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DR. LAURA FOUASSIER (Orcid ID : 0000-0001-6377-5610)

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ZEB1 promotes cholangiocarcinoma progression through tumor dedifferentiation and tumor-stroma paracrine signaling

Cindy Lobe^{1,*}, Marie Vallette^{1,*}, Ander Arbelaiz¹, Ester Gonzalez-Sanchez^{1,2,3}, Laura Izquierdo^{3,4}, Anna Pellat¹, Nathalie Guedj⁶, Corentin Louis⁷, Valérie Paradis⁶, Jesus M Banales^{3,4,5}, Cédric Coulouarn⁷, Chantal Housset^{1,8}, Javier Vaquero^{1,2,3,9,#,§} & Laura Fouassier^{1,#,§}

1-Sorbonne Université, Inserm, Centre de Recherche Saint-Antoine (CRSA), Paris, France.

2-TGF- β and Cancer Group, Oncobell Program, Bellvitge Biomedical Research Institute (IDIBELL), Barcelona, Spain.

3-National Biomedical Research Institute on Liver and Gastrointestinal Diseases (CIBERehd), Instituto de Salud Carlos III, Madrid, Spain.

4-Department of Liver and Gastrointestinal Diseases, Biodonostia Health Research Institute - Donostia University Hospital, University of the Basque Country (UPV/EHU), San Sebastián, Spain.

5-IKERBASQUE, Basque Foundation for Science, Bilbao, Spain.

6-Service d'Anatomie Pathologique Hôpital Beaujon, F-92110 Clichy, France; INSERM, UMR 1149, Centre de Recherche sur l'Inflammation, F-75018 Paris, France.

7-Inserm UMR 1241, Univ Rennes 1, Rennes, France.

8-Assistance Publique-Hôpitaux de Paris (AP-HP), Saint-Antoine Hospital, Reference center for inflammatory biliary diseases and autoimmune hepatitis, ERN Rare-Liver, Paris, France.

9-LPP, Sorbonne Université, CNRS, Ecole Polytech., Univ. Paris-Sud, Observatoire de Paris, Université Paris-Saclay, PSL Research University, 4 Place Jussieu, 75252 Paris, France

* Share co-first authorship

Share co-senior authorship

§ Authors to whom any correspondence should be addressed

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Contact information:

Laura Fouassier, Ph.D.

Sorbonne Université, Faculté de Médecine, Site Saint-Antoine,
27 rue Chaligny, 75571 Paris cedex 12, France.

E-mail: laura.fouassier@inserm.fr Phone: +33-698774001

Javier Vaquero, Ph.D.

TGF-Beta and Cancer Group, IDIBELL,

Gran Via de L'Hospitalet, 199, 08908 Barcelona, Spain.

E-mail: jvaquero@idibell.cat Phone: +34-626569867

List of Abbreviations: cancer-associated fibroblasts (CAF), cancer stem cell (CSC), cholangiocarcinoma (CCA), conditioned media (CM), connective tissue growth factor (CTGF), epidermal growth factor (EGF), epithelial-to-mesenchymal transition (EMT), EMT-inducing transcription factors (EMT-TF), epithelial splicing regulatory protein 1 (ESRP1), extrahepatic CCA (eCCA), hepatic stellate cells (HSC), hepatocyte growth factor (HGF), insulin growth factor 2 (IGF2), interleukin (IL), intrahepatic CCA (iCCA), platelet derived growth factor subunit B (PDGFB), reactive oxygen species (ROS), small hairpin RNAs (shRNAs), the cancer genome atlas (TCGA), transforming growth factor beta (TGF- β), tumour-associated macrophages (TAM).

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Abstract

Background & aims

ZEB1 is a transcription factor that promotes metastatic and stem cell features, which has been associated with poor prognosis in cholangiocarcinoma (CCA), a desmoplastic cancer enriched in cancer-associated fibroblasts (CAF). Here, we aimed to define ZEB1 regulatory functions in malignant and stroma compartments of CCA.

