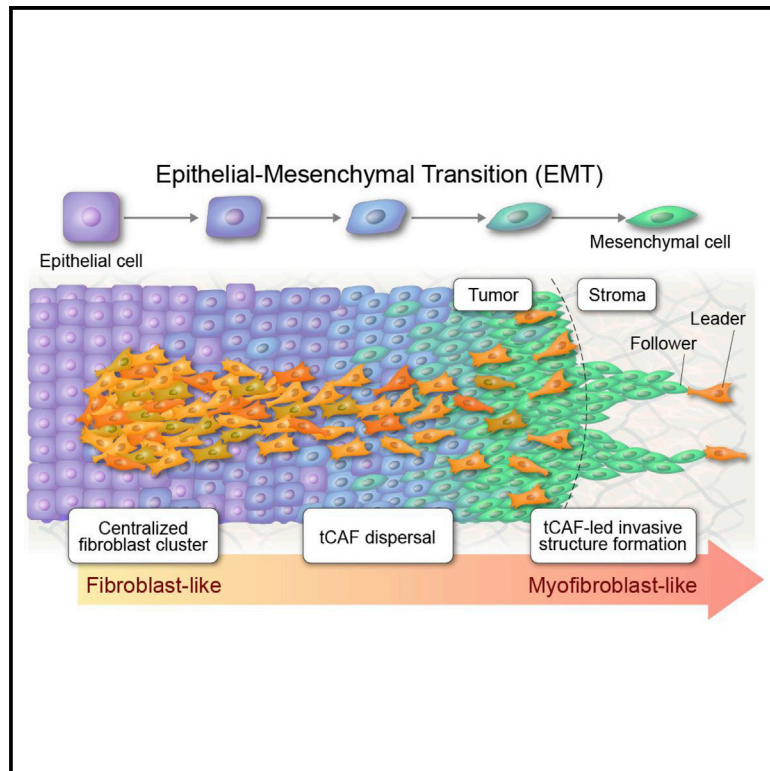


Contextual cues from cancer cells govern cancer-associated fibroblast heterogeneity

Graphical abstract



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In brief

Bota-Rabassedas et al. show that EMT in lung adenocarcinoma cells activates a secretory process that governs CAF heterogeneity and, in turn, sensitizes lung adenocarcinoma cells to pro-metastatic signals from CAFs. Thus, EMT positions lung adenocarcinoma cells at the apex of a signaling hierarchy in the tumor microenvironment.

Highlights

- Epithelial-mesenchymal transition (EMT) of cancer cells governs CAF heterogeneity
- EMT sensitizes cancer cells to pro-metastatic signals from CAFs
- EMT initiates formation of CAF-led invasive structures
- EMT triggers CAF repulsion



Article

Contextual cues from cancer cells govern cancer-associated fibroblast heterogeneity

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SUMMARY

Cancer cells function as primary architects of the tumor microenvironment. However, the molecular features of cancer cells that govern stromal cell phenotypes remain unclear. Here, we show that cancer-associated fibroblast (CAF) heterogeneity is driven by lung adenocarcinoma (LUAD) cells at either end of the epithelial-to-mesenchymal transition (EMT) spectrum. LUAD cells that have high expression of the EMT-activating transcription factor ZEB1 reprogram CAFs through a ZEB1-dependent secretory program and direct CAFs to the tips of invasive projections through a ZEB1-driven CAF repulsion process. The EMT, in turn, sensitizes LUAD cells to pro-metastatic signals from CAFs. Thus, CAFs respond to contextual cues from LUAD cells to promote metastasis.

INTRODUCTION

Accumulation of a densely fibrotic, immunosuppressive tumor stroma facilitates metastasis and is correlated with a worse clinical outcome in advanced epithelial malignancies (Kalluri, 2016; Werb and Lu, 2015). Cancer-associated fibroblasts (CAFs) are phenotypically heterogeneous mesenchymal cells that originate from diverse cell types and are a major source of cytokines and extracellular matrix molecules that enhance tumor cell-invasive activity, recruit vasculature, and suppress anti-tumor immunity

(Gascard and Tlsty, 2016; Kalluri, 2016). In line with these findings, single-cell RNA sequencing studies have identified functional differences between CAFs and have shown that extracellular matrix production and immunoregulatory functions segregate to distinct CAF populations (Bartoschek et al., 2018; Elyada et al., 2019). Pharmacologic and genetic approaches to target CAFs in preclinical models result in widely disparate effects that range from tumor suppression to tumor promotion (Feig et al., 2013; Kraman et al., 2010; Loeffler et al., 2006; Olive et al., 2009; Özdemir et al., 2014; Provenzano et al., 2012; Su



et al., 2018). Clearly, the molecular underpinnings of CAF heterogeneity must be better understood before CAF-targeting approaches can be tested in cancer patients.

There is a growing appreciation that cancer cells function as primary architects of the tumor microenvironment (Li and Stanger, 2019; Prager et al., 2019). Oncogenic mutations and epigenetic events in cancer cells upregulate immunoregulatory molecules and activate cytokine secretion, leading to the creation of an immunosuppressive milieu marked by the presence of pro-tumorigenic lymphoid and myeloid cell populations and reductions in anti-tumor CD8⁺ T cells (Li et al., 2018; Spranger and Gajewski, 2018). Cancer stem cells, which have the capacity to self-renew and undergo a reversible epithelial-to-mesenchymal transition (EMT), are capable of immune evasion and creating and maintaining tumor-promoting myeloid and T cell populations in the tumor microenvironment (Prager et al., 2019; Zhou et al., 2015). In lung adenocarcinoma (LUAD) patients, EMT features are correlated with advanced disease and worse clinical outcome (Larsen et al., 2016). Based on this conceptual framework, here, we postulated that contextual cues from LUAD cells govern CAF heterogeneity.

RESULTS

CAF heterogeneity in LUAD

To test our hypothesis, we studied CAF heterogeneity in *Kras*^{LA1} mice, which develop LUAD from somatic activation of a latent *Kras*^{G12D} allele (Johnson et al., 2001). *Kras*^{LA1} mice harbor a Thy-1⁺ CAF population (hereafter called tCAFs) that, in co-culture with *Kras*^{LA1}-derived LUAD cells, generates a cytokine-rich collagenous matrix and promotes LUAD cell invasion (Pankova et al., 2016; Roybal et al., 2011). We isolated tCAFs from *Kras*^{LA1} mice (Figure 1A) and subjected them to multiple annealing and dC-tailing-based quantitative single-cell RNA sequencing (MATQ-seq), a highly sensitive single-cell RNA-seq assay (Sheng et al., 2017). After excluding non-fibroblastic cells based on expression levels of cell-type-specific markers (Table S1), we identified two distinct tCAF clusters (Figure 1B) that were distinguishable based on 980 differentially expressed genes (Table S2). Quality control metrics showed that clustering was not related to batch-to-batch variability or sequencing depth (Figure S1). By gene set enrichment analysis (GSEA), cluster 1 was enriched in, among other terms, “EMT,” “inflammatory response,” “hypoxia,” “glycolysis,” “myogenesis,” and “angiogenesis” (Figure 1C). These terms are features of activated fibroblasts (Kalluri, 2016). In contrast, cluster 2 was enriched in “Notch signaling” and “PI3K/AKT/mTOR signaling” (Figure 1C). Thus, tCAFs contained two distinct subpopulations.

To assess whether clusters 1 and 2 are generated by contextual cues from LUAD cells, we co-cultured tCAFs with LUAD cells derived from *Kras*^{LA1};Trp53^{R172H} mice (Gibbons et al., 2009). These murine LUAD cell lines are positioned at either end of the EMT spectrum and display variable metastatic properties (high or low) following injection into syngeneic, immunocompetent mice (Gibbons et al., 2009). Highly metastatic LUAD cell lines undergo a reversible EMT through mutual antagonism between the EMT-activating transcription factor zinc finger E-box binding homeobox 1 (ZEB1) and microRNA-200 (miR-200) family members

(Ahn et al., 2012; Gibbons et al., 2009; Tan et al., 2017, 2018; Yang et al., 2011). We generated 80-cell aggregates containing tCAFs alone or tCAFs and LUAD cells (30:50 ratio) that are mesenchymal and highly metastatic (344SQ cells) or epithelial and poorly metastatic (393P cells). After 48 h, tCAFs were isolated from the multicellular aggregates by flow sorting and subjected to bulk-cell RNA-seq (Figure 1D). tCAFs acquired distinct transcriptomic features in co-culture with 344SQ cells or 393P cells (Figure 1E; Table S2). A gene expression signature derived from 344SQ-co-cultured tCAFs was more highly correlated with cluster 1 than cluster 2 (Figure 1F), whereas a gene expression signature derived from 393P-co-cultured tCAFs was more highly correlated with cluster 2 than cluster 1 (Figure 1G). Thus, clusters 1 and 2 were distinguishable based on signatures tCAFs acquired in co-culture with LUAD cells at either end of the EMT spectrum.

LUAD cells at either end of the EMT spectrum govern tCAF heterogeneity

Based on GSEA of upregulated genes, the terms “TGF- β signaling,” “EMT,” “angiogenesis,” and “glycolysis” were higher in co-cultured than mono-cultured tCAFs (Figures 2A, left bar graph and S2A). Comparisons between tCAFs co-cultured with 344SQ cells or 393P cells revealed sharp differences; among other terms, “hypoxia,” “interferon- γ ,” “TNF- α signaling via NF- κ B,” “IL-2 STAT5 signaling,” and “inflammatory response” were higher in 344SQ-co-cultured tCAFs, whereas “angiogenesis,” “EMT,” “myogenesis,” “Notch signaling,” and “Wnt β -catenin signaling” were higher in 393P-co-cultured tCAFs (Figure 2A, center bar graph), which led us to postulate that the EMT state of LUAD cells influences the phenotypic properties of tCAFs. To test this hypothesis, we generated multicellular aggregates containing tCAFs and 344SQ cells that have been subjected to small hairpin RNA (shRNA)-mediated ZEB1 depletion, which induces a shift toward an epithelial state (Tan et al., 2017), and found that GSEA hallmarks upregulated in tCAFs co-cultured with ZEB1-deficient 344SQ cells overlap with those upregulated in 393P-co-cultured tCAFs (Figure 2A, center and right bar graphs), which supported our hypothesis.

In multicellular aggregates, cells can communicate through direct contact or soluble factor exchange. Both means of communication are thought to drive the outgrowth of myofibroblastic and pro-inflammatory CAFs in pancreatic cancer (Biffi et al., 2019; Dror et al., 2016; Mitra et al., 2012). To address the role of soluble factor exchange, we introduced tCAFs and LUAD cells (344SQ or 393P) into Boyden chambers separated by a porous filter. Based on bulk-cell RNA-seq, tCAFs acquired distinct transcriptomic features in co-culture with 344SQ cells and 393P cells (Figures 2B and S2B; Table S2). Concordance between the GSEA hallmarks that tCAFs acquired in Boyden chambers (Figure 2B) and multicellular aggregates (Figure S2A) was higher in tCAFs co-cultured with 344SQ cells than 393P cells (55% versus 26%, respectively), which suggests that soluble factor exchange played a more central role in tCAF reprogramming by 344SQ cells than by 393P cells.

