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Evolution of Placental Invasion and Cancer Metastasis are Causally Linked

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Data and Materials Availability: data for the human-cow transcriptome comparison with and without co-culture with trophoblast cells is available under GSE136299 in the Gene Expression Omnibus (GEO) database of NCBI, <https://www.ncbi.nlm.nih.gov/geo/>. The comparative fibroblast gene expression data is available under PRJNA564062 under SUB6229748 and SUB6264591 on the Sequence Read Archive (SRA) of NCBI, <https://www.ncbi.nlm.nih.gov/sra>.

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Abstract

Among mammals, placental invasion is correlated with vulnerability to malignancy. Animals with more invasive placentation (e.g. humans) are more vulnerable to malignancy. To explain this correlation, we propose the hypothesis of *Evolved Levels of Invasibility*: the evolution of invasibility of stromal tissue affects both, placental and cancer invasion. We provide evidence for this hypothesis using an *in vitro* model. We find that bovine endometrial and skin fibroblasts are more resistant to invasion than their human counterparts. Gene expression profiling identified genes with high expression in human but not in bovine fibroblasts. Knocking down a subset of them in human fibroblasts leads to stronger resistance to cancer cell invasion. Identifying the evolutionary determinants of stromal invasibility can provide significant insights to develop rational anti-metastatic therapeutics.

Placental invasion into the maternal endometrium of the uterus exhibits substantial similarities to early cancer dissemination into stroma¹⁻⁴. These similarities have inspired the hypothesis of antagonistic pleiotropy^{5,6}. According to this hypothesis, trophoblasts evolved the capacity to invade into the endometrium leading to invasive placentation. These mechanisms can become reactivated in cancer cells, leading to a predisposition to metastasis. This hypothesis implies that cancer malignancy should be limited to placental mammals, where invasive placentation first evolved. This prediction, however, is inconsistent with the fact that opossums, with ancestrally non-invasive placenta^{7,8}, get invasive skin cancers⁹. Here we explore an alternative scenario in which stromal cells of the uterus evolved to either resist or permit invasion, determining the outcome of placental invasiveness⁹.

The likelihood that the evolution of the stromal environment is driving the evolution of cancer malignancy is enhanced by the fact that the molecular mechanisms employed by cancer cells to metastasize are shared with other biological processes. For instance, the mechanisms regulating gastrulation, wound healing, extravasation by leukocytes etc., are shared with both trophoblast and cancer invasion^{6,10,11}. This implies that invading cancer cells employ mechanisms that evolved much earlier than placental invasion, and therefore, the evolution of invasive placentation *per se* cannot be responsible for the origin of malignant cancer. It is important to note, however, that the invasiveness of the placenta continued to evolve after its origin. Placental invasion reverted to a non-invasive phenotype in several lineages of placental mammals, as well as evolved into an even higher degree of invasiveness in the great apes, which includes humans¹²⁻¹⁴. A complete loss of placental invasion evolved in the hoofed mammals, such as cows and horses and their relatives, and it is exactly in those animals that have lower malignancy rates for a variety of cancers⁹. In contrast, humans with very invasive placentas are highly vulnerable to melanoma malignancy.

Based on the arguments outlined above, we advance the hypothesis that evolutionary changes in the permissiveness, or resistance of uterine stromal cells to placental invasion are mechanistically linked to the vulnerability to cancer malignancy⁹. We term this hypothesis *Evolved Levels of Invasibility* (ELI) (Figure 1A), and experimentally test the ELI hypothesis using an *in vitro* model (Figure 1B). We demonstrate that human fibroblasts are more permissive to invasion by trophoblasts as well as cancer cells compared to their bovine counterparts. We then identify factors responsible for the resistance to invasion paving a way for therapeutic interventions. This study highlights how investigating evolutionarily processes may lead to the identification of therapeutic targets, pointing to the clinical potential of evolutionary analysis.

Results

Collective cell invasion behavior can be modelled in an ECM-mimetic co-culture system

To test the ELI hypothesis, we developed a method for measuring collective cell invasion (CCI) into a lawn of stromal cells. The CCI into stroma is a complex process^{15,16,17}. During infiltration and spread, invading cells interact with stroma, which modulates their invasive capacity, while experiencing contact guidance from the underlying Extracellular Matrix (ECM) fibers^{1,18,19}. Based on these considerations, we sought to mimic the stromal invasion process through a patterned co-culture of stromal and invasive cells on structured adhesion substrata. This approach was validated in previous cancer-stroma interaction studies^{20,21}. The extent of invasion was measured as the average distance over which invasive cells have penetrated the stromal monolayer. The platform also enabled the sensitive measurement of cell invasion (Figure 1C).

We first compared the spread of malignant (1205Lu) and non-malignant (WM35) melanoma cells into a monolayer of human skin fibroblasts (BJ5ta) on substrata reproducing the nano-topographic features of aligned ECM fibers, or on flat surfaces. Aligned matrix fibers are a common feature in many tissues²². We found that mimicking aligned ECM matrix microenvironment allowed malignant 1205Lu melanoma cells to more extensively invade into skin fibroblast monolayer than non-malignant WM35 cells (Figure 1D–E, S1A). Furthermore, when 1205Lu and WM35 cells were compared on a flat surface, no statistically significant difference could be detected. In contrast, a significant difference between 1205Lu and WM35 was detectable in the ECM-mimetic platform (Figure 1E, S1B, C), indicating that this assay has higher sensitivity in measuring stroma invasion compared to experiments on a flat surface.

Bovine endometrial stroma is resistant to trophoblast invasion

To explore the causes of species differences in placental invasion, we measured the invasion of a human choriocarcinoma (J3) cell line into human endometrial stromal fibroblasts (hESFs), and compared them to bovine trophoblasts (F3) invading into bovine endometrial stromal fibroblasts (bESFs). At 48 hours J3 cells invaded deeply into hESF layer, while the F3 invasion into bESFs was much more limited (Figure 1F–G, Supplementary Movies). Upon closer inspection we found that human J3 cells formed invasive forks, which propelled

invasion into hESFs, while the bovine F3 showed no invasive forks (Figure 1F, G, H). These results recapitulate the *in vivo* species differences during embryo implantation^{23–26}.

In human placenta, extravillous trophoblasts (EVTs) constitute the most invasive cell type^{27,28}. We therefore measured the invasion of human chorioblastoma cell line BeWo, the EVT cell line HTR8 and the bovine F3 cells invading into the layer of human or bovine stromal fibroblasts in all combinations (Figure 2A). BeWo cells, though being a carcinoma cell line that have been extensively used as model of human cyto-trophoblast cells²⁹. The results showed that while hESF were invaded by all trophoblast cell lines, both human and bovine, bESF were more resistant to invasion (Figure 2B). The largest differences were found in the case of F3 invasion; with F3 invading rapidly into hESF monolayers, but are nearly completely halted by bESFs. This cross-species invasion experiments confirmed that the limited invasion in bESF is, to a large extent, a property of the stroma cells rather than the trophoblast cells (Figure 2B). In summary, the degree of trophoblast invasion is mostly controlled by the identity of the stromal cells rather than the invasive capacity of trophoblast cells, as assumed in the ELI hypothesis.

Bovine skin fibroblasts resist melanoma invasion

The findings above leave open the possibility that the observed effects could be specific to the fetal-maternal interface rather than reflecting species differences in stromal invasibility in general. In humans, melanoma invasion into the surrounding stroma is a strong predictor of malignancy³⁰. Would stromal compartments in other tissues than the uterus, e.g., the skin, show similar species-dependent properties as the uterine stromal cells? To address this question, we investigated whether gene expression profiles of human and bovine skin fibroblasts are more similar to those of their corresponding endometrial fibroblasts in the same species than they are to their homologous cell type in the other species.

The null hypothesis of independent transcriptome evolution of cell types predicts that corresponding (homologous) cell types should be more similar to each other than they are to a different cell type in the same species³¹. This is because, for instance, the human and bovine lineages diverged more recently than skin and endometrial fibroblasts have differentiated in evolution. This is necessarily true, because otherwise corresponding cell types would not be homologous³². On the other hand, if gene expression profiles in skin and endometrial fibroblasts co-evolved, the cells from the same species could be more similar than to the corresponding/homologous cell in the other species. The ELI hypothesis predicts that, in each species, endometrial fibroblasts and skin fibroblasts share the same level of invasion resistance because their gene expression profiles co-evolved, such that selection for higher or lower endometrial invasiveness has a parallel effect on the invasibility of skin fibroblasts. Therefore, we expect that the stromal cell types from the same species are more similar to each other than they are to corresponding cell types in the other species³³.

To evaluate this expectation, we performed RNA sequencing on bovine and human skin fibroblasts, and compared them to human and bovine endometrial fibroblasts. Principal component analysis showed that the gene expression distance between skin fibroblasts and ESFs of the same species was less than the distance between corresponding cell types of different species (Figure 2C). The observed pattern is consistent with the hypothesis that

gene expression in endometrial and skin fibroblasts indeed co-evolved due to pleiotropic effects of mutations on transcriptional regulation^{31,34}. In other words, the biology of skin and endometrial fibroblasts can be expected to be similar within species even though they differ between species.

We then measured invasion of malignant human melanoma cells (A375) into the skin fibroblasts from human (BJ5ta), and bovine (bSkFb), and found that bSkFb indeed resisted A375 invasion more strongly than human skin fibroblasts (Figure 2D). This greater invasion resistance of bSkFb was supported by a similarly high resistance of these cells to invasion by WM35, 1205Lu and SKMe128 melanoma cell lines (Figure 2E). A similar assay with bovine melanoma cells was precluded owing to lack of bovine melanoma cell lines³⁵. Overall, these results indicate that bovine skin fibroblasts can resist cell invasion better than their human counterparts.