Approach and Results

Bioinformatic and immunohistochemical analyses were performed to determine correlations between ZEB1 and markers of progressiveness in human intrahepatic CCA (iCCA). Gain/loss of function models were generated in CCA cells, and liver myofibroblasts, as a model of CAF. Conditioned media (CM) was used to unravel tumor-stroma interplay. *In vivo* experiments were performed using xenograft CCA model. ZEB1 expression in tumor cells of human iCCA was associated with undifferentiated tumor and vascular invasion. *In vitro*, ZEB1 promoted epithelial-mesenchymal transition and stemness in tumor cells leading to cell migration and spheroid formation. *In vivo*, ZEB1-overexpressing CCA cells formed larger tumors with more abundant stroma. *CCN2/CTGF* expression was increased in tumor cells from ZEB1-overexpressing xenografts and correlated with ZEB1 expression in human tumors. *In vitro*, CM from ZEB1-overexpressing tumor cells or recombinant CTGF induced myofibroblast proliferation. ZEB1 was also expressed by CAF in human CCA and its expression correlated with *CCN2* in myofibroblast and CCA stroma. In mice, co-transplantation of CCA cells with ZEB1-depleted myofibroblasts reduced CCA progressiveness compared to CCA cells/ZEB1-expressing myofibroblasts. Furthermore, ZEB1 controls the expression of paracrine signals (i.e. HGF and IL6) in tumor cells and myofibroblasts.

Conclusions

ZEB1 plays a key role in CCA progression by regulating tumor cell-CAF cross-talk, leading to tumor dedifferentiation and CAF activation.

1-Introduction

Cholangiocarcinoma (CCA) includes a heterogeneous group of malignancies that develop from the epithelial layer of the biliary tract.(1, 2) CCA is the second most common primary liver tumor after hepatocellular carcinoma, and its incidence is rising worldwide.(1, 2) Late diagnosis often compromises surgical resection which is the only effective therapeutic option, that is applicable in only 20% of cases. Patients ineligible for surgery undergo a palliative treatment with a combination of gemcitabine and cisplatin (GEMCIS).(1-3) High recurrence rates after tumor resection and chemoresistance contribute to the very poor prognosis of CCA.(1, 2) Therefore, it is essential to investigate the molecular mechanisms involved in the pathogenesis of CCA, still largely unknown, to identify new potential targets for therapy.

Epithelial-to-mesenchymal transition (EMT) is a dynamic and reversible process by which cells lose epithelial characteristics and acquire mesenchymal properties.(4, 5) This process plays a major role in the formation of metastatic carcinoma cells,(4, 5) the acquisition of stem cell properties(4, 5) and the induction of chemoresistance against a variety of chemotherapeutic drugs and targeted therapies.(4-7) Diverse extracellular signals promote EMT by increasing the expression of EMT-inducing transcription factors (EMT-TF) from the 3 major families: SNAI (SNAIL and SLUG), TWIST (TWIST1 and TWIST2) and ZEB (ZEB1 and ZEB2).(4, 5) ZEB1, that belongs to the Zinc finger E-box binding protein family, is able to recognize E-box sequences in the promoter of certain genes, acting as a transcriptional repressor (*i.e.* E-cadherin) or activator (*i.e.* vimentin).(8) In human, ZEB1 becomes aberrantly expressed in different types of cancer, both in tumor and stromal cells, *i.e.* cancer-associated fibroblasts (CAF)(9), tumor-associated macrophages (TAM),(10) and endothelial cells.(11) Of note, ZEB1 expression in both tumor and stromal cells has been linked to early recurrence and poor prognosis in human pancreatic ductal adenocarcinoma.(12) In this cancer, which displays a strong desmoplastic stroma, ZEB1 expression in CAF supports cancer progression via paracrine signaling with tumor cells.(13) Similarly, CCA is characterized by a dense desmoplastic stroma enriched in CAF that derive either from hepatic stellate cells (HSC) or portal myofibroblasts, and participate in tumor progression(14, 15) and chemoresistance.(16)

ZEB1 has been associated with poor prognosis in human intrahepatic CCA (iCCA).(17) Yet, little is known regarding its cellular expression and functions in CCA. Thus, the present study aimed to define the expression of ZEB1 in tumor cells and CAF of CCA and its potential function in the interplay between these two cell types by using preclinical models and human samples of CCA. Here, we describe a new dual function of ZEB1 in CCA. Our data demonstrate that ZEB1 promotes CCA tumor cell dedifferentiation through the acquisition of an EMT/cancer stem cell

(CSC) phenotype, along with an upregulation of CTGF that contributes to the development of a profibrogenic microenvironment. Moreover, ZEB1 impacts the cross-communication between malignant cells and stromal myofibroblasts by regulating the expression of HGF and IL6, leading ultimately to CCA progression.