EMT sensitizes LUAD cells to pro-metastatic signals from tCAFs

To assess how tCAF reprogramming influences LUAD progression, we generated orthotopic LUADs by intra-thoracic injection

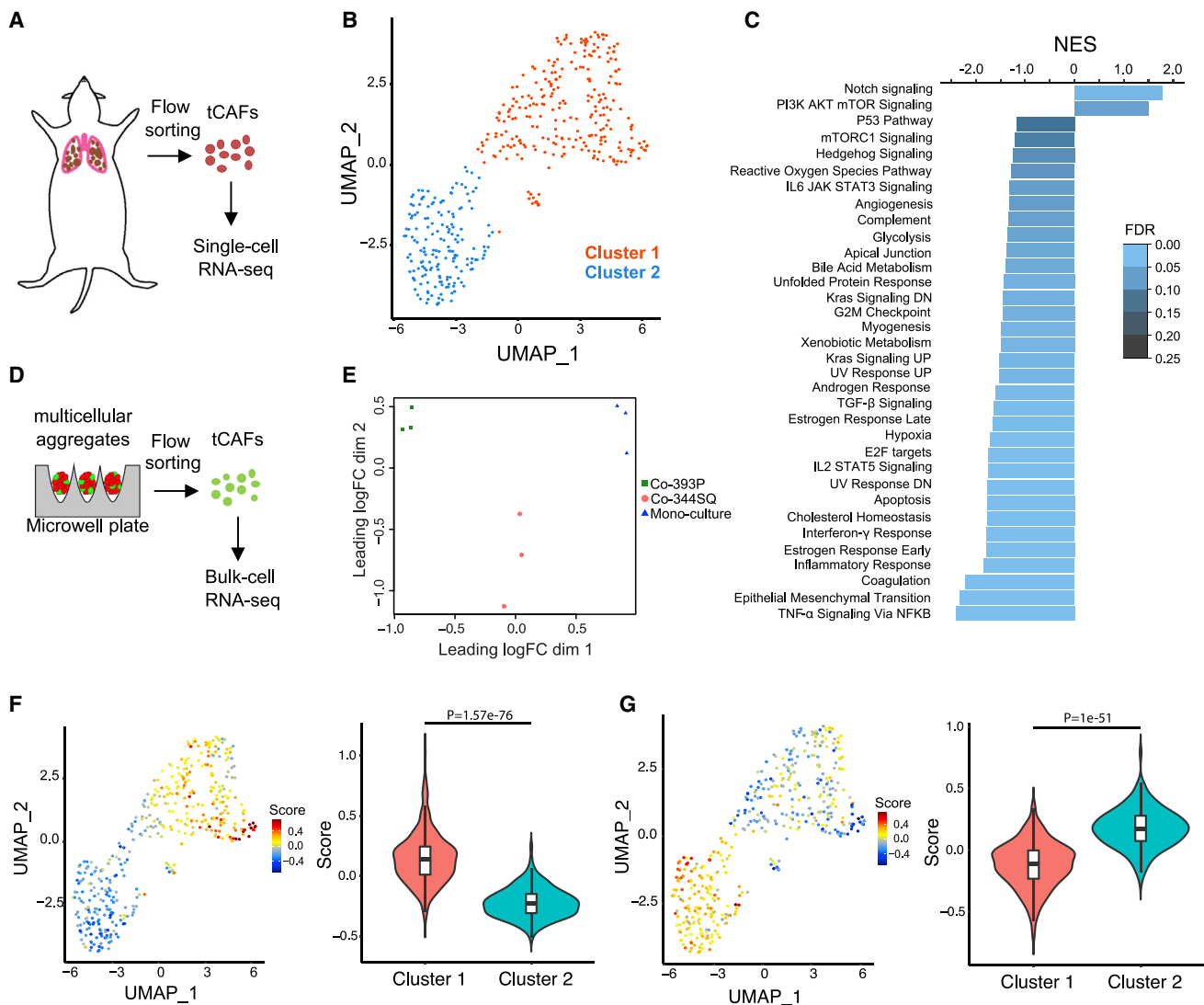


Figure 1. LUAD cells shape tCAF heterogeneity

(A) tCAFs isolated by flow sorting from lung tissues in *Kras*^{LA1} mice were subjected to single-cell RNA sequencing. (B) Principal-component analysis (PCA) plot of murine tCAFs ($n = 428$) subjected to single-cell RNA sequencing. (C) GSEA of upregulated genes in tCAFs clusters. Normalized enrichment scores (NESs) represented by bar length. False discovery rate (FDR) values color-coded. (D) Multicellular aggregates generated in microwell plates were incubated for 48 h and subjected to flow sorting to isolate tCAFs for bulk-cell RNA sequencing. $n = 3$ biological replicates per condition. (E) PCA plot of tCAFs from triplicate multicellular aggregate preparations (dots) subjected to bulk-cell RNA sequencing. (F and G) Correlations between single-cell RNA-sequencing data and expression signatures from tCAFs co-cultured with 344SQ cells (F) or 393P cells (G) are depicted with correlation plots (left) and violin plots (right).

of LUAD cells alone (344SQ or 393P, 10^6 per mouse) or in combination with tCAFs (5×10^5 of each) into syngeneic, immunocompetent mice (Figure 3A) and found that tCAFs enhanced the metastatic activity of orthotopic tumors generated by 344SQ cells but not 393P cells (Figures 3B and 3C). Based on measurements of primary tumor sizes (Figure 3B), tCAF-induced metastatic activity was not related to increased 344SQ tumor growth. To test the hypothesis that EMT sensitizes LUAD cells to pro-metastatic signals from tCAFs, we repeated the experi-

ment on ZEB1-deficient 344SQ cells and found that tCAFs did not enhance metastasis under those conditions (Figure 3D), which supported our hypothesis.

High ZEB1 levels in LUAD cells drive the formation of tCAF-led invasive structures

Notably, the co-injected tCAFs were present in regional lymphatics and distant metastatic deposits (Figures 3E and 3F), which led us to reason that tCAFs and LUAD cells have the

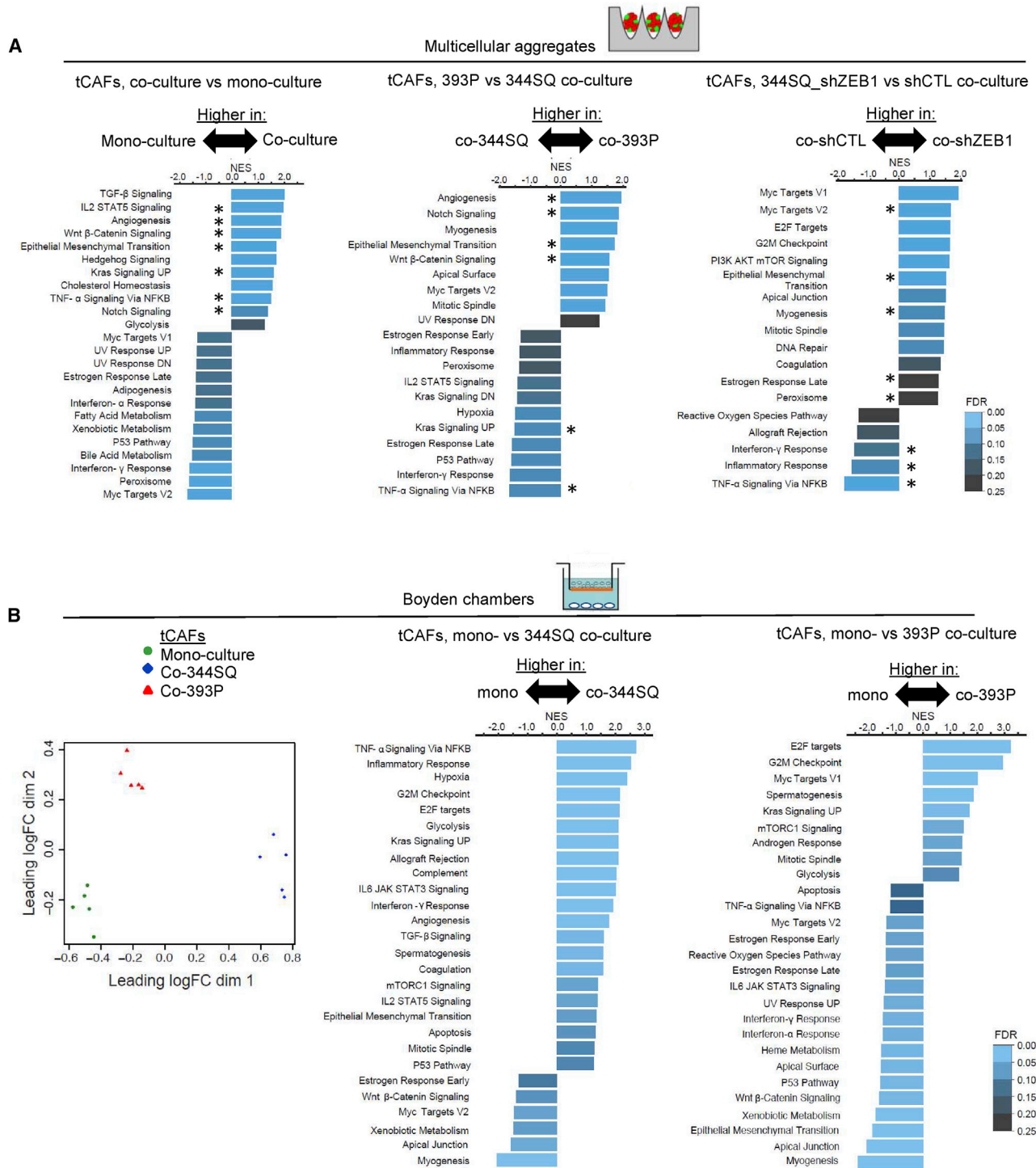


Figure 2. LUAD cells at either end of the EMT spectrum shape tCAF heterogeneity

(A) GSEA of upregulated genes in tCAFs that were mono- or co-cultured (left), co-cultured with 344SQ cells or 393P cells (center), or co-cultured with ZEB1-deficient (shZEB1) or ZEB1-replete (control shRNA [shCTL]) 344SQ cells (right) in multicellular aggregates. NES represented by bar length. FDR values color-coded. Hallmarks shared between co-cultures are indicated (asterisks).

(B) PCA plot of tCAFs from Boyden chamber preparations (dots) subjected to bulk-cell RNA sequencing. GSEA of upregulated genes in tCAFs that were mono-cultured (mono-) or co-cultured (co-) with 393P cells or 344SQ cells in Boyden chambers. NES represented by bar length. FDR values color-coded. n = 5 biological replicates per condition.

