Original research

A GATA6-centred gene regulatory network involving HNFs and ΔNp63 controls plasticity and immune escape in pancreatic cancer

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INTRODUCTION

Molecular taxonomy of tumours harbours great potential for the development of personalised medicine. In the case of pancreatic ductal adenocarcinoma (PDAC), molecular classification has revealed the existence of multiple subtypes with distinctive biological and clinical behaviour and likely specific vulnerabilities. Four PDAC...
taxonomies were proposed until now, differing in the number of subgroups and nomenclature.1–4 All classifications identified a PDAC subtype with loss of cell identity features, associated with significantly worse patient survival. This subtype, called ‘quasi-mesenchymal’,2 ‘basal-like’,3 ‘squamous’,1 ‘pure basal’8 or ‘basal A/B’9 showed rather homogeneous gene expression profiles across classifications.6 We will refer to this PDAC subtype as ‘basal’. A better understanding of the molecular drivers of this aggressive PDAC subtype would improve patients’ management in the context of an almost invariably lethal malignancy.

The transcription factor GATA6, a crucial regulator of acinar cell differentiation7 and suppressor of KRasG12D-driven tumorigenesis in mice,11 is highly expressed in the classical subtype2 and silenced through promoter methylation in squamous tumours.1 Recently published data indicated that the basal phenotype is driven by broad epigenomic reprogramming, especially at super-enhancers, controlled by ΔNp63.11–13 The shorter isoform of the TP63 transcription factor marking the basal layer of stratified epithelia. Interestingly, GATA6 itself was identified as being controlled by a superenhancer lost in basal patient-derived cells14 and EZH2-driven epigenetic silencing of GATA6 is involved in PDAC de-differentiation,15 suggesting that loss of GATA6 might be embedded in the basal programme rather than driving it.

Here, we aimed at elucidating whether loss of GATA6 is the cause or the consequence of the basal phenotype in PDAC. By combining the analysis of patient-derived samples and transcriptomics datasets, in vitro experiments with PDAC cells, and a next-generation KRasG12D-driven mouse PDAC model where Gata6 was deleted at late stages of tumorigenesis, we show that GATA6 loss is necessary, but not sufficient, for the appearance of a basal programme in PDAC. Concomitant downregulation of HNF1A and HNF4A is required for the full phenotypic switch. Additionally, Gata6 loss in preneoplastic lesions (PanINs) favoured the development of metastases in mice, possibly by promoting plasticity and immune escape of tumour cells. We demonstrate that an epithelial/progenitor transcriptional network acts as a barrier against tumour progression, and provide a molecular link between the basal gene programme in vivo and the metastatic spread in PDAC.

METHODS AND MATERIAL
All relevant methods and materials can be found in online supplemental file.

RESULTS
GATA6 loss is necessary for the expression of the basal programme
We analysed five PDAC transcriptomic datasets with molecular classification, which revealed that GATA6 expression was consistently lower in the poorly differentiated subtypes (quasi-mesenchymal2 p=0.008, basal-like3 p=5.54e-10, squamous1 p=1.57e-11, pure-basal8 p<2e-16, basal A/B9 p<0.0001) (figure 1A). Since ΔNp63 was suggested to drive the basal transcriptional programme in PDAC,11 12 we explored its relationship with GATA6 in 4/5 of the datasets (the Collisson was excluded due to low sample size) plus the TCGA PAAD dataset. TP63 expression was negatively correlated with GATA6 expression in 4/5 datasets (figure 1B, online supplemental figure 1A). Additionally, ΔNp63-target genes12 were significantly enriched among those upregulated in GATA6low tumours (bottom quartile) in 3/5 datasets and showed a tendency in the remaining 2 (figure 1C, online supplemental figure 1B), supporting that the ΔNp63-dependent programme is induced when GATA6 is lost. We showed previously that GATA6 loss in PDAC associates with ectopic expression of the basal marker KRT14 in a small collection of patient-derived samples.9 We measured GATA6, TP63 and KRT14 expression with immunohistochemistry (IHC) in an independent larger set of 60 formalin-fixed paraffin-embedded tissues from PDAC resections (figure 1D). GATA6 expression was lost in >10% of tumour cells in 23/60 patients (38.3%). In addition, KRT14 was exclusively expressed in GATA6low tumours (16/23, 69.6% p=1.64e-09) and TP63 expression was detected in 14/23 (60.1%) GATA6low and 10/37 (27%) GATA6high tumours (p=0.014) (online supplemental figure 2A). Of note, the GATA6low/TP63low tumours only had small foci of TP63-positive cells, which, on more detailed analysis, were found to be located in metastatic lesions containing GATA6-negative cells in 9/10 cases (online supplemental figure 2B, black arrowhead). Moreover, we compared the frequency of basal phenotypes between the top and bottom GATA6 expression quartiles (GATA6high, GATA6low) in the five PDAC datasets with classification and observed only 2/207 GATA6high/Basal cases (online supplemental figure 1C). These data strongly indicate that GATA6 loss is necessary for the expression of the basal phenotype.