Human and bovine fibroblasts respond differently to trophoblast co-culture

To understand the genetic underpinnings of ELI in bovine stromal cells, we collected RNA sequencing data from human and bovine endometrial fibroblasts, with and without co-culture with the respective trophoblasts. We focused on differentially expressed genes in stromal cells, both basally and upon interaction with species-specific trophoblasts. Endometrial fibroblasts of both species were labeled with the DiI dye, and co-cultured with equal number of unlabeled trophoblast cells, HTR8 and F3 for human and bovine cells respectively (Figure S3A). Co-cultures were maintained for 72 hours, and the cells were sorted using fluorescence activated cell sorting (FACS)²⁰. Henceforth, the bovine cells will be referred to as bESFs and bESF^{co-F3} and human ESFs, hESFs and hESF^{co-HTR8}.

A large number of genes were differentially expressed in bovine vs. human endometrial fibroblasts (Figure S3B), and ESFs responded strongly to co-culture with their cognate trophoblasts cells (Figure S3C–D). Furthermore, human and bovine ESFs responded differently to trophoblast co-culture (Figures 3A–B, Figures S3E–F). Human and bovine endometrial stromal cells differed substantially with respect to the number of genes affected by co-culture, with human cells displaying a larger number of differentially expressed genes (5,349 genes changed at $p < 0.01$, $FDR=3.1 \times 10^{-3}$) compared to bovine cells (3,101 genes at $p < 0.01$, $FDR=7.01 \times 10^{-3}$). Gene set analysis revealed genes belonging to chemotactic activity, cell motility, and metastasis ontologies at higher relative abundance in hESF^{co-H8} compared to bESF^{co-F3} (Figure 3B). These results suggest that species differences in invasibility may be caused by differential gene expression among the stromal cells in response to trophoblast cells.

We found that human and bovine ESFs expressed markedly different sets of chemokine ligands (Figure 3C) and chemokine receptors (Figure 3D), although both were enriched in transcripts associated with angiogenesis. Genes showing elevated expression in hESFs included Fibroblast Growth Factors (FGFs), Vascular Endothelial Growth Factors (VEGFs), Semaphorins, members of the Transforming Growth Factor (TGF) family, as well as NRPs and *ROBO1*, whereas bESFs showed high expression of Endothelin (*END1*), Plasminogen activator (*PLAU*), Thrombopoeitin (*THPO*) and notably, Transforming Growth Factor beta 2 (*TGFB2*). In contrast, genes belonging to ontologies GO_Adherens Junction and Endothelial

Barrier did not show systematic expression differences in favor of either hESF or bESF (Figure S4A–B). Finally, the genes related to GO_Fibroblast Migration tend to be expressed at lower levels in bESF vs. hESF (Figure S4C). These data point to the possibility that stromal response to trophoblast co-culture varies strongly between the species for genes that could play a role in the interaction between invading trophoblast cells and stromal cells, as opposed to genes which regulate basal stromal integrity. Differential expression of these genes may affect the invasibility phenotype in either species.

These analyses suggested that differences in the interactions between the stromal and invading cells were drivers of ELI in human and bovine. For instance, transforming Growth Factor (TGFB), and WNT signaling pathways are mediators of cancer stromal interaction^{36–41} and thus we explored the effect of trophoblast co-culture on the activation of these signaling pathways in order to see whether there are parallels between cancer-stroma and trophoblast-endometrial stroma interactions. Expression of genes in non-canonical WNT and TGFB pathways were indeed higher in hESFs compared with bESFs (Figure 3E, F). Ingenuity pathway analysis showed that both signaling pathways were differentially activated in bovine and human ESFs (Figure S4E). Although co-culture with trophoblasts resulted in reduction in expression of genes belonging to TGFB pathway for hESFs, they continued to remain many-fold higher when compared in bESFs. While TGF ligands TGFB1, and TGFB3, as well as receptors of the TGFB family, e.g. ACVR1 were highly expressed in hESFs, negative regulators like TGF β 2 and inhibin-A were expressed at higher levels in bESFs. Similarly, we observed that downstream transcription factors Smad 2, –3 and –5, as well as the co-SMAD, SMAD4, were expressed at higher levels in hESFs compared to bESFs (Figure 3E). Similarly, many members of the WNT signaling network displayed higher expression in hESF vs bESF (Figure 3F, S4D). These results supported a role for paracrine signaling between heterotypic cells during the invasion processes in both, the uterus and in tumor lesion, prompting us to further explore the role of the signaling networks in stromal invasion.

Selected gene knockdown in human fibroblasts increases resistance to melanoma invasion

We hypothesized that modulating the differentially expressed genes in human stromal cells could induce them to become more resistant to invasion in human cells. We focused on genes related to WNT and TGFB signaling. Specifically, we selected 16 genes from the TGFB and WNT pathways that had higher expression in hESF or were upregulated in hESF in co-culture with trophoblast cells (Figure 4A). We then targeted these genes using a battery of siRNAs in hESFs and BJ5ta to test whether they could modulate stromal invasibility.

Comparison of each siRNA-transfected stromal cell population was made with appropriate untransfected controls, as well as with cells transfected with control non-targeting siRNA (Figure S5A–B). HTR8 invasion into hESF was significantly reduced after 24 hours in eight out of 16 genes (Figure 4B). A375 invasion into BJ5ta skin fibroblast monolayer was observed for 18 hours and 36 hours after gene knockdown. Nine of these gene knockdowns significantly decreasing invasion into the skin fibroblast monolayer (Figure 4C), including members of WNT superfamily, TGFB ligands, as wells as less established targets and

effectors of WNT signaling, e.g. STARD7⁴², LPIN1^{43–47}, and YAP1⁴⁸. Furthermore, we detected a weak but significant correlation in the increase of stromal resistance to invasion following gene knockdowns in both hESF and BJ5ta ($r^2=0.25$, $p = 0.02$), indicating that gene silencing enhances resistance to invasion in a similar manner in both human endometrial and skin fibroblasts (Figure 4D). Moreover, the average response for all gene knockdowns was an increase in resistance to invasion (Figure 4E). These results suggested that human skin fibroblasts could be induced to resist melanoma invasion and that knowledge of gene expression in cow fibroblasts can tell us how to achieve this.

Opposite evolutionary trends of vulnerability to malignancy in humans and bovines

In order to assess the evolutionary history of the expression of the genes identified above as important for ELI, we cultured the endometrial stromal fibroblasts from rabbit, rat, guinea pig, cat, horse and sheep to extend our analysis (Figure 4F). This taxon sample represents two clades of eutherian mammals, the Euarchontoglires, and the Laurasiatheria, together forming the clade of Boreoeutheria. We plotted the phylogenetic tree of these eight species⁴⁹, along with the expression of these genes in skin fibroblasts compared to the inferred boreoeutherian ancestor ($TPM_{species} - TPM_{ancestor}$).

We found, surprisingly, that humans evolved higher expression levels of invasibility enhancing genes (*TGFB1*, *ACVR1*, *DDR2*, *LPIN1*, *CD44*, *MMP1*), compared to the inferred boreoeutherian ancestor. The expression levels of these genes in human skin fibroblasts is higher than that of rabbit, rat and guinea pig. All of these species have hemochorial placentation and thus the higher expression level of these genes in humans likely evolved in the primate lineage. The limited data on a selected group of genes, therefore, supports the hypothesis of Evolved Levels of Invasibility (ELI) amongst mammals, and points to the possibility of increased stromal vulnerability in humans, and decreased stromal vulnerability in bovines.

Discussion

Mammalian species differ in their potential for tumorigenesis, as well as their vulnerability to cancer metastasis^{9,50}. The comparative biology of cancer incidence across different animals has identified species specific tumor suppressor mechanisms explaining the variation in occurrence of tumors between species^{51,5253,5455,56}. In a recent review, Constanzo and collaborators made a convincing case for a model where cancer progression in humans includes the re-activation of embryonic gene expression normally controlling placenta development and immune evasion⁵⁷. Here, we focus on differences across species of the rates of cancer malignancy rather than differences in rates of tumorigenesis. For instance, melanoma does occur in bovines and equines, but remains largely benign^{58–60}, while it is highly malignant in humans. This correlates with the phenotype of the fetal-maternal interface, i.e. the degree of placental invasion during pregnancy.

siRNA-guided knockdown in human stromal fibroblasts revealed genes that impart stromal invasibility by melanoma and trophoblast cells (Figure 4G). These genes include TGFB ligands, consistent with results from colorectal cancer³⁹. Other genes enhancing stromal invasibility are members of the non-canonical WNT pathway, as well as WNT signaling

modulators, e.g. *LPIN1*⁶¹, by directly regulating β -Catenin levels, e.g. *YAP/TAZ*⁴⁸, or by being regulated by β -catenin induced transcription, e.g. *CD44*⁶². Of these, YAP1, a Hippo pathway target promoting tumor growth⁶³, can also be regulated by non-canonical WNT signaling, and can in turn inhibit canonical WNT signaling⁴⁸. Both TGFB^{64,65} and non-canonical WNT signaling^{66,67} are known to affect tumor progression and metastasis. TGFB1 and TGFB3, expressed at relatively higher levels in hESF, are also reported to regulate β -catenin activity, indicating that human stromal vulnerability to invasion may be influenced by paracrine signaling to invading cells through secreted ligands. Also of interest, many of the genes identified in our comparative gene expression screen have connections to metabolic regulation. These include genes encoding LIPIN1 which regulates triglyceride metabolism,⁶⁸ and MGAT5, known to regulate glucose uptake in tumor cells⁶⁹.

The results presented here show that species differences in malignancy rates may, in part, be caused by species differences in invasibility of stromal cells. We found that bovine stroma evolved lower invasibility likely via decreased paracrine signaling through TGFB and WNT pathways. In particular, these results argue that TGFB secretion, and high non-canonical WNT signaling in stromal cells are causal factors explaining the high vulnerability of human stromal tissues to cancer invasion, at least in the case of melanoma. Comparative transcriptomic data across multiple additional species further suggests that the human lineage evolved higher expression of these genes enhancing tumor and trophoblast invasion and thus likely have evolved higher malignancy rates than in the common ancestor of eutherian mammals.