2-Materials and methods

ZEB1 gain and loss of function cell models

Stable ZEB1 inhibition in SK-ChA-1, HuCCT1 R and in liver myofibroblasts (hTERT-HSC cells) was performed by lentiviral transduction with specific small hairpin RNAs (shRNAs). Plasmids containing sequences targeting ZEB1 (shRNA-ZEB1) or scramble (shRNA-Control) designed by GenecopoeiaTM (Maryland, USA) were used to produce Δ U3 SIN lentivirus in the Viral and Gene Transfer Vectors Platform of Necker (Paris, France). Briefly, cells were infected and maintained in presence of lentivirus for 24h. Then, lentiviruses were removed, and fresh media was added to maintain cells in culture another 48h until mCherry expression was observed. Positive transduced cells were detected by positive fluorescence signal and they were selected by treatment with lethal doses of puromycin for 72h to eliminate non-infected cells. Due to incomplete efficiency of puromycin selection in SK-ChA-1 and HuCCT1 R, 2 clones of each cell type were isolated and characterized throughout the study. The cell transduction was performed with level 2 precautions according to biosafety regulation standard ISO 15190:2003.

ZEB1 experimental overexpression was performed by retroviral infection. Plasmids coding for the murine open reading frame of Zeb1 or Mock (control), were used to produce retrovirus in the same platform, following the protocol previously described by Dr. Julie Caramel (Cancer Research Center, Lyon, France).(18) For retroviral transduction, the same protocol was applied to the HuCCT1 cells. In this case, puromycin was effective and cell populations were used in the study.

HuCCT1 resistant cells to erlotinib (HuCCT1 R) were generated through a process of slowly escalating exposure to erlotinib (from 1 to 20 μ M), an EGFR tyrosine kinase inhibitor, as previously described.(16)

For clarification purposes, along the manuscript cell lines or murine models derived from injection of cells with low ZEB1 expression will be represented by white bars, while those with high ZEB1 expression will be represented by black bars.

Further information about the materials and methods is available in the Supplementary Information.

3-Results

3.1- ZEB1 expression in malignant cells is associated with tumor differentiation and vascular invasion in human CCA

To examine ZEB1 expression, we performed an immunostaining in a set of tissue samples from 45 subjects with iCCA. While ZEB1 expression was absent from cholangiocytes of adjacent liver tissue (Figure 1A, yellow arrow), 20% of iCCA displayed a nuclear expression of ZEB1 in cancer cells (Figure 1A, black arrows). ZEB1 was also observed in stromal cells (Figure 1A, red arrows). In cancer cells, ZEB1 expression positively associated with tumor differentiation (Figure 1B) and vascular invasion (Figure 1C). Accordingly, the expression of ZEB1 in a panel of 8 human CCA cell lines (including iCCA and extrahepatic CCA (eCCA) cell lines) was inversely correlated with the expression of E-cadherin and *ESRP1*, two epithelial markers, and positively with that of vimentin, a mesenchymal marker, at protein (Figure 1D) and mRNA levels (Supplementary Figure 1). Furthermore, ZEB1 is the EMT-TFs that best correlated both negatively with the expression of epithelial markers and positively with the mesenchymal marker (Supplementary Figure 1). Interestingly, CCA cell lines with negligible ZEB1 expression are derived from more differentiated human tumors, than the cell lines with high ZEB1 expression that were obtained from undifferentiated tumors.(19-24) Accordingly, transcriptomic expression analyses showed a strong negative correlation between *ZEB1* and both *CDH1* (encoding E-cadherin) and *ESRP1* in human CCA from Rennes and TCGA cohorts, and a strong positive correlation between *ZEB1* and *VIM* (encoding vimentin) in both datasets (Figure 1E).

3.2- ZEB1 regulates CCA cell dedifferentiation by inducing epithelial-to-mesenchymal transition and stemness phenotype

To decipher the molecular mechanisms by which ZEB1 regulates the differentiation of cancer cells, we modulated ZEB1 expression in CCA cells by different approaches. In HuCCT1 cells, the murine form of Zeb1 was overexpressed as previously described,(18) and, in SK-ChA-1 cells, ZEB1 was downregulated by shRNA (Figure 2A-C). Additionally, HuCCT1 cells resistant to erlotinib (HuCCT1-R, previously generated in our laboratory),(16) which overexpressed ZEB1 as a cause of resistance, were also modified to downregulate ZEB1 (Figure 2A-C). In the 3 cellular models, genetic manipulation of ZEB1 did not induce a compensatory alteration at the mRNA or protein level of other EMT-TFs (Supplementary Figure 2). Interestingly, ectopic expression of ZEB1 in HuCCT1 induced a decrease of the epithelial markers *CDH1* and *ESRP1* and an upregulation of the mesenchymal marker *VIM* (Figure 3A-C), whereas ZEB1 downregulation had opposite effects in SK-ChA-1 and HuCCT1-R (Supplementary Figures 3A-C and 4A-C). Because

