borohydride were obtained from Sigma-Aldrich (Steinheim, Germany). 8 M guanidine hydrochloride (GuHCl) was obtained from Thermo Fisher Scientific. Peptide N-glicosidase F (PNGase F) was purchased from Roche Diagnostics (Mannheim, Germany). Endoglycosidase I (EGCase I, recombinant clone derived from Rhodococcus triotomea and expressed in Escherichia coli), 10x EGCase I buffer (500 mM HEPES, 1M NaCl, 20 mM DTT and 0.1% Brij 35, pH 5.2), glycobuffer 1 (50 mM sodium acetate, 5mM CaCl₂, pH 5.5), and α2–3 neuraminidase S were purchased from New England BioLabs Inc., (Ipswich, MA, USA). Dithiothreitol (DTT), HPLC SupraGradient acetonitrile (ACN) was obtained from Biosolve (Valkenswaard, The Netherlands) and other reagents and solvents such as chloroform, methanol, ethanol, 2-propanol, and glacial acetic acid were from Merck (Darmstadt, Germany). MultiScreen® HTS 96 multwell plates (pore size 0.45 µm) with high protein-binding membrane (hydrophobic Immobilon-P PVDF membrane) were purchased from Millipore (Amsterdam, The Netherlands), conical 96-well Nunc plates from Thermo Scientific (Roskilde, Denmark). The 50 mg TC18-reverse phase (RP)-cartridges were from Waters (Breda, The Netherlands). Ultrapure water was used for the all preparations and washes, generated from a Q-Gard 2 system (Millipore, Amsterdam, The Netherlands).

Rabbit anti-human GALNT3 and anti-human GCNT3 (Atlas Antibodies, Stockholm, Sweden) antibodies (Abs) were used for detection of GALNT3 and GCNT3, respectively. Mouse anti-human galectin (Gal)-1 (71), and anti-Tn monoclonal antibodies were kindly provided by Dr. RD Cummings (Boston, USA), goat anti-human Gal-4 was purchased from R&D Systems (Minneapolis, MN) and anti-sialyl Lewis A/CA 19-9 monoclonal antibody was from LifeSpan Biosciences (Seattle, WA). Goat anti-human actin (Santa Cruz Biotechnology, Heidelberg, Germany) was used as an endogenous control. Secondary Abs and proteins used were goat anti-rabbit Alexa Fluor 488 (AF 488), anti-mouse AF 488 and rabbit anti-goat AF 488, streptavidin AF 488 (Molecular Probes, Invitrogen, Carlsbad, CA), goat anti-human Fc (Jackson Immunoresearch, Suffolk, UK) and IRDye 680 anti-goat and IRDye 800CW anti-rabbit IgG (LI-COR Biosciences, Lincoln, Nebraska). Roti-Block blocking buffer was acquired from Roth (Karlsruhe, Germany) and Calceine AM from Molecular Probes. Recombinant human Gal-1 (72) was kindly provided by Dr. Rabinovich. Production of biotinylated recombinant human Gal-3 was described in van Stijn et al. (73) and recombinant human Gal-4 was purchased from R&D. Recombinant C-type lectins [MGL-Fc (74), DC-SIGN-Fc (75), DCIR-Fc (76), and Dectin1-Fc] consist of the extracellular domains of the lectins fused to the human immunoglobulin IgG1-Fc tail. To construct Dectin1-Fc, the extracellular domains of human Dectin-1(a) (amino acids 65-241) were amplified using total RNA from immature DCs and fused at the C-terminus.

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to human IgG1-Fc in the Sig-pIgG1-Fc vector. Dectin-1-Fc was produced by transient transfection of HEK-293T cells.

Biotinylated lectins from peanut (PNA) and Helix pomatia (HPA) were purchased from Sigma-Aldrich (Zwijndrecht, The Netherlands). Lectins from Aleuria aurantia (AAL), Lotus tetragonolobus (LTA), Maackia amurensis (MAA; mixed MAA-I and MAA-II), soybean agglutinin (SBA), Sambucus nigra agglutinin (SNA), Ulex europaeus I (UEA-1), Wistera floribunda (WFA), and Wheat germ agglutinin (WGA) were bought from E-Y Laboratories (San Mateo, CA).

