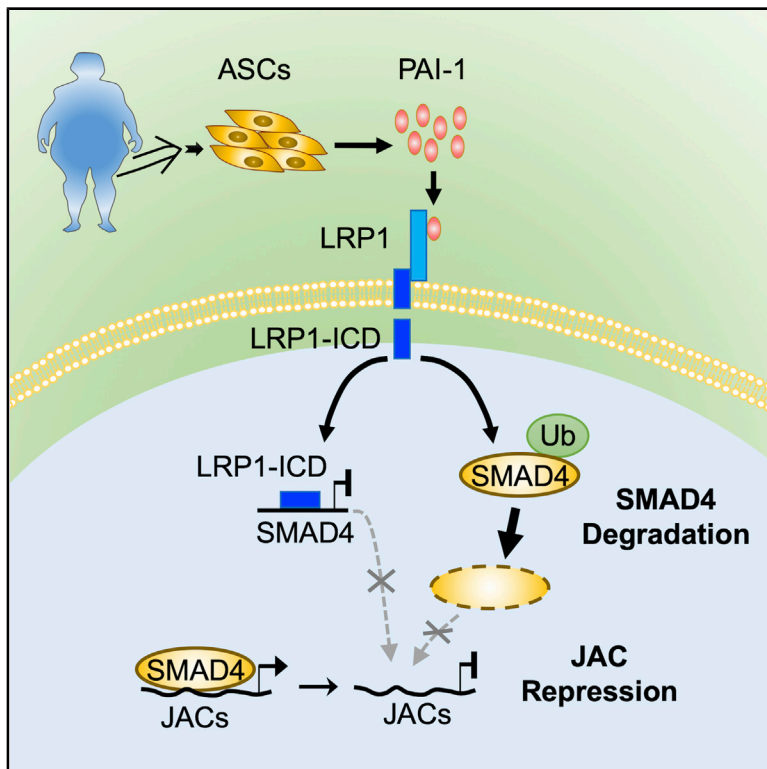


PAI-1-Dependent Inactivation of SMAD4-Modulated Junction and Adhesion Complex in Obese Endometrial Cancer

Graphical Abstract



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In Brief

Lin et al. demonstrate that PAI-1 secreted by adipose stromal cells interacts with LRP1 to repress TGF- β /SMAD4-regulated genes linked to cellular junction and adhesion complexes. This action disrupts cell-cell communication and facilitates the development of obesity-driven endometrial cancer.

Highlights

- ASC infiltration in endometrial tumor microenvironments positively correlates with BMI
- ASC-secreted PAI-1 influences heterogeneous endometrial epithelial cell subpopulations
- PAI-1-LRP1-modulated SMAD4 downregulation leads to JAC repression *in vitro*
- Repressive JAC is present in cancer cell subpopulations of persons with obesity



Article

PAI-1-Dependent Inactivation of SMAD4-Modulated Junction and Adhesion Complex in Obese Endometrial Cancer

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SUMMARY

While plasminogen activator inhibitor-1 (PAI-1) is known to potentiate cellular migration via proteolytic regulation, this adipokine is implicated as an oncogenic ligand in the tumor microenvironment. To understand the underlying paracrine mechanism, here, we conduct transcriptomic analysis of 1,898 endometrial epithelial cells (EECs) exposed and unexposed to PAI-1-secreting adipose stromal cells. The PAI-1-dependent action deregulates crosstalk among tumor-promoting and tumor-repressing pathways, including transforming growth factor β (TGF- β). When PAI-1 is tethered to lipoprotein receptor-related protein 1 (LRP1), the internalized signaling causes downregulation of SMAD4 at the transcriptional and post-translational levels that attenuates TGF- β -related transcription programs. Repression of genes encoding the junction and adhesion complex preferentially occurs in SMAD4-underexpressed EECs of persons with obesity. The findings highlight a role of PAI-1 signaling that renders ineffective intercellular communication for the development of adiposity-associated endometrial cancer.

INTRODUCTION

Type 1 endometrioid endometrial cancers are often low grade and early stage, arising from the epithelial lining of the uterus (Kost et al., 2016; Onstad et al., 2016). There has been an increase in the incidence of this cancer type for the past two decades, due in part to the increasing rate of obesity (Onstad et al., 2016; Rota et al., 2016; Soliman et al., 2005). Of all cancers showing a positive correlation between body mass index (BMI) and incidence, endometrial cancer exhibits the strongest association with obesity (Sahoo et al., 2018). While it has long been theorized that the endometrial microenvironment aberrantly influences gene expression profiles and consequential neoplastic output of epithelial cells, the contribution of adiposity to this tumorigenic process has not been well elucidated (Onstad et al., 2016; Soliman et al., 2005). During weight gain in humans and mouse models, accumulation of visceral white adipose tissues, including the parametrial fat pad in proximity to the uterus, releases adipokines and hormones to disrupt local and systemic lipid homeostasis (Chusyd et al., 2016; Zhang et al., 2010).

In addition to these circulating factors, adipose tissues disseminate adipocytes, immune cells, and other cell types into the lymphatic and vascular systems (Escobedo and Oliver, 2017; Ghosh et al., 2014). One of these cellular components

that cannot be overlooked is adipose stromal cells (ASCs), which are adipocyte progenitors (Zhang et al., 2010). When released from adipose tissues, ASCs target distal organ sites through chemokine gradients (Zhang et al., 2016). Upon infiltrating into the tumor microenvironment, these progenitors can differentiate into myofibroblasts to remodel extracellular matrix architecture (Seo et al., 2015). Embedded ASCs are also known to stimulate angiogenesis and release cytokines to promote inflammatory responses (Carmeliet, 2003; O'Sullivan et al., 2018). Removal of ASCs with pro-apoptotic peptides reduces tumor vascularization and growth in mouse models (Daquinag et al., 2017). These actions of ASCs in the microenvironment play a role in cancer development.

Aside from having an indirect influence on tumorigenesis through fibroblastic and vascular differentiation, emerging evidence indicates that ASCs have a direct effect on tumor cells through paracrine signaling (Freese et al., 2015; Ishii et al., 2018). For example, ASCs activate C-C motif chemokine ligand 5 (CCL5) signaling of cancer cells through interactions with chemokine receptors to enhance their proliferative and metastatic potential (Kauts et al., 2013). Another ASC-secreted adipokine is leptin bound to its surface receptor to promote epithelial-to-mesenchymal transition in cancer cells (Strong et al., 2015). In addition, ASCs may enhance anti-inflammatory differentiation



of immune cells in the tumor microenvironment (O'Sullivan et al., 2018). While these findings confirm a role of ASCs in cancer development and progression, the mechanistic aspects of ASC contribution to endometrial tumorigenesis are still poorly understood.

We have recently shown that the plasminogen activator inhibitor-1 (PAI-1) is a major adipokine secreted by ASCs, exerting paracrine influences on endometrial epithelial cells (EECs) *in vitro* (Polusani et al., 2019). While PAI-1 is a serine protease inhibitor well known for modulating extracellular matrix proteins (Czekay et al., 2011), our initial findings suggest that PAI-1 acts as an oncogenic ligand transducing intracellular signals that deregulate transcriptional programs of EECs (Polusani et al., 2019). This paracrine action may bring about heterogeneous alterations among different epithelial cell populations with pro-tumorigenic effects (Polusani et al., 2019). Prompted by these findings, we conducted single-cell RNA sequencing (scRNA-seq) and analyzed signaling pathways and cell-fate trajectories of heterogeneous cellular subpopulations after EEC exposure to PAI-1-secreting ASCs *in vitro*. We then mechanistically determined how PAI-1 signaling is internalized through protein-protein interactions, leading to disruption of transforming growth factor β (TGF- β)/SMAD-modulated transcription. To validate these transcriptomic findings *in vivo*, a single-cell mass cytometry (i.e., cytometry by time-of-flight [CyTOF]) marker panel was built to analyze proteomic profiles of primary endometrial tumor subpopulations and to assess intra- and inter-tumor heterogeneity associated with obesity. Here, we set out to demonstrate that PAI-1 initiates signaling cascades altering a transcriptional program in EECs, in addition to its known role in cell-surface proteolytic regulation.

RESULTS

ASC Infiltration in Tumor Microenvironments of Persons with Obesity

Consistent with previous publications (Lachance et al., 2006; Soliman et al., 2005), *in silico* analysis of The Cancer Genome Atlas (TCGA) endometrial cancer cohort demonstrated that obesity occurs more frequently in endometrioid (or type 1) endometrial cancers (Figure 1A). In these cancers, patients diagnosed at ages <50 and 50–59 years tended to have higher BMIs than older patients (>60 years old) (Figure 1B). In our South Texas patient population, the obese group comprised a large proportion of patients in the <50-years age group (Table S1). To control for this age group, embedded tumor samples from persons without obesity (BMI < 30 kg/m²) and persons with obesity (BMI \geq 30 kg/m²) were examined for ASC infiltration within the endometrial tumor microenvironment. We conducted an immunofluorescent analysis of tissue microarrays in 35 primary endometrial tumors and 6 non-cancerous age matched controls using the white adipose tissue 7 (WAT7) peptide that binds specifically to the surface of ASCs (Table S1) (Daquinag et al., 2017, 2016). The tissue sections from tumors of persons without obesity or non-cancerous controls showed negative or low WAT7 staining, while an increase in the number of WAT7 stained granules was seen in the endometrial stroma of persons with obesity and cancer (Figure 1C). Importantly, this increase was positively corre-

lated with BMI, consistent with a process of infiltration of WAT7-positive ASCs into the tumor microenvironment of persons with obesity ($p < 0.0001$; Figure 1D). We then assessed which factors are commonly secreted by ASCs *in vitro* using a multiplex assay of adipokines and cytokines (Polusani et al., 2019). The result showed that PAI-1 appeared to be the most abundant adipokine (i.e., 2 orders of magnitude greater than the other 10 factors) detected in 10 different ASC-conditioned media (ASC-CM; Figure S1). While increased levels of secreted PAI-1 were not correlated with the BMI status of women from whom these ASC samples were derived (data not shown), we suggest that more ASCs are accumulated in the endometrial microenvironment of women with increasing levels of obesity.

ASCs Engender PAI-1-Dependent and PAI-1-Independent Expression of Target Genes

To determine how ASC-secreted PAI-1 influences epithelial transcriptomes, we first exposed the immortalized EEC line EME6/7t to ASC-CM in cell culture for 24 h (Figure 2A), simulating ASC paracrine actions *in vitro*. Consistent with our previous findings (Polusani et al., 2019), a pro-tumorigenic phenotype was induced in this exposure model, including enhanced cellular invasion, proliferation, and mobility (Figures S2A and S2B, left, and S2C and S2D). The finding was independently confirmed in primary, un-immortalized EECs (Figures S2A and S2B, right). Then, we conducted scRNA-seq of EME6/7t EECs exposed to ASC-CM and stratified scRNA-seq profiles of 1,269 single cells (i.e., 669 control and 600 ASC-CM-exposed EME6/7t cells) (Figures 2A and 2B). Principal component analysis displayed cellular clustering of control and ASC-CM groups based on the most variable set of expressed genes (Figures S2E–S2G). The dimensionality reduction algorithm, t-distributed stochastic neighbor embedding (t-SNE), was employed to visualize distribution aggregates of these EECs, yielding a total of six subpopulations (Figure 2B). Subpopulations #1 and #2 were predominantly present in the control group, while two major *de novo* subpopulations, #5 and #6, appeared after the ASC-CM exposure (Figure 2C). This dynamic change in population size was moderately affected in two minor subgroups, #3 and #4, following the exposure (Figure 2C).

To further determine whether PAI-1 secreted by ASCs had an effect on these subpopulation changes, we additionally treated ASC-CM-exposed EME6/7t cells with the small-molecule inhibitor tiplaxtinin (Tip), known to bind and cleave this adipokine (Elkhdah et al., 2004). The concentration of Tip used in the present study was 50 μ mol/L, which had no known adverse cytotoxic effect in EME6/7t cells, based on our previous work or in other cell lines (Crandall et al., 2006; Polusani et al., 2019). When the additional scRNA-seq data of 629 cells in the ASC-CM+Tip group were integrated into those of the control and ASC-CM groups, we identified a total of 970 target genes whose expression patterns were directly influenced by PAI-1. There were four different PAI-1-mediated regulatory types within the two *de novo* subpopulations, #5 and #6 (Figures 2D and 2E). In type 1, PAI-1 initiated the upregulation of 331 genes that were partially reversed by Tip in subpopulation satellite #5a but were reversible by the PAI-1 inhibition in the other satellite, #5b (see *ECL2* as an example in Figure 2F, top). In contrast, the

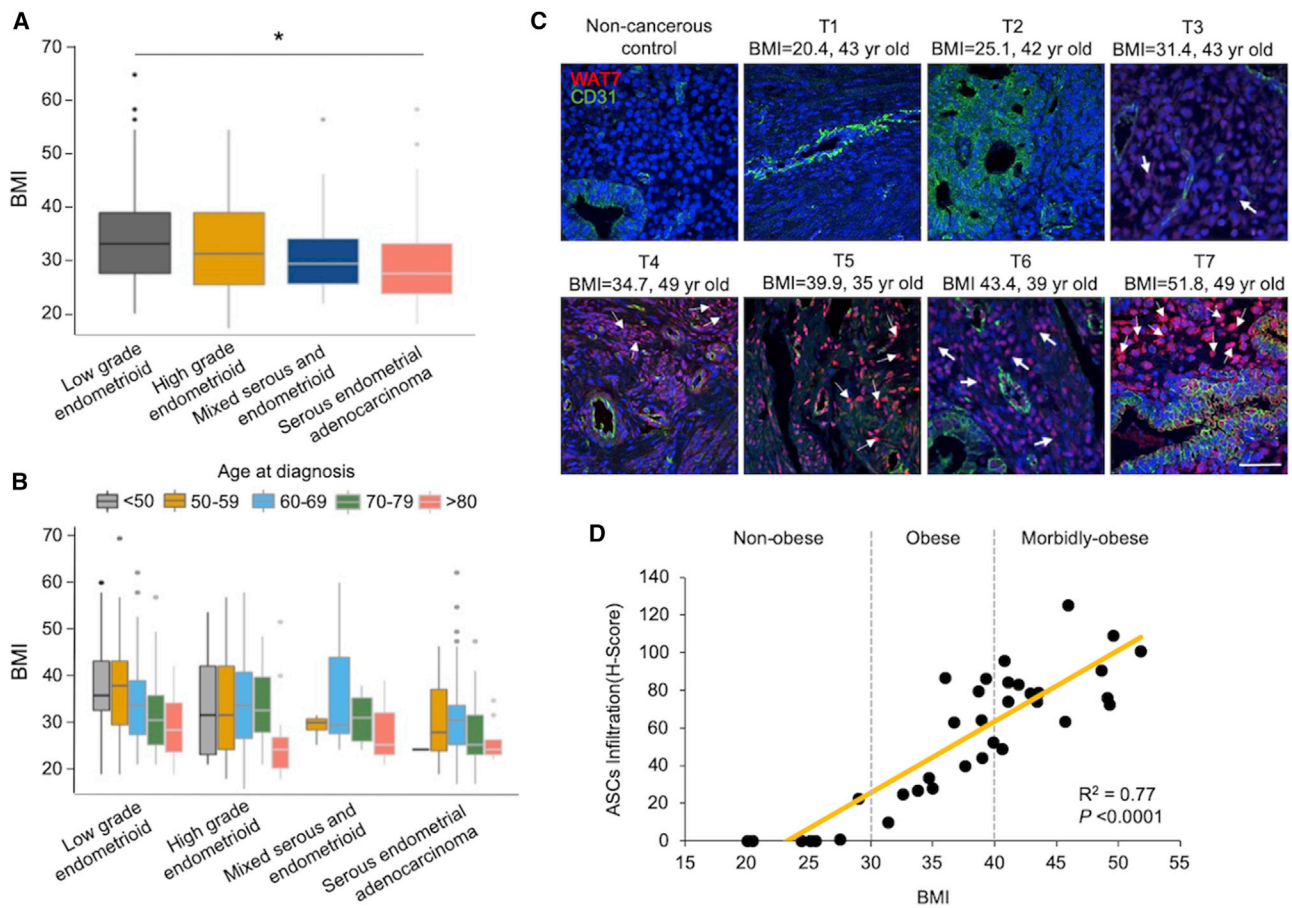


Figure 1. ASC Infiltration in Endometrial Tumor Microenvironments of Persons with Obesity

(A) In the TCGA endometrial cancer cohort, patients with low-grade endometrioid tumors had higher BMIs relative to patients with serous endometrial adenocarcinoma. * $p < 0.05$.

(B) In low-grade endometrioid tumors, patients in the <50 and 50–59 age groups tended to have higher BMIs than older patients. This age difference was not evident in the serous subtype.

(C) Representative sections of tissue microarrays immunofluorescently stained with the white adipose tissue 7 (WAT7) homing peptide, a known marker specific for ASCs (pink/red; see white arrows), CD31 for endothelial cells (green), and DAPI for nuclei (blue). Scale: 70 μ m.

(D) Scoring of ASC infiltration shows a direct association with BMI ($p < 0.0001$).