EMT-TFs are linked to the acquisition of stemness features,(8) we evaluated the impact of ZEB1 in the regulation of two CSC markers, CD44 and CD24, allowing the quantification of a population (CD44^{high}/CD24^{low}) that shares EMT/CSC traits.(16, 25). Accordingly, overexpression of ZEB1 caused an upregulation of CD44 and a marked downregulation of CD24 (Figure 3D), whereas ZEB1 knockdown had the opposite effect (Supplementary Figures 3D and 4D). These data were confirmed by flow cytometry analysis, indicating that high ZEB1 expression concurred with an increase of CD44^{high}/CD24^{low} cell population (Figure 3E) and inversely when ZEB1 is down-regulated (Supplementary Figures 3E and 4E).

These phenotypic modifications were functionally translated by an increased migration, colony and sphere formation in HuCCT1 Zeb1-overexpressing cells (Figure 3F-H) and by a mirroring effect in SK-ChA-1 and HuCCT1-R cells downregulated for ZEB1 (Supplementary Figures 3F-H and 4F-H), compared with the corresponding controls. Altogether, these observations indicate that ZEB1 expression in CCA cells confers high plasticity leading to cell dedifferentiation.

3.3- ZEB1 induces the generation of larger CCA tumors with abundant stromal content

To determine the biological relevance of ZEB1 in CCA cells, we generated CCA tumors into immunocompromised mice by engrafting cancer cells that overexpressed ZEB1 (HuCCT1-Zeb1). CCA cells overexpressing Zeb1 generated larger tumors (Figure 4A-B), histologically characterized by a prominent desmoplastic stroma (Figure 4C) with higher collagen deposition (Figure 4D), compared with control tumors. In this model, engraftment of human cancer cells gave rise to tumors with murine stromal cells, which we took advantage of to analyze gene expression in cancer cells and stromal cells using human and murine specific primers, respectively. In this context, the expression of murine *Acta2* (α -Sma), a marker of CAF, and *Col1a1* (collagen 1) and *Col4a1* (collagen 4), extracellular matrix (ECM) components produced by CAF, was increased in the stroma of tumors developed from CCA cells overexpressing Zeb1 (HuCCT1-Zeb1) (Figure 4E) in agreement with the histological observations (Figure 4C-D). These observations agreed with our previous findings in resistant CCA tumors (developed from HuCCT1-R cells that overexpress ZEB1), in which the stroma content was also increased.(16) To understand how the stromal compartment expanded upon transplantation of Zeb1-overexpressing tumor cells (HuCCT1-Zeb1), we evaluated the expression of key pro-fibrogenic factors, *TGFB1* (encoding transforming growth factor beta 1, TGF- β 1), *CCN2* (encoding connective tissue growth factor, CTGF) and *PDGFB* (encoding Platelet Derived Growth Factor Subunit B, PDGFB) in the tumoral human compartment, corresponding to CCA cells. The expression of *TGFB1* and *PDGFB* in CCA cells, was unchanged in tumors generated from the cells overexpressing Zeb1, whereas a significant increase in *CCN2* mRNA levels was observed

(Figure 4F). The analysis of resistant CCA tumor samples overexpressing ZEB1, obtained from our previous work(16) revealed an even stronger increase in *CCN2* expression, compared to WT (Figure 4G). Protein analyses confirmed the upregulation of CTGF expression in ZEB1 overexpressing tumors (Figure 4 H-I). These results suggest that in tumor cells, ZEB1 positively regulates *CCN2*/CTGF, which boosts the stromal expansion of CCA.