Cells and Cell Culture
PaTu-S and PaTu-T cell lines were purchased at DSMZ culture bank (Braunschweig, Germany), following the manufacturers protocol. The quality of the mRNA was established by measurement of the A260/A280 ratio and in-gel agarose visualization. Microarray screening was performed by the Consortium for Functional Glycomics (CFG) (http://www.functionalglycomics.org), using the GlycoV4 oligonucleotide array, a custom Affymetrix GeneChip (Affymetrix, CA, USA) designed for the CFG. This array includes ~1,260 human probe-ids related to glyco-genes. The data were normalized using RMA Express 1.0 (http://rmaexpress.bmbolstad.com) with quantile normalization, median polish, and background adjustment. Differential expression was calculated using the Limma package in R software. Fold change and standard errors were estimated by fitting a linear model for each gene and empirical Bayes smoothing was applied to the standard errors. The adjusted \( p \)-value is the \( p \)-value adjusted for multiple testing using the Benjamini and Hochberg’s method (77) to control the false discovery rate of 0.1 or less. The transcripts identified as differentially expressed were those with adjusted \( p \) < 0.1 and a difference in fold change > 1.5. Cluster Analysis and TreeView Visualization Software was used to visualize the microarray data. Both programs are integrated for analysis and visualization of complex microarray experiments data. Both software programs were developed by Michael Eisen at the Michael Eisen’s lab in the Howard Hughes Medical Institute (HHMI) at University of California at Berkeley (UCB) and the Lawrence Berkeley National Lab (LBNL) USA.

Microarray Analysis
Cells were harvested at 60% confluency using trypsin-EDTA (Gibco, Invitrogen). mRNA was isolated using the RNeasy Mini kit (Qiagen, Venlo, The Netherlands), following the manufacturer's guidelines. cDNA was synthesized with Reverse Transcription System kit (Promega, San Luis Obispo, CA). Part of the oligonucleotides used were previously described Garcia-Vallejo et al. (78). The remaining oligonucleotides used in this study were designed using software from http://www.ncbi.nlm.nih.gov/nuccore, and were synthesized by Invitrogen (Table 1). PCR reactions were performed as described Belo et al. (22) and relative mRNA abundance was calculated from CT values from the target vs. the endogenous reference gene glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (78).

Western-Blotting
Shortly, 150 µg total proteins of whole cell lysates were separated by SDS-PAGE and the proteins were transferred to nitrocellulose membranes (Whatman Protran, Sigma). Blots were incubated with anti-GALNT3 and GCNT3 Abs (0.1 µg/ml) and anti-Actin Abs (0.2 µg/ml, as a loading control), followed by incubations with the secondary IRDye Abs (0.07 µg/ml). Western-blot analysis was performed using LI-COR Odyssey systems scanner and software.

Glycosyltransferase Activity
Cells were lysed in TSM buffer containing protease inhibitors by five sonication cycles of three sec each on ice after which Triton X-100 (0.5% final concentration) was added and cell lysates were further incubated on ice for 20 min. Lysates were centrifuged at 1000 × g at 4°C for 5 min and supernatant was used in both T-synthase and GALNT assays. T-synthase activity was measured as described in Ju et al. (79), using GalNAc-α-(4-MU) (1,000 µM) as an acceptor, UDP-Gal (500 µM) as a donor, and the cell extracts as enzyme source. Cell extracts from human Jurkat T cells were used as negative control. For the standard curve measurements, free 4-MU was used. Reaction mixtures were incubated for 60 min at 37°C and the final fluorescent product was measured on a FLUOstar Spectrofluorimeter (excitation 360 nm and emission at 460 nm).

GALNT assays were carried out in a volume of 50 µl reaction mixture (pH = 7.0), containing 0.1 M sodium cacodylate, 10 mM MnCl₂, 4 mM ATP (to counteract enzymatic degradation of the nucleotide sugar), 0.5% Triton X-100, 0.25 mM UDP-[³H]-GalNAc (4,15 Ci/mol), 1 mM acceptor peptide [PVPPTPPSPTPPSPTPPS, comprising the IgA hinge region (80)] and PTTPPTITTTTVTPTPTPTGQT, representing MUC2 tandem repeat region, and 5 µl cell lysate as enzyme source. After a 4 h incubation at 37°C, the radioactive products were isolated using Sep-Pak C-18 cartridges (Waters, Milford, MA) as described Palcic et al. (81). Incorporation of radioactivity in the peptides was assessed using a liquid scintillation MicroBeta2 Plate Counter 2450 (Perkin Elmer, USA). Control assays lacking the acceptor were carried out to correct for incorporation into endogenous acceptors.