Our data support the ELI hypothesis, suggesting that differences in stromal gene expression between species are causal in determining both, the degree of embryo implantation as well as stromal resistance to early cancer dissemination. Epitheliochorial placentation has evolved multiple times from the pre-existing invasive placentation, suggesting evolutionary advantages of non-invasive placentation^{13,14,70} may drive corresponding differences in malignancy resistance. We further found that apes may also have evolved increased stromal receptivity to trophoblast invasion, and correlatively have evolved higher vulnerability to cancer malignancy. As extravillous trophoblasts have evolved recently in great apes, the endometrial stroma may also have increased invasibility to accommodate the more invasive trophoblast types⁷¹. Our ELI hypothesis, and its experimental validation, suggests that studying the genetic basis for evolved resistance to invasion can identify factors determining the stromal control of cancer progression and identify novel therapeutic targets.

Materials and Methods

Cell Sourcing

The study used a variety of cell sources for comparative analyses. Human ESFs were immortalized by the Charles Lockwood lab at Yale, and obtained from Gil Mor group⁷². F3 cells were previously established by Pfarrer group at the University of Veterinary Medicine, Hannover, Germany^{73,74}. J3, HTR8/SVNeo, BeWo, and BJ5ta cells were obtained from ATCC.

bSkFb bovine (*Bos Taurus*) skin fibroblasts, as well as those from guinea pig (*Cavia porcellus*), sheep (*Ovis aries*), pig (*Sus scrofa*), and rabbit (*Oryctolagus cuniculus*) were isolated from the skin tissue of the face of a cow from Bristol Butchers. A small piece of skin was harvested, hair removed, and the sample was washed in PBS and cut into strips approximately 1.0 cm². Dermis was separated from epidermis by enzymatic digestion (30 minutes in 0.25% trypsin at 37°C, followed by Dissociation Buffer (1 mg/ml collagenase, 1 mg/ml Dispase, 400 µg/ml DNaseI) for 45 minutes at 37°C). Epidermis was removed, and 2 mm pieces were cut from the dermis, and transferred to a 12-well plate and covered with media. Fibroblasts emerged from the explants, and grew to confluency in growth media. Extra tissue was removed, and cells subcultured for a few times using a cell scraper.

Bovine ESFs were isolated as follows: Uterine tissues were harvested from each species and primary ESF were isolated by enzymatic digestion. Uterus fragments, 2–3 mm in size, were created using scalpel, and digested with 0.25% trypsin-EDTA for 35 min at 37°C, followed by Dissociation Buffer (1 mg/ml collagenase, 1 mg/ml Dispase, 400 µg/ml DNaseI) for 45 minutes at 37°C. Cell clumps were homogenized by passage through a 22-gauge syringe followed by passage through a 40 µm nylon mesh filter to remove remaining clumps. A single cell suspension was obtained from the lysate, transferred to fresh growth medium and cultured in T25 flasks. To facilitate enrichment of fibroblasts versus epithelial cells, media was exchanged in each well after fifteen minutes in order to remove floating cells that had not yet attached while stromal fibroblasts had attached. Cells were grown to confluency and sub-passaged by scraping the cells off the surface to be split into two T25 flasks. At confluency cells were split through one more round of differential attachment. We used immunohistochemistry to test for abundance of vimentin (Santa Cruz, sc-6260) and cytokeratin (Abcam, ab9377) to validate fibroblast subtype in the isolated cells.

hESF, bSkFb and bESFs were tested for potential mycoplasma contamination, and were found to be free from it.

Cell culture

Human ESFs were grown in phenol-red free DMEM/F12 with high glucose (25 mM), supplemented with 10% charcoal stripped calf serum (Hyclone), and 1% antibiotic/antimycotic (Gibco). Decidualization was induced in ESFs with 0.5 mM 8-Br-cAMP (Sigma), and 0.5 mM of progesterone analog medroxy-progesterone acetate (MPA) for 96 hours in DMEM supplemented with 2% charcoal-stripped calf serum (Hyclone). BJ5ta (ATCC) cells were cultured in 80% DMEM and 20% MEM supplemented with 10% FBS, 1% antibiotic/antimycotic, and 0.01 mg/ml hygromycin. F3 cells were obtained from Dr. Pfarrer's group, and were cultured in DMEM with high glucose, supplemented with 10% FBS, 1% antibiotic/antimycotic (Gibco). J3 cells were cultured in αMEM supplemented with 10% FBS, 1% antibiotic/antimycotic (Gibco), while HTR8/SVNeo (ATCC) were cultured in RPMI-1640 supplemented with 10% FBS, and 1% antibiotic/antimycotic (Gibco).

Fabrication of poly-urethane acrylate mold

Photoresist was spin coated on silicon wafers, and electron-beam lithography was used to nanopattern the wafers (JBX-9300FS, JEOL). After the photoresist was developed, exposed silicon was etched by deep-reactive ion etcher (STS ICP Etcher) resulting in formation of submicron parallel ridges. Residual photoresist was removed using ashing, and diced into silica masters for subsequent replica molding.

UV curable poly-urethane (PUA) was drop-wise dispensed onto the silicon master previously prepared, and contacted with polyethylene terephthalate (PET) film. Application of UV ($\lambda = 200\text{--}400\text{ nm}$, 100 mJ/cm^2) for 1 minute was used to cure PUA, after which the mold was peeled off with a pair of tweezers and cured overnight under UV to terminate the residual acrylate groups. The process resulted in PUA mold of $\sim 50\text{ }\mu\text{m}$ thickness.

Fabrication of nanotextured substrate

Previously prepared PET mold was used as a replica mold to transfer the topographic pattern on glass substrate using the technique of capillary force lithography. Briefly, glass substrate was cleaned using NaOH (0.1 M for 1 hour), washed with DIH_2O , and dried under nitrogen stream. Primer (phosphoric acrylate and propylene glycol monomethyl ether acetate in a ratio of 1:10) was spin-coated on the coverslip as a thin layer, and baked for 20–30 minutes at 70°C . $150\text{ }\mu\text{l}$ of PEG-DA (molecular weight = 575) precursor was dispensed dropwise on the primed cover slip and the PET mold was placed reversibly. After the PUA precursor filled the submicron cavities by capillary action, the substrate was cured in UV ($\lambda = 250\text{--}400\text{ nm}$, 100 mJ/cm^2) for 1 minute. The mold was peeled after polymerization using a pair of tweezers. The substrate resultant from the method consisted of nanotextured grooves with an expected elastic modulus of $\sim 70\text{ MPa}$. The substrate was cured again for an hour under UV to terminate residual active acrylate groups.

Cell patterning for stromal invasion assay

For cell patterning, we used a stereolithographic plastic mold to create a polydimethylsiloxane (PDMS) stencil. PDMS stencil was cast by mixing monomer and cross-linker in a ratio of 10:1, and cured at 80°C for 4 hours by placing on a pre-designed plastic mold. Stencils were washed using isopropyl alcohol and dried using N_2 stream, placed on the nanotopographic substrate coated with laminin ($25\text{ }\mu\text{g/ml}$), and the device kept in vacuum to allow air under the stencil to be removed to avoid chance of leakage. DiI-labeled invasive cells were seeded on the stencil at a density of $10^6\text{ cells}/100\mu\text{l}$, and allowed to attach and polarize for 8 hours. Unattached cells were washed off twice with PBS, and the stencil removed carefully using a pair of blunt-end tweezers. Unlabeled stromal cells were seeded at a density of $10^6\text{ cells}/100\mu\text{l}$ to attach to the area previously covered by PDMS stencil. After 6 hours of attachment, unattached cells were washed off, and the substrate mounted for live cell microscopy. To ensure selection of definitive fibroblasts, and to avoid selection of trophoblasts taking up dye from dead stromal cells and occasional heterotypic cell fusion events, we only collected DiI^{high} cells. All experiments were conducted in similar culture conditions, by 1:1 mixture of the media in either comparable conditions.

Live Cell Fluorescence Microscopy

Phase-contrast and epifluorescence microscopy was performed using Leica DMi8 model, with an HC PL Fluotar 10x/0.30 Dry objective, and images were acquired using Leica LAS X Software, and processed using Fiji Image Analysis Software. Lumencor SpectraX was used as a source of light for excitation, and routed through a Rhodamine excitation filter cube consisting of Excitation 546 nm/ 10 nm (band-pass filter), Dichroic 560 nm (long-pass filter), and Emission 585 nm/ 40 nm (band-pass filter), and acquired using Andor EMCCD iXon Ultra 888 camera.

Invasion Analysis

Acquired sequential images for each condition were analyzed using Fiji Image Analysis Software after contrast enhancement. DiI labeled invasive cell fronts (trophoblasts or melanoma cells) were identified manually for each time point. Area occupied by DiI⁺ cells was measured at each time point. Total invasion was calculated as follows:

$$\delta\text{Area}(t) = \text{Area}(t) - \text{Area}(t_0)$$

Extent of invasion per unit measurement of interface was determined by dividing δArea by the length of initial intercellular interface

$$\langle \delta\text{Area}(t) \rangle = \frac{\delta\text{Area}(t)}{L(\text{interface})}$$

Fluorescence Assisted Flow Sorting

Cells were detached from the substrate using TrypLE solution (Gibco), quenched with excess medium, and washed thrice with phosphate-buffered saline (PBS). Isolated cells were suspended in 1% AlbuMAX (Gibco) dissolved in PBS and sorted using fluorescence assisted flow sorting (FACS). FACS was performed using BD FACSARIA II using PE.Cy5 channel, and analysis performed using FACSDiva 6.0. In order to increase the purity of EStFs collected after the co-culture experiments, and to account for possible uptake of dye from dead or dying stromal cells by the trophoblasts, as well as for occasional cell-fusion events, we only sorted DiI^{hi} cells. Sorted cells were collected directly in RNaselect to limit RNA degradation. Even cells that were not in co-culture were subjected to the same sorting protocol.

siRNA Transfection and Characterization

For gene silencing experiments, stromal cells (BJ5ta, or hESFs) were cultured in 12 well plates transfected with at least two siRNAs (shown in table below) at a concentration of 50 nmoles per well. Lipofactamine RNAiMAX was used to transfect siRNAs, obtained from IDT.