3.4- ZEB1 regulates *CCN2* transcription in CCA cells leading to myofibroblast expansion

The transcriptional activation of *CCN2*/CTGF by ZEB1 has been described in breast cancer cells through its cooperation with YAP signaling,(26) which is active in CCA cells.(27) Accordingly, analysis of several CCA cell lines showed that *CCN2* expression correlated with *ZEB1* expression (Supplementary Figure 5A). Evaluation of genetically manipulated CCA cells in which ZEB1 is either overexpressed or down-regulated, demonstrated that the mRNA expression of *CCN2* concurred with that of *ZEB1*, showing a strong upregulation of *CCN2* expression in Zeb1-overexpressing cells (HuCCT1-Zeb1) and a downregulation in ZEB1-depleted CCA cells (SK-ChA-1- and HuCCT1 R-shZEB1 cells), compared with controls (Figure 5A). At the protein level, CTGF expression also concurred with ZEB1 in HuCCT1 Control/Zeb1 and HuCCT1 R-shRNA cell lines (Figure 5A). Despite mRNA level difference in SK-ChA-1, there were no differences of CTGF at the protein level (Figure 5A), hypothesizing the existence of unknown posttranslational mechanisms in this cell line. Regarding *TGFB1* expression, no differences were observed (Supplementary Figure 5B) corroborating our *in vivo* findings (Figure 4F and 4G). Transcriptomic analysis of human CCA showed a strong positive correlation between *CCN2* and *ZEB1* expressions in the Rennes Cohort and a similar trend in the TCGA cohort (Figure 5B). To evaluate the effect of CTGF on stromal cells, human liver myofibroblasts (hTERT-HSC cell line), used as a model of CAF, were exposed to recombinant CTGF (Figure 5C) or to conditioned media (CM) from Zeb1-overexpressing cells (HuCCT1-Zeb1) in which *CCN2* is upregulated (Figure 5D). In both cases, the proliferation of myofibroblasts was increased, compared with the controls (Figure 5C-D), indicating a pro-proliferative effect of CTGF on myofibroblasts, consistent with its role in hepatic fibrosis.(28, 29) Furthermore, addition of a neutralizing antibody against CTGF to CM of HuCCT1-Zeb1 reduces significantly the effect on myofibroblasts viability (Figure 5D). Contrarily, CTGF had no effect on CCA cell proliferation (Supplementary Figure 5C) nor on the expression of EMT markers and pro-inflammatory cytokines (Supplementary Figure 5D). Finally, ChIP analyses demonstrated that ZEB1 upregulates *CCN2* expression by directly binding its promoter (Figure 5E), as previously demonstrated in breast cancer.(26)

Finally, to extend our findings on the relationship between *ZEB1* and *CCN2* to other cancer types, *ZEB1* and *CCN2* expression was analyzed both in human cell lines and tissues using the

Broad-Novartis Cancer Cell Line Encyclopaedia database that includes 1156 cell lines (Supplementary Figure 6A) and the TCGA datasets (Supplementary Figure 6B). We confirmed that *ZEB1* and *CCN2* are interrelated in other cancers, suggesting a ubiquitous mechanism of regulation.

3.5- ZEB1 is expressed by CAF and regulates their activation status

CCA is characterized by a dense desmoplastic stroma rich in CAF, which have an important role in tumor progression.⁽¹⁴⁾ Our observation of ZEB1 expression in stromal cells from CCA patients (Figure 1A, red arrows), led us to further investigate the role of ZEB1 in CAF. First, immunostaining of ZEB1 and α -SMA in human iCCA samples, showed a co-expression of both proteins in stromal areas (Figure 6A, red arrows), indicating that α -SMA expressing CAF also express ZEB1. Of note, some stromal cells showed ZEB1 expression in absence of α -SMA (Figure 6A, black arrows), suggesting that other stromal populations may express ZEB1. Furthermore, transcriptomic analysis of laser microdissected tissue from stromal areas of human iCCA showed a strong correlation of *ZEB1* with *ACTA2* and *COL4A1*, and with the pro-fibrotic factors *TGFB1* and *CCN2* (Figure 6B), but not with *PDGFB* (Supplementary Figure 7A). Accordingly, analysis of xenografts derived from ZEB1-overexpressing CCA cells (HuCCT1-Zeb1 and HuCCT1-R), which display a more prominent and activated stroma, showed an upregulation of *Zeb1* in stroma (*i.e.* mouse gene, Figure 6C-D) together with that of *Acta2*, *Col4A1* (Figure 4E and ref (16)), *Tgfb1*, *Ccn2* and *Pdgfb* (Supplementary Figure 7 B-C). To validate *in vitro* the regulation of these factors by ZEB1, ZEB1 was stably downregulated in hepatic myofibroblasts (Figure 6E), resulting in decreased expression of *TGFB1*, *CCN2*, *PDGFB*, *ACTA2* and *COL4A1*, (Figure 6F). ChIP analyses confirmed the direct regulation of *CCN2* by ZEB1 also in human liver myofibroblasts (Supplementary Figure 7D). Downregulation of *CCN2* in hTERT-HSC by siRNA was insufficient to modify the expression of *TGFB1*, *PDGFB* and *COL4A1* and only a small although significant reduction of *ACTA2* was observed in cells (Supplementary Figure 7E), indicating that ZEB1 effects are not dependent on CCN2 expression or at least not exclusively. Altogether, these results provide evidence of the role of ZEB1 in the activation of stromal CAF.