See also [Figure S1](#) and [Table S1](#).

downregulation of 428 genes by PAI-1 in type 2 was reversible by Tip in subpopulation satellite #5a, but partially reversed by Tip in subpopulation satellite #5b (see *RPL6* in [Figure 2F](#), bottom). However, these mixed regulatory modes in subpopulation #5 were less frequently seen in subpopulation #6. While the expression of 378 upregulated genes could be reversed by Tip in subpopulation satellites #6a, #6b, and #6c (i.e., type 3; see *STMN1* in [Figure 2G](#), top), this reversal was less evident in 370 PAI-1-downregulated genes in subpopulations #6b and #6c (i.e., type 4; see *TPT1* in [Figure 2G](#), bottom). From this single-cell expression analysis, it became apparent that PAI-1 initiates context-dependent gene regulation in diverse EME6/7t subpopulations.

To trace cellular origins of the two major *de novo* subpopulations, #5 and #6, pseudotime analysis was used to order cell trajectories based on their transcriptomic similarities ([Haghverdi et al., 2016](#)). After time tracking of subpopulation #5, we

observed a putative movement from #1/#2 to #3 and then to #5 after exposing EME6/7t cells to ASC-CM. The inhibition by Tip partially reversed the movement back to control subpopulations #1/#2 or #3, suggesting that PAI-1 had a transient influence on target gene expression in subpopulation #5 (see examples of two genes, *SLNF11* and *ATF3*, in [Figure 2H](#)). In contrast, gene regulation in subpopulation #6 was less dependent on PAI-1. Although the cellular origin of subpopulation #6 could similarly be traced back to #1/#2 after the ASC-CM exposure, these cells were less responsive to the PAI-1 inhibition (see also expression patterns of *IFITM2* and *VIM* in [Figure 2I](#)). Corroborating the aforementioned gene heatmaps, the pseudotime analysis suggests that PAI-1-dependent regulatory modes appear prominent in subpopulation #5, while other factors secreted by ASC-CM are more involved in gene regulation for subpopulation #6. Alternatively, PAI-1 triggers an irreversible transcriptional process in

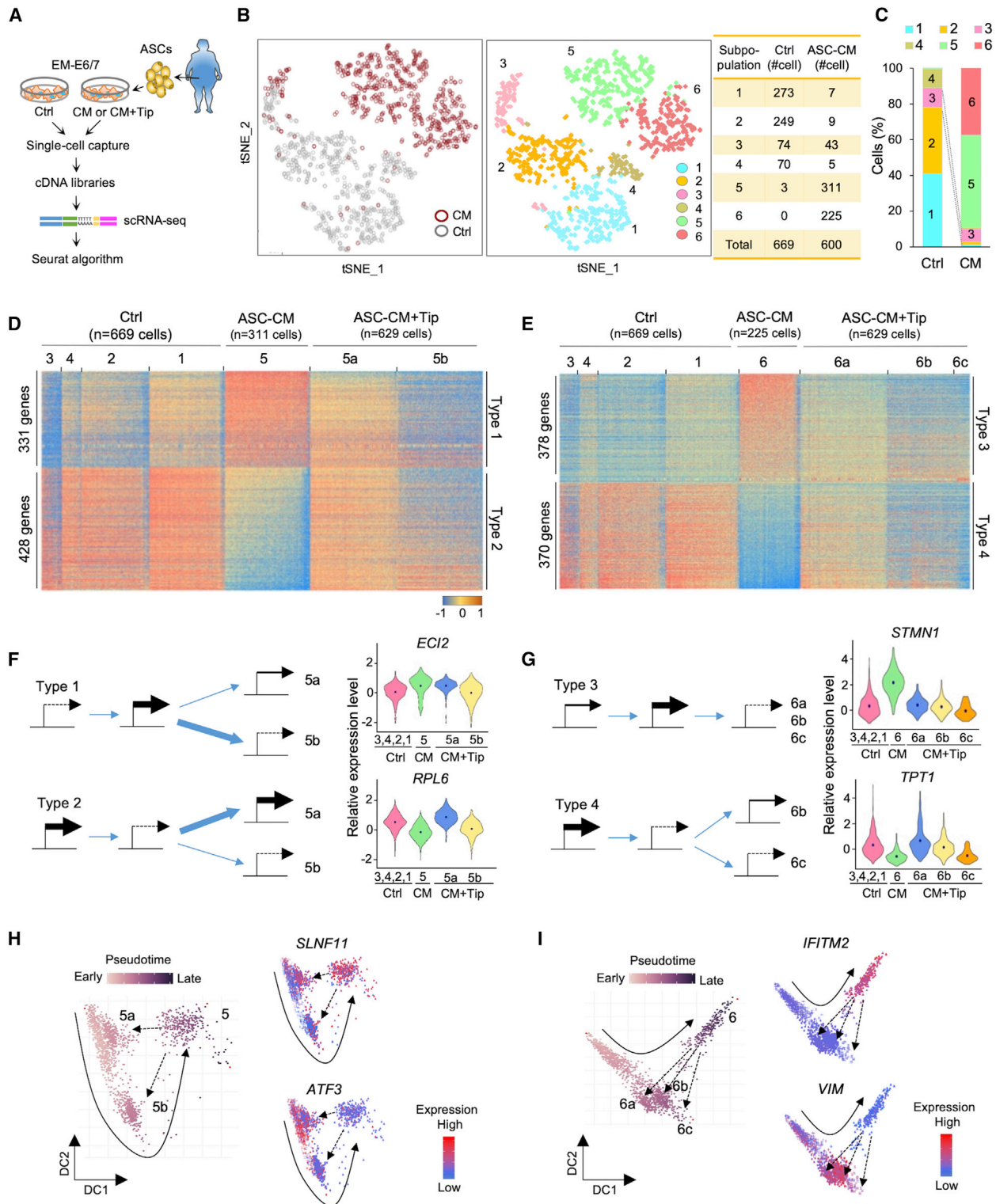


Figure 2. Single-Cell Transcriptome Profiling of Endometrial Epithelial Subpopulations Exposed to ASC-CM

(A) Schematic of study design of the immortalized EEC line EME6/7t exposed to control media and conditioned media of adipose stromal cells (ASC-CM) in the presence or absence of the PAI-1 inhibitor Tip, and subsequent unsupervised clustering analysis of EEC subpopulations based on scRNA-seq analysis. (B) tSNE scRNA-seq plot of EME6/7t cells with or without ASC-CM exposure (left) and six cell clusters from 1,269 individual cell transcriptomes (right). (C) Bar plots showing the proportions of clustered cell subpopulations in each treatment.

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type 3, especially for those undergoing epigenetic repression, suggested by our recent finding (Polusani et al., 2019).

PAI-1 Deregulates TGF- β /SMAD-Mediated Junction and Adhesion Complex

To determine how PAI-1 alters intracellular signaling activities, we performed network analysis of 13 prominent oncogenic signaling pathways (Liberzon et al., 2015) using scRNA-seq data. When aligning these pathways based on their PAI-1 dependence, we found the top six signaling pathways—TGF- β /SMAD, Notch, MYC, E2F targets, p53, and tumor necrosis factor alpha (TNF- α) via NF- κ B—underwent reversible repression by PAI-1 in subpopulation #5 (Figure 3A). This transient repression was similarly observed in five of these six pathways in subpopulation #6. The exception is the upregulation of E2F targets, which is a key pathway to highlight the differential impact of PAI-1 on subpopulations #5 and #6. The remaining seven signaling pathways appeared to be upregulated after ASC-CM exposure, but this activation was less dependent on PAI-1.

We additionally found enrichment of five pathways associated with cellular complex consisting of gap junction, tight junction, focal adhesion, apical surface, and apical junction (collectively termed junction and adhesion complex [JAC]; Figure 3B) (Aoki-Kinoshita and Kanehisa, 2007; Liberzon et al., 2015). The results also showed PAI-1-dependent influences on JAC in the satellites of subpopulations #5 and #6 (Figure 3B). The JAC response to ASC-CM exposure was plastic and could be reversed by Tip in #5a, #6a, and #6c (Figure 3B). On the other hand, the Tip treatment led to activation of JAC in #5b and #6b, likely attributed to a compensatory mechanism. Correlation matrix analysis further revealed that the JAC expression was positively associated with TGF- β /SMAD signaling in control subpopulations #1 and #2, but this relationship was uncoupled by PAI-1 in subpopulations #5 and #6 exposed to ASC-CM (see gap-junction and apical surface pathways that are most negatively affected in Figure 3C).

To narrow down which target loci contribute to adiposity-related endometrial cancer, we first screened 484 JAC genes listed in the Molecular Signatures Database (Figure 3D) (Aoki-Kinoshita and Kanehisa, 2007; Liberzon et al., 2015). A total of 382 of these genes had significantly different expression levels between endometrial tumors and adjacent normal controls in the TCGA cohort ($n = 522$; $p < 0.05$). Among these, 101 genes were also differentially expressed in ASC-CM-exposed cells based on our scRNA-seq data. Fifty-eight genes were determined to be BMI associated when their increased or decreased expression levels in tumors were stratified by obesity status (Fig-

ure 3D). These genes were either downregulated ($n = 36$) or upregulated ($n = 22$) in ASC-CM-exposed subpopulation #5 and/or #6, and their differential expression status could be partially reversed by PAI-1 inhibition (Figure S3). Moreover, the deregulated gene profiles occurred more frequently in the endometrioid endometrial subtype (or type 1) and in patients with clinical stage I cancers when we unbiasedly correlated these expression levels with clinicopathological parameters of TCGA endometrial tumors (Figure 3E). Although we did not reach statistical significance of $p < 0.05$ among the five age groups categorized in the 36 downregulated candidate genes, we observed that upregulated expression of the 22 candidate genes in the 50–59 age group is higher than in other older patient groups ($p < 0.05$; Figure 3E). The result suggests that PAI-1-mediated expression changes of candidate JAC genes are capable of inducing endometrial neoplasms.

PAI-1 Modulates JAC Expression by Interacting with LRP1 to Downregulate SMAD4

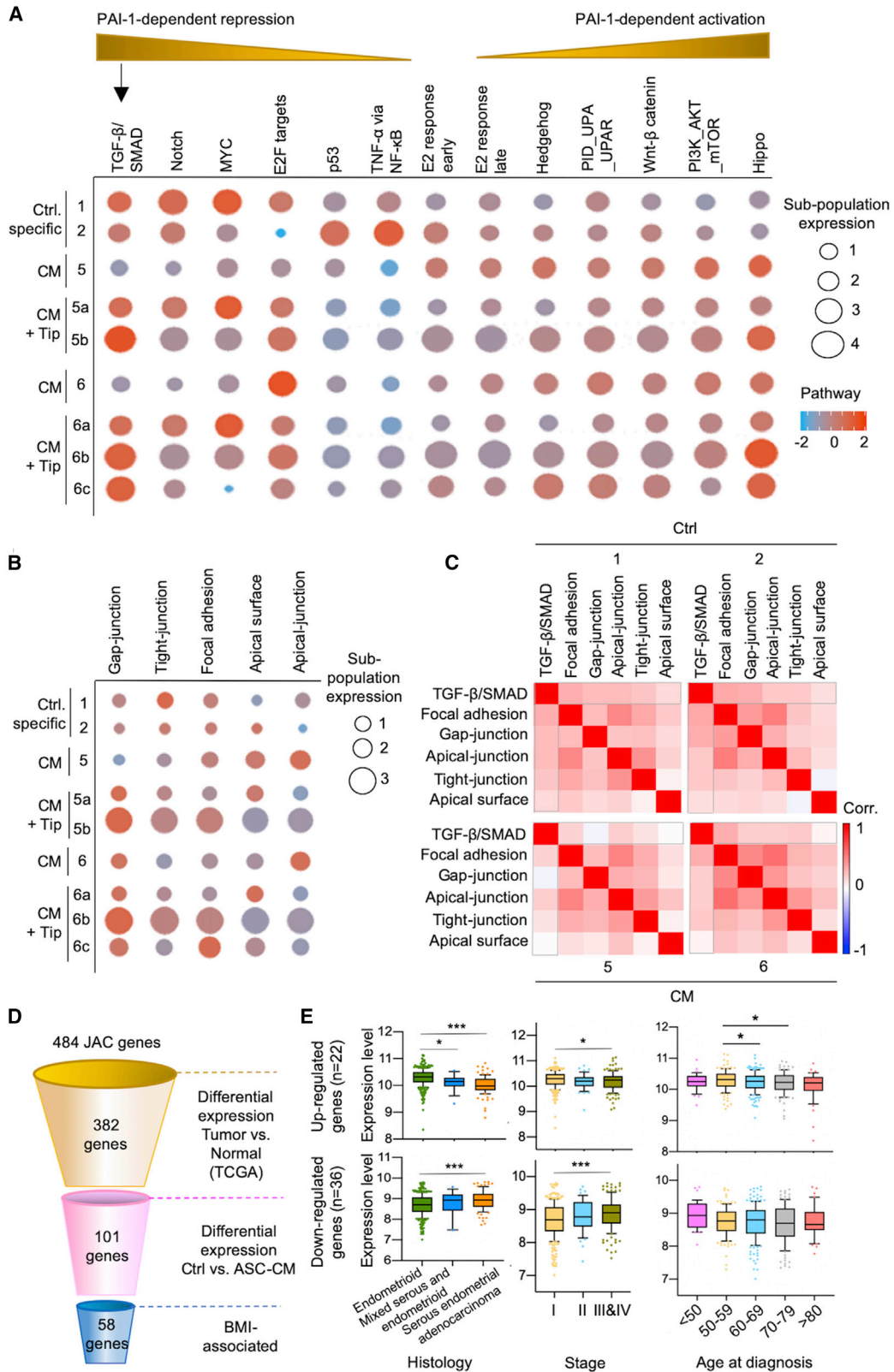
Based on the correlation between PAI-1 regulatory action and reduced TGF- β /SMAD signaling (Figure 3A), we examined the potential regulation of the co-SMAD, SMAD family member 4 (SMAD4), by PAI-1. While PAI-1 activities include interactions with the urokinase plasminogen activator pathway (uPA-uPAR), PAI-1 also has a strong affinity to the low-density lipoprotein receptor-related protein 1 (LRP1) to transduce signals for downstream transcriptional modulation (Degryse et al., 2004; Polusani et al., 2019; Wei et al., 2019). As these published observations strongly implicate LRP1 as a modulator of PAI-1 transcriptional activity, we mechanistically determined how the interaction of PAI-1 with LRP1 influences SMAD4 levels, using proximity ligation assay (PLA). The data demonstrate an increase in the interaction between PAI-1 and LRP1 on the cell surface of ASC-CM-exposed EME6/7t cells relative to control cells (Figures 4A and 4B). Further, this interaction was attenuated by pretreating cells with LRP1 antagonist receptor-associated protein (RAP). When PLA was conducted on cells treated with Triton-X detergent to increase membrane permeability, we similarly observed increased intracellular interactions between PAI-1 and LRP1 in ASC-CM-exposed EME6/7t cells (Figure 4C). This internalization is known to produce the cleaved intracellular domain (ICD) of LRP1 that can be translocated into the nucleus upon signaling activation (Xian et al., 2017; Yamada et al., 2019). Therefore, chromatin immunoprecipitation (ChIP)-qPCR was performed to determine if ICD-LRP1 binds to the SMAD4 promoter in EME6/7t cells exposed to ASC-CM. The result showed a 5- to 6-fold increase in the LRP1-ICD binding to the promoter, which was abrogated

(D and E) Heatmaps showing single-cell gene expression patterns from each treatment in subpopulations #5 (D) and #6 (E).

(F and G) Schematic showing four types of PAI-1-related modes of transcriptional regulation in subpopulation #5 (F) or #6 (G). In types 1 and 2, genes were altered by ASC-CM and partially reversed by adding Tip (F). In type 3, ASC-CM induced expression, which was reversed by Tip; on the other hand, ASC-CM repressed gene expression, but PAI-1 inhibition had little effect on restoring expression in type 4 (G). The thickness of blue arrows represents differential levels of gene regulatory expression. Violin plots showing expression distributions of representative genes (dots refer to the average) for the four regulatory types.

(H and I) Pseudotime trajectory of EME6/7t cells derived from subpopulation #5 and related satellites (i.e., #5a and #5b) (H) or subpopulation #6 and related satellites (i.e., #6a, #6b, and #6c) (I). Representative genes show upregulation (*SLNF11*) or downregulation (*ATF3*) due to ASC-CM exposure (#5) and their responses to Tip (see #5a and #5b). Genes show upregulation (*IFTM2*) or downregulation (*VIM*) due to ASC-CM exposure and their responses to Tip (see #6a, #6b, and #6c). A solid arrow represents the time trajectory of cellular subpopulations. A dotted arrow represents the trajectory of EME6/7t cells in response to ASC-CM exposure.

See also Figure S2.



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by RAP treatment (Figure 4D). Acting as a transcription repressor, LRP1-ICD subsequently downregulated *SMAD4* gene expression in ASC-CM-exposed cells (Figure 4E). The expression of *SMAD4* was restored in exposed cells additionally treated with Tip or RAP.