Gene Name	siRNA Duplex Name	Duplex Sequence 5'	Duplex Sequence 3'
LPIN1	Hs.Ri.LPIN1.13.1	GCGUCUACUUGGAUGACCUCACAGA	GCCGCAGAUGAACCUACUGGAGUGUCU
	Hs.Ri.LPIN1.13.2	AAGCAUUUUUUGUUCAAGAAACAGA	UCUUCGUAAAAACAAGUUCUUUGUCU
RAC2	Hs.Ri.RAC2.13.1	CCGUGUUUGACAACUAUUCAGCCAA	GUGGCACAAACUGUUGAUAGUCGGUU
	Hs.Ri.RAC2.13.2	GGACUUUUGCUAUUGCAAUAGAAA	GUCCUGAAAACGAUACGUUUUUCUUU
MGAT5	Hs.Ri.MGAT5.13.1	CUCUGACUUGUUGAGUAAUCAGUCA	GUGAGACUGAACCAUCUUAUAGUCAGU
	Hs.Ri.MGAT5.13.2	CAUGACAGCUUAUGAUCUGAAGAAA	CAGUACUGUCGAUACUAGACUUCUUU
WNT11	Hs.Ri.WNT11.13.1	AGCCAAUAAACUGAUGCGUCUACAC	GUUCGGUUAUUUGACUACGCAGAUGUG
	Hs.Ri.WNT11.13.2	AAGGAAACCGACAAGAUUAAAAATA	GUUUCCUUUGGCUGUUCUAAUUUUUAU
WNT5B	Hs.Ri.WNT5B.13.1	CGGUUUCUCUCUGACAUAUAAUGCC	AUGCCAAAGAGAGACUGUAAUUUACGG
	Hs.Ri.WNT5B.13.3	GAGGAAGCUGGCCAAUUGUACCAG	GUCUCCUUCGCACGGUUAACAUGGUC
STARD7	Hs.Ri.STARD7.13.2	GGUAUCUCACUGAGCUAAUCUGGAA	AGCCAUAGAGUGACUCGAUUAGACCUU
	Hs.Ri.STARD7.13.3	GUCCCCACAAGUCAUUUGAUGAGAA	GGCAGGGGUGUUCAGUAAACUACUCUU
YAP1	Hs.Ri.YAP1.13.1	GGUCAGAGAUACUUCUAAAUCACA	GACCAGUCUCUAUGAAGAAUUUAGUGU
TGFB1	Hs.Ri.TGFB1.13.1	CAGCAACAAUCCUGGCGAUACCTC	AUGUCGUUGUUAAGGACCGCUAUGGAG
	Hs.Ri.TGFB1.13.2	CUACUGUAGUAGAUCAUUUUATG	GUGAUGACAUCAAUUCUAGAUAAAUAAC
TGFB3	Hs.Ri.TGFB3.13.1	CAACUUAGGUCUAGAAUCAGCATT	AGGUUGAAUCCAGAUUUUAGUCGUAA
	Hs.Ri.TGFB3.13.2	AAGUAUGAAUUAUACUCUAAAATC	UGUUCAUACUUAUAAUGAGAGUUUAG
ACVR1	hs.Ri.ACVR1.13.1	GACUUUGACCAAAAUUGAUAAUCC	UUCUGAAACUGGUUUUAAACUUAUAAAGG
	hs.Ri.ACVR1.13.2	CGCUACGGGGAAAUGCAUUUUCTT	UUGCGAUGCCCCUUUACGUAAAAGAA
FOXO1	hs.Ri.FOXO1.13.1	GAUGUUUCUGACUAAUCUAAAUCA	CACUACAAGAGCUGAUUAGAAUUUAGU
	hs.Ri.FOXO1.13.2	GGAAUUCUCAACCUUCUCUCAUCA	AACCUAUUAGAGUUGGAAGAGAGUAGU
Twist1	hs.Ri.TWIST1.13.1	UUCUGAUAGAAGUCUGAACAGUUGT	GUAAGACUACUUCAGACUUGUCAACA
	hs.Ri.TWIST1.13.2	CUAUUUUAAAUGGUAACAAUCAGA	GAGAUAAAUUUUAACCAUUGUUAGUCU
NCOA3	hs.Ri.NCOA3.13.1	CUGCCAUCUUAGUGAUUUGACAA	UAGACGGUUAGAUAUCUUAACUGUU
	hs.Ri.NCOA3.13.3	CCAGUGGAAUUGGUGAUUCUGAATT	CAGGUCACCUUAACCACUAAGACUAAA
ERBB3	hs.Ri.ERBB3.13.1	GCCAUCUUCGUCUAGUUGAACUATA	AACGGUAGAAGCAGUACAACUUGAUAU
	hs.Ri.ERBB3.13.2	CACUGUACAAGCUCUACGAGAGGTG	CUGUGACAUUGUUCGAGUAGCUCUCCAC
DDR2	hs.Ri.DDR2.13.1	GUCAGUACACCAUCUGAAGUUTA	GACAGUCAUUGUGUUAGACUUCAAAU
	hs.Ri.DDR2.13.3	CCACUCCAUCUGGACAUUUAUGAA	ACGGUGAGGUAGACCUGUAAAUUACUU
BMP4	hs.Ri.BMP4.13.1	CAGUCCUUGAGGAUAGACAGAUATA	CCGUCAGGAACUCCUUAUCUGUCUUAU
	hs.Ri.BMP4.13.2	CCUUGUUUUCUGUCAAGACACCATG	ACGGAACAAAAGACAGUUCUGUGGUAC
MMP1	hs.Ri.MMP1.13.1	AGCAGACAUCAUGAUUCUUUUGTC	GUUCGUCUGUAGUACUUAAGAAAACAG
	hs.Ri.MMP1.13.3	GUCAACCUUGUUUCUACUGUUUUAT	UUCAGUUGGAACAAAGAUAGACAAAUA
CD44	hs.Ri.CD44.13.1	GCUCUGAGCAUCGGAUUUGAGACCT	UCGAGACUCGUAGCCUAAACUCUGGA

RNA isolation and qRT-PCR

RNA was isolated using RNeasy micro kit (QIAGEN, Germany) and resuspended in 10 μ L water. SuperScript IV VILO Master Mix (ThermoFisher, USA) was used to synthesize cDNA from 1 μ g mRNA and ezDNase was used to remove DNA after reverse transcription. Primers from Integrated DNA technologies (IDT, USA) were used for qRT-PCR reaction using SybrGreen (ThermoFisher, USA) to evaluate the efficiency of siRNA mediated knockdown. B2M and RPL0 were used as internal reference genes, and both were found to be consistent within samples after knockdown. qRT-PCR was performed in Quantstudio3 (ThermoFisher, USA) and relative quantitation was performed by comparative C_t .

RNA Sequencing

Agilent Bioanalyzer 2100 was used to determine RNA quality and RIN number of over 8 was observed in the samples. TruSeq RNA Library from illumina was used to prepare mRNA library. These libraries were sequenced on illumina HiSeq to generate 30–40million reads per sample (Single end 75 bp reads) and a high Q score was observed ($Q>30$) for the sequenced data. Alignment of reads and gene specific analysis was performed in Partek® Flow® software, version 5.0 Copyright ©; 2018 Partek Inc., St. Louis, MO, USA. Human cells data was aligned to reference genome (hg38) using STAR alignment (in Partek Flow) and bovine data was aligned using Bos Taurus assembly UMD_3.1.1/bosTau8. Quantification was performed using quantify to annotation model (Partek E/M) using Ensemble Transcripts release 85 (humans) or Ensemble-UMD3.1 gene annotation model. Normalization was performed using TPM (Transcripts Per Million⁷⁵). Gene level comparison and statistical analysis was performed using ANOVA and significance of (adjusted) $p=0.05$ or less was considered for analysis. Gene lists comparison was performed by obtaining Ontology/pathway specific list from Gene Ontology⁷⁶ or Kyoto Encyclopedia of Genes and Genomes(KEGG⁷⁷)