3.6 Depletion of ZEB1 in myofibroblasts reduces CCA progressiveness

Next, we evaluated the *in vivo* relevance of ZEB1 expression in human liver myofibroblasts on CCA growth. CCA cells were co-engrafted with liver myofibroblasts depleted for ZEB1 (hTERT-HSC-shZEB1) or not (hTERT-HSC-shControl). We used EGI-1 CCA cells because they have negligible expression of ZEB1 (to avoid interference with ZEB1 expressed by stromal cells) and their tumorigenic potential is the highest among our CCA cell lines. Transplantation of CCA cells with liver myofibroblasts depleted for ZEB1 generated smaller tumors compared to the controls (Figure 7A). One concern of transplanting human liver myofibroblasts in immunocompromised mouse is their replacement by resident mouse fibroblasts during tumorigenesis. Indeed, mRNA expression of fibroblast related genes from human origin (*ACTA2*, *COL1A1* and *COL4A1*) was very low in the tumors (Supplementary Figure 8A) and similar to that of xenografts derived from implantation of EGI-1 cells alone.⁽¹⁶⁾ Interestingly, analysis of the murine counterparts showed a downregulation of *Zeb1*, *Tgfb1*, *Acta2*, *Col4a1*, and a trend for *Ccn2* and *Pdgfb* expression in the stromal compartment of tumors developed from CCA cells with ZEB1-depleted myofibroblasts, compared with the controls (Figure 7B), indicating that, in addition of promoting tumor growth, human myofibroblasts could educate the recruited murine stroma at an early time point. On the contrary, no changes in profibrotic factors were observed in the tumor compartment (Supplementary Figure 8B).

Finally, to evaluate the direct effect of liver myofibroblasts on tumor cells, CCA cells were exposed to CM from ZEB1-depleted myofibroblasts or the controls and CCA cell viability was analyzed. CM from control myofibroblasts (hTERT-HSC-shRNA-Control) strongly increased the viability of EGI-1 and HuCCT1 cells, compared with serum-free media. In contrast, CM from ZEB1-depleted myofibroblasts showed a weak effect on the viability of EGI-1 cells and no effect on HuCCT1 cells (Figure 7C). Changes in cell viability were accompanied by the activation of pro-survival intracellular signaling pathways. CM from control myofibroblasts activated STAT3 and AKT in CCA cells, while CM from ZEB1-depleted myofibroblasts had very low or no effect on these signaling pathways (Figure 7D). No differences were observed in the status of ERK1/2 (Figure 7D). No changes in mRNA level of EMT markers and pro-inflammatory cytokines were observed in CCA cells treated with any of the CM from myofibroblasts (Supplementary Figure 9). Altogether, these results suggest that ZEB1 in myofibroblasts regulates their interplay with CCA cells, that depends on growth factors secreted by CAF.

3.7- ZEB1 impacts the interplay between CCA tumor cells and stromal myofibroblasts

Our study suggests that ZEB1 regulates the interplay between CCA cells and CAF to promote CCA progression. Therefore, as a proof of concept, we selected four molecules with major roles in CCA progression that could play a role in ZEB1-regulated interplay: insulin growth factor 2 (IGF2),(16) epidermal growth factor (EGF),(30) hepatocyte growth factor (HGF)(31) and interleukin 6 (IL6).(32) We analyzed the expression of these factors in our preclinical models, including i/ Zeb1-overexpressing cells (HuCCT1-Zeb1 cells vs control cells) and mouse CCA tumors generated from these cells, ii/ CCA cells resistant to erlotinib that overexpressed ZEB1 (HuCCT1-R vs WT cells) and tumors generated from these cells, and iii/ ZEB1-depleted myofibroblasts (hTERT-HSC-shZEB1 vs control cells) and tumors generated from EGI-1 cells with myofibroblasts (Figure 8 and Supplementary Figure 10).