In search of other regulatory mechanisms, we observed upregulation of *NEDD4-like E3 ubiquitin (NEDD4L)* and *ubiquitin-conjugating enzyme E2 I (UBE2I, formerly UBC9)* genes based on our scRNA-seq data, suggesting a role of PAI-1-LRP1 in the proteasome pathway (Figure S4A). The protein products of both genes are known to bind to SMAD2/3/4 complexes, which may cause ubiquitin-mediated degradation of TGF- β -associated transcription factors (Lin et al., 2003; Wang et al., 2011; Yilmaz et al., 2018). The PLA result of membrane-permeated EME6/7t cells confirmed this PAI-1-mediated degradation by ubiquitination of SMAD4, which could be abrogated by the proteasome inhibitor MG132, as well as inhibited by Tip or RAP treatments (Figure 4F). The data from capillary western immunoassays independently supported the role of PAI-1 in mediating SMAD4 degradation in ASC-CM-exposed EME6/7t cells (Figures 4G and S4B).

To verify the direct impact of SMAD4 on JAC gene expression, we generated stable knockdown clones of EME6/7t cells transfected with *SMAD4* shRNA (sh*SMAD4*) or control (shCtrl) that showed efficient suppression of *SMAD4* after the knockdown (Figure S5). The expression of 11 JAC genes was then determined in control and sh*SMAD4* cells in the presence or absence of ASC-CM by qRT-PCR (Figure 5A). Supporting the scRNA-seq findings, exposure of EME6/7t cells to ASC-CM or attenuation of TGF- β /SMAD signaling by *SMAD4* knockdown resulted in expression downregulation of these genes (Figure 5A). Moreover, ChIP-qPCR assays indicated that SMAD4 binds to their corresponding promoters (Figure 5B). With the exception of *ADAM15*, SMAD4 promoter binding was decreased in the 10 remaining loci in EME6/7t cells exposed to ASC-CM. Taken together, these results demonstrate that PAI-1-LRP1-modulated SMAD4 downregulation occurs at both the transcriptional and the post-translational levels. As a result, tumor-suppressive activities of TGF- β /SMAD signaling are attenuated, leading to deregulation of JAC genes in EECs.

Low Expression of SMAD4 and JAC Is Linked in Cell Subpopulations of Persons with Obesity

To confirm the expression profiling data and examine the effects of obesity and PAI-1 on JAC expression, we conducted CyTOF profiling of 16 protein factors in nine primary endometrial tumors (Table S2) and three sets of EME6/7t cells (i.e., control, ASC-CM,

and ASC-CM plus Tip) (Figures S6A and S6B). The 11 BMI-associated JAC factors were chosen (Figure 3D; Polusani et al., 2019) because of their critical involvements in cell-cell adhesion and extracellular matrix communication, including growth arrest-specific 1 (Gas1), adhesion molecule with Ig like domain 2 (AMIGO2), syndecan-3 (SDC3), disintegrin and metalloproteinase domain-containing protein 15 (ADAM15), connexin 43 (Cx43), zonula occludens-1 (ZO1), zonula occludens-2 (ZO2), rho-associated protein kinase 1 (ROCK1), signal transduction adaptor protein paxillin (Paxillin), tyrosine protein kinase Fyn (Fyn), and cellular inhibitor of apoptosis protein 1 (cIAP1) (see functional descriptions of these factors in Figure S6C). Three TGF- β /SMAD signal components—GFR-2, SMAD2, and SMAD4—were also included to determine whether PAI-1-mediated repression of this signaling activity occurred in the samples analyzed. Additional markers for immune, stromal, and endothelial cells were used to gate EECs from non-epithelial components in primary tumors (Figure S6B). EECs were positively identified by two epithelial markers, EpCAM and CK8/18 (Figure S6B).

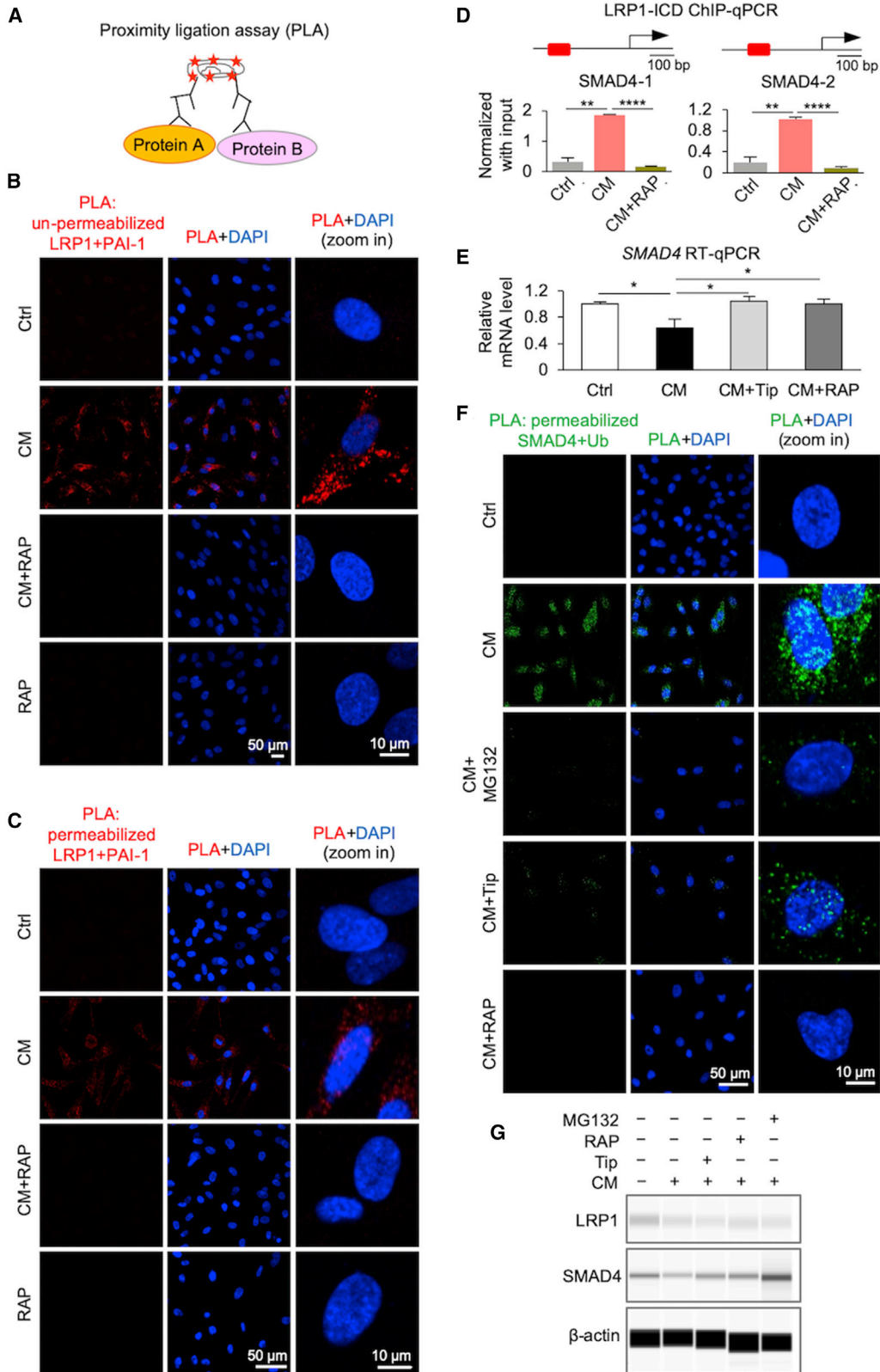
The tSNE plots of the 16 protein factors displayed 27 cellular subpopulations derived from a total of 74,048 single EECs (Figures 6A and 6B). Consistent with the scRNA-seq findings of EME6/7t cells, various degrees of JAC protein repression were seen in the ASC-CM-exposed subpopulations #4, #6, and #7 relative to those of control subpopulations #8, #9, and #10 (Figure 6C). Except for ADAM15, Fyn, and cIAP1, JAC repression could be reversed by Tip treatment in subpopulations #1, #2, and #3. This PAI-1-dependent repression was similarly observed in TGF- β /SMAD signal components SMAD2 and SMAD4 in EME6/7t subpopulations (Figure 6D). However, the remaining subpopulations derived from primary tumors showed highly heterogeneous expression patterns of these protein markers.

To further explore how PAI-1-mediated repression of TGF- β /SMAD4 signaling influences intra-tumor and inter-patient heterogeneity in endometrial cancer, we classified the aforementioned 27 cell subpopulations into five categories (i.e., I, II, III, IV, and V) on the basis of (1) expression levels of SMAD4 ($p < 0.0001$; Figure 7A), (2) expression levels of 11 candidate JAC markers ($p < 0.0001$; Figure 7B), and (3) subpopulation size (Figure 7C). Violin plot profiling of SMAD4 levels further showed that decreasing expression of this TGF- β mediator is usually correlated with reduced JAC expression in EEC subpopulations (Figure 7). In general, EEC subpopulations derived from EME6/7t cells exhibited higher expression levels of SMAD4 and JAC

Figure 3. PAI-1-Dependent Repression of TGF- β /SMAD Signaling Deregulates JAC Expression

(A and B) Network analysis delineated PAI-1-dependent signaling pathways (A) as well as JAC (B), based on exposure to ASC-CM \pm Tip.
(C) Correlation matrix heatmaps showing the association between cell signaling and JAC pathways. Bracketed areas indicate a decrease in correlation between TGF- β /SMAD and JAC due to ASC-CM exposure (subpopulations #5 and #6) compared to control (subpopulations #1 and #2).
(D) Schematic of the stepwise filtering used for JAC gene selection. PAI-1-dependent JAC genes were filtered using the TCGA endometrial cancer cohort, based on differential expression patterns between adjacent normal control and tumors ($p < 0.05$) and BMI as described in Method Details. This stepwise analysis identified putative BMI-associated JAC genes ($n = 58$).
(E) The 58 BMI-associated PAI-1 dependent genes were divided into up- ($n = 22$) and downregulated ($n = 36$) groups. Expression profiles of patients in TCGA endometrial cancer cohort were classified according to histology, clinical stage, and age. Downregulated JAC genes showed lower expression in the endometrioid subtype compared to the serous subtype, whereas upregulated genes showed higher expression levels in the endometrioid subtype (left). This correlation was also seen in early-stage disease compared to stage III/IV disease (middle). The relationship between gene expression and age at diagnosis was less evident (right). * $p < 0.05$, *** $p < 0.001$.

See also Figure S3.



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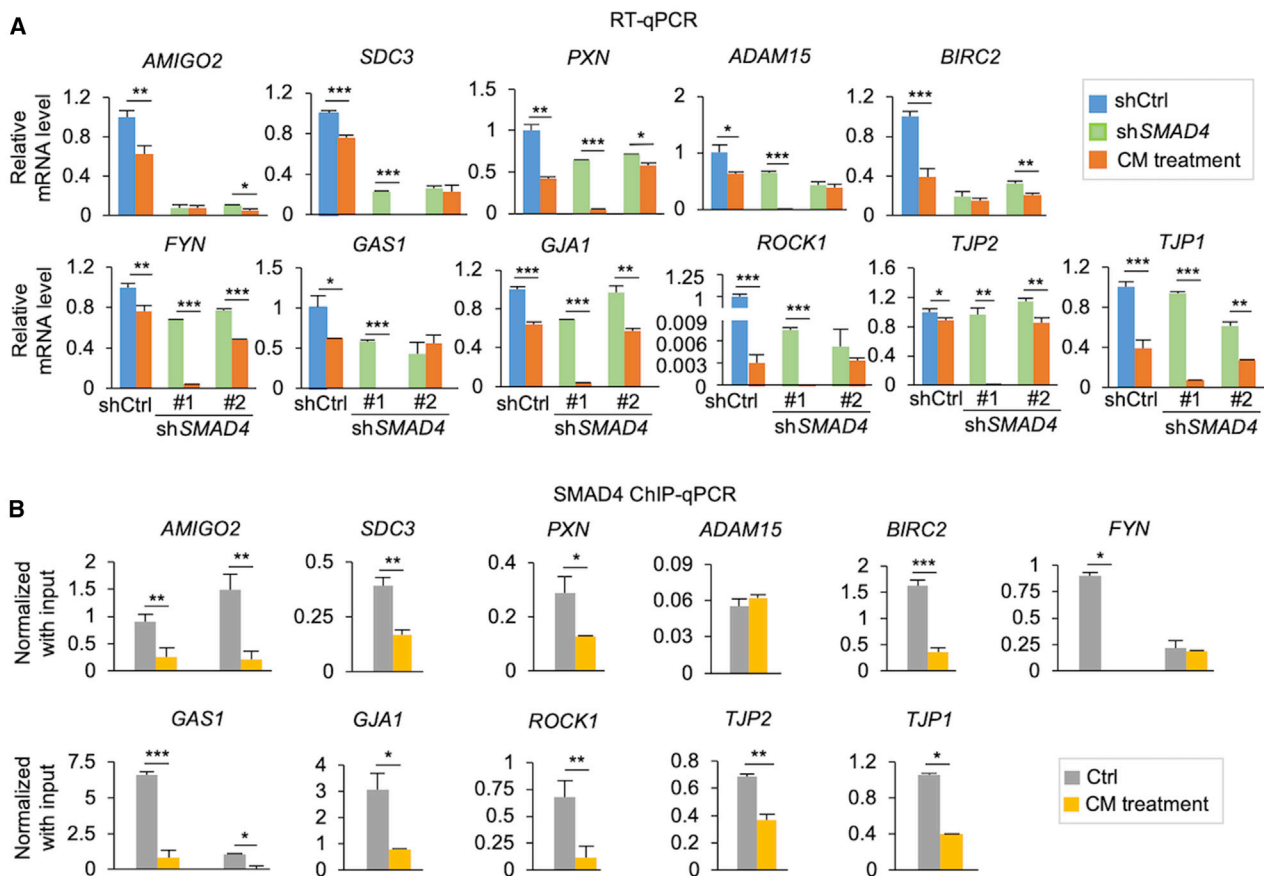


Figure 5. TGF- β /SMAD4-Mediated Downregulation of Candidate JAC Genes

(A) Bar plots showing mRNA expression levels of selected PAI1-regulated JAC genes by qRT-PCR in the two shSMAD4 knockdown clones, compared to shCtrl in the presence or absence of ASC-CM.

(B) ChIP-qPCR analysis of SMAD4 binding to promoter regions of the JAC genes in control (Ctrl) cells and ASC-CM-exposed cells.

Data represent mean \pm SD, n = 3 technical replicates; *p < 0.05, **p < 0.01, ***p < 0.001, Student's t test. See also Figure S5.

markers (Figure S7; predominantly in categories IV and V) than those of primary tumors (Figure S7; predominantly in categories I, II, and III). Interestingly, we found that low SMAD4-expressing

subpopulations (categories I and II) were more frequently associated with JAC reduction in endometrioid cancer patients with increased BMI (Figure 7B and 7C).

Figure 4. PAI-1-LRP1 Interactions Downregulate SMAD4 Levels via Transcription Repression and Ubiquitin-Mediated Degradation

(A) PLA utilizes a pair of oligonucleotide-conjugated secondary antibodies that bind corresponding antibodies targeting each interacting protein partner. The proximity of these two protein partners allows circular DNA amplification of the oligonucleotides template upon *in situ* ligation. Fluorescent probes complementary to this amplicon engender a detectable signal via fluorescence microscopy. Plasma membrane permeabilization was used for intracellular protein interactions. (B and C) PLA indicating protein-protein interactions of PAI-1 and low-density LRP1 on the cell surface (B) or intracellular (C) due to ASC-CM exposure in the presence or absence of the LRP1 inhibitor RAP. Representative micrograph images showing positive signal (red) for PAI-1/LRP1 interactions. DAPI was used to stain nuclei (blue).

(D) ChIP-qPCR analysis of LRP1-ICD binding to two promoter regions of the SMAD4 gene in control (Ctrl) cells and CM-exposed cells treated and untreated with RAP (5 μ g/mL). Data represent mean \pm SD, n = 3 technical replicates; **p < 0.01, ***p < 0.001.

(E) qRT-PCR analysis of mRNA expression levels of SMAD4 gene in control cells and CM-exposed cells treated and untreated with Tip (50 μ M) or RAP (5 μ g/mL). Data represent mean \pm SD, n = 3 technical replicates; *p < 0.05.

(F) PLA demonstrating protein-protein interactions of SMAD4 and ubiquitin due to ASC-CM exposure in the presence or absence of the following inhibitors: Tip for PAI-1, RAP for LRP1, and protease inhibitor MG132. Micrograph images showing positive PLA signal (green) for SMAD4-ubiquitin interactions. DAPI was used to stain nuclei (blue).

(G) Protein expression of LRP1 and SMAD4 by capillary western immunoassays. Endometrial epithelial EME6/7t cells were treated with control media, ASC-CM, and ASC-CM plus Tip, or with pre-treatment with LRP1 antagonist RAP or protease inhibitor MG132. The result shown is a representative of three independent experiments.

See also Figure S4.