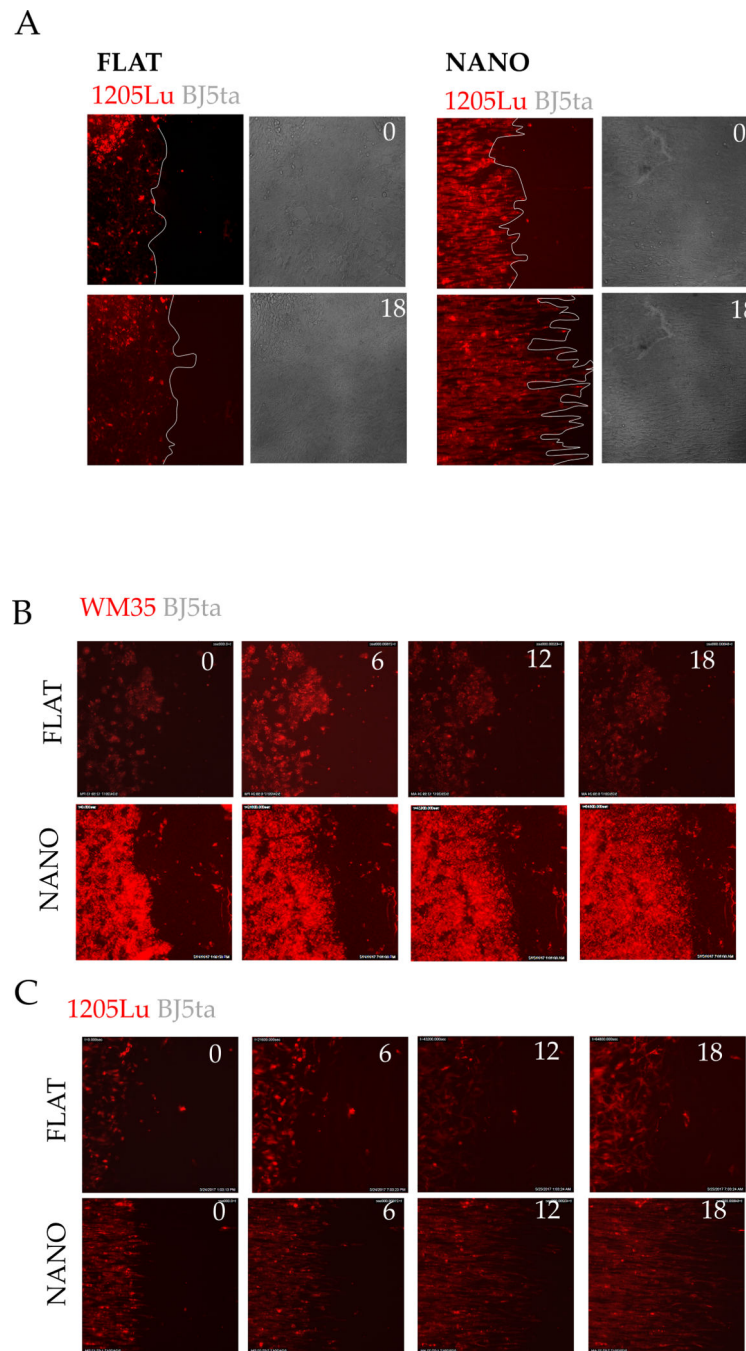
Comparative transcriptome analysis

Raw sequence reads (single-ended 75bps) for endometrial stromal fibroblasts from rabbit (*Oryctolagus cuniculus*), rat (*Rattus norvegicus*), guinea pig (*Cavia procellus*), cat (*Felis catus*), horse (*Equus caballus*), and sheep (*Ovis aries*) were aligned to reference genome assemblies OryCun2.0, Rnor_6.0, Cavpor3.0, Felis_catus_8.0, Equcab2, and Oar_v3.1 respectively using Tophat2 (v2.1.1). Read counts for all genes were calculated using HTSeq (v0.6.1p1, https://htseq.readthedocs.io/en/release_0.11.1/index.html) with Python (v2.7) according to Ensembl gene annotation release 92 (horse) or release 89 (other species). Then transcripts per million (TPM) were calculated to estimate relative mRNA abundance⁷⁵. To make gene expression levels comparable across species, we further normalized TPMs such that one-to-one orthologs have the same sum across species. The ortholog tables in this analysis were downloaded from Ensembl using BioMart tool. The ancestral state reconstruction for gene expression was performed on square root TPMs using function ‘ace’ from R package ‘APE’^{78,79}.

Two genes that have been found to influence invasibility in our knockdown experiments (Figure 4B and 4C), YAP1 and MMP1, have one-to-many or no orthologs in some species in our analysis. YAP1 has two orthologs in cow, ENSBTAG00000039307 (TPM=183.4) and

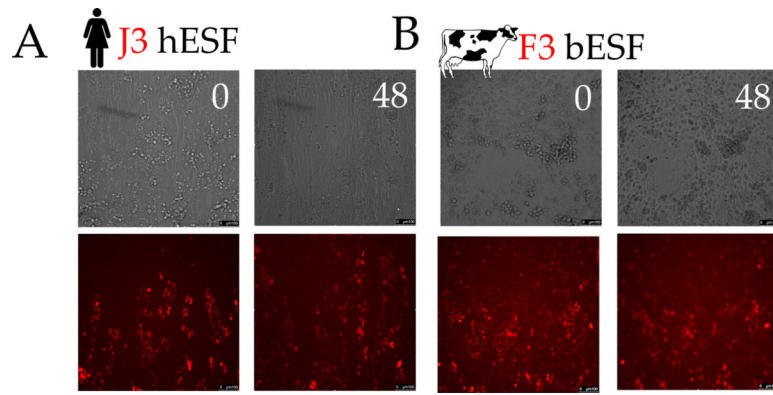
ENSBTAG00000047406 (TPM=96.3), and no ortholog in guinea pig. MMP1 has two orthologs in cow, ENSBTAG00000015818 (TPM=0.053) and ENSBTAG00000048029 (TPM=0.645), and three orthologs in rat, ENSRNOG00000055895 (TPM=0), ENSRNOG00000008881 (TPM= 0.157) and ENSRNOG00000032353 (TPM= 0.698). In these cases, the orthologs with the highest TPM were utilized to perform the ancestral state reconstruction.

Extended Data



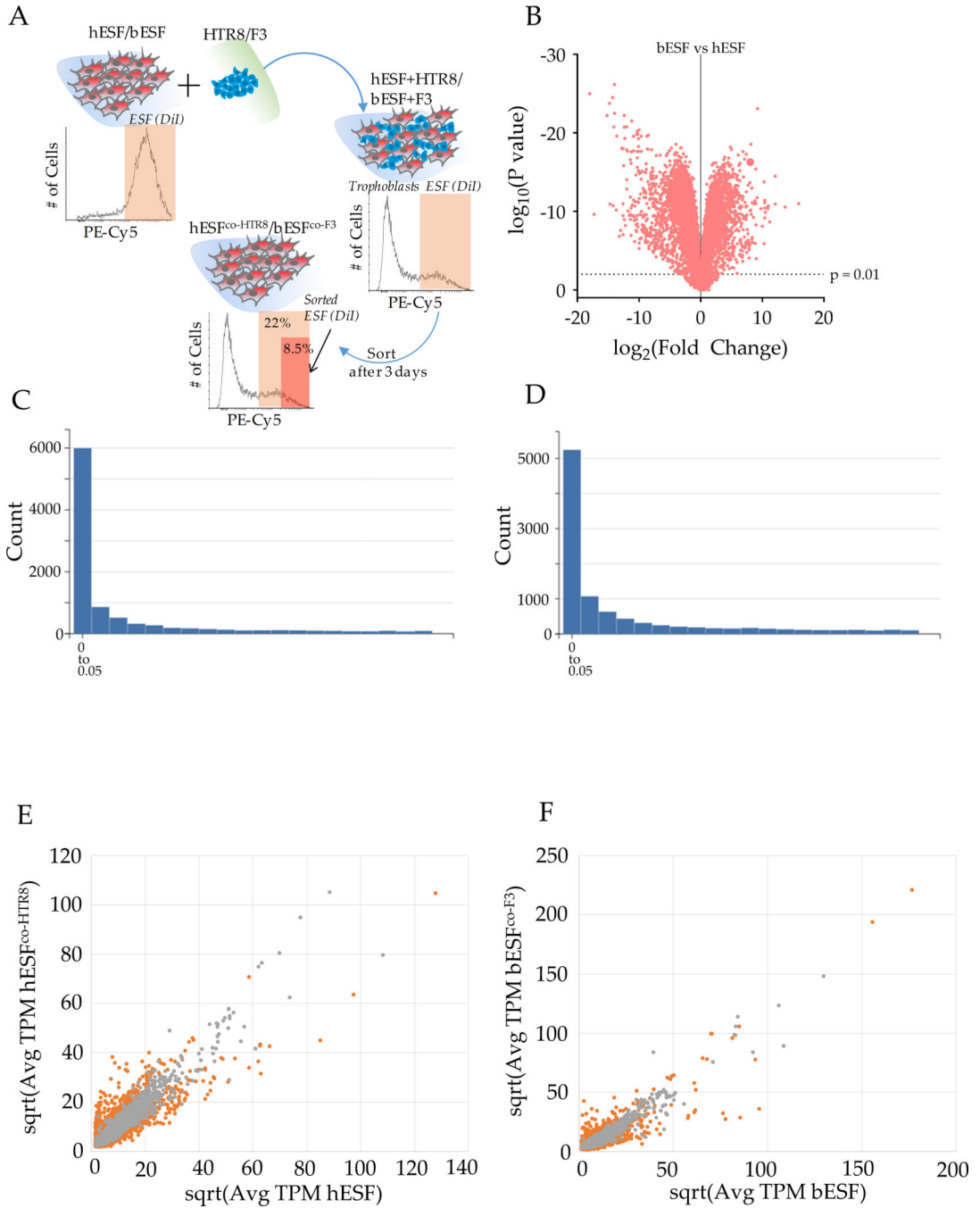
Extended Data Figure 1. Nanotextured stromal invasion assay quantitatively and sensitively measures collective intrusion of cells into stroma.

(A) Time stamped images at 0 and 18 hours showing extent of invasion by DiI labeled 1205Lu cells (red) into BJ5ta stromal fibroblast monolayer (unlabeled) on flat and nanotextured substrata; Quantification shown in Fig. 1D. (B-C) Sensitive measurement of differences in invasion by nanotextured stromal invasion platform; Time stamped images at 0, 6, 12, and 18 hours showing extent of invasion by DiI labeled non-malignant WM35 (B), and malignant 1205Lu (C) cells into BJ5ta monolayer shown in both flat, and nanotextured cell patterned assay; Quantification shown in Fig. 1E.



Extended Data Figure 2. Human and bovine trophoblasts exhibit different invasive potential into endometrial stroma.

