Tipifarnib, a potential adjuvant that unlocks therapeutic resistance by normalizing microenvironment

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Introduction

More than 1.5 million new cancer cases are diagnosed in US every year [1]. Unlike the normal cells that cooperate with each other to form an ecosystem within the tissue, the cancer cells break the rules by bypassing the normal proliferation, breaching the basement membrane, disseminating into circulation and finally establishing distant colonization [2]. To initiate this malignant cellular program, cancer cells hijack a variety of existing cellular mechanisms that are critical to normal tissue development and regeneration [3]. Despite the increasing knowledge on these molecular regulators leveraged by cancer cells to fulfill the aberrancy, most advanced cancers are still incurable, indicating the therapies to treat cancer have not kept up. In fact, the majority of current therapeutic regimens are designed to target cancer cells. However, the neighbors of cancer cells – especially the connective tissue or stroma – are far from passive bystanders [4]. Tumor-associated stromal cells including endothelial cells, fibroblasts and immune cells can also behave abnormally in tumor microenvironment and contribute to disease progression and therapeutic resistance [2]. An important hallmark of cancer is the dysregulation of angiogenesis, which leads to abnormal vascular network within the tumor that is filled by dilated, tortuous, and hyperpermeable vessels [5]. It has been reported that normalization of tumor vasculature by inhibiting the pro-angiogenic activity of vascular endothelial growth factor (VEGF) can improve the efficacy of chemotherapy and immune checkpoint blockade therapy [6-8]. In this short communication, we extend the concept of “vascular normalization” to “microenvironment normalization”. The underlying hypothesis is that alteration of cancer-associated stroma abates the drug delivery and therefore limits the efficacy of therapeutic regimen. Thus, normalizing tumor-associated stromal components through molecular or pharmacologic perturbation might provide a novel opportunity for cancer treatment by enhancing the efficacy of conventional therapy and immune infiltration, in addition to antiangiogenesis therapy (Figure 1).

Results

We first ask which cancer type can be the best model to test microenvironment normalization hypothesis. We think the ideal model should have high abundance of stroma components. To estimate the percent of stromal cells within the tumor, we used ESTIMATOR [9], a computational tool that uses gene expression data as input and predicts the tumor purity, to estimate the abundance of stromal cells. Across 22 different tumor types in TCGA database, we found that the pancreatic tumors are of the highest stromal scores (Figure 2a), which is consistent with previous studies [10]. Importantly, the high stroma scores correlate with poor prognosis in pancreatic cancer patients (Figure 2b). It is also notable that the progression of pancreatic cancer is accompanied by the increase of stroma, indicating that the stromal cells within malignant tissues may have evolved to cope with tumor progression (Figure 2c). Based on these observations, “microenvironment normalization” may bring the great therapeutic benefit for pancreatic cancer patients.

We next investigate which stroma-related pathways are enriched in pancreatic adenocarcinoma tissues versus normal tissues. We found the hydrolase activity is substantially increased in pancreatic adenocarcinoma (Figure 3a, b). Indeed, in pancreatic cancers, the serine hydrolase RBBP9 has been found to promote anchorage-independent growth in vitro and tumor-initiation in vivo [11]. In addition, hyaluronidase inhibitor PEGPH20 has been developed to reduce the “desmoplastic reaction” in the stage IV metastatic pancreatic cancer and is currently in the Phase II clinical trial. However, since hydrolase activity is also essential to diverse cellular processes such as myofibroblast and fetal development, systemic inhibition of hydrolase activity can cause side effect on other healthy tissues. Therefore, we queried the L1000CDS² database to search for other compounds that can reverse the hydrolase activity (Figure 4a). L1000CDS² is a dataset that integrates the LINCS L1000 small molecule expression profiles and is under development by Ma'ayan Lab at the Icahn School of Medicine at Mount Sinai [12]. The L1000CDS² dataset will search for the small molecules that can potentially reverse the custom defined gene signature. We found that the top candidate compounds target the Ras signaling (Figure 4b). Indeed, more than 50% of the pancreatic tumors carry Ras mutation (Figure 4c). And inhibiting Ras signaling in cancer cells alone may be an effective strategy to treat pancreatic cancers. Based

Figure 1. Microenvironment normalization hypothetical model. Hypothetical model describing that normalizing microenvironment can increase the efficacy of therapeutic intervention and improve prognosis outcome.

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Figure 2. Pancreatic cancer as the model to test microenvironment normalization hypothesis. a) Comparison of stroma and immune scores across 22 tumor types in TCGA dataset. The box and whisker plots showing the abundance of stromal cells, and the color key representing the percent of immune cells within the tumor. b) Kaplan-Meier curves showing that high stromal abundance in pancreatic tumors is associated with poor survival outcome. c) Comparison of stroma scores among normal, neoplasm and adenocarcinoma tissues in pancreas.

Figure 3. Pancreatic Cancer Progression is Accompanied by the Increased Hydrolase Activity. a) Heat map showing the top 15 gene signatures enriched in adenocarcinoma versus normal pancreatic tissues. Column-side annotation shows the types of the samples. b) Volcano plot showing the differential expression of the genes (n=1,327) in Regulation of Hydrolase Activity (GO:0051336). The expression difference between adenocarcinoma and normal tissues is plotted on the x axis (log scale), and the false discovery rate (adjusted significance) is plotted on the y axis (-log scale). Upregulated and downregulated genes in adenocarcinoma are labeled with red and blue, respectively.
on our bioinformatics prediction, suppressing Ras signaling may also normalize microenvironment and increase therapeutic efficacies, which may present promising strategies for potential synergy with chemotherapy and immune checkpoint therapy, respectively (Figure 4d).

Tipifarnib [13], a potent and specific farnesyltransferase inhibitor, ranks among the top compounds that can reverse the hydrolase activity in pancreatic cancer (Figure 4b). Upon treatment with Tipifarnib, the prenylation of the CxxX tail motif in Ras kinase will be blocked, which prevents Ras from binding to the membrane upon activation. As a result, Tipifarnib prominently abates the proliferation of Ras mutant cells. However, the activity of membrane-free cytoplasmic Ras is not suppressed by Tipifarnib, indicating Tipifarnib monotherapy may not be effective. This might also partially explain why Tipifarnib was suspended in Phase III clinical trials [14].

Discussion

Despite the ineffectiveness of Tipifarnib monotherapy, Tipifarnib now provides the opportunity to address the questions associated with the “microenvironment normalization” hypothesis. How does the hydrolase activity change in pancreatic cancer stroma after Tipifarnib treatment? And, how does the treatment of Tipifarnib alter pancreatic cancer microenvironment, such as hypoxia, and drug delivery efficiency? It is also notable that immune checkpoint blockade therapy works poorly for pancreatic cancer, presumably due to the low immune infiltration efficiency. It will be interesting to investigate whether the combination of anti-PD1/anti-CTLA with Tipifarnib can enhance immune cytotoxic and forge the innovative combinatorial intervention to pancreatic cancer.

Methods

TCGA RNA-seq data were downloaded from UCSC Cancer Genome Browser (https://genome-cancer.ucsc.edu/) and used as input for the R package of ESTIMATOR to estimate the stroma percent. Gene signature were calculated using single sample Gene Set Enrichment Analysis (ssgGSEA) on the server of GenePattern (https://genepattern.broadinstitute.org/gp). In silico drug screening was performing using the online graphical user interface of L1000CDS2 (http://amp.pharm.mssm.edu/L1000CDS2)

References

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