O-Glycan Release and Purification From PaTu-S and PaTu-T Cell Lines
We have previously described a high-throughput polystyrylhexylidene difluoride (PVDF)-membrane-based 96-well plate for N-glycans release (24). Here, a sequential N-glycan release followed by β-elimination was established to achieve reduced O-glycans derived from each cell line. After desalting and purification, the O-glycans were subjected to analysis on
PGC nano-LC-ESI-MS/MS (82). Cell pellets with ∼2 x 10^6 cells from three biological replicates of PaTu-S and PaTu-T were washed with PBS and suspended in 100 µl H_2O. A high-throughput release of N-glycans has been performed first to avoid contamination during O-glycan release as previously described Holst et al. (24). After removal of N-glycans, the O-glycan release via β-elimination, desalting and PGC clean-up was carried out in a high-throughput method using a PVDF membrane-based protocol adapted from Jensen et al. (82). Briefly, 60 µl of 0.5M NaBH_4 in 50 mM KOH were applied onto the PVDF membrane in each well of 96-well plates after rewetting the PVDF membrane by 3 µl of methanol. Plates were shaken for 30 min on a horizontal shaker and incubated in a moisturized, sealed box as an incubation chamber within an oven for 16 h at 50°C. After incubation and cooling to RT, released O-glycans were recovered by centrifugation at 1,000 x g for 2 min into 96-well collection plates. The wells were rewet by 3 µl of methanol and washed by 50 µl of H_2O with 10 min incubation steps on a horizontal shaker prior to centrifugation at 1,000 x g for 2 min. This wash and centrifugation steps were performed three times, and the washes were pooled to the collection plates.

Prior to desalting of O-glycans, the collected samples were concentrated to ∼30 µl under vacuum in an Eppendorf Concentrator 5301 (Eppendorf, Hamburg, Germany) at 30°C for 2 h. Subsequently, 3 µl of glacial acetic acid was added to neutralize the reaction followed by a brief centrifugation to collect the sample at the bottom of the well. In the next step, desalting of O-glycans was performed on cation exchange solid phase extraction (SPE) which consisted of 80 µl of AG50W-X8 resin slurry deposited onto OroChem 96-well filter plates (Perfect Pure, Millipore) as previously described Jensen et al. (82). Glycan samples were loaded onto the plates, and flow-through was collected into a collection plate after centrifugation at 1,000 x g for 2 min. Glycan alditols were eluted with 50 µl of methanol twice. The combined flow-through and eluate were pooled and dried under vacuum in an Eppendorf Concentrator at 30°C. Subsequent drying steps with 100 µl of methanol were performed thrice to remove residual borate. As the last step, the carbon SPE clean-up of O-glycans was performed following the same procedure with adjustment to a 96-well filter plates (82). Briefly, 80 µl of PGC slurry was loaded onto OroChem 96-well filter plates, and a two-step preconditioning were performed with 100 µl of 80% ACN in H_2O with 0.01% TFA followed by 100 µl of H_2O with 0.01% TFA. The columns were washed 3 x 50 µl of H_2O with 0.01% TFA after loading samples. O-glycans were subsequently eluted by 40 µl of 60% ACN in H_2O with 0.01% TFA and dried in concentrator. The purified glycan alditols were re-suspended in 10 µl of water prior to PGC nano-LC-ESI-MS/MS analysis.

**Glycosphingolipid Extraction and Purification by RP-SPE**

Extraction of GSLs from PaTu-S and PaTu-T cell lines was performed in triplicate using glass centrifuge tubes. Cells were harvested (2 x 10^6) and washed three times in 1 ml of water followed by centrifugation at 2,000 x g for 30 min. The supernatant was removed and replaced by 200 µl of water. The cell samples were vortexed for 5 min and sonicated (Ultrasonic Cleaner, VWR) for 30 min. Chloroform (550 µl) was added to the samples followed by 15 min sonication. Methanol (350 µl) was added to the cell pellets and incubated for 4 h with shaking at room temperature. The upper phase containing GSLs was collected after centrifugation at 2,700 x g for 20 min. Then, 400 µl of chloroform/methanol (2:1, v/v) was added, followed by adding 400 µl of methanol/water (1:1, v/v). After sonication and centrifugation, the upper phase was collected and pooled to the previous sample. The process of adding methanol/water (1:1, v/v), sonication, centrifugation and removing upper phase was repeated for another two times. In each replicate, the upper phase was collected and replaced by the same volume of methanol/water (1:1, v/v). The combined upper phases were dried under vacuum in an Eppendorf Concentrator 5301 (Eppendorf, Hamburg, Germany) at 30°C.