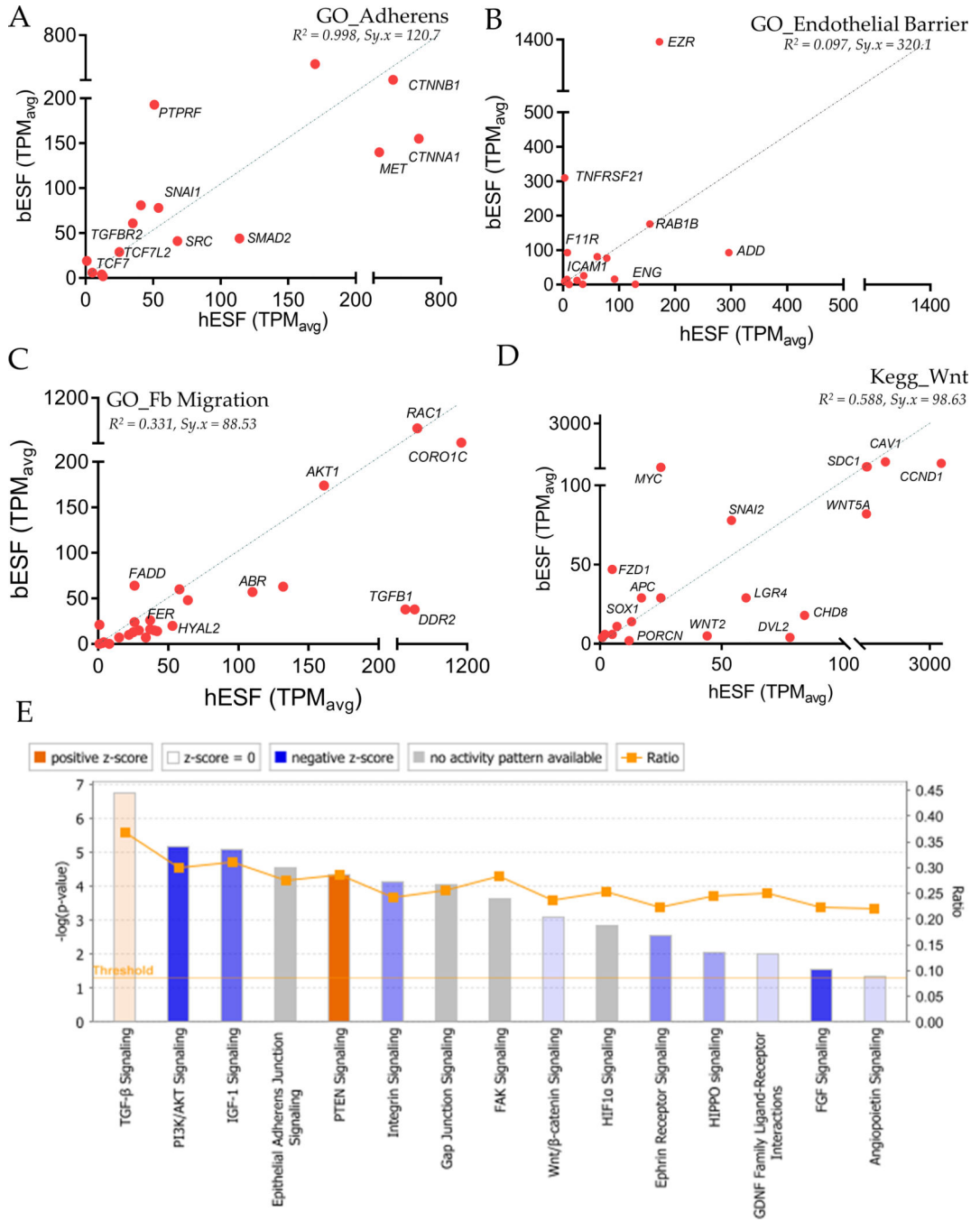
Time stamped images showing extent of invasion of human choriocarcinoma derived trophoblasts, J3 (red) (A), and bovine trophoblasts, F3 (red) (B) into their respective endometrial stromal fibroblast monolayers at 0, and 48 hours.



Extended Data Figure 3. Human and bovine endometrial stromal fibroblasts respond differently to co-culture with trophoblasts.

(A) Experimental plan to isolate endometrial cells after co-culture with respective trophoblast; (B) Volcano plot showing fold change in genes between bovine vs human endometrial stromal fibroblasts, along with their significance depicted in p value. (C) P-value distribution of t-tests comparing human ESFs with and without co-culture with HTR8 trophoblast cells. (D) P-value distribution of t-tests comparing bovine ESFs with and without co-culture with F3 trophoblast cells. In both C and D, note that thin right hand tail of the distribution, which indicates that a large number of genes are differentially expressed in

response to the presence of the corresponding trophoblast cells. (E-F) Scatter plots showing relative TPM values of genes between (E) hESF and hESF co-cultured with HTR8, and (F) bESF and bESF co-cultured with F3; Red dots refer to individual gene transcripts abundance significantly different in between the compared conditions.



Extended Data Figure 4. Human and bovine endometrial stromal fibroblasts do not appear to differentially respond to their respective trophoblasts by regulating cell-cell adhesion. Gene ontology analysis of individual genes belonging to GO_Adherens (A), GO_Endothelial Barrier (B), GO_Fibroblast Migration (C), and Kegg ontology for Wnt Signaling (D) for hESF and bESF shown with their relative TPM values; Also are shown the coefficient of determinant, R2 for the linear regression between TPM values of hESF and bESF in a given gene-sets, along with the standard deviation of the residuals, Sy.x. (E) Ingenuity Pathway Analysis of selected signaling pathways differentially activated in hESF

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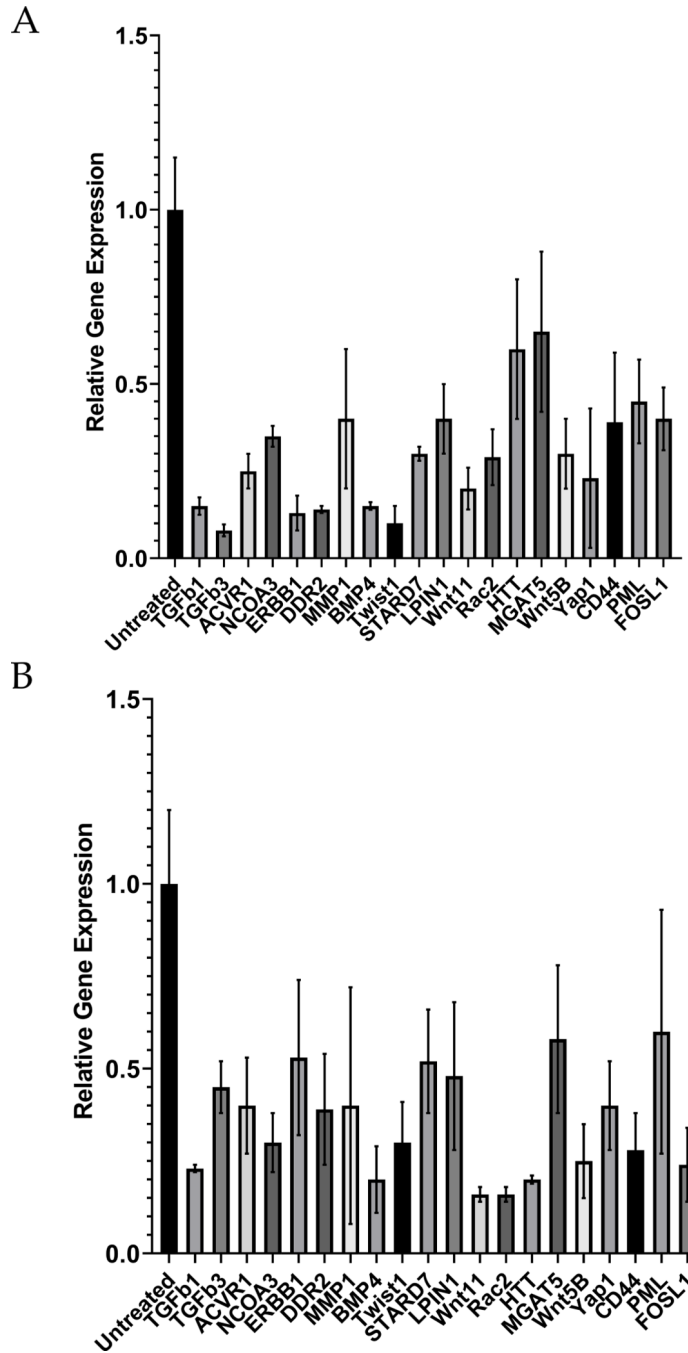
and bESF; Shown are $-\log(p\text{-values})$ of genes in the respective pathways differentially activated, while the color depicts the z-score.

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Extended Data Figure 5. siRNA induced gene knockdown reduces transcript levels in hESFs and BJ5ta cells.