To understand the hierarchical relationship between GATA6 and ΔNp63 in the regulation of the basal phenotype, we overexpressed GATA6 in BxPC3, a PDAC cell line with high levels of ΔNp63.11 12 GATA6 overexpression led to a 40% reduction of ΔNp63 protein (p=4.15e-04) and 30% reduction of the mRNA (p=0.03) (figure 1E, online supplemental figure 2D). Consistently, using reverse transcription followed by qPCR (RT-qPCR) we observed the upregulation of classical (HNF4A, CDH1, FOXA1) and downregulation of basal markers (ΔNp63, KRT14, KRT5, FAT2, S100A2, PTHLH); KRT14 was strongly reduced both at mRNA (p=6.3e-06) and protein level (p=1.42e-06) (figure 1E,F, online supplemental figure 2D). Intriguingly, we did not observe clear changes in proliferation, migration or Matrigel invasion in vitro (not shown). Conversely, ΔNp63 and KRT14 proteins were slightly induced in PaTu8988S cells (classical) after GATA6 knock-down (online supplemental figure 2E). GATA6 expression in PaTu8988S shrG6 cells was similar to the basal level in BxPC3 cells, while GATA6 overexpression in the latter was close to the endogenous expression in PaTu8988S cells (online supplemental figure 2C) indicating that the range of GATA6 expression on experimental manipulation remains within endogenous physiological levels.

A reanalysis of published RNA-Seq data12 showed that TP63 knock-out in BxPC3 cells significantly induced GATA6 expression (adj. p=0.02) while ΔNp63 overexpression in PaTu8988S cells only resulted in a small, not significant, decrease in GATA6 mRNA (online supplemental figure 2F). ChIP-Seq from the same publication showed a TP63 peak downstream of GATA6 transcription start site (not shown), possibly indicating a direct repression. The lack of a GATA6 peak in the vicinity (<10 kb) of the ΔNp63 TSS in PaTu8988S cells,12 suggests indirect regulation.
Figure 1  GATA6 loss is necessary for the expression of the basal-like programme. (A) Analysis of GATA6 mRNA expression in pancreatic ductal adenocarcinoma (PDAC) datasets with transcriptomics-based molecular classification. (B) Correlation between TP63 and GATA6 mRNA expression in the indicated PDAC datasets. (C) Enrichment of the gene set ΔNp63 target genes among the genes upregulated in GATA6low versus GATA6high tumours in the indicated datasets. (D) Representative images of GATA6low and GATA6high PDAC. H&E and immunohistochemical stainings for TP63, GATA6 and KRT14. Scale bar=200 µm. (E) Expression of GATA6, ΔNp63 (red arrowhead) and KRT14 in control (Ctrl) and GATA6-overexpressing (G6) BxPC3 cells, analysed by western blotting of whole protein lysates. Vinculin was used as loading control. (F) Expression of a set of classical and basal genes in GATA6-overexpressing BxPC3 cells compared with Ctrl cells, measured by RT-qPCR. Results are shown as mean±SD of at least n=3 biological replicates. *p>0.05. (G) Expression of GATA6, ΔNp63 (red arrowhead) and KRT14 in control (shCtrl) and GATA6-silenced (shG6) PaTu8988S cells, analysed by western blotting of whole protein lysates. GAPDH was used as loading control. FRD, false discovery rate. NES, normalised enrichment score.
The intersection between TP63 ChIP-Seq peaks in BxPC3\textsuperscript{12} and GATA6 ChIP-Seq peaks in PaTu8988S\textsuperscript{9} showed limited overlap (0.8% of GATA6 peaks, 10.7% of TP63 peaks), suggesting that the two transcription factors control separate programmes in basal versus classical cells. Interestingly, only 1 out of 3802 GATA6 distal peaks (>50 Mb from a TSS) was located on regions identified as 'Squamous elements' by Somerville\textit{et al}\textsuperscript{12} (online supplemental figure 2G). These data indicate that, while important, neither GATA6 loss nor ΔNp63 expression is sufficient to drive a full phenotypic switch in PDAC cells and support the involvement of a cooperative model of transcriptional regulation.