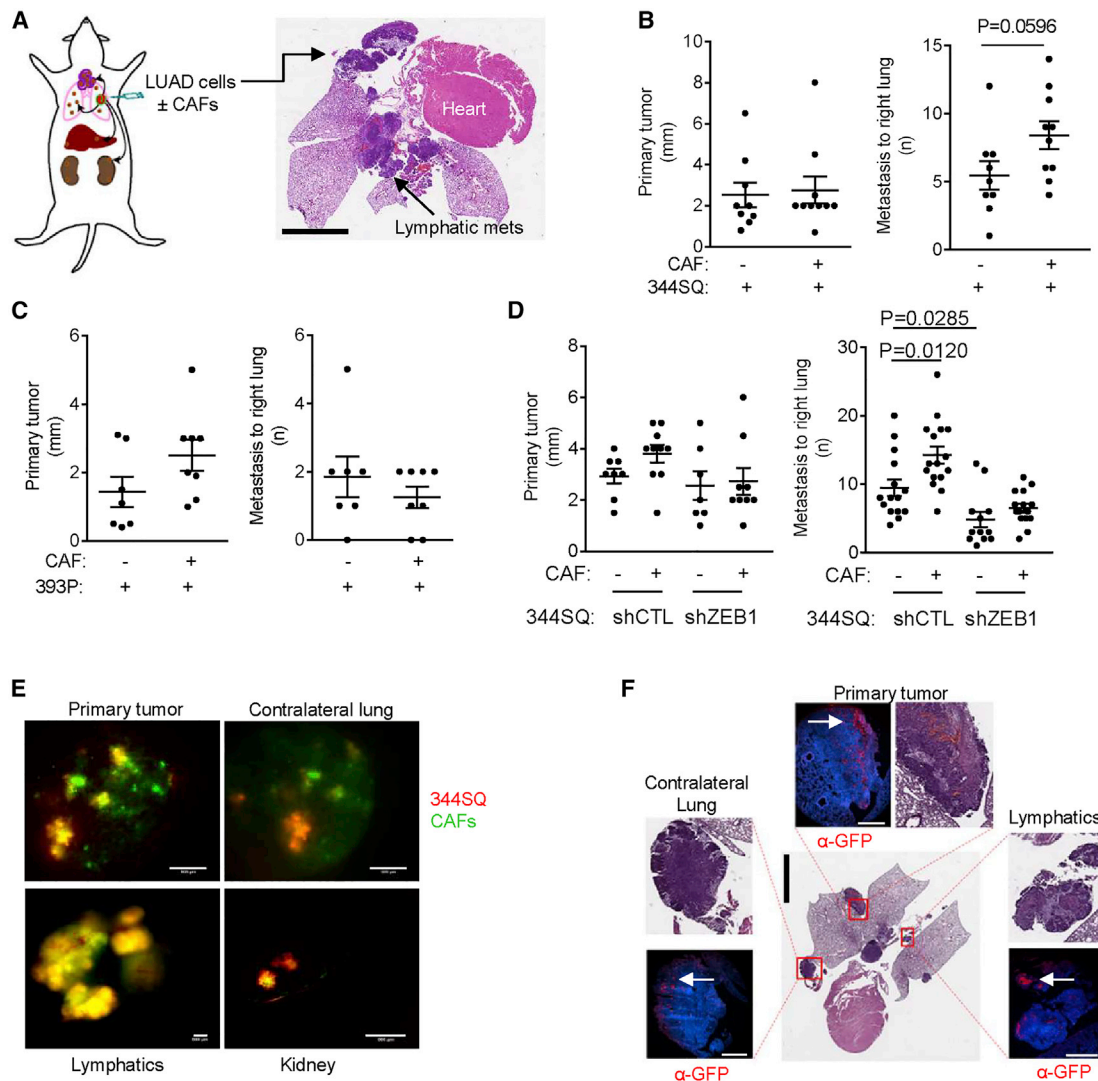


Figure 3. EMT sensitizes LUAD cells to pro-metastatic signals from tCAFs

(A) Orthotopic LUADs generated by intra-thoracic injection of LUAD cells alone or in combination with tCAFs in syngeneic, immunocompetent mice. Arrows point to sites of primary tumor and mediastinal lymph node metastasis. Scale bar, 4 mm.

(B–D) Primary tumor diameters and numbers of contralateral lung metastases per mouse (dots) injected with 344SQ cells (B), 393P cells (C), or ZEB1-deficient (shZEB1) or ZEB1-replete (shCTL) 344SQ cells (D) alone (–) or in combination with tCAFs (+). n = at least 8 mice per condition. p values, 2-sided t test, 1-way ANOVA test.

(E) Merged fluorescence micrographs of fluorescently tagged 344SQ cells (red) and tCAFs (green) in primary tumor and metastases to contralateral lung, mediastinal nodes, and kidney. Intact tissues imaged immediately after resection. Scale bars, 500 μ m. n = 2 mice.

(F) Micrographs of primary tumor and metastases to contralateral lung and mediastinal nodes in a mouse co-injected with fluorescently tagged 344SQ cells and tCAFs. Boxed regions are shown at higher magnifications (insets). Consecutive sections stained with hematoxylin and eosin or fluorescently tagged anti-GFP antibody (α -GFP) to identify co-injected tCAFs (arrows). Nuclei counterstained with DAPI (blue). Scale bars, 4 mm (central image), 500 μ m (fluorescent micrographs). n = 2 mice.

capacity to generate organized, invasive structures. To examine this possibility, we performed live-cell imaging on multicellular aggregates in 3-dimensional (3D) collagen gels (Figure 4A) and identified invasive projections containing a leading tCAF and LUAD cells following collectively behind (Figure 4B; Video S1). By time course studies, tCAF-led invasive structure formation was initiated by a tCAF that migrated radially to the aggregate periphery, exited the aggregate, and was followed by a LUAD

cell that maintained contact with the leading tCAF (Figure 4C; Video S2). Treatment with a matrix metalloproteinase inhibitor at a dose that does not inhibit cell proliferation (Figure S3) reduced the numbers of invasive projections (Figure 4D), suggesting that invasion required proteolytic activity. These findings recapitulate features of “leader-follower” structures reported by Gaggioli et al. (2007). tCAF-led invasive projections were significantly more numerous in aggregates containing mesenchymal

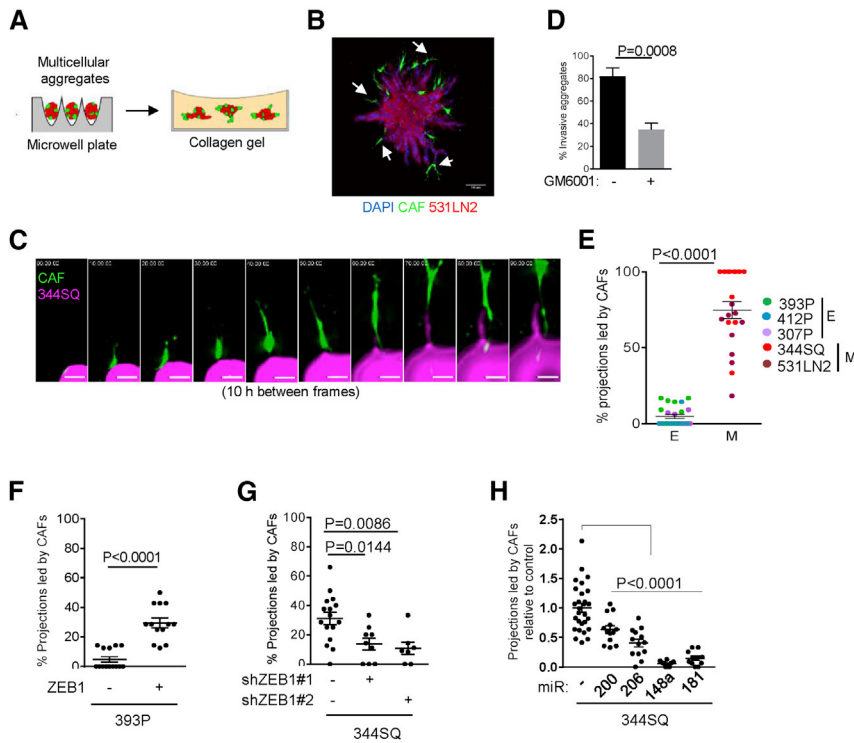


Figure 4. ZEB1 drives tCAF-led invasive structure formation

(A) Multicellular aggregates generated in microwell plates were transferred to collagen gels for imaging of invasive structure formation.

(B) Confocal micrograph of a multicellular aggregate containing tCAF-led invasive structures (arrows). Scale bar, 100 μ m.

(C) Montage of confocal micrographs from live-cell imaging of a multicellular aggregate. tCAF (green), 344SQ cells (magenta). Scale bars, 15 μ m.

(D) Percentage of aggregates that develop invasive projections in the presence or absence of protease inhibitor GM6001 (20 μ M). n = at least 25 aggregates per condition. p values, Fisher's exact test.

(E) Percentages of total projections led by tCAFs per multicellular aggregate (dots). n = at least 6 aggregates per cell line. Epithelial (E) or mesenchymal (M) LUAD cell lines. p values, 2-sided t test.

(F–H) Percentages of total projections led by tCAFs per multicellular aggregate (dots). 393P cells stably transfected with ZEB1 (+) or empty vector (–). n = at least 13 aggregates per condition (F). 344SQ cells stably transfected with ZEB1 shRNA (+) or scrambled control shRNA (–). n = at least 7 aggregates per condition. (G) 344SQ cells stably transfected with miRs or empty vector (–). (H) n = at least 11 aggregates per condition. p values, 2-sided t test, 1-way ANOVA test.

than epithelial LUAD cell lines (Figure 4E). In gain- and loss-of-function studies, ZEB1 enhanced tCAF-led invasive structure formation (Figures 4F and 4G). Based on reports that ZEB1 promotes LUAD metastasis by relieving key mediators from miR-dependent silencing (Ahn et al., 2012; Gibbons et al., 2009; Tan et al., 2017, 2018; Yang et al., 2011), we ectopically expressed ZEB1-silenced miRs (miR-200, miR-206, miR-148a, or miR-181) in 344SQ cells and found that the numbers of tCAF-led invasive structures sharply decreased (Figure 4H). Thus, miR silencing in LUAD cells plays a key role in tCAF-led invasive structure formation.

LUAD cells repel tCAFs through a ZEB1-driven secretory process

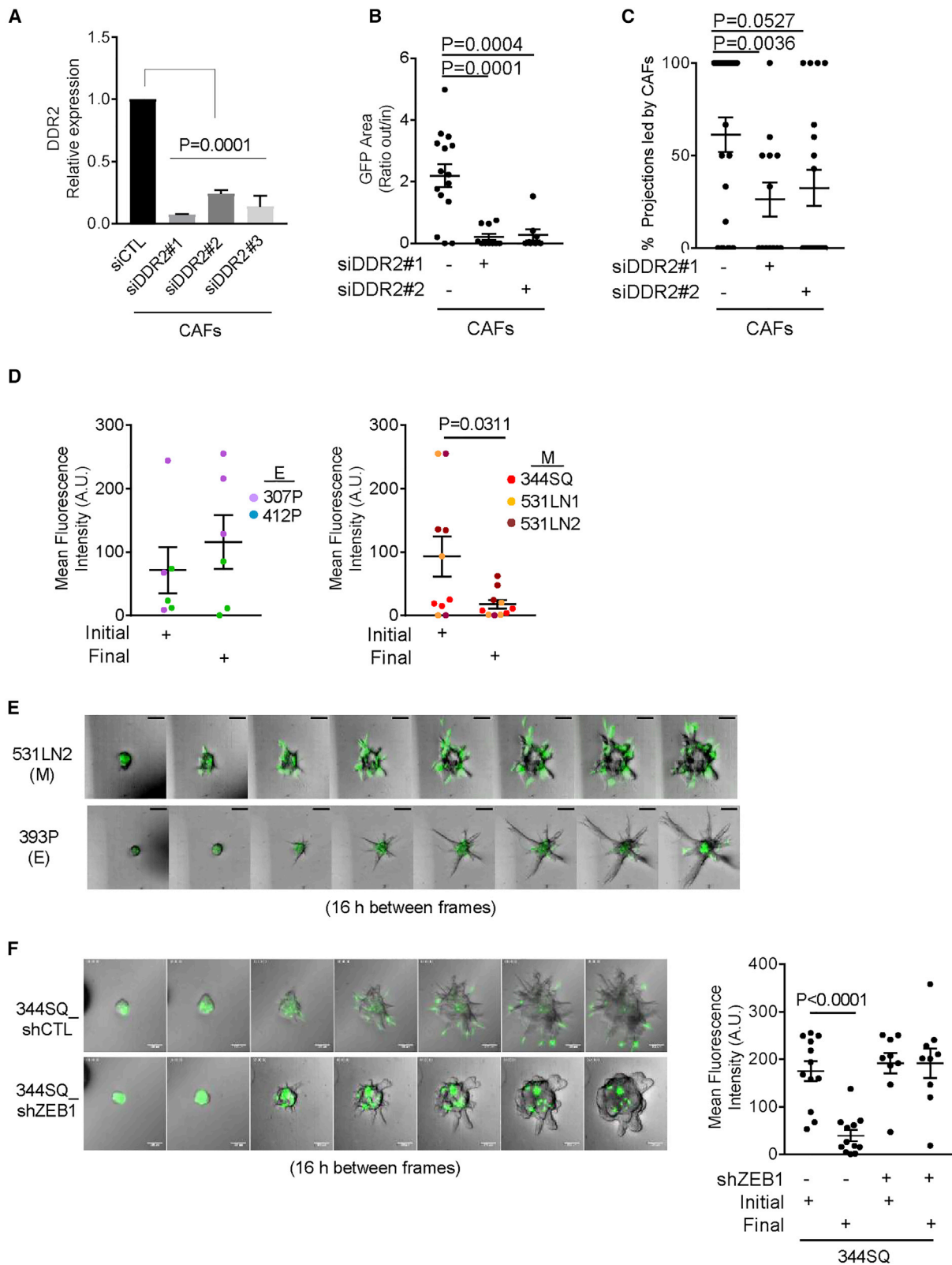
To assess the importance of tCAF motility in invasive structure formation, we subjected tCAFs to small interfering RNA (siRNA)-mediated depletion of the DDR2 collagen receptor (Figure 5A), which promotes CAF motility (Bayer et al., 2019), and found that tCAF migration to the aggregate periphery and tCAF-led invasive structure formation were reduced (Figures 5B and 5C). To determine whether tCAF motility was related to the EMT state of LUAD cells, we quantified radial tCAF movement in aggregates containing LUAD cells at either end of the EMT spectrum and found that movement was greater in aggregates containing mesenchymal than epithelial LUAD cells (Figures 5D and 5E; Video S3) and was impaired by shRNA-mediated ZEB1 depletion in 344SQ cells (Figure 5F). To determine whether high ZEB1 expression causes LUAD cells to repel tCAFs, we assessed the ability of tCAFs and LUAD cells to mix across a scratch wound and found that tCAFs mixed to a greater extent with epithelial than mesenchymal LUAD cells (Figure 6A;