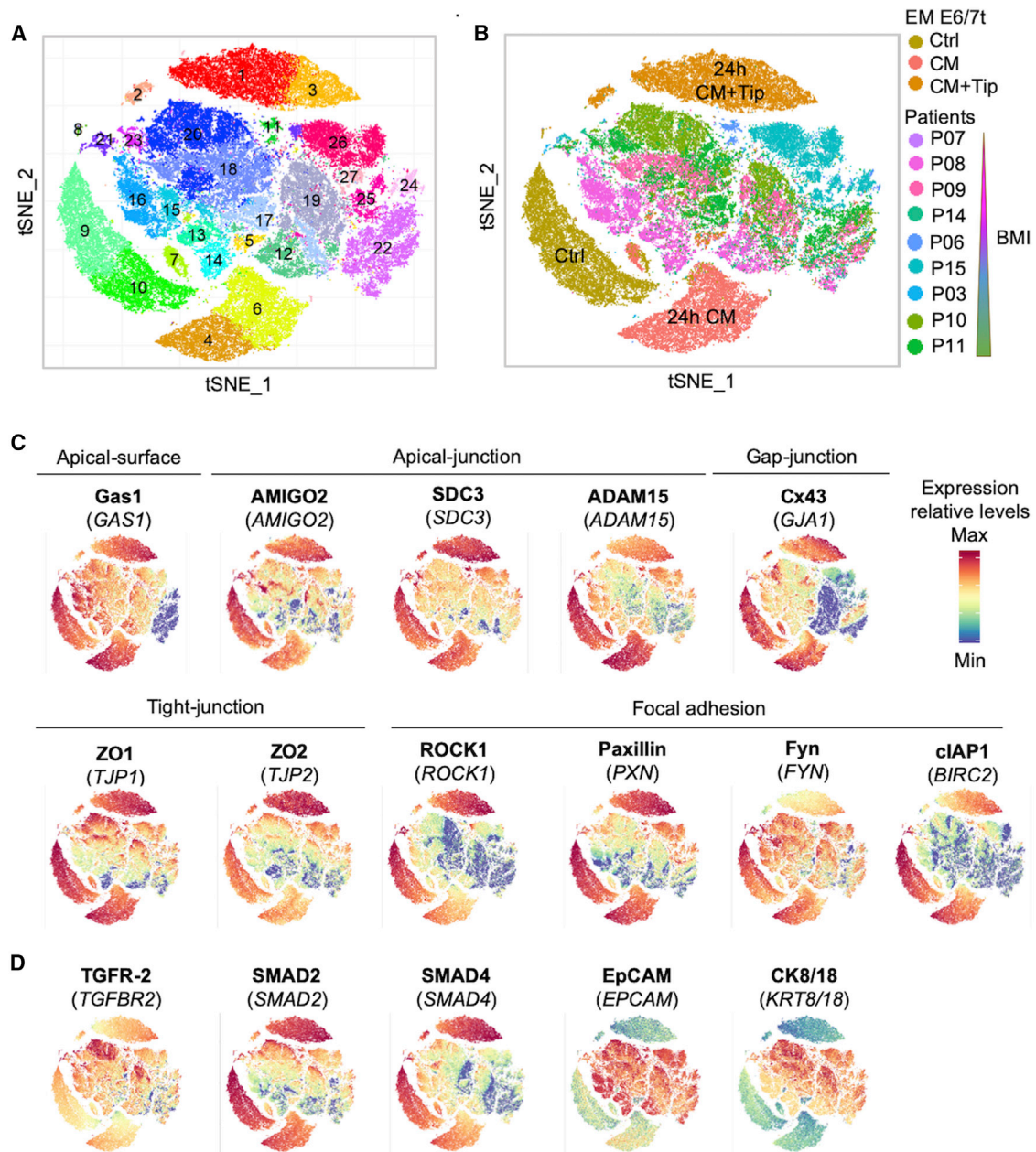


Figure 6. Single-Cell Proteomic Analysis of Cellular Subpopulations in EME6/7t EECs and Primary Tumors

(A and B) tSNE plots of mass cytometry (CyTOF) analysis visualizing 74,048 epithelial cells, which were divided into clusters ($n = 27$) based on their proteomic profiles (A). These cells included immortalized EEC line EME6/7t (exposed to ASC-CM \pm Tip) and primary endometrial tumors of patients with known BMIs (B). (C) tSNE-based phenographs showing abundance and cellular distributions of JAC proteins. (D) tSNE-based phenographs showing EpCAM and CK8/18 as epithelial cell markers and TGFR-2, SMAD2, and SMAD4 as TGF- β factors. See also [Figure S6](#) and [Tables S2](#) and [S3](#).

When further aligning these EEC subpopulations with tumor histology, we found that patients with endometrioid endometrial cancers (i.e., P09, P14, P06, P15, P03, P10, and P11) tended to have larger proportions of low SMAD4-expressing cells (e.g., categories I and II) and smaller proportions of high SMAD4-expressing cells (e.g., categories III and IV) than patients with

papillary serous carcinomas (i.e., P07 and P08) ([Figure 7D](#); [Table S2](#)). The average expression of 4 of the 11 JAC markers analyzed (i.e., ADAM15, ROCK1, Cx43, and Gas1) was also decreased in the endometrioid group relative to the papillary serous group ($p < 0.0001$; [Figure 7E](#)). A similar trend of reduced expression of both SMAD4 and JAC markers was seen in obese and

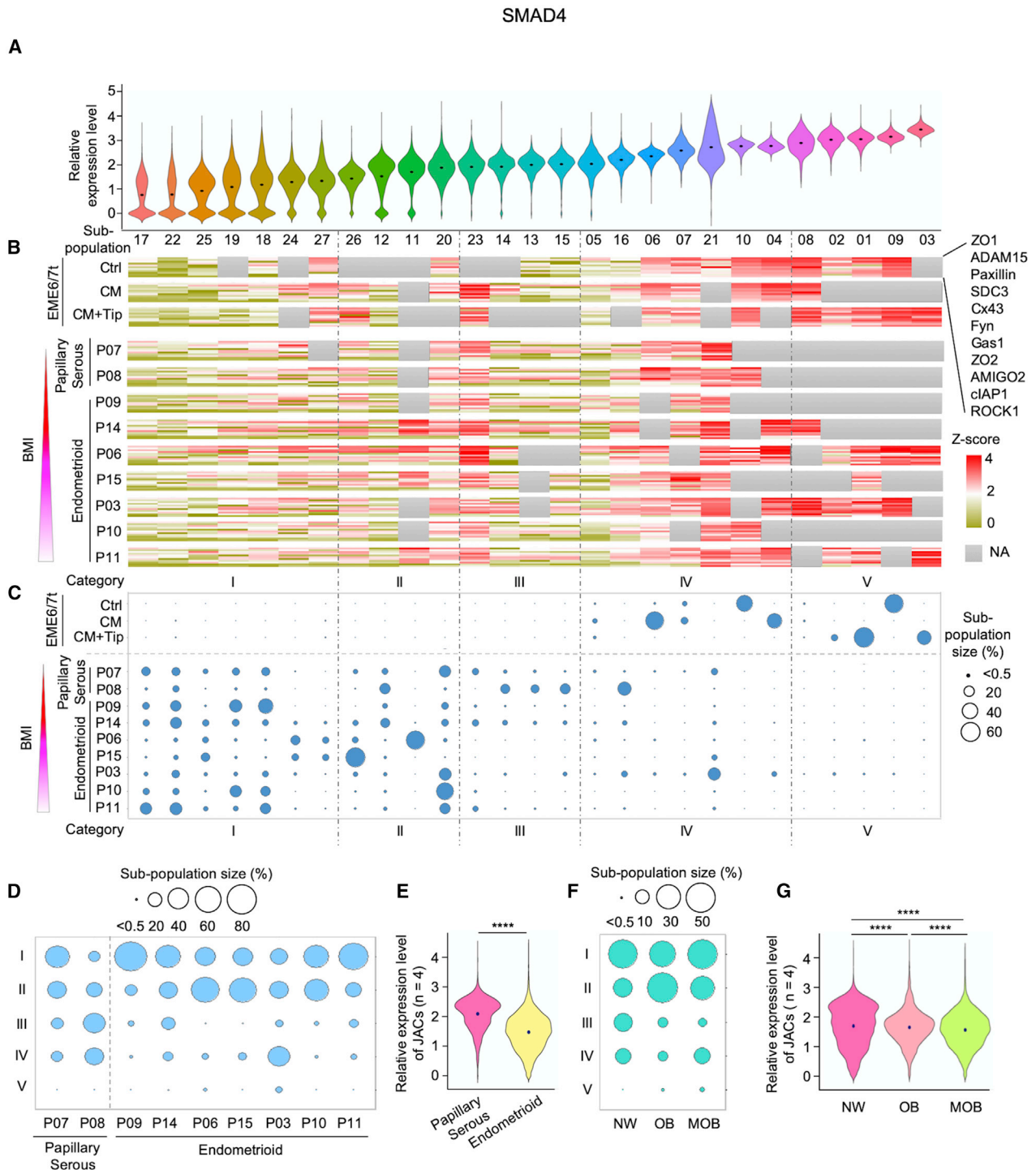


Figure 7. Stratification of Endometrial Cell Subpopulations Based on SMAD4 and JAC Expression Levels and Patients' BMIs

(A–C) Expression profiles of SMAD4 and JAC based on CyTOF were classified into five categories: I, II, III, IV, and V. Violin plots show ordered SMAD4 expression level (low to high) and distribution in the individual subpopulations (A). Expression heatmaps of JAC were aligned according to SMAD4 expression levels (in descending order) in different cellular subpopulations (B). Area of circle denotes relative cell proportion of a subpopulation within each sample and ordered in the aforementioned five categories in cellular subpopulations of EME6/7t lines and primary tumors (C). Tumors were ordered from top to bottom based on increased patients' BMIs and histology classes.

(legend continued on next page)

morbidity obese groups relative to the normal weight group ($p < 0.0001$; Figures 7F and 7G). This CyTOF finding further corroborates the notion that SMAD4-modulated JAC repression observed *in vitro* recapitulates a transcriptional regulatory program *in vivo*, particularly in endometrioid carcinomas.

DISCUSSION

ASCs derived from adipose tissues, including the parametrial fat near the uterus, can be mobilized via chemokine actions to the endometrial microenvironment and promote epithelial tumorigenesis (Ma et al., 2016; Zhang et al., 2010; Zhang and Kolonin, 2016). These adipocyte progenitors exert paracrine actions by secreting abundant PAI-1 and other factors that may alter signaling responses in nearby EECs. To simulate this event *in vitro*, we exposed EECs to ASC-CM for downstream analyses of gene and protein expression. Furthermore, transcriptomic profiling and pathway enrichment analysis identified TGF- β /SMAD as a key pathway negatively regulated by PAI-1, which could diminish TGF- β -mediated tumor suppressor activity. While the link between hyperplastic development and tumorigenesis is not directly addressed by the current *in vitro* exposure model, future studies will include EECs derived from endometrial hyperplasia to further assess the role of PAI-1 in promoting endometrial tumorigenesis. Long-term chronic ASC exposure may engender obesity-associated transcriptional repression profiles. These tumorigenic effects may be initiated early in pre-menopausal persons with obesity, resulting in the observed increased incidence of premenopausal endometrial hyperplasia and type 1 cancer. Alternatively, the PAI-1 tumorigenic effects may manifest later in life, resulting in the development of typical postmenopausal endometrial cancer (Gressel et al., 2015; Nevadunsky et al., 2014; Onstad et al., 2016; Wise et al., 2016). In the current study, ASC infiltration was associated positively with BMI in a premenopausal patient cohort (<50 years old). A limitation of the study was the absence of a postmenopausal study cohort. Future studies will examine the association of ASC infiltration in a large cohort of both premenopausal and postmenopausal patients, balanced BMI, and other clinical variables. In addition, a limited number of tumor samples, yet reflective of our patient population, was used in the current CyTOF analysis. These studies support the TCGA transcriptomic data that lower SMAD4 and corresponding JAC expression occurred in tumors from higher-BMI type 1 patients. Because of the extensive time needed to acquire a large set of prospective fresh tumor samples for CyTOF analysis, we will expand the patient cohort in future studies.

As a serine protease inhibitor, PAI-1 is a well-known antagonist of the plasminogen activator system (Czekay et al., 2011). However, a growing body of evidence indicates that PAI-1 interacts with signal transducer LRP1 and becomes internalized to regulate oncogenic transcription programs, resulting in epithelial-to-mesenchymal transition, cytoskeletal reorganization, and cellular migration and invasion (Czekay et al., 2011; Degryse et al., 2004; Xian et al., 2017; Yamada et al., 2019). Our present findings add to the list of PAI-1-mediated tumorigenic events by describing two functions of PAI-1-LRP1 interactions in deregulating TGF- β /SMAD signaling. First, PAI-1 mobilizes LRP1-ICD to bind the SMAD4 promoter for transcription repression. Second, the PAI-1-LRP1 signaling independently enhances ubiquitin-mediated SMAD4 degradation. These combined effects of transcriptional and post-translational regulation can greatly attenuate tumor-suppressive functions of TGF- β /SMAD signaling.

Among the PAI-1-dependent JAC factors we identified in the present study that are under the regulatory control of SMAD4, ADAM15, ROCK1, Cx43, and Gas1 are integral components involved in cell-cell adhesion, extracellular matrix (ECM) interaction, and inter-cellular communication (Jin et al., 2015; Parsons et al., 2010; Vicente-Manzanares et al., 2009). Consistently, turnover of focal adhesion factors is associated with a loss of integrin interactions with ECM substrates, leading to enhanced cellular motility (Nagano et al., 2010; Webb et al., 2002). Other studies showed that cell-ECM interactions are stabilized by ROCK1, whereas a loss of ROCK1 and internalization of integrins facilitate cell-cell detachment (Parsons et al., 2010; Vicente-Manzanares et al., 2009). In addition, cell-cell adhesion not only regulates, but is also stabilized by gap junctions, which are intercellular channels allowing for direct electro-chemical communication between adjacent cells (Evans and Martin, 2002; Kanaporis et al., 2008). The major gap-junction structural protein in the endometrium Cx43 is downregulated by PAI-1 in ASC-exposed EECs. These effects enhance cellular motility and facilitate cellular invasiveness (Polusani et al., 2019; Stuelten et al., 2018). Collectively, the data suggest that the deregulation of JAC by PAI-1 is a mechanism for enhancing pro-tumorigenic changes in obesity-driven endometrial cancer.

Overall, our studies outline mechanistic insights that ASC-secreted PAI-1 induces transcriptomic and proteomic programs altering tumor-suppressive TGF- β signaling activities and JAC gene expression, which are associated with decreased cellular and ECM interactions. These negative transcriptional regulatory effects are mediated in part by the suppression of SMAD4 via PAI-1-LRP1 interactions. The current study demonstrates a paracrine effect of tumor-infiltrating and PAI-1-secreting ASCs

(D) Balloon plots of subpopulation distributions of endometrial tumors in the five categories based on histology and BMI. Circle area indicates the size of each subpopulation. Categories IV and V were associated with relatively high expression levels of SMAD4 and JAC, whereas low levels of SMAD4 and JAC were noted in categories I, II, and III.

(E) Violin plots showing the average expression levels of four JAC markers—ADAM15, ROCK1, Cx43, and Gas1—based on patients' tumor histology: papillary serous and endometrioid.

(F) Balloon plots of subpopulation distributions in the five categories stratified by BMI into three groups: normal weight (NW), obesity (OB), and morbid obesity (MOB).

(G) Violin plots showing the average expression levels of ADAM15, ROCK1, Cx43, and Gas1, based on three BMI groups: NW, OB, and MOB. Lower JAC expressions corresponded with patients with higher BMIs. **** $p < 0.0001$.