Both *IGF2* and *EGF* expression increased in ZEB1-overexpressing cells (HuCCT1-Zeb1 and HuCCT1 R) compared with their controls (Supplementary Figure 10A-B). Similarly, in ZEB1-depleted myofibroblasts, *IGF2* and *EGF* expression was strongly downregulated compared to the controls (Supplementary Figure 10A-B). However, once implanted in mice no change was observed in the tumor (Supplementary Figure 10A-B), suggesting that the complexity of *in vivo* systems may influence ZEB1 regulation of these factors.

HGF was found overexpressed in Zeb1-overexpressing cells (HuCCT1-Zeb1) and resistant cells (HuCCT1-R) compared to the controls (Figure 8A). Interestingly, *in vivo* *HGF* was overexpressed in both human (reflecting expression in cancer cells) and murine (reflecting expression in stromal cells) compartments of CCA tumors (Figure 8A). On the contrary, no change in the expression of *HGF* was observed in cultured myofibroblasts and *in vivo* in the co-transplantation xenograft model (Figure 8A). The increased *HGF* expression in the stroma of mice tumors derived from ZEB1-overexpressing HuCCT1 cells suggested that ZEB1 in malignant cells induces *HGF* expression in stromal cells through paracrine signal. To test this hypothesis, we exposed liver myofibroblasts (hTERT-HSC) to CM from Zeb1-overexpressing cells (HuCCT1-Zeb1) or the controls. Indeed, CM from HuCCT1-Zeb1 cells upregulated *HGF* expression in myofibroblasts compared to control CM (Figure 8B). However, neither direct treatment with exogenous CTGF nor the addition of a neutralizing antibody against CTGF to the CM of HuCCT1-Zeb1 cells modified the *HGF* expression in hTERT-HSC cells, indicating that other factors upregulate *HGF* (Supplementary Figure 11A-B). Transcriptomic analysis of human CCA showed a strong and positive correlation between *ZEB1* and *HGF* expressions in both the Rennes and the TCGA cohorts, which was confirmed in other cancer types using public data bases (Supplementary Figure 12A-C). Further ChIP assays demonstrated the direct binding of ZEB1 to *HGF* promoter (Supplementary Figure 12D).

The only statistical change observed for *IL6* was a strong downregulation in myofibroblasts depleted for ZEB1 (hTERT-HSC-shZEB1) compared to controls, that was also observed in the stromal compartment of CCA xenografts (Figure 8C). This specific regulation of *IL6* in the liver myofibroblasts suggested that ZEB1 may regulate *IL6* in CAF but not in malignant cells. To address this possibility, we treated ZEB1-depleted myofibroblasts (hTERT-HSC-shZEB1) or the controls with TGF- β , the classical molecule used to activate myofibroblasts and we monitored the expression of *ZEB1* and *IL6* genes. *ZEB1* expression was upregulated in response to TGF- β in control myofibroblasts and as expected, this response was completely abolished in ZEB1-depleted cells, confirming the efficiency of our shRNA system (Figure 8D). In agreement with our hypothesis, *IL6* response overlapped with that of *ZEB1*, being upregulated by TGF- β in control myofibroblasts, but abolished in cells depleted for ZEB1, confirming ZEB1 regulation of *IL6* in liver myofibroblasts (Figure 8D). In human CCA, *IL6* showed a strong correlation with ZEB1 expression in both the Rennes and the TCGA cohorts (Supplementary Figure 13A). Similarly, this correlation was confirmed in other human cancers from the Broad-Novartis Cancer Cell Line Encyclopaedia and the TCGA datasets (Supplementary Figure 13B-C).