Videos S4 and S5). In gain- and loss-of-function studies, ZEB1 impaired LUAD cell mixing with tCAFs (Figures 6B and 6C).

Cancer cells secrete cytokines and extracellular matrix molecules that can attract or repulse cells in the tumor microenvironment (Lu et al., 2012). To identify factors secreted by 344SQ cells that mediate tCAF repulsion, we performed liquid chromatography-mass spectrometry on conditioned medium samples isolated from LUAD cells (344SQ or 393P) and tCAFs in monoculture or co-culture. Of the 1,296 total proteins identified, numerous collagens, matricellular proteins, and proteases were present at higher levels in 344SQ cells (Figure S4), including inter-alpha-trypsin inhibitor heavy chain 2 (ITIH2), a secreted protein that stabilizes hyaluronan, which reduces the elasticity of tumor tissue and increases interstitial fluid pressure (Whatcott et al., 2011). ITIH2 mRNA levels were higher in 344SQ cells than 393P cells and were upregulated by ectopic ZEB1 expression in 393P cells (Figure 6D). tCAFs mixed to a greater extent with ITIH2-deficient than ITIH2-replete 344SQ cells (Figures 6E and 6F), which suggests that LUAD cells repel tCAFs through a ZEB1-dependent secretory process.

DISCUSSION

Metastasis requires a high degree of tumor cell plasticity as manifested by the capacity to switch between migratory modes, evade immune surveillance, and maintain viability under stressful conditions (Egeblad et al., 2010; Odenthal et al., 2016). In one working hypothesis, EMT confers the invasive and stem-ness properties necessary for cancer cells to dissolve junctional complexes, intravasate into vascular structures, and colonize the metastatic niche (Chockley and Keshamouni, 2016; Krebs



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et al., 2017; Puisieux et al., 2014). The findings presented here support a paradigm that incorporates the tumor microenvironment into the regulatory network of EMT-activating transcription factors and a model in which cancer cells reside at the apex of a signaling hierarchy in the tumor microenvironment (Li and Stanger, 2019).

Reports on CAF heterogeneity have primarily used droplet-based single-cell RNA-seq technologies to interrogate large numbers of fibroblastic cells (Lambrechts et al., 2018). Here, we used a highly sensitive RNA sequencing platform to explore heterogeneity within an immunophenotypically defined (Thy-1⁺) CAF population. To deconvolve tCAF complexity, we probed single-cell RNA-seq data with expression signatures identified in co-cultured and mono-cultured tCAFs and found that tCAFs could be distinguished based on these signatures. Thus, comparative studies on defined CAF populations are feasible in the context of a highly sensitive single-cell RNA-seq method, and insight into the molecular underpinnings of tCAF complexity provided the reaction coordinates needed for complexity deconvolution.

The viability of circulating cancer cells is higher if they are incorporated into stromal fragments that provide an early growth advantage to cancer cells in the metastatic niche, which is the basis for the current belief that cancer cells bring their own “soil” to the metastatic niche (Duda et al., 2010). We found that tCAFs localized in the metastatic niche and generated tCAF-led invasive structures in response to ZEB1-driven repulsive forces from LUAD cells (Figure 6F), which builds on evidence that CAF-led invasive structures are stabilized by mechanically active heterotypic contacts that generate force and enable invasion (Labernadie et al., 2017). In line with our findings, EMT activates contact inhibition of locomotion to guide cell migration during embryonic development (Scarpa et al., 2015; Stramer and Mayor, 2017).

Collagen fibers secreted by CAFs create a stiff, fibrotic tumor stroma that promotes cancer cell invasion (Bayer et al., 2019; Corsa et al., 2016; Erdogan et al., 2017; Yamauchi et al., 2018). Selective antagonists of DDR2 and collagen-binding integrins ($\alpha 1\beta 1$, $\alpha 2\beta 1$, $\alpha 10\beta 1$, $\alpha 11\beta 1$) inhibit the pro-metastatic activity of CAFs (Grither and Longmore, 2018; Raab-Westphal et al., 2017). Based on the findings presented here, clinical trials designed to inhibit CAF motility through the use of selective inhibitors of collagen receptors are warranted as part of the ongoing efforts to develop CAF-targeting therapies (Kalluri, 2016; LeBleu and Kalluri, 2018).

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.celrep.2021.109009>.

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Figure 5. Radial tCAF movement precedes tCAF-led invasive structure formation

- (A) Quantitative PCR analysis of DDR mRNA expression in tCAFs transfected with DDR2 or control (CTL) siRNAs. Values expressed relative to control siRNA (siCTL). n = 3 replicates per condition. p values, 1-way ANOVA test.
- (B) Migratory properties of DDR2-deficient and DDR2-replete tCAFs in multicellular aggregates expressed as a ratio of extra:intra-aggregate GFP fluorescence per aggregate (dot). tCAFs transfected with siDDR2 (+) or scrambled control siRNA (siCTL) (–) are indicated. n = at least 8 aggregates per condition. p values, 1-way ANOVA test.
- (C) Percentages of total projections led by tCAFs per multicellular aggregate (dots). tCAFs transfected with siDDR2 (+) or siCTL (–). n = at least 13 aggregates per condition. p values, 1-way ANOVA test.
- (D) tCAFs in the center of aggregates (dots) containing epithelial (E) or mesenchymal (M) LUAD cell lines were quantified at T = 0 (initial) and 4d (final). n = 3 movies per cell line. Loss of fluorescence over time indicative of tCAF radial movement. LUAD cell lines color-coded. p values, 2-sided t test.
- (E and F) Montages of confocal micrographs from live-cell imaging of multicellular aggregates. tCAFs (green). Scale bars, 100 μ m. Scatterplot (F) shows GFP fluorescence at T = 0 (initial) and 4d (final) in the center of aggregate bodies (dots). Loss of fluorescence over time indicative of tCAF radial movement. n = at least 9 aggregates per condition. p values, 1-way ANOVA test.

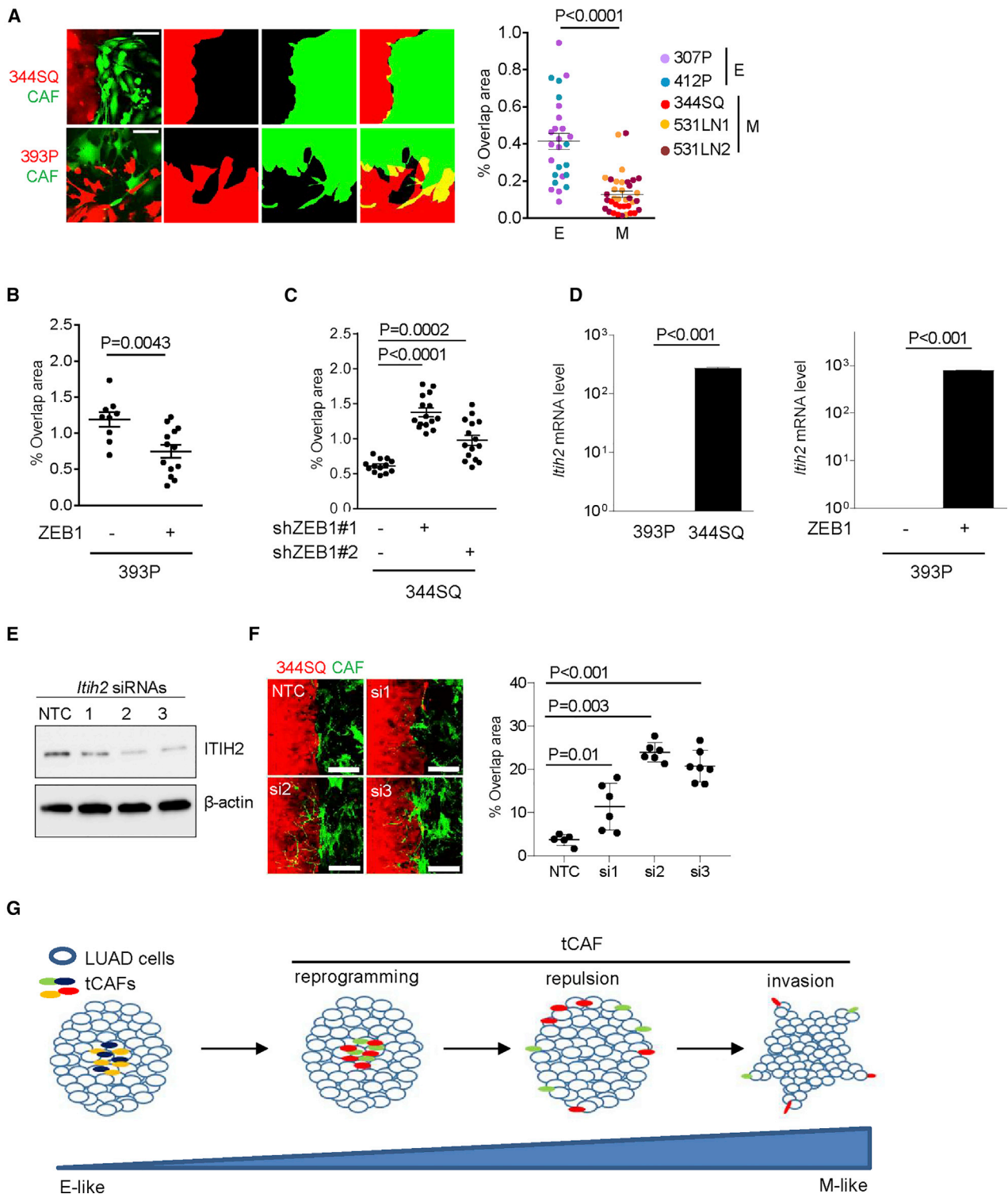


Figure 6. LUAD cells repel tCAFs through a ZEB1-driven secretory program

(A) Micrographs (far-left panels) of the leading front of a scratch wound containing tCAFs (green) and LUAD cells (red). Masks generated for each channel (2nd and 3rd panels) were overlaid (far-right panels). Scale bar, 100 μ m. Mixing was quantified based on the overlapping area per field (dots) in scratch wounds containing epithelial (E) or mesenchymal (M) LUAD cell lines (color-coded). n = 3 biological replicates per cell line. p values, 2-sided t test.