See also Figure S7.

on tumorigenic events in the epithelial compartment. Importantly, we provide evidence that this effect is heightened in persons with obesity. Epidemiologic studies have shown a positive correlation between BMI and the incidence of breast, ovarian, colon, and endometrial cancers (Sahoo et al., 2018). As the obesity epidemic widens, it is expected that the incidence of obesity-mediated cancers will rise sharply. We recommend that the role of PAI-1-mediated mechanisms outlined in this study be studied in other obesity-associated malignancies. Future studies will also examine whether PAI-1-LRP1 initiates a transcriptomic program shift from SMAD4- to STAT1-mediated gene transactivation and whether the long-term loss of SMAD4 leads to epigenetic silencing of target loci (Degryse et al., 2004). The expression signatures mediated by PAI-1 will need to be evaluated further in adiposity-linked endometrial clinical samples for tumorigenic development. Therefore, our current data provide the rationale to pursue translational studies for the development of proteomic signatures that evaluate the risk of endometrial cancer development in women with obesity.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

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AUTHOR CONTRIBUTIONS

Conceptualization, T.H.-M.H., N.B.K., and L.-L.L.; Methodology, T.H.-M.H., N.B.K., L.-L.L., and E.R.K.; Software, L.-L.L., C.-L.L., X.T., and C.-N.H.; Validation, L.-L.L., C.-M.W., and N.L.; Formal Analysis, L.-L.L., C.-L.L., and C.-P.W.; Investigation, L.-L.L., P.V., M.G.K., and A.C.D.; Resources, T.H.-M.H., N.B.K., and E.R.K.; Writing – Original Draft, L.-L.L., N.B.K., and T.H.-M.H.; Writing – Review & Editing, all authors; Funding Acquisition and Supervision, T.H.-M.H. and N.B.K.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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REFERENCES

- Angerer, P., Haghverdi, L., Büttner, M., Theis, F.J., Marr, C., and Buettner, F. (2016). destiny: diffusion maps for large-scale single-cell data in R. *Bioinformatics* 32, 1241–1243.
- Aoki-Kinoshita, K.F., and Kanehisa, M. (2007). Gene annotation and pathway mapping in KEGG. *Methods Mol. Biol.* 396, 71–91.
- Butler, A., Hoffman, P., Smibert, P., Papalexi, E., and Satija, R. (2018). Integrating single-cell transcriptomic data across different conditions, technologies, and species. *Nat. Biotechnol.* 36, 411–420.
- Carmeliet, P. (2003). Angiogenesis in health and disease. *Nat. Med.* 9, 653–660.
- Chen, T.J., and Kotecha, N. (2014). Cytobank: providing an analytics platform for community cytometry data analysis and collaboration. *Curr. Top. Microbiol. Immunol.* 377, 127–157.
- Chen, H., Lau, M.C., Wong, M.T., Newell, E.W., Poidinger, M., and Chen, J. (2016). Cytofkit: a bioconductor package for an integrated mass cytometry data analysis pipeline. *PLoS Comput. Biol.* 12, e1005112.
- Chusyd, D.E., Wang, D., Huffman, D.M., and Nagy, T.R. (2016). Relationships between rodent white adipose fat pads and human white adipose fat depots. *Front. Nutr.* 3, 10.
- Crandall, D.L., Quinet, E.M., El Ayachi, S., Hreha, A.L., Leik, C.E., Savio, D.A., Juhan-Vague, I., and Alessi, M.C. (2006). Modulation of adipose tissue development by pharmacological inhibition of PAI-1. *Arterioscler. Thromb. Vasc. Biol.* 26, 2209–2215.
- Czekay, R.P., Wilkins-Port, C.E., Higgins, S.P., Freytag, J., Overstreet, J.M., Klein, R.M., Higgins, C.E., Samarakoon, R., and Higgins, P.J. (2011). PAI-1: An integrator of cell signaling and migration. *Int. J. Cell Biol.* 2011, 562481.
- Daquinag, A.C., Tseng, C., Zhang, Y., Amaya-Manzanares, F., Florez, F., Dabbin, A., Zhang, T., and Kolonin, M.G. (2016). Targeted proapoptotic peptides depleting adipose stromal cells inhibit tumor growth. *Mol. Ther.* 24, 34–40.
- Daquinag, A.C., Dabbin, A., Snyder, B., Wang, X., Sahin, A.A., Ueno, N.T., and Kolonin, M.G. (2017). Non-glycanated decorin is a drug target on human adipose stromal cells. *Mol. Ther. Oncolytics* 6, 1–9.
- Degryse, B., Neels, J.G., Czekay, R.P., Aertgeerts, K., Kamikubo, Y., and Loskutoff, D.J. (2004). The low density lipoprotein receptor-related protein is a motogenic receptor for plasminogen activator inhibitor-1. *J. Biol. Chem.* 279, 22595–22604.

- Dobin, A., Davis, C.A., Schlesinger, F., Drenkow, J., Zaleski, C., Jha, S., Batut, P., Chaisson, M., and Gingeras, T.R. (2013). STAR: ultrafast universal RNA-seq aligner. *Bioinformatics* 29, 15–21.
- Elokda, H., Abou-Gharbia, M., Hennen, J.K., McFarlane, G., Mugford, C.P., Krishnamurthy, G., and Crandall, D.L. (2004). Tiplaxtinin, a novel, orally efficacious inhibitor of plasminogen activator inhibitor-1: design, synthesis, and pre-clinical characterization. *J. Med. Chem.* 47, 3491–3494.
- Escobedo, N., and Oliver, G. (2017). The lymphatic vasculature: its role in adipose metabolism and obesity. *Cell Metab.* 26, 598–609.
- Evans, W.H., and Martin, P.E. (2002). Gap junctions: structure and function (Review). *Mol. Membr. Biol.* 19, 121–136.
- Freese, K.E., Kokai, L., Edwards, R.P., Phillips, B.J., Sheikh, M.A., Kelley, J., Comerci, J., Marra, K.G., Rubin, J.P., and Linkov, F. (2015). Adipose-derived stem cells and their role in human cancer development, growth, progression, and metastasis: a systematic review. *Cancer Res.* 75, 1161–1168.
- Ghosh, S., Hughes, D., Parma, D.L., Ramirez, A., and Li, R. (2014). Association of obesity and circulating adipose stromal cells among breast cancer survivors. *Mol. Biol. Rep.* 41, 2907–2916.
- Gressel, G.M., Parkash, V., and Pal, L. (2015). Management options and fertility-preserving therapy for premenopausal endometrial hyperplasia and early-stage endometrial cancer. *Int. J. Gynaecol. Obstet.* 131, 234–239.
- Haghverdi, L., Büttner, M., Wolf, F.A., Büttner, F., and Theis, F.J. (2016). Diffusion pseudotime robustly reconstructs lineage branching. *Nat. Methods* 13, 845–848.
- Hammes, L.S., Korte, J.E., Tekmal, R.R., Naud, P., Edelweiss, M.I., Valente, P.T., Longatto-Filho, A., Kirma, N., and Cunha-Filho, J.S. (2007). Computer-assisted immunohistochemical analysis of cervical cancer biomarkers using low-cost and simple software. *Appl. Immunohistochem. Mol. Morphol.* 15, 456–462.
- Ishii, K., Takahashi, S., Sugimura, Y., and Watanabe, M. (2018). Role of stromal paracrine signals in proliferative diseases of the aging human prostate. *J. Clin. Med.* 7, 68.
- Jin, S., Martinelli, D.C., Zheng, X., Tessier-Lavigne, M., and Fan, C.M. (2015). Gas1 is a receptor for sonic hedgehog to repel enteric axons. *Proc. Natl. Acad. Sci. USA* 112, E73–E80.
- Kanaporis, G., Mese, G., Valiuniene, L., White, T.W., Brink, P.R., and Valiunas, V. (2008). Gap junction channels exhibit connexin-specific permeability to cyclic nucleotides. *J. Gen. Physiol.* 131, 293–305.
- Kauts, M.L., Pihelgas, S., Orro, K., Neuman, T., and Piirsoo, A. (2013). CCL5/CCR1 axis regulates multipotency of human adipose tissue derived stromal cells. *Stem Cell Res. (Amst.)* 10, 166–178.
- Kost, E.R., Valente, P.T., Lynch, B.A., Krishnegowda, N.K., Hertz, A.M., Hall, K.L., Riddle, N.D., and Tekmal, R.R. (2016). Clinical and pathologic features of hispanic endometrial cancer patients with loss of mismatch repair expression. *Int. J. Gynecol. Cancer* 26, 1129–1136.
- Kyo, S., Nakamura, M., Kiyono, T., Maida, Y., Kanaya, T., Tanaka, M., Yatabe, N., and Inoue, M. (2003). Successful immortalization of endometrial glandular cells with normal structural and functional characteristics. *Am. J. Pathol.* 163, 2259–2269.
- Lachance, J.A., Everett, E.N., Greer, B., Mandel, L., Swisher, E., Tamimi, H., and Goff, B. (2006). The effect of age on clinical/pathologic features, surgical morbidity, and outcome in patients with endometrial cancer. *Gynecol. Oncol.* 101, 470–475.
- Lambrechts, D., Wauters, E., Boeckx, B., Aibar, S., Nittner, D., Burton, O., Bassez, A., Decaluwé, H., Pircher, A., Van den Eynde, K., et al. (2018). Phenotype molding of stromal cells in the lung tumor microenvironment. *Nat. Med.* 24, 1277–1289.
- Liberzon, A., Birger, C., Thorvaldsdóttir, H., Ghandi, M., Mesirov, J.P., and Tamayo, P. (2015). The Molecular Signatures Database (MSigDB) hallmark gene set collection. *Cell Syst.* 1, 417–425.
- Lin, X., Liang, M., Liang, Y.Y., Brunnicardi, F.C., Melchior, F., and Feng, X.H. (2003). Activation of transforming growth factor-beta signaling by SUMO-1 modification of tumor suppressor Smad4/DPC4. *J. Biol. Chem.* 278, 18714–18719.
- Ma, T., Luan, S.L., Huang, H., Sun, X.K., Yang, Y.M., Zhang, H., Han, W.D., Li, H., and Han, Y. (2016). Upregulation of CC chemokine receptor 7 (CCR7) enables migration of xenogeneic human adipose-derived mesenchymal stem cells to rat secondary lymphoid organs. *Med. Sci. Monit.* 22, 5206–5217.
- Nagano, M., Hoshino, D., Sakamoto, T., Kawasaki, N., Koshikawa, N., and Seiki, M. (2010). ZF21 protein regulates cell adhesion and motility. *J. Biol. Chem.* 285, 21013–21022.
- Nevadunsky, N.S., Van Arsdale, A., Strickler, H.D., Moadel, A., Kaur, G., Levitt, J., Girda, E., Goldfinger, M., Goldberg, G.L., and Einstein, M.H. (2014). Obesity and age at diagnosis of endometrial cancer. *Obstet. Gynecol.* 124, 300–306.
- O’Sullivan, J., Lysaght, J., Donohoe, C.L., and Reynolds, J.V. (2018). Obesity and gastrointestinal cancer: the interrelationship of adipose and tumour micro-environments. *Nat. Rev. Gastroenterol. Hepatol.* 15, 699–714.
- Onstad, M.A., Schmandt, R.E., and Lu, K.H. (2016). Addressing the role of obesity in endometrial cancer risk, prevention, and treatment. *J. Clin. Oncol.* 34, 4225–4230.
- Parsons, J.T., Horwitz, A.R., and Schwartz, M.A. (2010). Cell adhesion: integrating cytoskeletal dynamics and cellular tension. *Nat. Rev. Mol. Cell Biol.* 11, 633–643.
- Polusani, S.R., Huang, Y.W., Huang, G., Chen, C.W., Wang, C.M., Lin, L.L., Osmulski, P., Lucio, N.D., Liu, L., Hsu, Y.T., et al. (2019). Adipokines deregulate cellular communication via epigenetic repression of gap junction loci in obese endometrial cancer. *Cancer Res.* 79, 196–208.
- Rota, M., Rumi, F., Bagnardi, V., Dal Maso, L., Zucchetto, A., Levi, F., La Vecchia, C., and Tavani, A. (2016). Modelling body mass index and endometrial cancer risk in a pooled-analysis of three case-control studies. *BJOG* 123, 285–292.
- Sahoo, S.S., Lombard, J.M., Ius, Y., O’Sullivan, R., Wood, L.G., Nahar, P., Jaaback, K., and Tanwar, P.S. (2018). Adipose-derived VEGF-mTOR signaling promotes endometrial hyperplasia and cancer: implications for obese women. *Mol. Cancer Res.* 16, 309–321.
- Seo, B.R., Bhardwaj, P., Choi, S., Gonzalez, J., Andresen Eguiluz, R.C., Wang, K., Mohanan, S., Morris, P.G., Du, B., Zhou, X.K., et al. (2015). Obesity-dependent changes in interstitial ECM mechanics promote breast tumorigenesis. *Sci. Transl. Med.* 7, 301ra130.
- Soliman, P.T., Oh, J.C., Schmeler, K.M., Sun, C.C., Slomovitz, B.M., Gershenson, D.M., Burke, T.W., and Lu, K.H. (2005). Risk factors for young premenopausal women with endometrial cancer. *Obstet. Gynecol.* 105, 575–580.
- Strong, A.L., Ohlstein, J.F., Biagas, B.A., Rhodes, L.V., Pei, D.T., Tucker, H.A., Llamas, C., Bowles, A.C., Dutreil, M.F., Zhang, S., et al. (2015). Leptin produced by obese adipose stromal/stem cells enhances proliferation and metastasis of estrogen receptor positive breast cancers. *Breast Cancer Res.* 17, 112.
- Stuelten, C.H., Parent, C.A., and Montell, D.J. (2018). Cell motility in cancer invasion and metastasis: insights from simple model organisms. *Nat. Rev. Cancer* 18, 296–312.
- Tanaga, K., Bujo, H., Zhu, Y., Kanaki, T., Hirayama, S., Takahashi, K., Inoue, M., Mikami, K., Schneider, W.J., and Saito, Y. (2004). LRP1B attenuates the migration of smooth muscle cells by reducing membrane localization of urokinase and PDGF receptors. *Arterioscler. Thromb. Vasc. Biol.* 24, 1422–1428.
- Van Der Maaten, L., and Hinton, G. (2008). Visualizing data using t-SNE. *J. Mach. Learn. Res.* 9, 2579–2625.
- van Tijn, P., de Vrij, F.M., Schuurman, K.G., Dantuma, N.P., Fischer, D.F., van Leeuwen, F.W., and Hol, E.M. (2007). Dose-dependent inhibition of proteasome activity by a mutant ubiquitin associated with neurodegenerative disease. *J. Cell Sci.* 120, 1615–1623.
- Vicente-Manzanares, M., Choi, C.K., and Horwitz, A.R. (2009). Integrins in cell migration—the actin connection. *J. Cell Sci.* 122, 199–206.
- Wang, J., Huo, K., Ma, L., Tang, L., Li, D., Huang, X., Yuan, Y., Li, C., Wang, W., Guan, W., et al. (2011). Toward an understanding of the protein interaction network of the human liver. *Mol. Syst. Biol.* 7, 536.

- Webb, D.J., Parsons, J.T., and Horwitz, A.F. (2002). Adhesion assembly, disassembly and turnover in migrating cells – over and over and over again. *Nat. Cell Biol.* 4, E97–E100.
- Wei, X., Li, S., He, J., Du, H., Liu, Y., Yu, W., Hu, H., Han, L., Wang, C., Li, H., et al. (2019). Tumor-secreted PAI-1 promotes breast cancer metastasis via the induction of adipocyte-derived collagen remodeling. *Cell Commun. Signal.* 17, 58.
- Wickham, H. (2009). *ggplot2: elegant graphics for data analysis* (Springer).
- Wise, M.R., Jordan, V., Lagas, A., Showell, M., Wong, N., Lensen, S., and Farquhar, C.M. (2016). Obesity and endometrial hyperplasia and cancer in premenopausal women: A systematic review. *Am. J. Obstet. Gynecol.* 214, 689.e681–689.e617.
- Xian, X., Ding, Y., Dieckmann, M., Zhou, L., Plattner, F., Liu, M., Parks, J.S., Hammer, R.E., Boucher, P., Tsai, S., and Herz, J. (2017). LRP1 integrates murine macrophage cholesterol homeostasis and inflammatory responses in atherosclerosis. *eLife* 6, e29292.
- Yamada, M., Hayashi, H., Suzuki, K., Sato, S., Inoue, D., Iwatani, Y., Ohata, M., Yuan, B., and Takagi, N. (2019). Furin-mediated cleavage of LRP1 and increase in ICD of LRP1 after cerebral ischemia and after exposure of cultured neurons to NMDA. *Sci. Rep.* 9, 11782.
- Yilmaz, E., Gul, M., Melekoglu, R., Inci Coskun, E., Sahin, N., Gul, S., Bastermur, A.G., and Ciplak, B. (2018). Neural precursor cell-expressed developmentally down-regulated 4-like: a new biomarker in the pathophysiology of endometrial cancer. *J. Int. Med. Res.* 46, 3709–3716.
- Zhang, Y., and Kolonin, M.G. (2016). Cytokine signaling regulating adipose stromal cell trafficking. *Adipocyte* 5, 369–374.
- Zhang, Y., Bellows, C.F., and Kolonin, M.G. (2010). Adipose tissue-derived progenitor cells and cancer. *World J. Stem Cells* 2, 103–113.
- Zhang, T., Tseng, C., Zhang, Y., Sirin, O., Corn, P.G., Li-Ning-Tapia, E.M., Troncoso, P., Davis, J., Pettaway, C., Ward, J., et al. (2016). CXCL1 mediates obesity-associated adipose stromal cell trafficking and function in the tumour microenvironment. *Nat. Commun.* 7, 11674.

STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-CD45 conjugated to 89Yb (clone H130)	Fluidigm	Cat# 3089003B; RRID:AB_2661851
Anti-CD181 conjugated to 142 Nd (clone 8F1/CXCR1)	Fluidigm	Cat# 3142009B
Anti-CD31 conjugated to 144Nd (clone WM59)	Fluidigm	Cat# 3143016B
Anti-CD182 conjugated to 147Sm (clone 5E8/CXCR2)	Fluidigm	Cat# 3147010B
Anti-CD34 conjugated to 148Nd (clone 581)	Fluidigm	Cat# 3148001B; RRID:AB_2810243
Anti-CD200 conjugated to 149Sm (clone OX104)	Fluidigm	Cat# 3149007B
CK8/18 (<i>KRT8/18</i> , clone C51)	Fluidigm	Cat# 3174014A
CD10 (clone H110a)	Fluidigm	Cat# 3158011B
ZO2 (clone 3E8D9)	ThermoFisher Scientific	Cat# 37-4700; RRID:AB_2533321
SMAD2 (clone 31H15L4)	ThermoFisher Scientific	Cat# 700048; RRID:AB_2532277
Fyn (<i>FYN</i>)	ThermoFisher Scientific	Cat# PA5-14780; RRID:AB_2108443
Gas1 (<i>GAS1</i>)	Thermo Fisher Scientific	Cat# PA5-48298; RRID:AB_2633756
ZO1 (<i>ZO1</i> , clone 1A12)	ThermoFisher Scientific	Cat# 33-9100; RRID:AB_87181
SDC3 (<i>SDC3</i>)	R&D Systems	Cat# AF3539; RRID:AB_622125
SMAD4 (<i>SMAD4</i> , clone 253343)	R&D Systems	Cat# MAB2097; RRID:AB_2286470
TGFR2 (<i>TGFBR2</i>)	R&D Systems	Cat# AF241-NA; RRID:AB_354416
EpCAM (<i>EPCAM</i>)	R&D Systems	Cat# AF960; RRID:AB_355745
SDC3 (<i>SDC3</i>)	R&D Systems	Cat# AF3539; RRID:AB_622125
SMAD4 (<i>SMAD4</i>)	R&D Systems	Cat# AF2097; RRID:AB_355150
cIAP1 (<i>BIRC2</i>)	Abcam	Cat# ab226031
Paxillin (<i>PXN</i> , clone Y113)	Abcam	Cat# ab216652
ROCK1 (<i>ROCK1</i> , clone EP786Y)	Abcam	Cat# ab219587
ADAM15 (clone 23G9)	Novus Biologicals	Cat# MAB935-100
Ubiquitin	Novus Biologicals	Cat# NB300129; RRID:AB_2180545
PAI-1 (clone C-9)	Santa Cruz Biotechnology	Cat# sc-5297; RRID:AB_628154
CD31	Santa Cruz Biotechnology	Cat# sc-1506-R; RRID:AB_831096
Cx43	Cell Signaling Technology	Cat# 3512S; RRID:AB_1007667
Cx43 (<i>GJA1</i>)	Sigma	Cat# SAB4501175; RRID:AB_10746172
AMIGO2 (<i>AMIGO2</i>) (clone 30G9.1G2)	Kerafast	Cat# EBS001
LRP1	Cell Signaling Technology	Cat# 64099S; RRID:AB_2799654
LRP1-ICD	Abcam	Cat# ab92544; RRID:AB_2234877
β-actin	R&D Systems	Cat# MAB8929; RRID:AB_10716360
20X Anti-Rabbit HRP Conjugate	ProteinSimple	Cat# 043-426
Normal goat IgG antibody	Abcam	Cat# ab37373
Normal rabbit IgG antibody	ThermoFisher Scientific Pierce Magnetic ChIP Kit	Cat# 26157
Biological Samples		
Human endometrial tumor FFPE tissues	The University of Texas Health Science center	Protocol: HSC20180584HR
Human endometrial tumors for CyTOF	The University of Texas Health Science center	Protocol: HSC20180584HR
Chemicals, Peptides, and Recombinant Proteins		
WAT7 peptides	This paper	N/A
Cell-ID Cisplatin	Fluidigm	Cat# 201064

(Continued on next page)

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Cell-ID Intercalator-Ir	Fluidigm	Cat# 201192A
Tiplaxtinin (PAI-039)	Selleckchem	Cat# S7922
RAP	Millipore Sigma	Cat# 55-350-650UG
MG132	Sigma Aldrich	Cat# M7449-200UL
RIPA buffer	Thermo Fisher Scientific	Cat# 89901
Fix I Buffer	Fluidigm	Cat# 201065
Critical Commercial Assays		
MILLIPLEX MAP Human Adipocyte Magnetic Bead Panel - Endocrine Multiplex Assay	MilliporeSigma	Cat# HADCYMag-61K
Qubit dsDNA HS Assay Kit	Invitrogen	Cat# Q32854
C1 Single-Cell mRNA Seq HT 10-17 μ m IFC	Fluidigm	Cat# 101-4982
C1 Single-Cell mRNA Seq HT Reagent Kit v2	Fluidigm	Cat# 101-3473
Nextera® XT DNA Library Prep Kit	Illumina	Cat# FC-131-1096
Nextera® XT Index Kit v2 Set A	Illumina	Cat# FC-131-2001
Nextera® XT Index Kit v2 Set B	Illumina	Cat# FC-131-2002
Direct-zol RNA Microprep Kits	Zymo Research	Cat# R2062
High-Capacity cDNA Reverse Transcription Kit	Applied Biosystems	Cat# 4368814
LightCycler® 480 SYBR Green I Master	Roche	Cat# 04887352001
12-230 kDa Wes Separation Module, 8 × 25 capillary cartridges	ProteinSimple	Cat# SM-W004
Anti-Rabbit Detection Module for Wes, Peggy Sue or Sally Sue	ProteinSimple	Cat# DM-001
Anti-Mouse Detection Module for Wes, Peggy Sue or Sally Sue	ProteinSimple	Cat# DM-002
2nd Generation Packaging System Mix	Abcam	Cat# LV003
Duolink® <i>In Situ</i> Red Starter Kit Mouse/Rabbit	Sigma Aldrich	Cat# DUO92101
Duolink® <i>In Situ</i> Red Starter Kit Goat/Rabbit	Sigma Aldrich	Cat# DUO92105
Pierce Magnetic ChIP Kit	ThermoFisher Scientific	Cat# 26157
Cell-ID 20-Plex Pd Barcoding Kit	Fluidigm	Cat# 201060
Deposited Data		
Raw Single-cell RNA-seq data	This paper	GEO: GSE140130
Experimental Models: Cell Lines		
EME6/E7/TERT	Kyo et al., 2003	N/A
Adipose stromal cells	Coriell Institute	Cat# AG17870
Adipose stromal cells	Zen-Bio Inc.	Cat# ASC-F-SL Lot# ASC0051, ASC0053, ASC0060, ASC0061, ASC0065, ASC0064, Cat# ASC-F Lot# ASC032003, ASC050709B, Cat# ASC-F-D Lot# ASCD011906
Human endometrial epithelial cells (HEuEC)	Lifeline Cell Technology	FC-0078
Oligonucleotides		
SMAD4 shRNA	Sigma-Aldrich	Cat# TRCN0000040032
Primers for qRT-PCR primers, see Table S5	This paper	N/A
Primers for ChIP-qPCR primers, see Table S4	This paper	N/A
Recombinant DNA		
pLKO.1 puro	Addgene	Cat# 8453
Software and Algorithms		
ZEN blue software	Zeiss Company	N/A
Seurat algorithm	Butler et al., 2018	https://satijalab.org/seurat/

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Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
MeV	N/A	http://mev.tm4.org
R package: ggplot2	Wickham, 2009	https://ggplot2.tidyverse.org
R package: Rtsne	Van Der Maaten and Hinton, 2008	https://github.com/jkrijthe/Rtsne
Compass for SW (Version: 3.1.7)	ProteinSimple	N/A

RESOURCE AVAILABILITY

Lead Contact

Further information and requests for resources should be directed to and will be fulfilled by the Lead Contact, Tim Hui-Mng Huang (huangt3@uthscsa.edu).

Materials Availability

This study did not generate new unique reagents.

Data and Code Availability

The raw single-cell RNA-seq data have been deposited in NCBI GEO under accession number GSE140130. Original data of Figure 1C, 4B, 4C, and 4F have been deposited to Mendeley Data (<http://doi.org/10.17632/5ywmdd6fnd.1>).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Clinical Samples

Tumor tissues for tissue microarray and CyTOF analyses were collected following the IRB protocols at the University of Texas Health Science Center at San Antonio (Tables S1 and S2). The Cancer Genome Atlas (TCGA) Uterine Corpus Endometrial Carcinoma (UCEC) cohort was used to identify BMI-associated genes.

Cell Lines, Cell Culture and In Vitro Exposure Model

ASCs isolated from ten female subjects were purchased from Coriell Institute (Camden, NJ, Cat# AG17870) and Zen-Bio Inc. (Research Triangle Park, NC, Cat# ASC-F-SL Lot# ASC0051, ASC0053, ASC0060, ASC0061, ASC0065, ASC0064, Cat# ASC-F Lot# ASC032003, ASC050709B, Cat# ASC-F-D Lot# ASCD011906). ASCs were cultured in non-differentiating media (DMEM supplemented with 5% FBS) and grown on fibronectin-coated culture dishes. For ASC-condition media (ASC-CM) exposure model, cells were treated with ASC-CM of 48 hr (Polusani et al., 2019). Non-tumorigenic and immortalized endometrial epithelial cell (EEC) line EM-E6/E7/TERT-1 (designated as EME6/7t) were established and characterized by Kyo et al. (2003). Cell line authentication was carried out by STR analysis at ATCC. EECs were cultured in Dulbecco's Modified Eagle Medium: Nutrient Mixture F-12 (DMEM/F-12) supplemented with 10% fetal bovine serum (FBS) and 100 U/mL penicillin/streptomycin. Un-immortalized human endometrial epithelial cells (HEuEC) were obtained from Lifeline Cell Technology (Cat# FC-0078) and cultured in reproductive medium complete kit (Lifeline Cell Technology, Cat# LL-0068) according to the manufacturer's instructions. For scRNA-seq, CyTOF analysis and function assays, PAI-1 antagonist, tiplaxtinin (PAI-039, 50 μ M, Selleckchem, Cat# S7922) was used with ASC-CM for 24 hr (Polusani et al., 2019). Before ASC-CM treatment, LRP1 antagonist, RAP (5 μ g/mL, Millipore Sigma, Cat# 55-350-650UG) was used to treat cells for 30 min (Tanaga et al., 2004); Proteasome Inhibitor, MG132 (10 μ M, Sigma Aldrich, Cat# M7449-200UL) was used to treat cells for 3 hr (van Tijn et al., 2007).

Generation of SMAD4 Stable Knockdown Cells

SMAD4 short hairpin RNA (shRNA) lentiviral particles were produced using MISSION shRNA Bacterial Glycerol Stock SMAD family member (TRCN0000040032, MISSION shRNA Bacterial Glycerol Stock; Sigma-Aldrich) according to the protocol provided. The shRNA nontargeting control vector pLKO.1 puro (Addgene, Cat# 8453) was used as a negative control. 293T cells were co-transfected with 2nd generation packaging system mix (ABM, Cat# LV003) according to the manufacturer's instructions. The lentivirus was harvested 48 hr later, filtered and infected EECs for 24 hr. Stable cell clones were generated by replacing with complete media containing 2 μ g/mL puromycin and selected for 3 weeks.

METHOD DETAILS

Immunohistochemistry (IHC)

Sections of formalin-fixed paraffin-embedded tissues were analyzed by using antigen retrieval, washing, and blocking. Adipose stromal cells (ASCs) were stained with the WAT7 peptide and endothelial cells with CD31 antibody, as described previously (Daquinag

et al., 2017). Differences in the number of ASCs within the stromal compartment of the tumor sections were counted and scored as previously described by two investigators independently (Hammes et al., 2007).

Luminex Technology Multiplex Assays

Fresh conditioned media was collected from culture dishes, and debris/cells were removed by centrifugation and 0.45 μm filters. PAI-1 levels were measured along with other adipokines and cytokines ($n = 11$) by using MILLIPLEX MAP Human Adipocyte Magnetic Bead Panel - Endocrine Multiplex Assay (MilliporeSigma, Cat# HADCYMag-61K) following the manufactures instructions. The values were detected by Luminex® 200 multiplexing system (MilliporeSigma). Levels of PAI-1 and other cytokines/adipokines were calculated in reference to a standard curve and normalized to total proteins from ASCs in cell culture.

Matrigel Invasion Assay

A 24-well chamber plate with polycarbonate member, 0.8 μm pore size (Thermo Scientific, Cat# 140629) was utilized to evaluate cell motility. The inserts were coated with Matrigel to shape a matrix barrier. 5×10^4 cells were plated in the upper compartment and cultured in serum-free medium. The bottom chambers were filled with 600 μL ASC-CM or ASC's regular media. Cells were incubated at 37°C for 18 hr and then removed by rubbing the upper surface of the membrane. After fixed in 70% methanol for 30 min, stained with 0.1% crystal violet for 10 min, and washed, migrated cells on lower surface of membrane were counted in six randomly selected microscopic fields and quantified using ImageJ software.

Cellular Proliferation and Migration Assays

Proliferation and migration assays were conducted using a live-cell imaging system IncuCyte (Essen Bioscience). For cellular proliferation, 3,000 cells/per well were plated in quadruplicate in 96 wells plates. For migration assay, cells were plated in quadruplicate in 96 wells plates and scratched using a 96-pin WoundMaker of IncuCyte scratch system when cells were grown to confluence. Cell and wound images were automatically carried. The real-time kinetic data of proliferation and migration was calculated as phase object confluence and wound confluence, respectively, and analyzed by IncuCyte software system.

Single-cell RNA-seq

For single-cell capture, cells were loaded into a 10-17 μm Fluidigm C1 mRNA Seq HT Integrated Fluidic Circuit (IFC), capable of capturing up to 800 cells, and processed by the C1 Single-Cell Auto Prep System. Single-cell cDNA was generated in the C1 System in an automated fashion, according to manufacturer's protocols (Fluidigm). Single-cell cDNA samples were then subjected to tagmentation and amplification. Indices were added during the enrichment step using the Nextera XT Index Kit v2 (Illumina) to generate barcoded cDNA libraries. The cDNA libraries sequencing was performed in the NextSeq 500 sequencing system (Illumina), with each sample running in one lane of a 4 lanes flowcell. The loading amount was adjusted for each sample (1.8 pM pooled library + 20% Phix which adds diversity for sequencing quality) following manufacturer's instruction.

Single-cell RNA-seq Data Processing

The output of scRNA-seq data from NextSeq 500 was first demultiplexed using stand-alone API script (Fluidigm), and R2 reads were aligned to GRCh38 human genome in STAR aligner (Dobin et al., 2013). The expression levels were further quantified as FPKM values using RSEM. For data quality purposes, sequenced cells with more than 150,000 reads were analyzed ($n \geq 600$ cells). The average gene numbers of each treatment are Ctrl: 6,786, CM: 6,696, CM+Tip: 7,253. The average gene number of all single cells is 6,911. Seurat software was applied to assess the batch effect by canonical correlation analysis (CCA), cluster the cells by performing non-linear dimensional reduction, tSNE, based on the information of the principal component analysis (Butler et al., 2018). The differentially expressed genes ($n = 1,301$) across all clusters between the treatments were the same with and without CCA, indicating that the batch effect if any is minimal. The differentially expressed genes in each cluster were further identified using FindMarkers in Seurat.

scRNA-seq data z-score normalization was performed for expression analysis. Heatmaps, t-SNE and violin plots (R packages: Rtsne and ggplot2) were used to display the cluster maps of different cell treatments (Van Der Maaten and Hinton, 2008; Wickham, 2009). For the pseudotime analysis, z-score of each subpopulation was analyzed by the R package destiny, which produced a dimensionality reduction and a pseudotime prediction. The dimensionality of data (DC1 and DC2) and diffusion pseudotime (DPT) was calculated to display distribution maps of individual subpopulation and their expression patterns. In pathway enrichment analysis, gene lists were collected from Hallmark Gene Sets, and the distribution of different signaling pathways was plotted using R packages: reshape2 and ggplot2. For gene correlation matrix analysis, the correlation between oncogenic signaling pathways and cellular junction complexes was calculated using R package corplot.

In Situ Proximity Ligation Assay (PLA)

Protein-protein interactions in different EECs treatment groups were examined by PLA using Duolink® In Situ Red Starter Kit following the manufactures instructions (Sigma Aldrich, Cat# DUO92101 and # DUO92105). For the association assay between LRP1 and PAI-1, cells were permeabilized with (intracellular interaction) or without (cell surface interaction) 0.2% (v/v) Triton X-100 for 10 min after fixing. Cells were incubated with rabbit polyclonal anti-LRP-1 antibody (1:250, Cell Signaling Technology,

Cat# 64099S) and mouse monoclonal anti-PAI-1 antibody (1:200, Santa Cruz Biotechnology, Cat# sc-5297), which were tagged with anti-rabbit PLUS and anti-mouse MINUS probes. For the association assay between SMAD4 and ubiquitin, cells were permeabilized with 0.2% (v/v) Triton X-100 for 10 min after fixing and incubated with goat polyclonal anti-SMAD4 antibody (1:200, R&D Systems, Cat# AF2097) and rabbit polyclonal anti-ubiquitin antibody (1:200, Novus Biologicals, Cat# NB300129) which were tagged with anti-goat PLUS and anti-rabbit MINUS. The cells were then mounted with DAPI after washing and signal amplification with Duolink reagents. Cell images were acquired by the Zeiss LSM 710 confocal microscopy and ZEN blue software.

Chromatin Immunoprecipitation (ChIP)-qPCR

ChIP assays were performed using Pierce Magnetic ChIP Kit according to the manufacturer's protocol (ThermoFisher Scientific, Cat# 26157). All reagents used were included in the kit. Briefly, 4×10^6 cells were cross-linked using 1% paraformaldehyde for 5 min and reactions were quenched by glycine for 5 min at room temperature. After lysis and MNase digestion, DNA fragments in the cell lysates were sheared to size 300-500 bp by the Q800R3 Sonicator (20% amplitude, 10 s on, 20 s off for 8 min) (Qsonica). Then, cell lysates were incubated with an anti-SMAD4 antibody (R&D Systems, Cat# AF2097), anti-LRP1-ICD antibody (Abcam, Cat# ab92544), normal goat IgG antibody (Abcam, Cat# ab37373) or normal rabbit IgG antibody (Pierce Magnetic ChIP Kit, ThermoFisher Scientific, Cat# 26157) overnight at 4°C. The immunoprecipitated genomic DNA fragments were eluted and amplified using quantitative real-time PCR. Specific antibody signals were normalized to the input signals. Quantitative real-time PCR results were compared statistically using Student's t test. Primers used for qPCR analysis are listed in [Table S4](#).