**GATA6 cooperates with HNF1A and HNF4A to sustain the classical phenotype**

To identify the crucial molecular events downstream of GATA6 loss, we analysed the PanCuRx dataset, including the largest series of all-stages PDAC samples and thus better representing the PDAC patient population than datasets only including resectable tumours. We compared GATA6\textsuperscript{low}/Basal (n=41) versus GATA6\textsuperscript{low}/Classical (n=21) tumours. Gene set enrichment analysis (GSEA) revealed that HNF1A and HNF4A putative target genes were enriched among the upregulated transcripts in GATA6\textsuperscript{low}/Classical tumours (figure 2A). Accordingly, HNF1A and HNF4A mRNAs were significantly higher in GATA6\textsuperscript{low}/Classical tumours, compared with the basal ones (figure 2B and online supplemental figure 3). Similarly, KRT14\textsuperscript{pos} regions of patients’ tumours showed reduced HNF4A protein levels, compared with KRT14\textsuperscript{pos} regions (figure 2C). Of note, loss of GATA6 expression was broader and more pronounced in all samples with KRT14\textsuperscript{pos} regions, compared with HNF4A reduction.

The classical/progenitor AsPC1 and SUIT2 cells\textsuperscript{12} have lower GATA6 expression than PaTu8988S, while HNF1A and especially HNF4A are high, thus they represent a model for GATA6\textsuperscript{low}/Classical tumours (online supplemental figure 2B). HNF4A knock-down was enough to induce expression of ΔNp63 in AsPC1 and in two clones of SUIT2 cells, SUIT2-007 and SUIT2-028, supporting that HNF4A represents a barrier to the basal phenotype downstream of GATA6 (figure 2D).

We previously reported the most comprehensive epigenomics data available for a collection of basal and classical patient-derived xenografts (PDX)-derived cell lines.\textsuperscript{14} We reprocessed raw data to compare the epigenetic marks over GATA6, HNF1A

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**Figure 2** Concomitant loss of HNF1A and HNF4A is required for the expression of the basal phenotype after loss of GATA6. (A) Enrichment plot of the gene sets containing putative HNF1A and HNF4A target genes when comparing basal (low-BAS) and non-basal (low-CLA) GATA6\textsuperscript{low} tumours of the PanCuRx cohort. (B) Expression of HNF1A, HNF4A and GATA6 in the different groups of patients in the PanCuRx dataset. *p<0.05, **p<0.001, ***p<0.0001. (C) Representative images of KRT14 and HNF4A staining in a pancreatic ductal adenocarcinoma (PDAC) sample. Bottom images show KRT14\textsuperscript{pos}/HNF4A\textsuperscript{high} (brown box) and KRT14\textsuperscript{pos}/HNF4A\textsuperscript{low} (orange box) regions. Scale bar: 500 µM. (D) Expression of HNF4A and TP63 in AsPC1 and SUIT-2 on HNF4A knock-down. Vinculin was used as loading control. (E) H3K27me3 distribution along the HNF4A locus in patient-derived xenografts-derived cell lines of the three categories. (F) The proposed model: GATA6 is the primary gatekeeper of the classical phenotype; HNF1A and HNF4A can block the full basal programme but, once lost, the ΔNp63-driven basal programme is fully expressed and drives PDAC progression toward metastasis. A negative feedback regulation driven by ΔNp63 might contribute to stabilise the basal phenotype. FDR, false discovery rate. MET, mesenchymal-to-epithelial transition; NES, normalised enrichment score.
and HNF4A loci in GATA6low/Basal (n=4), GATA6low/Classical (n=5) and GATA6high/Classical (n=5) cell lines. Notably, while we found a marked accumulation of the heterochromatin marker H3K27me3 on the HNF4A locus in GATA6low/Basal cells, the locus was not epigenetically silenced in GATA6low/Classical ones (figure 2E, online supplemental figure 4A). GATA6 showed a similar pattern but H3K27me3 was predominantly enriched upstream of the TSS (online supplemental figure 4A). HNF1A was not highly marked with H3K27me3 in GATA6low/Basal cells (online supplemental figure 4A).

H3K27ac and H3K4me3 patterns around the TSS of all the three genes were consistent with higher transcription in GATA6high/Classical and GATA6high/Classical cells, that is, enrichment of these two markers of active chromatin was low or absent in GATA6low/Basal, with the exception of 1.037 cells, while it was high in all other cells (online supplemental figure 4B,C). These data suggest that GATA6 and HNFs are epigenetically silenced in basal cells, while classical cells retain HNFs expression even when GATA6 is low. Importantly, although a subset of GATA6high/HNF4Alow and GATA6high/HNF4Alow tumours was present in all patient-derived datasets, none of those tumours was basal, further supporting that GATA6 is sufficient to maintain classical features and that the concomitant loss of GATA6 and HNFs is required for the basal phenotype to emerge.