(B and C) Mixing quantified in scratch wound assays containing tCAFs and 393P cells stably transfected with ZEB1 (+) or empty vector (-) (B) or 344SQ cells transfected with shZEB1 (+) or scrambled shRNA (-) (C). n = 3 biological replicates per condition. p values, 2-sided t test, 1-way ANOVA test.

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AUTHOR CONTRIBUTIONS

N.B.-R. designed, executed, and interpreted 3D culture, cell culture, *in vivo* experiments, and performed and interpreted confocal microscopy. P.B. assisted N.B.-R. with the co-culture model design and confocal microscopy and performed and interpreted the live-cell imaging. Y.N. and J.L. performed the single-cell RNA-seq studies. K.S. performed the mini-bulk RNA-seq experiments. W.C. and Y.N. performed the bioinformatic analysis of the RNA-seq data. C.Z. supervised the single-cell RNA-seq experiments and analysis. J.A., D.W.S., and J.S.M. generated the microwell plates used in the co-culture models. Y.X., J.W., and C.J.C. analyzed the RNA-seq data. M.W. assisted with the design of the co-culture model. X.L., X.T., H.-F.G., Y.-H.A., S.L., and J.Y. assisted with the studies on cell culture and *in vivo* models. J.J.A.F. assisted with RNA-seq data handling and organization. B.L.R. and D.L.G. performed the flow cytometric analyses on orthotopic lung tumors. G.D.L. provided useful input into the experimental design and interpretation of findings from the DDR2 experiments. E.R.P., J.R.-C., B.M., L.M.S., P.V., M.G.R., H.A.C., and I.I.W. provided the data on human LUADs that were removed from the manuscript during the review process. J.M.K. conceived and supervised the project and contributed to the design and interpretation of all of the experiments.

DECLARATION OF INTERESTS

D.L.G. serves on scientific advisory committees for AstraZeneca, GlaxoSmithKline, Sanofi, and Janssen; provides consultation to Ribon Therapeutics; and receives research support from Janssen, Takeda, and AstraZeneca. I.I.W. serves on advisory boards for Genentech/Roche, Bristol-Myers Squibb, Medscape, Astra Zeneca/Medimmune, HTG Molecular, Merck, GlaxoSmithKline, and MSD and receives research support from Genentech, Oncoplex, HTG Molecular, DepArray, Merck, Bristol-Myers Squibb, Medimmune, Adaptive, Adaptimmune, EMD Serono, Pfizer, Takeda, Amgen, Karus, Johnson & Johnson, Bayer, 4D, Novartis, and Perkin-Elmer (Akoya). G.D.L. has received financial support from Pfizer-CTI. J.M.K. has received consulting fees from Halozyyme. P.B. has received consulting fees from ExpertConnect.

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(D) Quantitative PCR analysis of Itih2 mRNA levels in 393P cells and 344SQ cells (left bar graph) or 393P cells stably transfected with ZEB1 (+) or empty vector (–) (right bar graph). Values expressed relative to 393P cells and 393P_vector cells, respectively. n = 3 replicates per condition. p values, 2-sided t test.

(E) Western blot assay showing itih2 knockdown efficiency in 344SQ cells transiently transfected with control siRNA (NTC) or Itih2 siRNA (#1–3). β -Actin used as loading control. Results shown are representative of triplicate experiments.

(F) Micrographs of tCAFs (green) and siRNA-transfected 344SQ cells (red) at the leading fronts of scratch wounds. Scatterplot shows the overlapping area per field (dots). Scale bar, 100 μ m. n = 3 replicates per condition. p values, 1-way ANOVA test.

(G) Schematic illustration of a working model.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-GFP	Novus Biologicals	Cat# NB600-308; RRID:AB_10003058
Anti-Itih2	Novus Biologicals	Cat# NBP2-31750
Anti-β-Actin	Cell Signaling Technology	Clone 13E5, Cat# 4970; RRID:AB_2223172
PE anti-mouse CD45	Biolegend	Cat# 103106; RRID:AB_312971
PE Rat IgG2a, κ Isotype Ctrl	Biolegend	Cat# 400508; RRID:AB_326530
PE anti-mouse CD31	Biolegend	Cat# 102508; RRID:AB_312915
PE Rat IgG2b, kappa Isotype Ctrl	Biolegend	Cat# 400608; RRID:AB_326552
Alexa Fluor® 488 anti-mouse CD326 (Ep-CAM)	Biolegend	Cat# 118210; RRID:AB_1134099
Alexa Fluor® 488 Rat IgG2a, κ Isotype Ctrl	Biolegend	Clone RTK2758; Cat# 400525
PE/Cy7 anti-mouse Ly-6A/E (Sca-1)	Biolegend	Cat# 108114; RRID:AB_493596
PE/Cy7 Rat IgG2a, κ Isotype Ctrl	Biolegend	Cat# 400522; RRID:AB_326542
CD90 / Thy1 antibody [G7]	Abcam	Cat# ab25322; RRID:AB_470438
Rat IgG2b kappa Isotype Control (eB149/10H5), APC-eFluor 780	Thermo Fisher Scientific	Cat# 47-4031-80; RRID:AB_1272021
Anti-rabbit IgG, HRP-linked Antibody	Cell Signaling Technology	Cat# 7074; RRID: AB_2099233
Donkey anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor 568	Thermo Fisher Scientific	Cat# A10042; RRID:AB_2534017
Alexa Fluor 568 Phalloidin	Thermo Fischer Scientific	Cat# A12380
Bacterial and virus strains		
Bacteria: Subcloning Efficiency DH5α Competent Cells	Thermo Fischer Scientific	Cat# 18265017
Biological samples		
Human Lung Adenocarcinoma tissue	UT MD Anderson Cancer Center	N/A
Human IPF tissue	The University of California San Francisco	N/A
Chemicals, peptides, and recombinant proteins		
DAPI	Thermo Fischer Scientific	Cat# R37606
Collagenase type I (CLSS-1 filtered)	Worthington Biochemical	Cat# LS004216
Dispase II	Roche	Cat# 04942078001
RBC buffer	Biolegend	Cat# 420301
Puromycin	InvivoGene	Cat#ant-pr
Blasticidin	InvivoGene	Cat# ant-bl-1
Matrigel	Corning	Cat# 356231
CellTracker Deep Red Dye	Fisher Scientific	Cat# C34565
Collagen R solution 0.4%	Serva	Cat# 47256.01
Protease inhibitor GM6001	MD Millipore	Cat# 364206-1MG
paraformaldehyde	Electron Microscopy Sciences	Cat# 15714-S
JetPRIME® transfection reagent	Polyplus	Cat# 114-15
Pluronic F-127	Sigma-Aldrich	Cat# p2443-250G
PCR-grade mineral oil	Sigma-Aldrich	Cat# No. M8662
RnaseOUT	Thermo Fischer Scientific	Cat# 10777019
Superscript III	Thermo Fischer Scientific	Cat# 18080093
Bst 2.0 WarmStart® DNA Polymerase	NEB	Cat# M0538S

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
AMPure XP beads	Beckman Coulter	Cat#. A63880
Duplex-specific Nuclease	Evrogen	Cat# EA003
KAPA HIFI Hotstart Ready Mix	KAPA Biosystem	Cat# KK2601

Critical commercial assays

RNeasy Mini Kit	QIAGEN	Cat# 74106
qScript cDNA SuperMix	QuantaBio	Cat# 101414-106
WST-1	Takara	Cat# MK400
SuperSignal West Femto Maximum Sensitivity Substrate	Thermo Fischer Scientific	Cat# 34096
Nextera DNA library prep kit	Illumina	Cat# FC-121-1030
NextSeq 500/550 High Output Kit v2.5 (150 Cycles)	Illumina	Cat# 20024907
NextSeq 500/550 Mid Output Kit v2.5 (150 Cycles)	Illumina	Cat# 20024904
NEBNext Poly(A) mRNA Magnetic Isolation Module	NEB	Cat# E7490
NEBNext Ultra II RNA library Prep Kit	NEB	Cat# E7770

Deposited data

Raw and analyzed data	This paper	GEO: GSE166480
NCBI Gene Expression Omnibus database	Tian et al., 2020	GEO: GSE136904
Mouse reference genome NCBI build 38, mm10	Genome Reference Consortium	https://www.ncbi.nlm.nih.gov/grc/mouse

Experimental models: cell lines

Mouse: 344SQ, 393P, 531LN1, 531LN2, 307P, 412P, and their transfected derivatives	Gibbons et al., 2009	N/A
Mouse: 344SQ_RFP	Padhye et al., 2019	N/A
Mouse: 344SQ_shCtL, 344SQ_shZEB1, 344SQ_mir206, 344SQ_mir148a	Tan et al., 2017	N/A
Mouse: 393P_vec, 393P_ZEB1, 344SQ_miR-181	Tan et al., 2018	N/A
Mouse: 344SQ_miR-200	Ahn et al., 2012	N/A
Mouse: CAFS-GFP	This paper	N/A
Mouse: 393_RFP	This paper	N/A

Experimental models: organisms/strains

Mouse: KrasLA1/+, 129/SV	The Jackson Laboratory	N/A
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Oligonucleotides

siRNA against murine DDR2	Sigma-Aldrich	SASI_Mm01_00106702
siRNA against murine DDR2	Sigma-Aldrich	SASI_Mm01_00106703
siRNA against murine DDR2	Sigma-Aldrich	SASI_Mm01_00106704
siRNA against murine Itih2	Sigma-Aldrich	SASI_Mm01_00067716
siRNA against murine Itih2	Sigma-Aldrich	SASI_Mm01_00067717
siRNA against murine Itih2	Sigma-Aldrich	SASI_Mm01_00067718
Universal siRNA negative control #2	Sigma-Aldrich	N/A
DDR2_F: TCATCCTGTGGAGGCAGTTCTG	Sigma-Aldrich	N/A
DDR2_R: CTGTTCACTTGGTGATGAGGAGC	Sigma-Aldrich	N/A
RLP32_F: GGAGAAGGTTCAAGGGCCAG	Sigma-Aldrich	N/A
RLP32_R: TGCTC CCATAACCGATGTTTG	Sigma-Aldrich	N/A
Itih2_F: ACCAGGACACATCCTCTCAGCT	Sigma-Aldrich	N/A
Itih2_R: CAGAACCTCCGAAGTAGTTGTGG	Sigma-Aldrich	N/A
MATQ-seq primer mix	Sheng et al., 2017	N/A

Recombinant DNA

pLVX-puro	Clontech	Cat# 632164
EF-pLenti6.3-GFP	Dr. Scott lab (BCM)	N/A

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Software and algorithms		
ImageJ	Schneider et al., 2012	https://imagej.nih.gov/ij/
GraphPad Prism 7.03	GraphPad	N/A
BD FACSDiva 6.1.3 software	BD Biosciences	N/A
NIS-Elements	Nikon	N/A
R Studio	R studio	https://www.r-project.org/
GSEA 4.0.3	Subramanian et al., 2005	https://www.gsea-msigdb.org/gsea/index.jsp
MATLAB R2019a	The MathWorks, Inc.	https://www.mathworks.com/products/matlab.html
Cutadapt 1.18	Martin, 2011	https://github.com/marcelm/cutadapt/
STAR 2.5.3a	Dobin et al., 2013	https://github.com/alexdobin/STAR
Samtools 1.7	Li et al., 2009	https://github.com/samtools/
htseq-count 0.10.0	Anders et al., 2015	https://pypi.org/project/HTSeq/
seqtk 1.2-r94	Shen et al., 2016	https://github.com/lh3/seqtk
gencode.vM10.annotation.gtf	N/A	https://www.gencodegenes.org/mouse/release_M10.html
edgeR 3.26.8	Robinson et al., 2010	https://bioconductor.org/packages/release/bioc/html/edgeR.html
Other		
Transwell Boyden chambers	Thermo Scientific	Cat# 141078
Glass-bottom 35mm dishes	Mattek	Cat# P35G-1.5-14-C
Glass bottom 24-well plate	Mattek	Cat# P24G-1.5-13-F
Culture-insert 2-well, 35 mm plate	Ibidi	Cat# 81176

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Jonathan M. Kurie (jkurie@mdanderson.org)

Materials availability

Mouse lines generated during this study are completely available upon request to the Lead Contact.