qRTPCR results showing the percentage knockdown of transcript levels in hESF (A) and BJ5ta (B), calculated using DDCt method, compared using GAPDH as housekeeping gene with scrambled siRNA as control. For both A, and B, n = 4 biological replicates.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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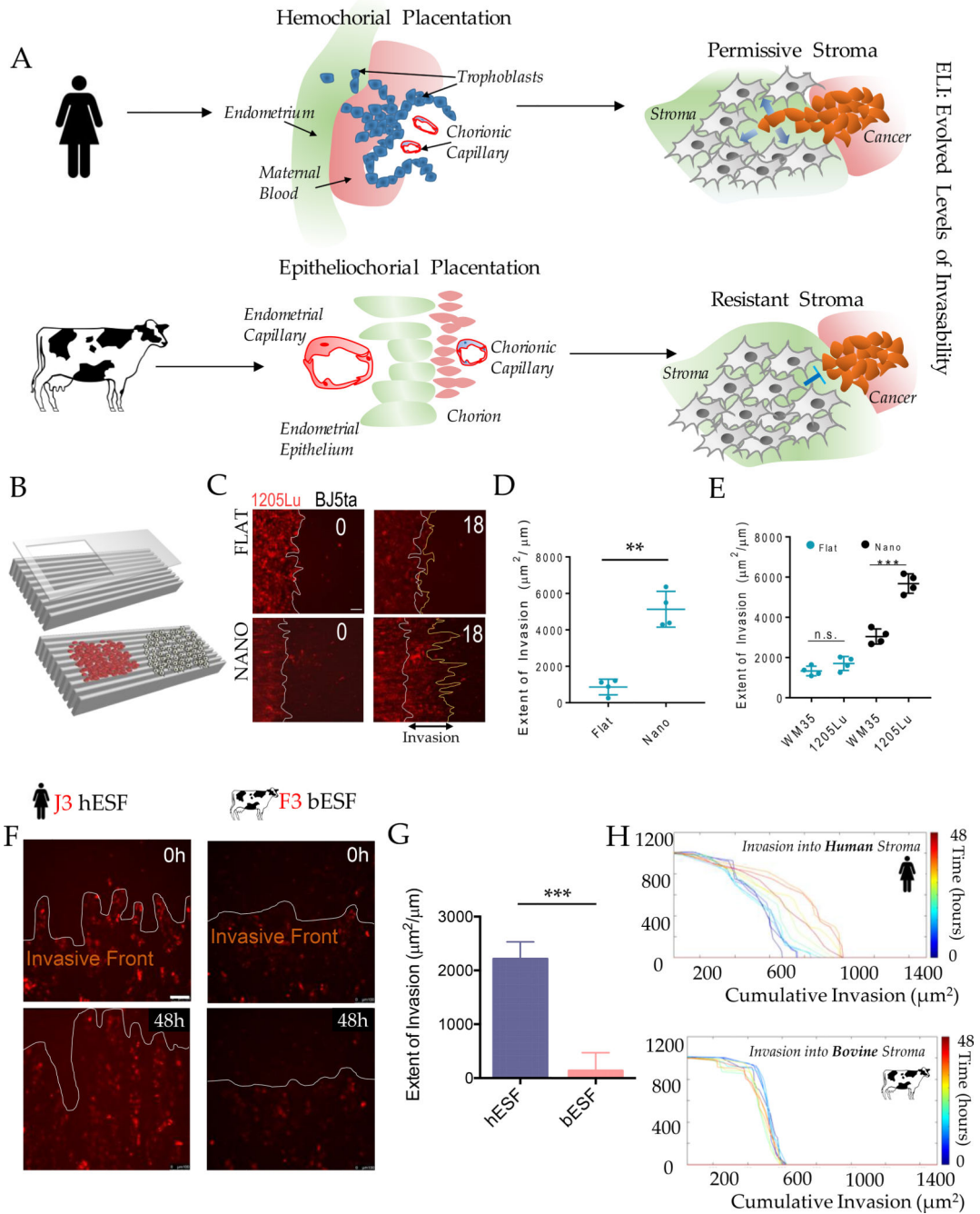


Figure 1. An experimental platform to test the hypothesis of Evolved Levels of Invasibility (ELI). (A) Schematic describing the ELI hypothesis. Placentation in humans is hemochorial, wherein the placental trophoblasts invade into the maternal stroma reaching the blood supply. In contrast, in cows and other boreoetherians, placentation has recently evolved to be epitheliochorial, where the trophoblast epithelium attaches to the endometrial epithelium, but does not invade the maternal interstitium. The ELI paradigm states that bovine stroma has evolved to resist invasion compared to human stroma, and therefore secondarily limits cancer metastasis. (B) Schematic showing a cell patterning nanotextured platform to

quantitatively and sensitively measure collective invasion into stroma; Stromal cells and invasive cells are patterned by a PDMS stencil into juxtaposed monolayers heterotypically interacting with each other, and imaged using live cell microscopy to observe collective cell invasion into the stroma. (C) Time course images showing invasion of 1205Lu malignant melanoma cells (red) into BJ5ta human skin fibroblasts (unlabeled) for 18 hours on a flat substrate vs a nanotextured substrate; Quantification of the extent of invasion per unit length of heterotypic intercellular interaction shown in (D). (E) Quantification of the extent of invasion of non-malignant WM35 and malignant 1205Lu melanoma into a monolayer of BJ5ta human skin fibroblasts on flat and nanotextured substrata. (F) Time course images showing invasion of human choriocarcinoma derived trophoblasts, J3 (red) into human endometrial stromal fibroblasts (unlabeled); and bovine trophoblasts, F3 (red) into bovine stromal fibroblasts (unlabeled) for 48 hours; See Figure S2 for phase-contrast and Supplementary Movies for dynamics of invasion; Quantification of the extent of invasion of trophoblasts into the respective stromal monolayer shown in (G). (H) Time course dynamic analysis showing cumulative invasion of J3 and F3 into respective species-specific endometrial stromal monolayers. In D, E, and G, $n = 4$ independent biological replicates; Statistical comparisons made using Student's t-tests **: $p < 0.01$, ***: $p < 0.001$; Error bars denote standard error of the mean (s.e.m.).

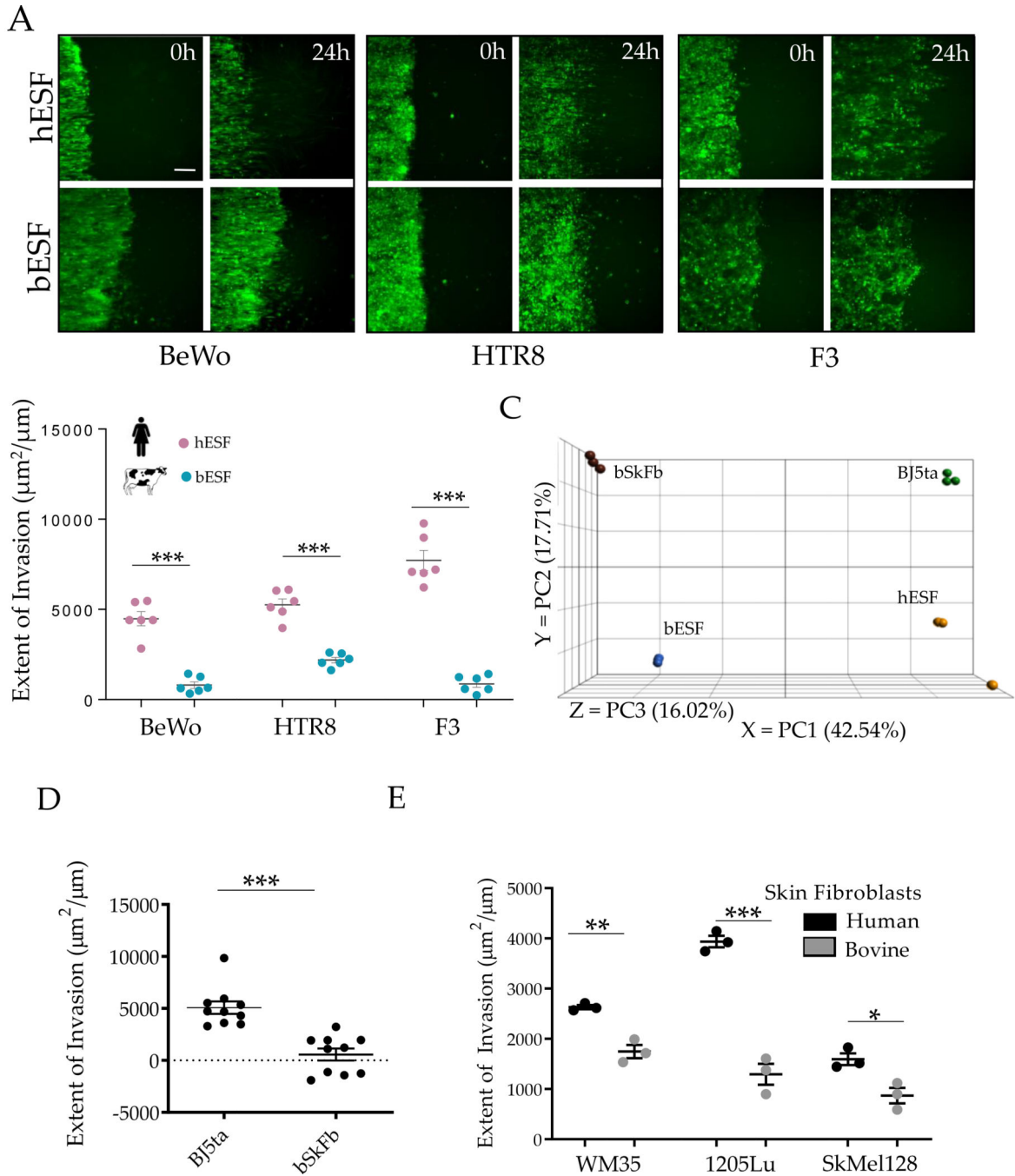


Figure 2. Bovine stroma resists trophoblast and melanoma invasion.

(A) Representative time course images of human trophoblasts BeWo, extravillous trophoblasts HTR8 (green) and bovine trophoblasts F3 (green) invading into endometrial stromal fibroblasts of the respective species for 18 hours; White traces show the boundary of the invasive fronts; Quantification of the extent of invasion for either trophoblasts shown in (B). (C) Principal Component Analysis (PCA) of the gene expression data from skin fibroblasts from human (BJ5ta) and bovine (bSkFb), and endometrial stromal fibroblasts from human (hESF) and bovine (bESF); 3 biological replicates were used for RNA

Sequencing. (D) Extent of invasion of malignant A375 cells into BJ5ta and bSkFb monolayers measured over 24 hours. (E) Extent of invasion of other well characterized human melanoma cell lines into BJ5ta and bSkFb monolayers after 18 hours of observation. In B, C, E, $n = 3$, and in D, $n = 8$ independent biological replicates; Statistical comparisons made using Student's t-tests *: $p < 0.05$, **: $p < 0.01$, ***: $p < 0.001$; Error bars denote standard error of the mean (s.e.m.).