Development of a next-generation mouse model to delete Gata6 in PanINs

We showed previously that Gata6 deletion at tumour initiation accelerates KRasG12D-driven pancreatic tumorigenesis. However, KRasG12D; Gata6−/− mice developed tumours that were generally well differentiated and Krt14-negative (not shown). To discriminate the effects related to tumour initiation from those related to tumour progression, we turned to a next-generation mouse model. For this purpose, we bred Gata6LoxLox mice27 with the dual recombinase mice harbouring the Pdx-1-Flp, Fsp-KrasG12D, Fsp-R26CreERT2 and R26Gale19 alleles, to generate KFC mice (KRas, Flp, Cre). This new model allows to uncouple the activation of KRasG12D expression and Gata6 deletion (figure 3A).

Flop-dependent recombination efficiency varied widely, ranging from <5% to >90% of the pancreas, and no malignant lesions were observed in pancreata having <30% of recombination, measured by IHC for the GFP reporter (not shown). We included in our analyses only mice where Flp-mediated recombination reached at least 30% of pancreatic epithelial cells. By 20 weeks of age, KFC mice developed throughout the pancreas multiple low-grade and high-grade PanIN lesions that stained positive for GFP (figure 3B,C). In Gata6 wt mice, Gata6 was detected in most epithelial cells. Occasionally, Gata6 expression was reduced in PanINs (figure 3C, black arrowhead) suggesting that spontaneous Gata6 loss might occur at this time point. We therefore administered tamoxifen (TMX) around 20 weeks of age to induce the deletion of Gata6 in KRasG12D-expressing cells and generate Gata6LateKO KFC mice. Successful TMX-induced recombination was verified by Gata6 IHC in all Gata6LateKO mice (figure 3D). GFP expression was retained in Gata6wt cells, suggesting that GFP is highly stable in pancreatic cells. Importantly, no recombination was detected in Gata6Galelox/FloX mice not receiving TMX, as assessed by Gata6 IHC (not shown), thus excluding leakiness. Mice were sacrificed at 65 weeks or when moribund. From a cohort of 82 mice, 43 were Gata6 LateKO and 39 were controls (Gata6Galelox/FloX); the latter included 28 Gata6wt and 3 Gata6wet mice receiving TMX and 8 Gata6elev/FloX mice not receiving TMX. Gata6Ctrl and Gata6LateKO mice developed highly heterogeneous tumours of widely varying sizes, and no significant difference in tumour size or density of Ki67 positive cells was observed (figure 3E–G). The experimental design did not allow for Kaplan-Meier survival analysis and we did not observe that Gata6LateKO mice became moribund significantly earlier than controls (figure 3H).

Gata6 loss in tumours favours the basal phenotype, metastases and lung tropism

We used Krt5 and Krt14 expression as a proxy for the basal phenotype in mouse PDAC, as the Tp63 staining did not give reliable results (not shown). Expression of both markers was highly concordant (p=4.64e-08, online supplemental figure 5A). The proportion of Krt5/14pos tumours was higher in Gata6LateKO mice than in Gata6Ctrl mice (28/44 (63.6%) versus 16/38 (42.1%)) (figure 4A). Importantly, among Gata6Ctrl mice, 15/16 Krt5/14pos tumours were Gata6neg (Gata6Low). When comparing tumours based on Gata6 expression, 43/63 (68.2%) Gata6neg tumours (Gata6LateKO+Gata6Loss) and only 1/19 Gata6pos (5.2%) were basal (Krt5/14pos) (p=3.4e-06, figure 4A). There was no significant difference in tumour grade between Gata6Ctrl and Gata6LateKO mice, while Gata6neg and basal tumours were significantly more often of grade 2 and 3 (online supplemental figure 5B, p=0.0132 and 0.034, respectively). This ultimately confirmed that Gata6 loss is necessary but not sufficient for the expression of the basal phenotype.

Patients with basal PDACs have worse outcome. Congruently, we observed that significantly more Gata6LateKO mice had clear signs of disease progression as reflected by significantly more metastases (30/43, 69.8%) than the Gata6Ctrl controls (8/39, 20.5%) (p=8.5e-06, figure 4B). Importantly, all 8 Gata6Ctrl mice with metastases had Gata6neg tumours (primary and metastatic). Gata6neg tumours were also more proliferative, as shown by Ki67 staining (p=2.93e-04, online supplemental figure 5C). These data indicate that Gata6 is an efficient suppressor of metastasis in murine KRasG12D-driven PDAC.