Data and code availability

The raw data of single cell RNA-seq and mini-bulk RNA-seq have been submitted to GEO: GSE166480. Sequencing depth and mapping can be found in [Table S1](#).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Animal husbandry and *in vivo* experiments

All mouse studies were approved by the Institutional Animal Care and Use Committee at The University of Texas MD Anderson Cancer Center. Mice received standard care and were euthanized according to the standards set forth by the Institutional Animal Care and Use Committee. To generate orthotopic lung tumors, 2-4 months old male littermates were randomly assigned to experimental groups. An incision was made in syngeneic, immunocompetent mice between the 2nd and 3rd ribs on the left side. Cells were directly injected into the left lung, and the wound was sutured. LUAD cells (RFP-tagged 344SQ, RFP-tagged 393P, 344SQ_shCTL or 344SQ_shZEB1) were injected alone (10⁶ cells) or in combination with GFP-tagged tCAFs (5x10⁵ of each cell type). Tissues were formalin-fixed and paraffin-embedded for histologic confirmation of tumor cells by hematoxylin and eosin staining and fluorescent antibody staining. Investigators were blinded to sample identities during analysis.

Cell lines

LUAD cells generated from *Kras*^{LA1/+}; *Tp53*^{R172H/+} male mice (344SQ, 393P, 531LN1, 531LN2, 307P, 412P, and their transfected derivatives) ([Gibbons et al., 2009](#)) were cultured in RPMI 1640 containing 10% FBS. Cells were maintained at 37°C in an incubator with a

humidified atmosphere containing 5% CO₂. Cells were passaged every 2-3 days or when they reached 80% confluence. tCAFs isolated from Kras^{LA1} male mice were cultured in DMEM with 10% FBS, 1% Pyruvate, and 1% Penicillin-Streptomycin, were passaged every 3-5 days or upon reaching 75% confluence, and were used for a maximum of 5 passages after thawing. All co-culture experiments and paired mono-culture controls were performed using a mix of both cell culture mediums, at a 1:1 ratio. 393_RFP cells were generated using the vector pLVX_mCherry (puromycin selection); the mCherry coding sequence was amplified by PCR and inserted into a pLVX-puro vector (Clontech, Cat#632164). GFP-transfected tCAFs were generated with the vector EF-pLenti6.3-GFP (Blasticidin selection). The remaining stably transfected cells are described elsewhere: 344SQ_RFP (Padhye et al., 2019); 344SQ_shCtL, 344SQ_shZEB1, 344SQ_mir206, 344SQ_mir148a (Tan et al., 2017); 393P_vec, 393P_ZEB1, 344SQ_miR-181 (Tan et al., 2018), and 344SQ_miR-200 (Ahn et al., 2012). For co-culture experiments, 50,000 tCAFs were seeded into the bottom chamber of a 0.4 μm Boyden chamber with or without 80,000 LUAD cells in the upper chamber. After 48h, tCAFs were collected from the bottom chamber and subjected to RNA extraction. For transient transfection, cells were transfected with jetPRIME® (Polyplus, Cat#114-15) transfection reagent following the manufacturer's suggested protocol.

METHOD DETAILS

Cell proliferation

Viable cell densities were quantified in sub-confluent monolayer culture conditions using water-soluble tetrazolium salt-1 (WST-1, Takara Cat#MK400) reagent as suggested by manufacturer's instructions (Takara).

Fibroblast isolation for single-cell RNA sequencing analysis

To obtain fresh fibroblasts for single cell RNA-seq, we resected whole lung tissues from age-matched (10-12 months-old) Kras^{LA1} mice (for tCAF isolation). The lungs were immediately perfused with 2% FBS in Hank buffered salt solution (FBS-HBSS) and dispersed into single-cell suspensions by immersion in 3 μg/mL of collagenase (Worthington Biochemical; Cat#LS004216) and Dispase II (Roche; Cat#04942078001) on a gentleMACS Dissociator (Miltenybiotec; Cat# 130-093-235) using the lung tissue dissociation programs (Lung_01 and Lung_02). Dispersed cells were filtered (70 μm and 40 μm), centrifuged, washed with FBS-HBSS, and subjected to red blood cell lysis by adding RBC buffer (Biolegend; Cat#420301). The remaining cells were centrifuged, washed, and counted. To isolate immunophenotypically defined tCAF populations (Thy-1⁺ Epcam⁻ CD45⁻ Sca-1⁻ CD31⁻) (Roybal et al., 2011), cells were stained for 45min on ice simultaneously with anti-Epcam-FITC (Biolegend; Cat#118210), anti-CD31-PE (Biolegend; Cat#102508), anti-CD45-PE (Biolegend; Cat# 103106), anti-Sca-1-PECy7 (Biolegend; Cat#108114), and anti-Thy1-APC (Abcam; Cat# ab25322). Control tubes containing 1x10⁵ cells were stained with single antibodies or no antibody. Cells were subjected to flow cytometry (FACS Aria II instrument, BD FACSDiva 6.1.3 software) using a 75 μm nozzle. Cells were gated with the following scheme: cells, singlets, live cells, negative for: FITC (Epcam), PE (CD31 and CD45), and PECy7 (sca-1), and positive for APC (Thy1). Live/dead marker (DAPI; Thermo Fischer Scientific; Cat#R37606) was added, and only live cells were placed individually in wells on a 96-well plate. The 96-well plate was kept at 4°C before and during sorting and contained the first enzymes of the RNA isolation protocol to minimize RNA degradation. Immediately after sorting, the plate was centrifuged for 30 s to mix the cells and enzymes. The remaining steps of the RNA sequencing protocol are described below.

RNA sequencing

We used a Bravo® Automated Liquid Handler to robotically perform MATQ-seq analysis (Sheng et al., 2017). To prevent evaporation during the reactions, we first added 10 μL of PCR-grade mineral oil (Sigma, Cat. No. M8662) to every well of a 96-well plate (Bio-rad, Cat. No. HSS9601). For cell lysis, 1 μL of lysis buffer containing 0.65 μL of 0.2% Triton X-100 ultrapure water (Thermo Fisher Scientific, 10777019), 0.2 μL of MATQ-seq primer mix (Sheng et al., 2017), 0.05 μL of dNTP (10mM each), 0.05 μL 0.1M DTT, and 0.05 μL RnaseOUT (Thermo Fisher Scientific, Cat. No. 10777019) were premixed and then added to each well using the Liquid Handler. The cell lysis plate was then briefly centrifuged. Individual tCAFs were flow sorted into 96-well plates. We lysed the cells with a 3-min incubation at 72°C on a 96-well PCR machine and then quickly transferred the plate to ice for 1 min. We then added 1 μL of reverse transcription mix containing 0.4 μL 5X FS buffer, 0.1 μL 0.1 M DTT, 0.05 μL RnaseOUT, 0.05 μL Superscript III (Thermo Fisher Scientific, Cat. No. 18080093), and 0.4 μL RNase-free H₂O to each well of the plate using the Liquid Handler. We performed all the following steps of MATQ-seq using the robot and a 96-well PCR machine for temperature ramping according to the protocol previously described (Sheng et al., 2017) with minor changes in enzyme amount. We also redesigned the second strand primer to create a 3-base mismatch at the 3' end of the primer for later library preparation. For mini-bulk cell RNA sequencing analysis of tCAFs isolated from multicellular aggregates, we used MATQ-seq as previously described (Sheng et al., 2017). As a quality control step, mapping rates from single-cell and bulk RNA sequencing were quantified (Table S1).

To prepare libraries, we diluted 20 ng of amplified product into 10 μL for each cell. Samples were heated to 95°C for 30 s to denature the cDNA. We then added double strand conversion mix containing 0.6 μL 10 μM barcoded primer, 2 μL 10X Thermopol Buffer, 0.5 μL 10 mM each dNTP, 0.3 μL Bst 2.0 WarmStart® DNA Polymerase (NEB, Cat. No. M0538S), and 6.6 μL PCR grade water to each sample. Twenty cycles of 60°C 20 s, 65°C 30 s were performed to convert the cDNA into double strand and to add Illumina adaptor and barcode to the samples. 5 μL of 50mM EDTA was then added to each sample. We then pooled the samples into a single tube or several tubes according to the assigned barcode. Pooled libraries were purified with 1.2x AMPure XP beads (Beckman Coulter, Cat.

No. A63880) into 20 μ L of water. We then used the Nextera DNA library prep kit (Illumina, Cat. No. FC-121-1030) to tag 50 ng of the libraries according to the protocol provided with the kit. We purified the tagged libraries with 1.2x AMPure XP beads. We then amplified the libraries according to the Nextera protocol using Illumina sequencing adapters. A size selection was then performed using AMPure XP beads to select cDNA ranging from 250bp to 600 bp. We removed the majority of the ribosomal cDNA with a treatment of Duplex-specific Nuclease (Evrogen, Cat. No. EA003) using 100 ng of the library. We then purified the library using 0.9x AMPure XP and amplified the library using KAPA HIFI Hotstart Ready Mix (KAPA Biosystem, Cat. No. KK2601). Libraries were then diluted and sequenced on a NextSeq 500 machine, using the NextSeq 500/550 High Output Kit v2.5 (150 Cycles) or NextSeq 500/550 Mid Output Kit v2.5 (150 Cycles) (Illumina, Cat. No. 20024907 and 20024904).