Before the purification of the GSLs using reverse phase (RP) SPE, the samples were dissolved in 100 µl methanol and vortexed for 10 min, followed by the addition of 100 µl water. TC18-RP-cartridges were prewashed with 2 ml of chloroform/methanol (2:1, v/v), 2 ml of methanol followed by equilibration with 2 ml methanol/water (1:1, v/v). The extracted GSLs were loaded to the cartridge 3 times and washed with 2 ml methanol/water (1:1, v/v). The GSLs were eluted from the column with 2 ml methanol and 2 ml chloroform/methanol (2:1, v/v). The samples containing the eluate were evaporated under nitrogen for 1 h and dried under vacuum in an Eppendorf Concentrator at 30°C.

**GSL-Glycan Release by EGCase I, Reduction, and Purification**

To release the glycans from the GSLs, a mixture of EGCase I (12 mU, 2 µl), EGCase I buffer (4 µl) and water (34 µl) (pH 5.2) was added to each sample and incubated for 36 h at 37°C. The released glycans were collected and loaded on TC18-RP-cartridges which had been preconditioned with 2 ml of methanol and 2 ml of water. The samples were washed with 200 ul of water and residual glycans were loaded to the cartridge. Then, 500 µl of water was added to the cartridge to wash the glycans from the column. The flow-through and wash fractions were pooled and dried in an Eppendorf Concentrator at 30°C.

The reduction was carried out with slight modifications following the same procedure as described in previous work (82). In brief, GSL-glycans were reduced to alditols in 20 µl of sodium borohydride (500 mM) in potassium hydroxide (50 mM) for 2 h at 50°C. Subsequently, 2 µl of 100% glacial acetic acid was added to neutralize the solution and quench the reaction. The desalting of GSL-glycans was performed as previously described Jensen et al. (82). Glycan alditols were eluted with 50 µl of water twice. The combined flow-through and eluate were pooled and dried under vacuum in an Eppendorf Concentrator at 30°C. The carbon SPE clean-up was performed and the purified glycan alditols were re-suspended in 20 µl of water prior to PGC nano-LC-ESI-MS/MS analysis.

**Exoglycosidase α2-3 Neuraminidase Treatment**

Briefly, O-glycans and GSL-glycans alditols were resuspended in 18 µl of Glycobuffer 1 (New England Biolabs, Ipswich, MA, USA), and split into two aliquots, one for exoglycosidase α2–3
neuraminidase digestion and the other one was used as a control. Two μL of α2-3 neuraminidase was added to the samples for digestion, and 2 μL of Glycobuffer 1 to the controls. The samples were incubated overnight at 37°C and cleaned by PGC SPE, as described earlier Jensen et al. (82).

**PGC Nano-LC-ESI-MS/MS Analysis of O-Glycans and GSL-Glycans Alditols**

The LC-MS/MS analysis of O-glycan and GSL-glycan alditols was performed on a Dionex Ultimate 3000 nano-LC system equipped with a Hypercarb PGC trap column (5 μm Hypercarb Kappa, 32 μm × 30 mm, Thermo Fisher Scientific, Waltham, MA) and a Hypercarb PGC nano-column (3 μm Hypercarb Kappa, 75 μm × 100 mm, Thermo Fisher Scientific, Waltham, MA) coupled to an amaZon speed ion trap mass spectrometer (Bruker Daltonics, Bremen, Germany). Mobile phase A consisted of 10 mM ammonium bicarbonate. Mobile phase B was 60% (v/v) acetonitrile/10 mM ammonium bicarbonate. To analyze glycans, 2 μl injections were performed and separation was achieved with a gradient of B (1–51% at 0.5%/min for O-glycan and 1–71% at 0.7%/min for GSL-glycans) followed by a 10-min wash step using 95% of B at a flow of rate of 0.6 μl/min. MS scans from m/z 340 to 1800 were recorded in enhanced mode using negative ion mode. MS/MS spectra were recorded using the top 3 highest intensity peaks.