Among the Gata6LateKO mice, 15/30 (50%) had only lung metastases, 4/30 (13.3%) had only liver metastases and 11/30 (36.7%) had both (figure 4C). This result differs from findings in patients, where the liver is the most common site of metastases.68 There is evidence that tumour cells are heterogeneous and must, in addition, be highly plastic to form metastases.11,22 This degree of plasticity influences the organotropism of PDAC metastatic cells, whereby cells that cannot fully revert the epithelial phenotype colonise preferentially the lungs.23 Among Gata6LateKO mice, liver metastases were significantly more often E-cadherinpos than lung metastases as detected by IHC (16/19, 84.2% E-cadherinpos LiMet; 2/30, 6.7% E-cadherinpos LuMet, p=3.69e-08) (figure 4D, online supplemental figure 5D). This observation is consistent with data showing undetectable E-cadherin in primary cells derived from a lung metastasis24 and might indicate that Gata6LateKO cells are more limited in their ability to efficiently reactivate the epithelial programme.

Gata6LateKO primary tumour cells are more proliferative and chemoresistant

We successfully established primary cell lines from Gata6pos (n=5), Gata6LateKO (n=15) and Gata6Loss (n=6) tumours. While Gata6pos and Gata6LateKO cell lines were homogeneously positive and negative for Gata6, respectively, Gata6pos lines displayed a more heterogeneous expression pattern (figure 5A). ΔNp63 mRNA was significantly higher in Gata6LateKO and Gata6Loss cells, and must, in addition, be highly plastic to form metastases.23 Among Gata6LateKO mice, liver metastases were significantly more often E-cadherinpos than lung metastases as detected by IHC (16/19, 84.2% E-cadherinpos LiMet; 2/30, 6.7% E-cadherinpos LuMet, p=3.69e-08) (figure 4D, online supplemental figure 5D). This observation is consistent with data showing undetectable E-cadherin in primary cells derived from a lung metastasis.24 and might indicate that Gata6LateKO cells are more limited in their ability to efficiently reactivate the epithelial programme.

Pancreas

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A next-generation mouse model for conditional Gata6 deletion. (A) Schematic representation of the alleles used to generate the Gata6^{LateKO} mouse model. (B) Representative H&E image of the pancreas of a 20-week Pdx1-Flp and KRas^{G12D} mouse. Scale bar: 2 mm. (C) Images showing H&E and expression of GFP and Gata6, in a magnified region of the pancreas shown in B (dotted square). Black arrowhead: cells with lower Gata6. Scale bar: 100 µM. (D) Representative images of a Gata6^{LateKO} pancreas after tamoxifen administration, showing H&E, GFP and Gata6 expression. Scale bar: 100 µM. (E) Representative images of the pancreas of two Gata6^{Ctrl} and two Gata6^{LateKO} mice with variable tumour size. (F) Quantification of the tumour area in Gata6^{Ctrl} (n=37) and Gata6^{LateKO} (n=42) mice. (G) Quantification of the Ki67-positive cells per high-magnification field in Gata6^{Ctrl} (n=22) and Gata6^{LateKO} (n=36) mice. (H) Age at necropsy for Gata6^{Ctrl} (n=38) and Gata6^{LateKO} (n=44) mice. Statistical significance for F–H was checked with Mann-Whitney U test.

(figure 5B) and a similar trend was observed for a set of basal markers (Runx3, S100a2 and Krt14) but not classical/progenitor markers (Pdx1 and Hnf4a, online supplemental figure 6A) indicating that these cells preserve some basal features in vitro. Gata6^{LateKO} and Gata6^{Loss} cells were significantly more proliferative than Gata6^{Ctrl} cells (figure 5C). The migratory capacity of KFC cells was highly variable and no statistical differences were observed, although some of the Gata6^{LateKO} and Gata6^{Loss} cells...
Figure 4  Gata6 loss in tumours leads to a basal-like phenotype and increased metastatic potential with lung-specific tropism. (A) Expression of Gata6 and Krt14 in representative Gata6Ctrl and Gata6LateKO pancreatic ductal adenocarcinomas, detected by IHC (left) and quantification of Krt5 and Krt14 expression in tumours classified either by genotype (Gata6Ctrl n=38 and Gata6LateKO n=44) or by Gata6 expression (Gata6pos n=19 and Gata6neg n=63). *p<0.05, ***p<0.001. Scale bar: 200 µm. (B) Representative H&E images of liver and lung metastases in a Gata6Ctrl and a Gata6LateKO mouse and quantification of metastasis occurrence in Gata6Ctrl (n=38) and Gata6LateKO (n=44) mice. **p<0.01. Scale bar: 5 mm left/centre, 100 µm right. (C) Distribution of metastases to the liver (LiMet) or to the lung (LuMet) in Gata6LateKO mice. (D) Quantification of E-cadherin IHC in Gata6LateKO liver (n=19) and lung (n=30) metastases, **p<0.01.
Pancreas showed high migratory potential (online supplemental figure 6B). No difference was observed in the invasive capacity in vitro (online supplemental figure 6C).