Multicellular aggregates

Multicellular aggregates were created in a 24-well plate containing 1700 laser-ablated microwells per well as described (Albritton et al., 2016). The microwells were passivated with 0.05% pluronic acid (Sigma-Aldrich; Cat#p2443-250G) for 1 h prior to seeding the cells. For invasion assays in collagen gels, 85,000 LUAD cells alone or in combination with 51,000 tCAFs were seeded per well and cultured for 48 h to generate aggregates, each containing 50 LUAD cells alone or 50 LUAD cells and 30 tCAFs. tCAFs used to generate the aggregates originated from a common pool. Aggregates were mixed with rat tail collagen I (Collagen R solution 0.4%, Serva; Cat#47256.01) to generate collagen gels with defined concentrations (2mg/ml), volumes (200 μ l/gel), and aggregate densities (35 aggregates/ gel). The gels were allowed to polymerize upside-down on a glass-bottom 35mm plate (Mattek; Cat#P35G-1.5-14-C) or a glass-bottom 24-well plate (Mattek; Cat#P24G-1.5-13-F) at 37°C for 30 min. The aggregates were cultured for up to 5 days. For protease activity inhibition, aggregates were treated with GM6001 (20 μ M, MD Millipore; Cat#364206-1MG) in the collagen gel, for 3 days. Aggregates were fixed with 4% paraformaldehyde (Electron Microscopy Sciences; 15714-S) for 20 min at room temperature (RT) for confocal microscopy. For bulk-RNA sequencing studies, GFP-tagged tCAFs were isolated from multicellular aggregates (cultured for 48h) by disrupting the aggregates using a 30^{1/2} gauge needle and isolating the tCAFs by flow sorting in the green channel (FACS Aria II instrument, BD FACSDiva 6.1.3 software) using a 75 μ m nozzle. Live/dead marker (DAPI, Thermo Fischer Scientific; Cat#R37606) was added. Live cells (at least 500 per replicate) were selected and subjected to RNA extraction protocol described elsewhere in this methods section.

Liquid chromatography-mass spectrometry of conditioned medium samples

For proteomic studies, conditioned medium samples were collected from microwells containing multicellular aggregates that had been in culture for 48 h. After collection, the conditioned medium samples were filtered, concentrated, and analyzed as previously described (Tan et al., 2020).

Microscopy

Images of fixed multicellular aggregates in 3D collagen gels were acquired with a Nikon A1 confocal microscope, 10x objective. After collecting z-slices, the volume was rendered into 2-dimensional projections by maximum intensity projection algorithm (ImageJ; Schneider et al., 2012), and invasive structures were analyzed. An invasive projection was defined as at least one visible LUAD cell protruding out of the aggregate. A tCAF-led invasive structure was defined as a projection containing at least 1 tCAF at the tip. Invasive projections were manually counted. Results were determined from at least 10 multicellular aggregates per condition, at least two collagen gels per condition, and at least two independent batches of aggregates. When possible, investigators were blinded to sample identity when performing these quantifications.

Live-cell imaging was performed in an automated wide-field microscope (Nikon Eclipse TiE) with 20x objective. For 3-dimensional image stacks, Z stacks were projected by using a sum of the slices algorithm. Invasive projections were identified after thresholding to remove aggregate bodies (ImageJ). Diameters were measured after thresholding (ImageJ) to remove invasive projections from the live-cell images. Results were determined from at least 3 aggregates per condition, at least two gels per condition, and two independent batches of aggregates.

Cell-cell repulsion

Multicellular aggregates were generated, transferred to collagen gels, and subjected to time-lapse movies. Fluorescence intensity in the center of the aggregate was measured at initial (T = 0) and final (T = 4d) time points by performing radial profile analysis (ImageJ) on tCAFs in the green channel. Results represent values from at least 3 biological replicates per condition. For scratch-wound closure assays, 15,000 LUAD cells and 6,000 tCAFs were seeded on either side of a 35 mm culture-2-well insert glass-bottom dish (Ibidi; Cat#81176). tCAFs and LUAD cells were distinguishable on the basis of fluorescent tags that included GFP in tCAFs and RFP or CellTracker deep red dye (Fisher Scientific; Cat#C34565) in LUAD cells. Cells were stained with CellTracker dye for 30 min at RT at a concentration of 1.5 μ M prior to seeding. The cells were allowed to reach confluence in each chamber, at which time the insert was lifted to initiate cell migration and wound closure. Cells were imaged when the insert was lifted (initial time point, T = 0) and at the final time point (T = 4 d) using a confocal microscope (Nikon A1, 10x objective). Leading edges of the migrating cells were imaged and analyzed for area of overlap using ImageJ. In brief, 16-bit raw images were converted to 8-bit, and a mask was generated for each channel, which comprised the total area of each cell type. The two channels were overlaid and converted to RGB. The overlapping

area was measured and adjusted for the initial area of the chamber as described (Carmona-Fontaine et al., 2008). Results represent values from at least 4 images per replicate, 3 biological replicates per condition.

To quantify extrusion of DDR2-deficient and β -replete tCAFs from multicellular aggregates, we quantified GFP outside and inside each aggregate body and used those values to calculate a ratio (outside/inside GFP area). For this purpose, z stack images of para-formaldehyde-fixed aggregates in collagen gels were acquired by confocal microscopy (10X objective). Z stacks were rendered into 2-dimensional images by using a maximum intensity projection algorithm (ImageJ), converted to 8-bit, thresholded, and segmented to generate masks of red and green channels, which were used to quantify LUAD cells and tCAFs, respectively, in the aggregate body. GFP areas within and outside the aggregate body mask were determined.

tCAF detection in distant metastases

For direct detection of fluorescently tagged cells, tumor-bearing organs were resected from mice bearing orthotopic LUAD, placed into a glass bottom dish, covered with 2% FBS in 1X phosphate-buffered saline, and imaged using a wide-field Nikon Eclipse TIE microscope (NIS-Elements Software, Nikon). As a second approach, 4 μ m tissue slices from FFPE tumor-bearing lungs were stained with anti-GFP primary antibody and a fluorescently-labeled secondary antibody (anti-rabbit 568) and imaged with a wide-field Nikon Eclipse TIE microscope (NIS-Elements Software, Nikon), 10x objective.

Western blotting

Cells were lysed with RIPA buffer (Cell Signaling Technology) containing protease inhibitors (Cell Signaling Technology). Cell lysates were separated on a 4%–20% Bis-Tris gel, transferred to nitrocellulose membranes using Trans-Blot Turbo Transfer System (Bio-Rad), and then incubated with primary antibodies and HRP-conjugated secondary antibodies (Cell Signaling Technology). Protein bands were visualized with SuperSignal West Femto Maximum Sensitivity Substrate (Thermo Fisher Scientific). Antibodies against ITIH2 (Novus Biologicals, #NBP2-31750) and β -actin (Cell Signaling Technology, #4967) were purchased.

QUANTIFICATION AND STATISTICAL ANALYSIS

For MATQ data analysis, we first trimmed the unique molecular identifiers (UMI) sequence using seqtk (Shen et al., 2016). STAR (Dobin et al., 2013) was used to map the trimmed reads to the transcriptome. We annotated the mapped reads using HT-seq with gene code GRCm38 as a reference. We then parsed the reads annotation to the untrimmed reads and counted the gene expression using UMI. The expression level of every gene was then normalized to the total UMI numbers of the sample. To identify non-CAFs cells, we performed first PCA and clustering analysis by using all cells as an explorative analysis. A small proportion of cells ($n = 32$) were spreading out on the PCA plot and most of them were outliers. To investigate the underlying source of these outliers, we performed PCA analysis on these 32 cells and found that these cells are separated into three branches on the PCA. By looking into the genes on PC1 and PC2, we identified several cell lineage markers including epithelial markers (Cdh1, Krt7, Krt18, Krt19) and hematopoietic lineage markers (Laptn5, Hoxa10) expressed in these 32 cells. Additionally, we checked the expression of three markers (Cdh1, Hoxa10, Laptn5) across all cells. The cells with the highest expression level of each marker were the outlier cells. These observations suggested these outlier cells were potentially non-CAFs. Principal component analysis and clustering analysis were performed by using ‘Seurat’ R package. Genes differentially expressed between tCAF clusters were identified by using ‘limma-trend’ function in ‘limma’ R package.

For bulk-cell RNA sequencing of mono-cultured and co-cultured tCAFs, STAR (Dobin et al., 2013) was used to map the reads to the transcriptome. Reads annotation was performed using HT-seq with gene code GRCm38 as a reference. Reads normalization was performed using EdgeR (Robinson et al., 2010). To identify significantly enriched pathways and functional hallmarks, gene set enrichment analyses were performed using GSEA 4.0.3 (Subramanian et al., 2005). Gene set shuffling was used for all bulk data. For differentially expressed gene analyses, EdgeR was used for tCAFs isolated from multicellular aggregates and limma-trend was used for tCAFs co-cultured with LUAD cells in Boyden chambers.

To determine whether tCAFs were distinguishable based on gene expression signatures they acquire in co-culture with highly or poorly metastatic LUAD cells, we first performed differential expression analyses on bulk RNA sequencing data from tCAFs co-cultured with highly or poorly metastatic LUAD cells. The genes with FDR < 0.05 were selected for expression signatures of each condition. We calculated the average gene expression Z-score of each signature in each tCAF analyzed by single-cell RNA sequencing. The Z-scores were compared between clusters 1 and 2 by Student's t test.