Powerful isomer separation of glycans on PGC was performed. Structures of detected glycans were studied by fragmentation analysis using MS/MS in negative mode as well as additional exoglycosidase α2-3 neuraminidase treatment (data not shown). Glycan structures were assigned on the basis of the known MS/MS fragmentation patterns in negative-ion mode (28–30), signal patterns evidence from the profiling spectra, and general glycochemical knowledge, with help of Glycoworkbench (83) and Glycomod (84) software. The total quantity of glycans released from the same number of PaTu-S and PaTu-T cells was calculated by summing peak area of all detected glycans. The relative abundances of total O-glycans and GSL-glycans were determined by normalizing the total peak area of all detected glycans in PaTu-S to 100%. Relative quantification of individual glycans was performed by normalizing the total peak area of all glycans within one sample normalizing it to 100%. Annotated glycan structures were used for calculations of structural glycan features. Relative abundances of specific glycan structures were grouped by summing relative abundances of each glycan multiplied by the number of epitopes per glycan. The glycans representing 95% of the relative abundance were identified and quantified. The bioinformatics analysis of O-glycans was performed following the standardized e-infrastructure, which has been submitted into the public repository UniCarb-DR (85).

**Flow Cytometry**

PaTu-S and PaTu-T cells were incubated with anti-glycan antibodies for 1 h at 37°C in TSM (20 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1 mM CaCl₂ and 2 mM MgCl₂), followed by a 30 min incubation with the fluorescent secondary antibody. Biotinylated lectins and human Gal-3 (5 μg/ml) were detected using Streptadivin AF 488 (Molecular Probes, Invitrogen). Recombinant human Gal-1 and human Gal-4 proteins (5 μg/ml) were incubated for 1 h at 4°C prior to antibody staining with anti-Gal-1 and anti-Gal-4 Abs, respectively. The lectins MGL-Fc (10 μg/ml), DC-SIGN-Fc (20 μg/ml), DCIR-Fc (100 μg/ml) and Dectin-1-Fc (20 μg/ml) were used in the presence or absence of 10 mM EDTA and detected using FITC-labeled anti-human-IgFc. Fluorescence was measured using a FACSscan flow cytometer (Becton Dickinson, Oxnard, CA), or a FACScalibur flow cytometer. CellQuest (BD Biosciences, San Jose, CA) or Summit software was used to determine the mean fluorescence intensity (MFI) of cell populations, after which the background fluorescence negative control (cells stained only with the secondary antibody) was subtracted from the absolute MFI values.

**Cell Adhesion Assay**

PaTu-S and PaTu-T were seeded onto a 96 well-plate at confluence. Cells were allowed to adhere overnight at 37°C + 5% CO₂ in DMEM complete medium. Binding of Calceine AM-labeled iDCs to PaTu-S and PaTu-T cells was measured in TSM with 1% BSA. Briefly, 40,000 iDCs in the presence or absence of EDTA (10 mM), were added to each well-containing either PaTu-T or PaTu-S cells and were allowed to bind for 90 min at 37°C. Subsequently, non-adherent cells were removed by gentle washing. Cells were first visualized using a Leica DMILM (Leica Microsystems Wetzlar GmbH, Rijswijk, The Netherlands) and subsequently lysed, whereafter fluorescence was measured using a FLUOstar Spectrofluorimeter (BGM Labtech, Offenburg, Germany).

**Data and Statistical Analysis**

Data are presented as mean ± SEM. Statistical analysis applied to the cell adhesion assay data was one-way ANOVA using Dunnett t-tests and for enzyme activity studies the bivariate student t-tests. Statistics on flow cytometry studies were performed using paired t-tests. Data were considered significant if p < 0.05.

**DATA AVAILABILITY STATEMENT**

The microarray datasets for this study can be found in the Functional Glycomics Gateway (http://www.functionalglycomics.org/glycomics/publicdata/microarray.jsp), Accession 2087.

**AUTHOR CONTRIBUTIONS**

TZ, AB, SV, LL, JZ, and PD performed the experiments. TZ, BT, MW, AB, and ID conceptually designed the work. TZ, MW, ID, and AB wrote the manuscript. All authors read and commented on the manuscript.

**FUNDING**

This work was supported by the Portuguese Foundation for Science and Technology [grant number SFRH/BD/44820/2008 to...
ACKNOWLEDGMENTS

We thank Lana Scharfer and Steven R. Head and the Consortium for Functional Glycomics (CFG) for performing the gene array, Tineke van der Pouw Kraan for help with the TreeView program, Hakan Kalay for the synthesis of the MUC2. Chunsheng Jing, Prof. Nicolas G. Karlsson, Stephanie Holst, Katarina Madunic, and Yoann Rombouts for fruitful discussions and assistance with data interpretation.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/onc.2020.00732/full#supplementary-material

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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