We then investigated the contribution of epigenetic changes to our observations, since this mechanism was shown to control the emergence of the basal phenotype in PDAC cells.\textsuperscript{11 13 17} ChIP-qPCR for H3K27ac, a marker of open chromatin, showed higher enrichment on the \(\Delta Np63\) promoter in Gata6\textsuperscript{LateKO} and Gata6\textsuperscript{Loss} cells (figure 5D). No significant differences were observed for the promoters of a subset of basal (\(\text{Runx3, S100a2, Krt14}\)) and classical (\(\text{Pdx1 and Hnf4a}\)) genes (online supplemental figure 6D). Based on this analysis, Gata6 deletion does not cause widespread remodelling of chromatin accessibility in KFC cells.

Finally, we evaluated the relationship between GATA6 status and the response of tumour cells to chemotherapeutic agents commonly used for the treatment of PDAC. Patients with low GATA6\textsuperscript{low} or basal-like PDAC respond worse to 5-FU-based adjuvant treatments.\textsuperscript{9 10} Consistently, Gata6\textsuperscript{LateKO} cell lines were significantly more resistant to 5-FU than Gata6\textsuperscript{pos} cells (figure 5E). In contrast to the findings in patients, however, Gata6\textsuperscript{LateKO} cells were also more resistant to gemcitabine (figure 5E). Gata6\textsuperscript{Loss} cells had a mixed behaviour and no clear conclusion could be drawn. These observations indicate that the KFC cell line panel generated in this study recapitulates some features of the human disease, including the high inter-patient heterogeneity.

GATA6 loss favors cell plasticity and immune escape

To pinpoint the mechanism underlying Gata6 basal-suppressive and metastasis-suppressive function, we performed RNAseq analysis of Gata6\textsuperscript{pos}, Gata6\textsuperscript{LateKO} and Gata6\textsuperscript{Loss} primary tumour cells. A recent multi-omics analysis of primary mouse PDAC cells

Figure 5 Primary cells from Gata6\textsuperscript{LateKO} tumours are more proliferative and chemo-resistant in vitro. (A) Representative immunofluorescence images of primary KFC tumour cells isolated from Gata6\textsuperscript{pos}, Gata6\textsuperscript{LateKO} and Gata6\textsuperscript{Loss} mice. Top: expression of Gata6 (green). Bottom: merged Gata6 (green), E-cadherin (red) and DAPI (blue) stainings. (B) Expression of \(\Delta Np63\) measured by retrotranscription + qPCR (RT-qPCR). (C) Proliferation of primary KFC cells from the indicated groups, represented as fold increase in cell number 48 hours after seeding. Gata6\textsuperscript{pos} n=4, Gata6\textsuperscript{LateKO} n=15, Gata6\textsuperscript{Loss} n=5. (D) H3K27ac enrichment at the promoter of \(\Delta Np63\), detected by Chip-qPCR in primary KFC cells. Data are represented as % of input chromatin. Gata6\textsuperscript{pos} n=4, Gata6\textsuperscript{LateKO} n=7, Gata6\textsuperscript{Loss} n=3. (E) Graphs representing the IC50 values measured for primary KFC cells on treatment with 5-FU and Gemcitabine in cytotoxicity assays. Gata6\textsuperscript{pos} n=4, Gata6\textsuperscript{LateKO} n=8, Gata6\textsuperscript{Loss} n=6. (B–E) Each dot represents the average value of at least three independent experiments for each tumour cell line. *p<0.05, **p<0.01.
identified two transcriptomics-based clusters: a mesenchymal cluster C1 (Mes-C1) and a more epithelial cluster C2 (Epi-C2), including three subclusters C2a, C2b and C2c. Reference cell lines from that analysis were included in our experiment. KFC cells could be assigned to three clusters: C1, C2a and C2b/c. All but one Gata6pos lines fell into cluster Epi-C2b/c and one was assigned to the Epi-C2a cluster. In contrast, all lines assigned to the Mes-C1 cluster were either Gata6LateKO or Gata6Loss cells, supporting the strong anti-EMT role of GATA6 (figure 6A, online supplemental figure 7A). Eight of the Gata6LateKO cell lines included in the RNAseq analysis were isolated from primary tumours that had metastasised. Interestingly, the Mes-C1 cluster included only cells from lung-tropic tumours, while the Epi-C2b/c cluster only included cells from tumours that also generated liver metastases (figure 6A). These data further supports that PDAC cells with a strong mesenchymal phenotype colonise preferentially the lungs.