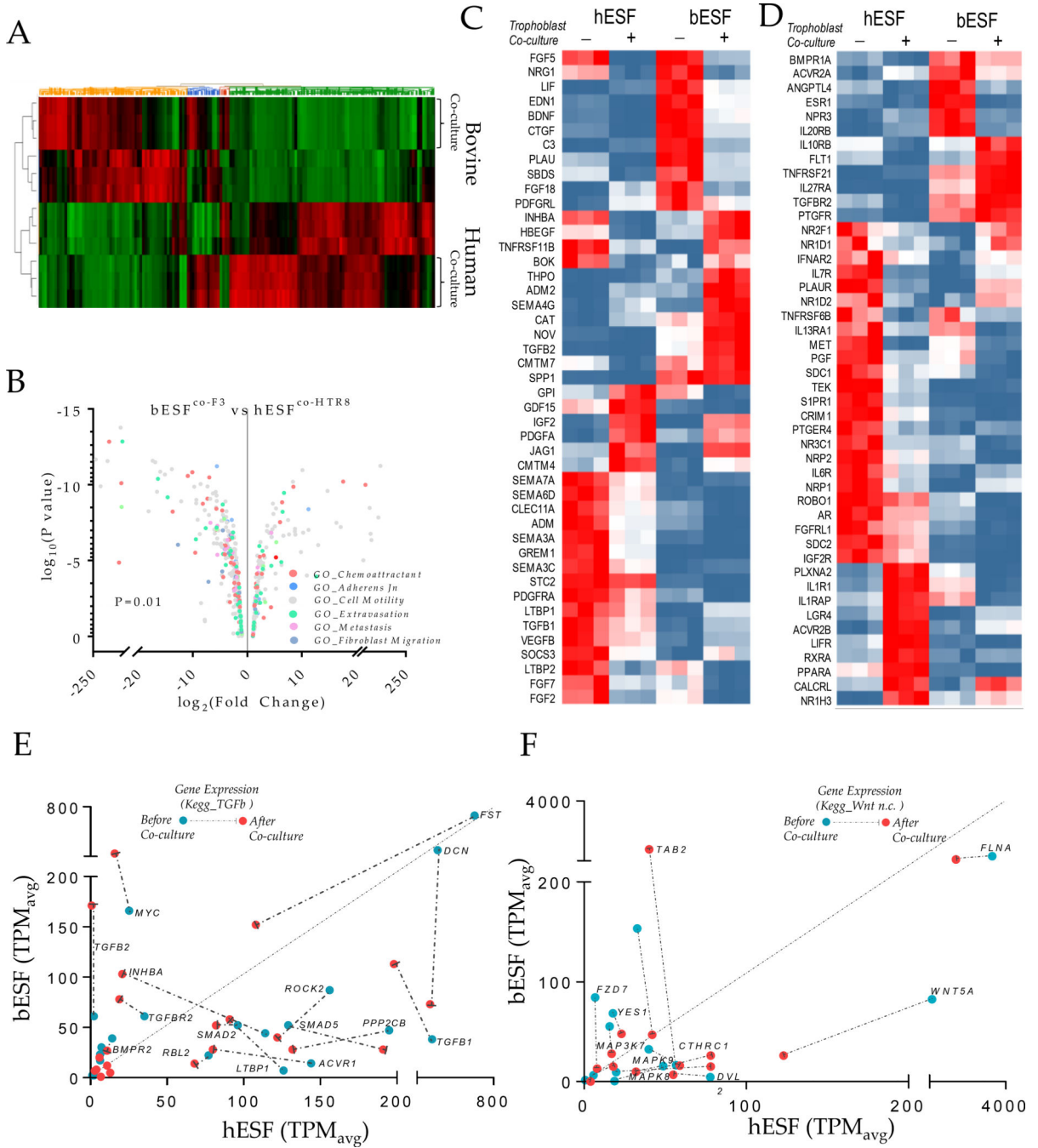


Figure 3. Transcriptomic analysis of bovine and human endometrial stromal fibroblasts reveals differential response to trophoblasts.

(A) Heat map of gene expression differences of human and bovine endometrial stromal fibroblasts with and without co-culture with HTR8 and F3 trophoblast cells respectively. (B) Volcano plot showing fold differences in the TPM values of bESFs vs hESFs co-cultured with the respective trophoblasts, color coded for relevant gene ontologies. (C-D) Heatmap showing expression of genes in hESFs and bESFs with or without co-culture with HTR8 and F3 respectively, belonging to the gene-sets chemokine (C) and chemokine receptors (D). TPM values of genes significantly different in hESFs and bESFs for KEGG pathways for

TGFB (E), and non-cannonical WNT signaling (F); For each gene, TPM values before and after co-culture are shown in green, and red dots respectively; Arrows show the change in TPM values due to trophoblast co-culture; Also are shown coefficient of determination, R^2 for the linear regression of TPM values between hESF and bESF, and the standard deviation of the residuals, $Sy.x$.

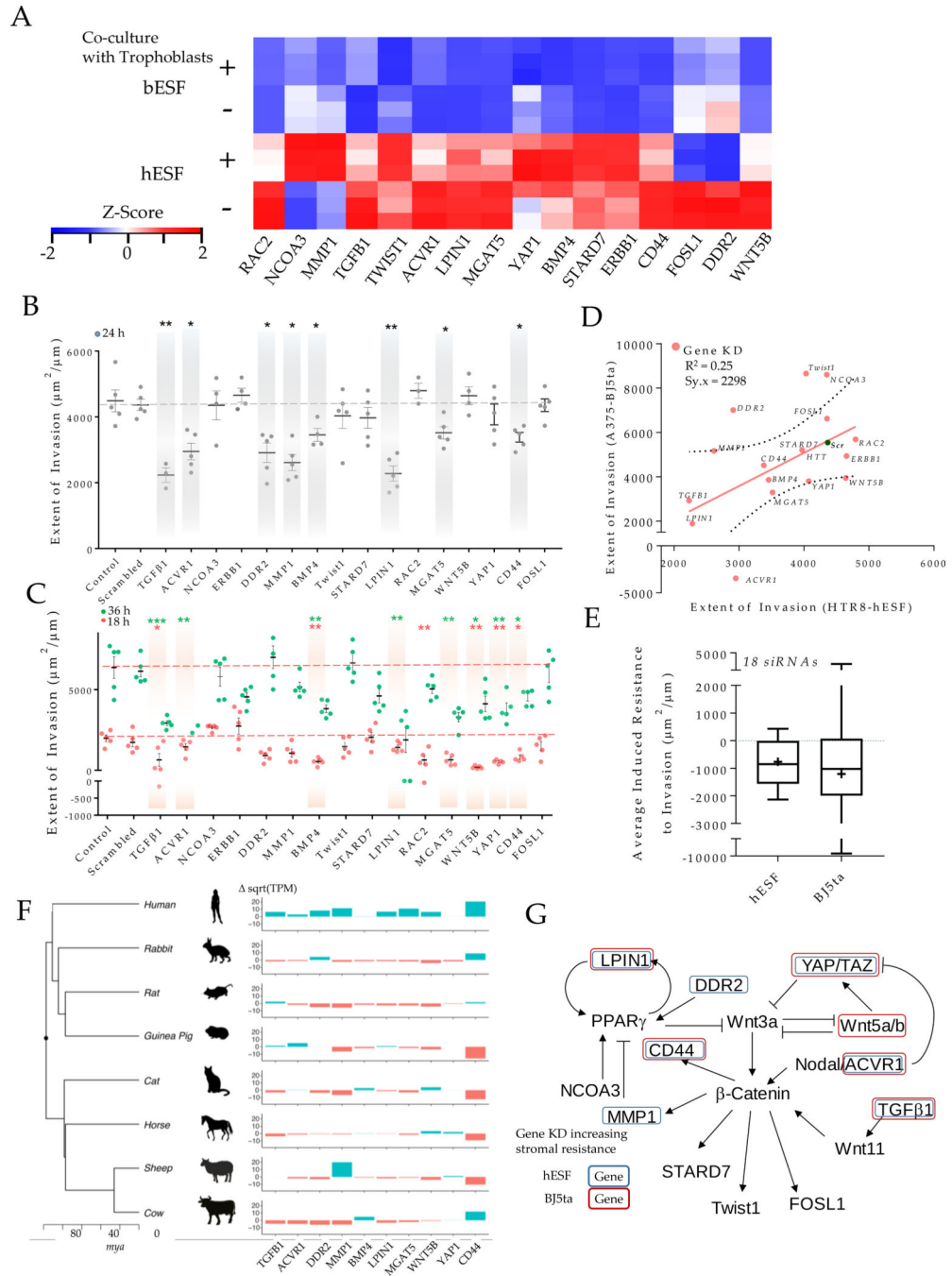


Figure 4. Induced resistance to invasion in human stroma by evolutionarily inspired gene silencing.

(A) Heatmap showing selected set of genes used for gene silencing in human stromal fibroblasts; Shown are z-score of TPM values for each transcript. (B-E) siRNA based gene knockdown in human stroma to induce expression similar to bESF increases resistance to invasion; (B) Extent of invasion of HTR8 cells into hESFs subjected to siRNA mediated gene silencing measured over 24 hours using nanotextured platform; (C) Extent of invasion measured for 18 hours (red), and 36 hours (green) using nanotextured platform for A375 invasion into BJ5ta subjected to siRNA induced silencing of individual genes; Each dot in B,

C reflects an individual invasion observation, Error bars show s.e.m., Statistical comparisons made with scrambled siRNA using Student-t-test; *: $P < 0.05$, **: $P < 0.01$, ***: $P < .0001$. (D) Correlation shown between the extent of invasion of HTR8 into hESF, and A375 into BJ5ta for each siRNA based gene silencing; (E) Average extent of invasion in hESF and BJ5ta for the above 20 gene silencing normalized to the extent of invasion observed for control scrambled siRNA knockdown in either human stromal cells; Whiskers show min to max values; horizontal bar is the median; average values are indicated by +, box is 25 to 75 percentile. (F) Relative transcript levels of the selected genes in various boreoetherians species compared to their most common ancestor; red colored bars denote decrease, and green bars show difference in $\sqrt{\text{TPM}}$ values vs the common ancestor; mya refers to million years ago. (G) Network of WNT signaling and its interaction with TGFB pathway showing previously known interactions, mapped with the gene silencing induced resistance in hESF, and BJ5ta stromal cells.