Reverse Transcription (RT)-qPCR

RNA was isolated by Direct-zol RNA Microprep Kits (Zymo Research, Cat# R2062) and reversed transcribed using a High-Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Cat# 4368814). Quantitative real-time PCR was performed using a LightCycler® 480 SYBR Green I Master (Roche, Cat# 04887352001) by LightCycler® 480 System (Roche). Each sample was tested at least in triplicate. Target gene expression levels were normalized based on the housekeeping ubiquitin (UBB) gene expression levels. Relative mRNA levels were calculated using the $2^{-\Delta\Delta Ct}$ method. PCR primers for specific gene amplification are listed in [Table S5](#).

Quantitative Capillary Western Immunoassay

Cells were lysed on ice using RIPA buffer (Thermo Fisher Scientific, Cat# 89901). The lysates were analyzed using the 12–230 kDa Wes separation module of the quantitative capillary Western immunoassay system (ProteinSimple). Anti-rabbit or anti-mouse detection module and anti-goat secondary antibodies were used for Wes per the manufacturer's recommendation (ProteinSimple). Primary antibodies used were as follows: LRP1 (1:25, Cell Signaling Technology, Cat# 64099S), SMAD4 (1:10, R&D Systems, Cat# AF2097), and β -actin (1:200, R&D Systems, Cat# MAB8929) as a housekeeping control. Data were analyzed and displayed in Compass for SW (Version: 3.1.7).

CyTOF Mass Cytometry

CyTOF staining panels are detailed in [Table S3](#). Antibodies were purchased from the Fluidigm or conjugated in-house according to the manufacturer's instructions (Fluidigm). Tumors were dissociated in digestion medium as previously described ([Lambrechts et al., 2018](#)), and then passed through 40 μ m filters into PBS at 4°C. The cells were centrifuged at 300 g for 5 min at 4°C and then depleted of erythrocytes by hypotonic lysis for 5 min at room temperature. Single-cell suspensions were incubated with Cell-ID Cisplatin (Fluidigm, Cat# 201064) to label dead cells to be excluded from analysis. Cells were fixed and permeabilized by Fix I Buffer (Fluidigm, Cat# 201065) and then patients' samples were barcoded by Cell-ID 20-Plex Pd Barcoding Kit following the manufacturer's instructions (Fluidigm, Cat# 201060). After washing, cells were incubated with the panel of antibodies for 1h at room temperature for cell staining. Cells were then fixed with MaxPar Fix and Perm Buffer (Fluidigm, Cat #201067) containing Cell-ID Intercalator-Ir 191/193 (Fluidigm, Cat #201192A) at 0.125 mM and 4°C overnight to stain the nuclei. After signal beads were normalized by EQ Four Element Calibration, cell events were analyzed by Helios mass cytometer (Fluidigm).

After raw data debarcoding and normalization by the CyTOF software (Version 6.7.1014, Fluidigm), cell gating was performed in Cytobank (<https://www.cytobank.org/>, Cytobank Inc.) ([Chen and Kotecha, 2014](#)). Cytokit ([Chen et al., 2016](#)) was used for qualitative assessments and visual illustrations by an automated clustering algorithm PhenoGraph. Using the R package Cytokit, data was clustered and represented in t-SNE maps. Heatmaps (MeV), violin plots (R package ggplot2), and balloon plots (R package ggplot2) were employed to display differences in expression and subpopulation size in different clusters. For pseudotime analysis, normalized CyTOF data was analyzed by R package Destiny (version 3.8) which produces dimensionality reduction and pseudotime prediction ([Angerer et al., 2016](#)). Samples from each patient were randomly assessed for 300 cells by R package. DC1 and DC2 and DPT (R package ggplot2) were calculated to plot diffusion maps.

QUANTIFICATION AND STATISTICAL ANALYSIS

Statistical analysis was conducted using Microsoft excel Office 365 and Duncan's multiple range test by R package. Mann-Whitney U tests were used to compare the expression of individual genes in tumor and normal samples in the Cancer Genome Atlas (TCGA)

Uterine Corpus Endometrial Carcinoma (UCEC) cohort. scRNA-seq and CyTOF data were normalized before signal or expression analysis. Data was presented as mean \pm SD (standard deviation) in the figure legends unless otherwise stated. Significance was determined by a Student's t test and p values are indicated in the figure legends. A $p < 0.05$ was considered as statistical significance. Duncan's multiple range test was employed to compare gene expression among different individuals with BMI in TCGA cohort.

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Supplemental Information

**PAI-1-Dependent Inactivation
of SMAD4-Modulated Junction and Adhesion Complex
in Obese Endometrial Cancer**

Li-Ling Lin, Edward R. Kost, Chun-Lin Lin, Philip Valente, Chiou-Miin Wang, Mikhail G. Kolonin, Alexes C. Daquinag, Xi Tan, Nicholas Lucio, Chia-Nung Hung, Chen-Pin Wang, Nameer B. Kirma, and Tim H.-M. Huang

Table S1: Patients information for tissue microarrays from endometrial tumors. Related to Figure 1.

Age	27-50 (mean=41.5, total, n=35)
Histology	Endometrioid (94%) Clear cell (3%) Carcinosarcoma (3%)
FIGO Stage	I (86%) II (6%) III & IV (8%)
BMI	20-29 (20%) 30-39 (37%) ≥ 40 (43%)

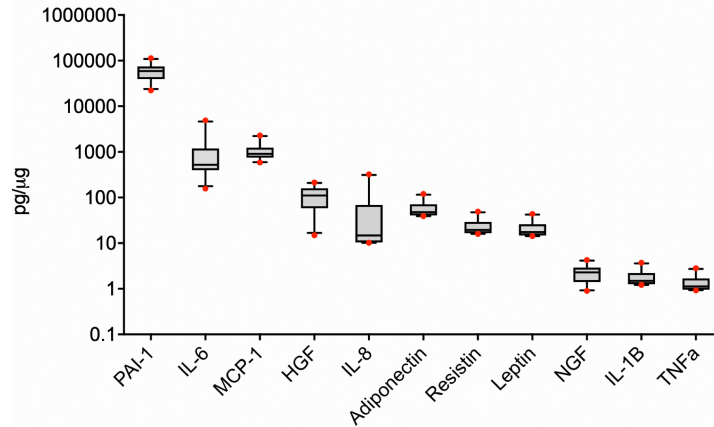


Figure S1: Abundant PAI-1 in conditioned media of adipose stromal cells. Related to Figure 1.

Conditioned media of 10 adipose stromal cell (ASC) lines were examined by FlexMap multiplexing assays for a panel of adipokines and cytokines. Secreted protein levels (pg), normalized to total proteins (μg) of cells in culture at the time of harvest, show about two orders of magnitude more abundance of PAI-1 compared to the other adipokines/cytokines.

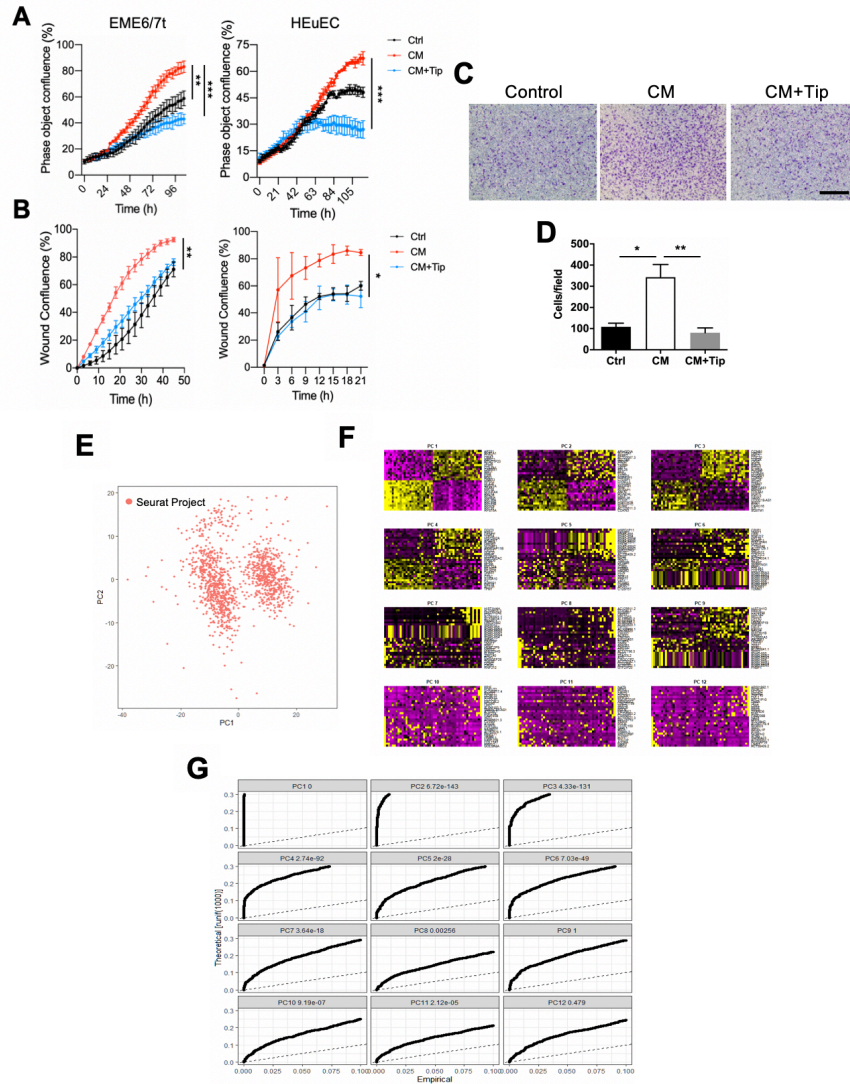


Figure S2: PAI-1-initiated pro-tumorigenic phenotype in endometrial epithelial cells. Related to Figure 2.

Immortalized EME6/7t endometrial epithelial cells (EECs) and an additional source of un-immortalized HEuEC primary cells were exposed to ASC-CM in the presence or absence of PAI-1 antagonist, Tip. Proliferation (**A**) and migration (**B**) rates in quadruplicate were determined using a live-cell imaging system IncuCyte. (**C-D**) Invasion assay of EME6/7t cells exposed to ASC-CM and CM+Tip, showing significantly enhanced invasive potential of EME6/7t cells after CM exposure, which was reversed by Tip. Scale: 500 μ m. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$. (**E**) Principal component analysis (PCA) utilizing the most variable genes. (**F**) Heatmaps showing the genes and cells ordered by PCA scores and used to select principal components. Yellow indicating high gene expression, and purple indicating low expression. (**G**) The JackStraw plots displaying p-values for each principal component. Dashed line, uniform distribution; Solid curve, the significant principal component.

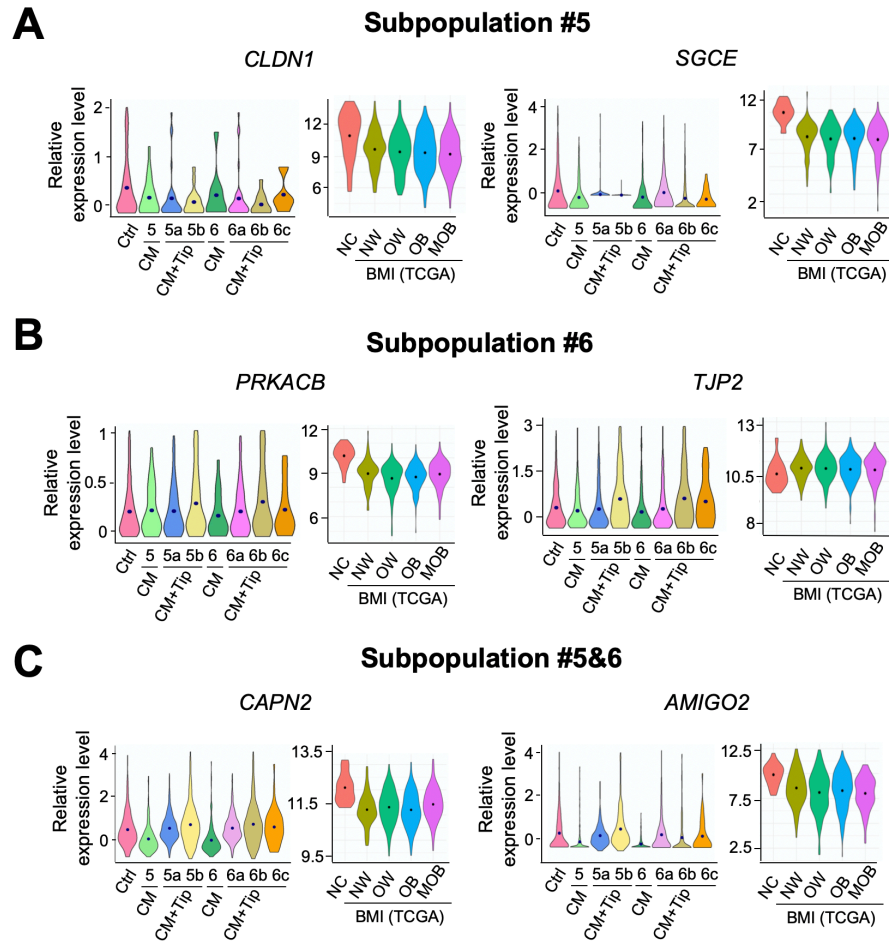


Figure S3: Representative BMI-associated JAC genes were regulated both in EEC subpopulations #5 and 6 after exposure to ASC-CM. Related to Figure 3.

Violin plots showing expression changes of (A) BMI-associated JAC genes corresponding to subpopulation #5, (B) BMI-associated JAC genes corresponding to subpopulation #6, (C) BMI-associated JAC genes that correspond to both subpopulations #5 and 6. The expression of genes in immortalized endometrial epithelial cells EME6/7t with or without exposure to ASC-CM (-/+ tiplaxtinin, Tip; plots on left) and in TCGA endometrial cancer cohort stratified by obesity status (plots on right). NC, non-cancer control; and cancer patients: NW, normal weight; OW, overweight; Ob, obese; MOB, morbidly obese.

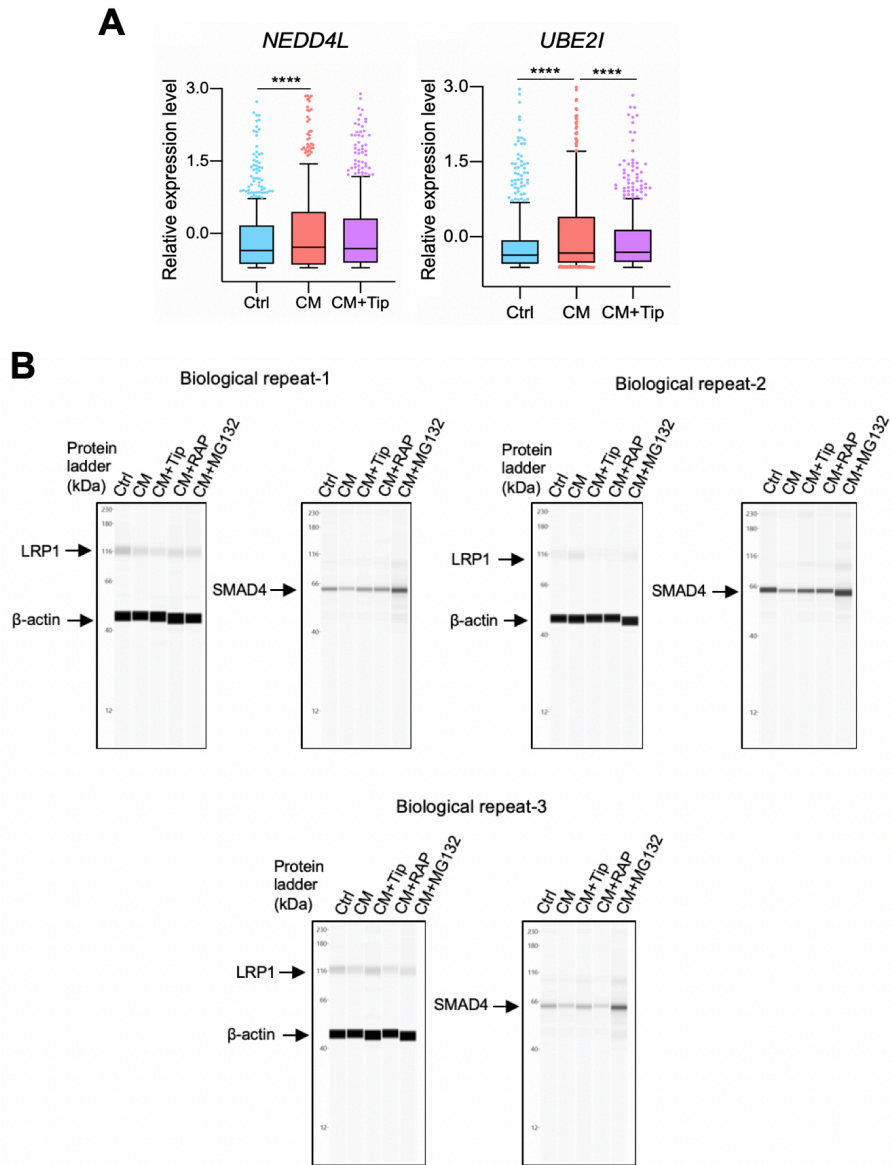


Figure S4: PAI-1 upregulates *NEDD4L* and *UBE2I* genes and suppresses SMAD4 protein expression but not LRP1 in endometrial epithelial cells. Related to Figure 4.