We performed GSEA with all possible comparisons. We found ‘EMT’ as the most highly enriched gene set among the genes upregulated in Gata6LateKO or Gata6Loss cells, supporting the enrichment of the ‘KRAS signaling_DN’ gene set among the upregulated genes. In contrast, the ‘Hypoxia’, ‘p53-pathway’ and ‘metabolism-related’ gene sets were downregulated (figure 6B, online supplemental figure 7B). Moreover, we also observed significant differences between Gata6LateKO and Gata6Loss cells (online supplemental figure 7B). Finally, when comparing Gata6LateKO cells with epithelial or mesenchymal features, EMT was clearly upregulated in Mes-Gata6LateKO, while two ‘MYC targets’ gene sets, cell cycle-related gene sets and DNA-repair-related ones were upregulated in Epi-Gata6LateKO (online supplemental figure 7D), confirming that Gata6LateKO cells are diverse. Therefore, heterogeneity in pancreatic cells is a defining characteristic not only of human but also of genetically engineered mice used to model this disease.

Interestingly, the major histocompatibility complex (MHC) class I genes H2-d1 and H2-k1 and the immunoproteasome gene Psmd8 were among the most significantly downregulated genes in the Gata6LateKO cells, suggesting that Gata6 loss might induce immune escape, thereby supporting higher metastatic potential.
Congruently, the gene set ‘MHCI Mediated Antigen Processing and Presentation’ was significantly enriched among genes down-regulated in the Gata6\textsuperscript{LateKO} cells (figure 6C). IHC analysis of GATA6\textsuperscript{low} patient tumours revealed a significantly decreased infiltration of CD8\textsuperscript{+} T cells compared with GATA6\textsuperscript{high} tumours (figure 6D). To expand these observations, we explored the available patient-derived datasets for evidence of GATA6 involvement in immune escape. The Puleo cohort showed the most consistent results: MHC I-mediated antigen processing and presentation and the estimated abundance of CD8\textsuperscript{+} T cells\textsuperscript{26} were significantly lower in GATA6\textsuperscript{flow} tumours (online supplemental figure 8A,B). Furthermore, the T cell checkpoint activator PDL-1 (encoded by the CD274 gene) was negatively correlated with GATA6 expression in 3/5 datasets (figure 6E and online supplemental figure 8C). Interestingly, CD274 and several genes related with antigen processing and presentation (PSMD8, PSMD9, B2M) had GATA6 peaks on the promoter in the ChIP-Seq we performed in PDAC cells,\textsuperscript{3} suggesting that GATA6 might directly regulate a subset of them. No significant decrease in tumour infiltration of CD8\textsuperscript{+} cells was observed in Gata6\textsuperscript{LateKO} mice compared with Gata6\textsuperscript{Ctrl}, but a trend was observed when comparing metastatic versus non-metastatic tumours (online supplemental figure 8D). Taken together, our data suggest that GATA6 loss in PDAC can facilitate immune escape, favouing metastasis.

**DISCUSSION**

Transcriptomic-based tumour taxonomy has revealed important differences and commonalities among PDACs. A detailed understanding of the molecular events driving the different phenotypes, particularly the highly aggressive basal one, will increase the chance of a successful translation of basic knowledge into clinical intervention.

Here, we show that loss of GATA6, a major regulator of epithelial identity, is necessary, but not sufficient, for the acquisition of the basal phenotype in patient-derived samples and in a next-generation mouse model in which Gata6 deletion was induced at the time of KRas\textsuperscript{G12D} -driven high-grade PanIN formation (Gata6\textsuperscript{LateKO}).

Multiple lines of evidence link GATA6 loss to the basal phenotype.\textsuperscript{1} Two studies on the mouse Kras\textsuperscript{G12D} organoid model revealed a significant decrease in the infiltration of CD8\textsuperscript{+} cells compared with GATA6 high patient tumours (online supplemental figure 8A,B). Furthermore, the T cell checkpoint activator PDL-1 (encoded by the CD274 gene) was negatively correlated with GATA6 expression in 3/5 datasets (figure 6E and online supplemental figure 8C). Interestingly, CD274 and several genes related with antigen processing and presentation (PSMD8, PSMD9, B2M) had GATA6 peaks on the promoter in the ChIP-Seq we performed in PDAC cells,\textsuperscript{3} suggesting that GATA6 might directly regulate a subset of them. No significant decrease in tumour infiltration of CD8\textsuperscript{+} cells was observed in Gata6\textsuperscript{LateKO} mice compared with Gata6\textsuperscript{Ctrl}, but a trend was observed when comparing metastatic versus non-metastatic tumours (online supplemental figure 8D). Taken together, our data suggest that GATA6 loss in PDAC can facilitate immune escape, favouing metastasis.