Supplemental information

**Contextual cues from cancer cells govern
cancer-associated fibroblast heterogeneity**

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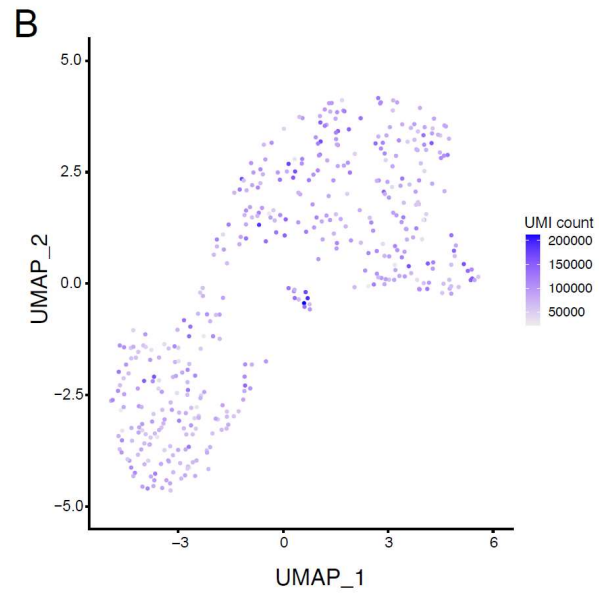
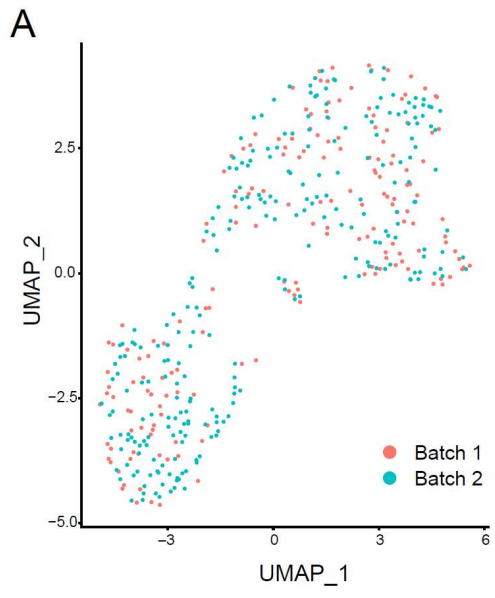
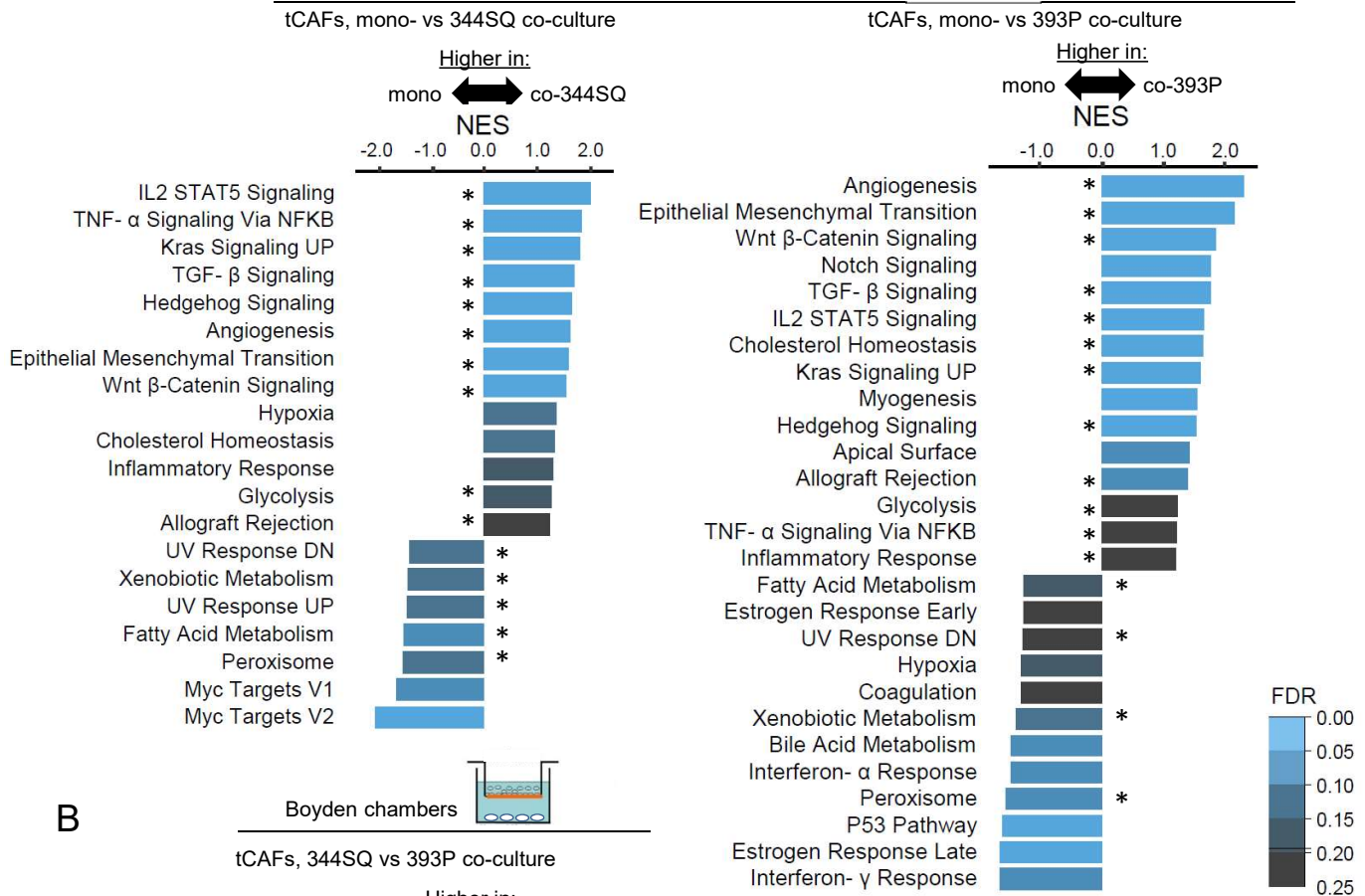
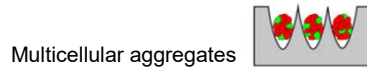
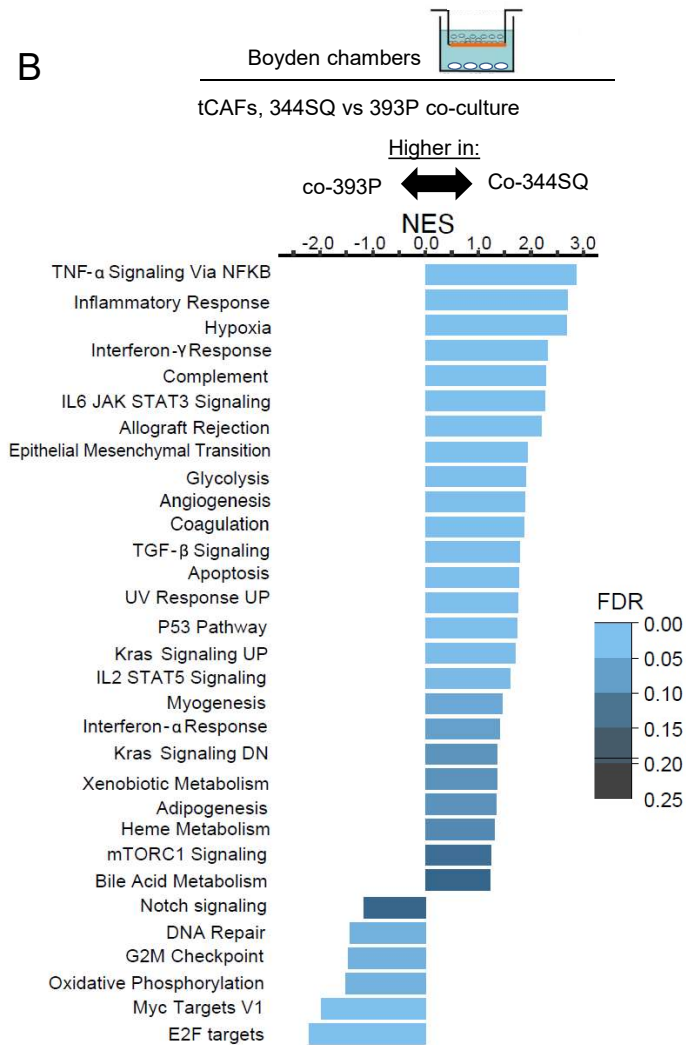


Figure S1. Single-cell RNA sequencing quality controls. Related to Figure 1. (A, B) PCA plots depict batch #1 and #2 (A) and Unique Marker Identifier (UMI) numbers in each cell (B).

A

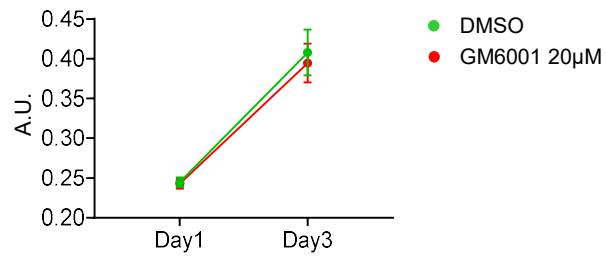


B



Supplementary Figure 2

Figure S2. LUAD cells at either end of the EMT spectrum shape tCAF heterogeneity. Related to Figure 2. (A) GSEA of upregulated genes in tCAFs that were mono-cultured (mono-) or co-cultured with 344SQ cells (co-344SQ) or 393P cells (co-393P) in multicellular aggregates. n= 3 biological replicates per condition. (B) GSEA of upregulated genes in tCAFs that were co-cultured with 344SQ cells (co-344SQ) or 393P cells (co-393P) in Boyden Chambers. NES represented by bar length. FDR values color-coded. n= 5 biological replicates per condition. Hallmarks that were shared between Boyden chambers and multicellular aggregates are indicated (A, asterisks).

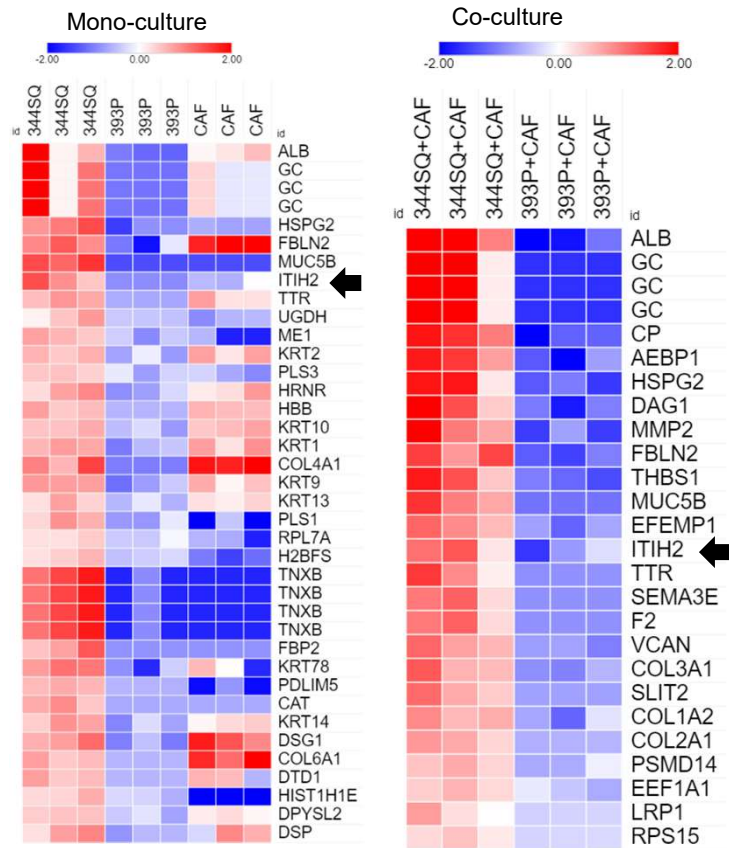


Supplementary Figure 3

Figure S3. Relative densities of tCAFs treated for indicated time periods with 20 μ M GM6001 (red) or vehicle dimethylsulfoxide (blue) determined by WST-1 assay. Related to Figure 4. A.U.: Absorbance units. n=3 biological replicates per condition.

A

LC/MS



B

Bulk-cell RNA-seq

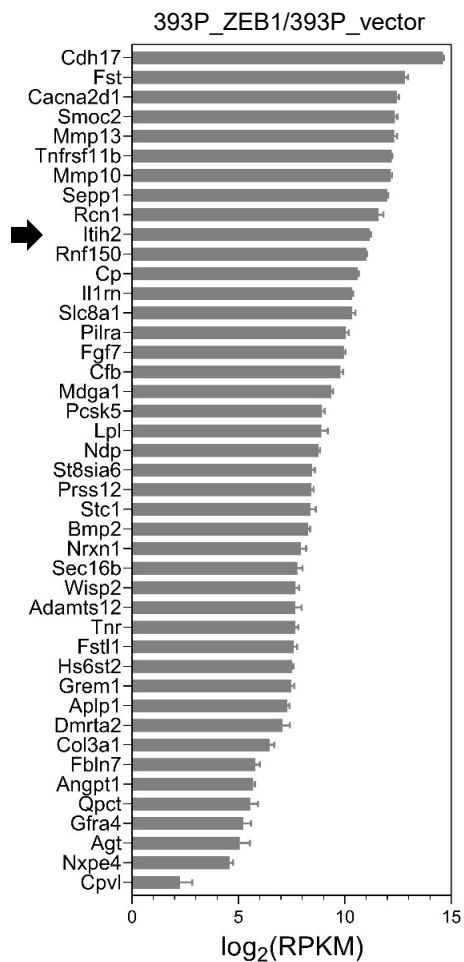


Figure S4. Identification of ZEB1-driven secreted factors. Related to Figure 6. (A) Heat map depiction of selected secreted factors identified by liquid chromatography-mass spectrometry analysis of conditioned medium samples from multicellular aggregates containing monocultured cells (left heat map) or co-cultured cells (right heat map). All proteins depicted had a P value < 0.05 (t-test). n=3 biological replicates per condition. (B) Relative levels of selected secreted factors identified by bulk-cell RNA sequencing (Tan et al., 2018). Values expressed as a ratio (393P_ZEB1/393P_vector). Log2FC: Log2 of fold change. n=3 biological replicates per condition