(A) Single-cell RNA-seq analysis reveals increased expression levels of *NEDD4L* and *UBE2I* in immortalized endometrial epithelial cells EME6/7t exposed to ASC-CM relative to cells additionally treated with a PAI-1 antagonist (Tip) or control EECs. **** $p < 0.0001$. (B) Capillary Western immunoassays showing protein expression levels of LRP1 and SMAD4 of EME6/7t cells in three independent experiments (three biological replicates).

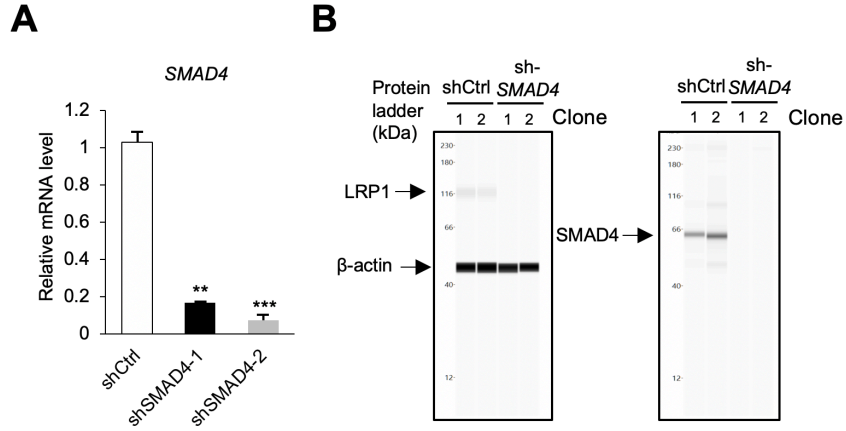


Figure S5: Expression levels of SMAD4 and LRP1 in two *SMAD4* table knockdown clones.

Related to Figure 5.

(A) Bar plots showing efficient *SMAD4* knockdown in EME6/7t cells in the two stable sh*SMAD4* clones relative to control shCtrl clone. ** $p < 0.01$, *** $p < 0.001$. (B) Capillary Western immunoassays showing protein expression levels of LRP1 and SMAD4 in the two stable shCtrl clones or two stable sh*SMAD4* clones of EME6/7t cells.

Table S2: Patients information for CyTOF analysis. Related to Figure 6.

Case#	Age	BMI	Ethnicity	FIGO Stage	FIGO Grade	Histology	Type
P07	61	20.5	African American	IB	3	Papillary serous	2
P08	56	21.7	non-Hispanic, white	IIIC2	3	Papillary serous	2
P09	72	28.0	Hispanic	IA	1	Endometrioid	1
P14	35	30.0	Hispanic	II	3	Endometrioid	1
P06	43	34.1	Hispanic	IA	1	Endometrioid	1
P15	56	34.7	non-Hispanic, white	IA	2	Endometrioid	1
P03	31	41.6	Hispanic	IA	1	Endometrioid	1
P10	53	48.1	Hispanic	IA	1	Endometrioid	1
P11	29	55.0	Hispanic	IB	1	Endometrioid	1

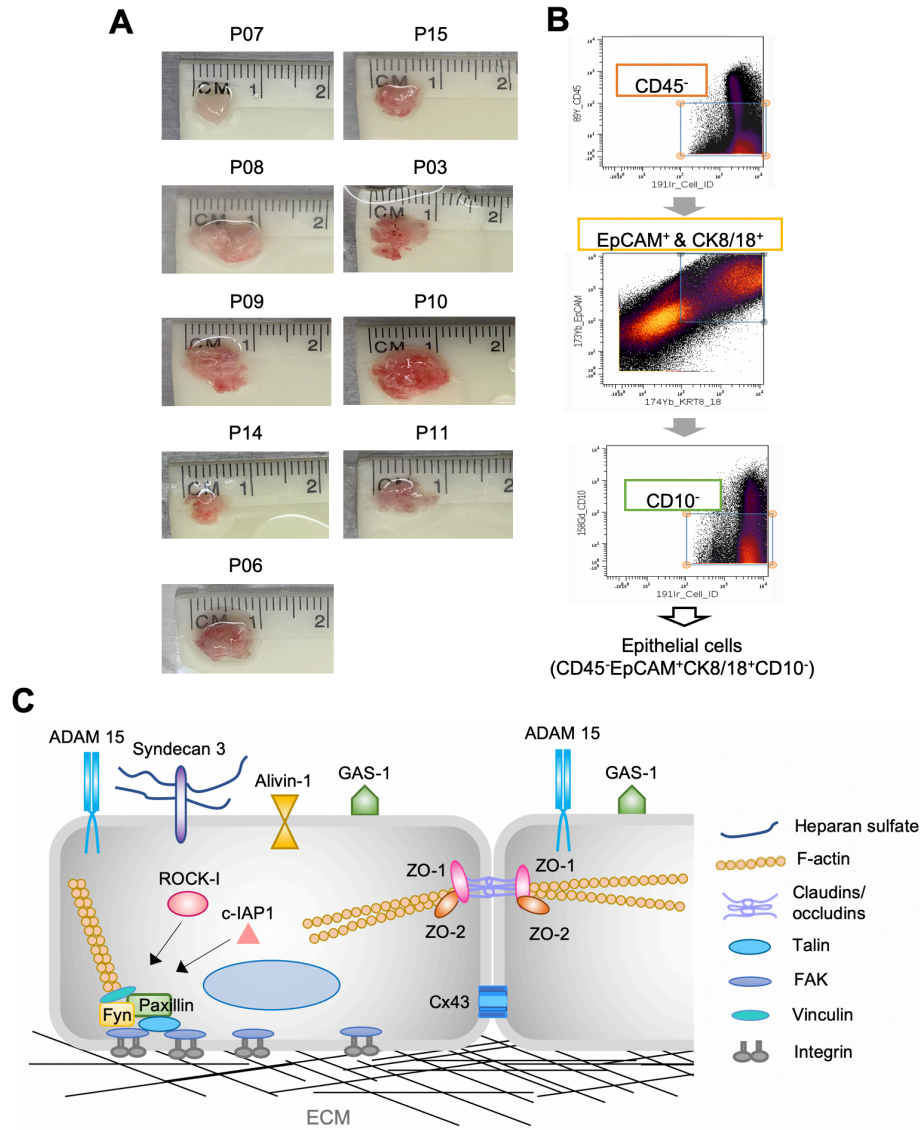


Figure S6: Isolating epithelial tumor cells from tissue of endometrial cancer patients and a schematic overview of JAC functions in epithelial cells. Related to Figure 6.

(A) Tumor appearance from endometrial cancer patients. (B) CyTOF gating strategies to analyze epithelial cells from endometrial tumors. (C) Apical surface protein, growth arrest-specific 1 (Gas1), apical junction (ADAM15, syndecan 3 (SDC3), and AMIGO2), and tight junction (zonula occludens (ZO)1 and ZO2) maintain epithelial polarity. Gap junction (connexin 43, Cx43) connects cell to cell and mediates intercellular communication. Rho-associated protein kinase 1 (ROCK1) and cellular inhibitor of apoptosis protein 1 (cIAP1) are focal adhesion regulators. Paxillin and Fyn associated with integrins adhere to the extracellular matrix (ECM).

Table S3: CyTOF antibody panel. Related to Figure 6.

Protein name	Gene name	Clone	Metal	Cellular functions
Gas1	<i>GAS1</i>		166Er	Cooperate with hedgehog signals
AMIGO2	<i>AMIGO2</i>	30G9.1G2	143Nd	Decrease cell adhesion/migration
SDC3	<i>SDC3</i>		141Pr	Cooperate with hedgehog signals
ADAM15	<i>ADAM15</i>	23G9	154Sm	Bind to $\alpha v\beta 3$ and $\alpha 5\beta 1$ integrin to mediate cell motility
Cx43	<i>GJA1</i>		162Dy	Assemble to gap-junction intercellular communication channels
ZO1	<i>TJP1</i>	1A12	169Tm	Interaction with F-actin and occludin in tight junction complexes
ZO2	<i>TJP2</i>	3E8D9	146Nd	Interaction with F-actin and occludin in tight junction complexes
ROCK1	<i>ROCK1</i>	EP786Y	161Dy	Cooperate with zonula adherens organization, RhoA signaling, and cortical tension
Paxillin	<i>PXN</i>	Y113	160Gd	As a docking protein to recruit signaling proteins to focal adhesions and coordinate downstream signaling
Fyn	<i>FYN</i>		153Eu	Associate with Src and focal adhesion kinase (FAK) to regulate cell migration
cIAP1	<i>BIRC2</i>		167Er	Control Rho GTPases signaling for focal adhesion formation
TGFR-2	<i>TGFBR2</i>		165HO	The ligand-binding receptor for TGF β signaling
SMAD2	<i>SMAD2</i>	31H15L4	152Sm	Form complex with SMAD3 or/and SMAD4 to induce TGF β signaling
SMAD4	<i>SMAD4</i>	253343	164Dy	A central mediator of TGF β signaling and induces a tumor suppressive effect
CK 8/18	<i>KRT 8/18</i>	C51	174Yb	Cytoskeletal proteins and be induced by endometrial cancer invasion
EpCAM	<i>EPCAM</i>		173Yb	As an epithelial tumor cell marker and associate with epithelial proliferation

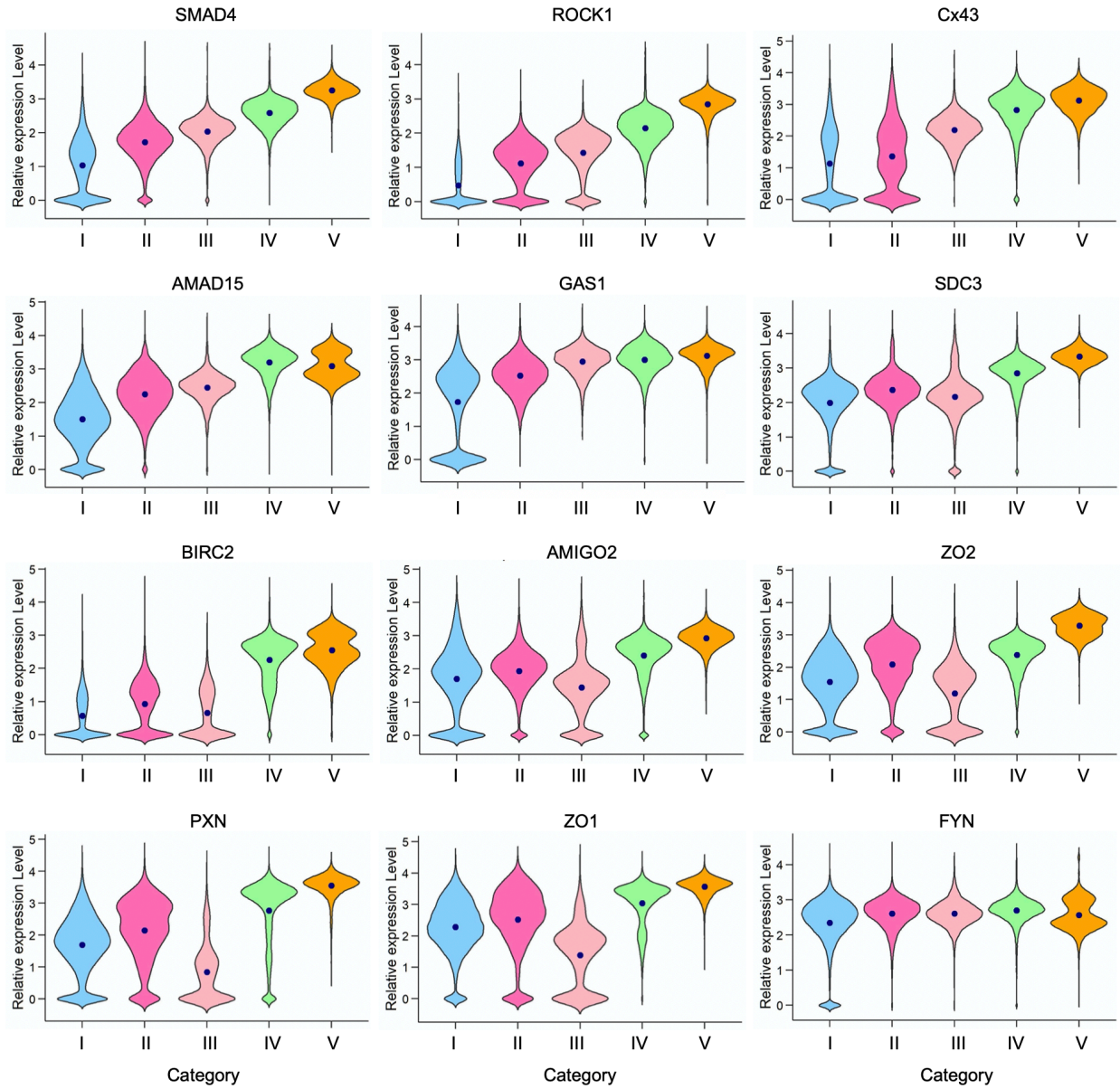


Figure S7: Violin plots showing expression patterns of SMAD4 and 11 proteins associated with junction and adhesion complex (JAC) within the five categories. Related to Figure 7.

Table S4: Primer sequences for ChIP-qPCR. Related to STAR Methods.

Gene	Sequence 5'	Sequence 3'
<i>SMAD4 #1</i>	GTGGCGCCTGTGATTTGCTTCT	CCTTGCCATGTGGTGCTCTCC
<i>SMAD4 #2</i>	CTTTGTTCCAGCCTCACTT	TAGTTACCTATTGCTGTATAACAAT CTC
<i>AMIGO2 #1</i>	CCAACTCTTCCCTCCCCT	TAACAAATCTGCCTGACCACA
<i>AMIGO2 #2</i>	TAATAATAATCCCAGAGCAGACCT	GGAAATGAGAGAGAGAGAACG
<i>SDC3</i>	TGCCAAAACAAGGGAGTC	GTGTGTCATTTCAAGGGTCTA
<i>PXN</i>	CACAGCCACCCTCCCTA	TGTTATTATTACCCAATTTCCGAGT
<i>ADAM15</i>	ATGACACACAATAGGGGC	TTCACATTCAGTTTGCTTCACGA
<i>BIRC2</i>	TGGAGAGGGCACAAACC	CAACTGGACGCTCATCG
<i>FYN #1</i>	GGTCTGATTTAGGGGGTTT	CTCAAGGTGCTGTCTCCA
<i>FYN #2</i>	AGCAAAAGTCTGGAGGAAG	CGAGGTTTTTTGGGGGT
<i>GAS1 #1</i>	AGCTTCCCACGGCAGACCAA	GGAGGCCCAGCACTAGGAGATTC
<i>GAS1 #2</i>	CGTGCCCATTTATTCCGTCTTCT	GCGCCAGGCATCGTTTTCA
<i>GJA1</i>	CCAAGTAGAGGCGTTCA	GGCTGGTAAATGTGGAAGG
<i>ROCK1</i>	GAGAGGTCCAAAGTCCAG	GTTTCGTTTCATTCATTCATTGTGT
<i>TJP2</i>	TTCGACCCCCGCCTGATTC	CCGCAACCCGCCCTGAG
<i>TJP1</i>	GGTCTCGAACTCCTGGGCTCAT	CAGTCCTCGTTGCACGGTAAT

Table S5: Primer sequences for real-time qPCR. Related to STAR Methods.

Gene	Sequence 5'	Sequence 3'
<i>AMIGO2</i>	TGCTGGTCTTTTGGTATCGTA	GCACGAAAGGAACCATTGA
<i>BIRC2</i>	GGTTTTTATTATGTGGGTCGC	GCTGTTCAAGAAGATGAGGA
<i>SDC3</i>	GGTCACACTGCTCATCTATC	GAACTCCTCCTGCTTGTC
<i>PXN</i>	CACCCAACAGCAGACAC	TTGAAATCCGACAGCGAAG
<i>FYN</i>	CCCAACTACAACAACCTCCAC	GCAAGGTCCCCGTATGAGA
<i>TJP1</i>	ACGGACCAGTTTTCTCG	TGGGTAGGGCTGTTTGT
<i>GAS1</i>	CCGCTACCTGACCTACT	TGTTCTCCTTGACCGACT
<i>GJA1</i>	GGTGACTGGAGCGCCTTAG	GCGCACATGAGAGATTGGGA
<i>ROCK1</i>	ATGAAGATGAATAAGGAAGGCA	AAATACCCCAACCGACC
<i>TJP2</i>	GGGAAGGTCGCTGCTATTGT	CTCTCGCTGTAGCCACTCC
<i>ADAM15</i>	GAGGGACACACTTTGGAGAA	GACCACCAAGCCTCTGA
<i>SMAD4</i>	CTCATGTGATCTATGCCCCGTC	AGGTGATACAACCTCGTTCGTAGT