While the in vivo findings and the correlations observed in patient-derived samples were highly consistent, in vitro modulation of GATA6 and ΔNp63 expression in cell lines yielded variable results suggesting highly context-dependent effects including roles for the stroma and the immune system. In particular, BxPC3 are KRAS wt, thus representing a rare subset of patients with PDAC. The different behaviour we observed in BxPC3 (in this work) and L3.6pl cells (in our previous work\textsuperscript{7}) after GATA6 overexpression might reflect the contribution of mutant KRAS. These differences might indicate that BxPC3 and L3.6pl cell lines do not faithfully represent the complexity of the basal-like PDACs. Indeed, only 3/6 PDX originally defined as basal-like\textsuperscript{14} shared the H3K27ac pattern of BxPC3 and L3.6pl.\textsuperscript{11} Our panel of primary mouse tumour cell lines represents a valuable tool for understanding the basal phenotype, adding to the PDX-derived cells described previously.\textsuperscript{14} Repression of basality is intrinsic in the epithelial phenotype governed by GATA6, HNFs, and likely other transcription factors. It remains to be determined whether the reconstitution of the lost repressive barriers would revert the phenotype, once it is established. Different levels of epigenetic regulation might determine such reversibility.

We additionally show that Gata6 loss dramatically increases the rate of metastasis, thus providing a molecular link between the basal programme and the metastatic potential of PDAC. A recently published transcriptomic dataset of PDAC encompassing all-stages confirmed that the basal phenotype is highly enriched among metastatic tumours.\textsuperscript{1} A thorough characterisation of primary cell lines isolated from mouse tumours revealed that Gata6 loss results in higher plasticity and possibly immune evasion, both characteristics of metastatic cells.

Cellular plasticity is one hallmark of metastatic cells. While EMT is required to initiate metastatic spread, the reverse process—MET—is necessary for the growth of metastases at distant sites and cells that cannot revert the EMT are not able to grow metastases in mouse models.\textsuperscript{21} The degree of plasticity seems to play an important role in defining the mode
of dissemination and the organotropism of metastases, with more epithelial-like cells forming metastases preferentially the liver and more mesenchymal-like cells favouring lung metastases. Gata6<sup>flx/flx</sup> mice preferentially developed lung metastases, mostly E-cadherin-negative, indicating that Gata6-KO cells might not be able to revert to a fully differentiated status. In patients, however, it is conceivable that Gata6 expression might be reactivated during tumour evolution or under the selective pressure of therapy.

Disseminating tumour cells must overcome multiple hurdles during their path to the metastatic site, among them the immune surveillance. Suppression of antigen processing and presentation is one immune evasion mechanism that tumour cells have hijacked from viruses. We observed that GATA6 loss in tumours from patients and mice decreased the expression of the antigen processing and presentation machinery and that infiltration of CD8<sup>+</sup>-positive T cells was reduced in GATA6<sup>flx</sup> tumours in patients, possibly indicating a more efficient immune evasion. Accordingly, Gata6 knockout favoured T cell-mediated tumour cell killing in an in vivo CRISPR screening, suggesting the immunogenicity gene programme is embedded within the GATA6-dependent epithelial cell identity programme. We reported similar findings for another master regulator of the epithelial cell identity, NR5A2, which actively inhibits an inflammatory gene expression programme in the normal pancreas. The polycomb repressive complex 2 (PRC) was recently shown to promote PDAC de-differentiation and metastasis through GATA6 repression and independent results indicate that PRC2 can silence the MHC class I antigen presentation pathway. These data point to an indirect role of GATA6 in modulating the antigen processing and presentation machinery, possibly through PRC2 and establish an unexpected link between identity maintenance programmes and immune recognition.

In summary, we show here that a GATA6-centred gene regulatory network functions as a gatekeeper of cell identity and blocks cell plasticity and immune evasion, thus providing a molecular link between the basal-like phenotype and metastasis. The Gata6<sup>flx/flx</sup> mouse model is therefore a valuable preclinical tool to study the most aggressive subtype of PDAC.

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**Contributors**

PM designed the study and secured funding, analysed the experiments and prepared the manuscript. BK performed most of the experiments and participated in writing the manuscript. NH managed the mouse colony and performed experiments. VI, SP and DH-B performed some experiments. JM performed some bioinformatics analyses. RO, SM and RR performed and analysed the RNA-Seq experiment. HPD and JS did the pathology assessments. MS and EG provided access to resection tissues. GAL and RAU performed the analysis of epigenetic data from PDX-derived cells. BS and DS provided the dual recombinase mouse strain. FXR provided the Gata6 flox mice and participated in the experimental design and manuscript preparation.

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**Competing interests**

None declared.

**Patient consent for publication**

Not required.

**Ethics approval**

The study was approved by the local ethics committee of the Medical University of Vienna (‘Ethikkommission’, protocol no. 1753/2014).

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**Data availability statement**

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**Supplemental material**

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