To functionally translate the importance of ZEB1 regulation over these factors, we performed viability studies on a panel of malignant cells and liver myofibroblasts (including those in which ZEB1 expression was genetically modified) by exposing the cells to HGF and IL6. These experiments show that HGF was able to increase the viability of almost all cell lines tested (Figure 8E), while IL6 was only able to potentiate the viability of HuCCT1 cells (both HuCCT1 Control and Zeb1) (Figure 8F). Nevertheless, the most interesting effect observed was that the modulation of ZEB1 expression leads to increase (in HuCCT1-Zeb1) or reduce (in hTERT-HSC-shZEB1) response to HGF (Figure 8E). A similar ZEB1-dependent modulatory response was observed when HuCCT1-Control and Zeb1 cells were exposed to CM of hTERT-HSC, and when hTERT-HSC-shControl and shZEB1 were exposed to CM of HuCCT1 (Supplementary Figure 14A-B). Furthermore, HGF promotes a moderate activation of hTERT-HSC-shControl cells, but has no effect in hTERT-HSC depleted for ZEB1 (Supplementary Figure 14C). However, mRNA expression of receptors for HGF (MET) and IL6 (IL6R and IL6ST, encoding GP130) did not explain the altered sensitivity to these soluble factors (Supplementary Figure 15A-C), indicating that modifications of other signaling mediators account for these effects.

Altogether, these results indicate that ZEB1 regulates not only the production of soluble signaling molecules (such as HGF) but their signaling pathways creating a signaling hub between tumor and stromal cells to boost CCA growth.

4 -Discussion

The final purpose of translational research is to extrapolate findings from preclinical models into new therapeutic targets, which are potentially targetable in clinical practice to improve patient management, in this case CCA patients. Following this strategy, we have used multiple cell culture models, animal experimentation with tumor xenografts and validated our findings by taking advantage of immunostaining and transcriptomic data from human CCA to pinpoint ZEB1 as an ideal therapeutic candidate against CCA for which new ways of treatment should be further explored.

ZEB1 is a transcription factor originally discovered as a mediator of the EMT process in embryonic development.(33) During the last decades it has acquired a prominent role in the biology of tumor cells in a variety of cancers by promoting tumor cell stemness, invasion and chemoresistance to an assortment of drugs.(8) Here we confirm a role for ZEB1 in the dedifferentiation and the acquisition of a more aggressive phenotype of CCA tumor cells. ZEB1 increased cell plasticity through modulation of epithelial (*i.e.* E-cadherin), mesenchymal (*i.e.* vimentin) and stemness (*i.e.* CD44/CD24) marker expression, providing them with migratory and stemness properties, which in human CCA was translated as a correlation of ZEB1 expression in tumor cells with poorly differentiated tumors and vascular invasion. These data open the way of the potential therapeutic usefulness of targeting ZEB1 in late stages of CCA, which could prevent cell dissemination by impairing EMT, and might have a profound impact in reducing cancer progression.

Recent evidence points to ZEB1 involvement in different aspects of tumor progression through its expression in several stromal cell types,(9-11, 13) although ZEB1 role in these cells is not yet fully understood. The desmoplastic nature of CCA stroma led us to focus a large part of our study on demonstrating the role of ZEB1 in the activation of stromal CAF. Nevertheless, we also found ZEB1 in areas of human iCCA that resemble tumor infiltrates (Supplementary Figure 16A) and vascular structures (Supplementary Figure 16B), indicating that ZEB1 is probably present also in immune cells and endothelial cells of CCA. Accordingly, major functions for ZEB1-expressing TAM and endothelial cells have been described in other cancers.(10, 11) This observation opens the door to further research to fully understand ZEB1 regulatory network over the different cell types present in CCA and how they interact to each other.

We also provide evidence that ZEB1 is at the interface of tumor-stroma cross-communication by regulating the expression of growth factors and proinflammatory cytokines that impact CCA progression.

CCN2/CTGF is a growth factor with a prominent role in the activation of HSC during liver fibrosis.(29) In accordance with previous observations that place ZEB1 as a regulator of CTGF,(26) we show that ZEB1 is an inducer of CCN2 expression at both sides of the CCA cell-CAF crosstalk by direct binding to its promoter. Modulation of ZEB1 resulted in a concomitant change of CCN2 expression in all genetically modified tumor and myofibroblast cells, even when these cell lines were implanted into immunocompromised mice to generate tumors. Moreover, ZEB1 and CCN2 expression correlated in human samples, translating our preclinical evidence into iCCA. Interestingly, our data suggest that this regulation occurs in both human and murine species, as shown by the ability of both murine (HuCCT1-Zeb1) and human (HuCCT1 R) ZEB1 to upregulate CCN2 in CCA cells. These observations show the high degree of evolutionary conservation of this regulatory mechanism, depicting the biological importance of ZEB1-CCN2 tandem in CCA and probably in other pathologies (including several types of cancer) as previously described(34, 35) and the strong correlation found in cancers from the TCGA cohort. Furthermore, CCN2 has been implicated in promotion of inflammation and angiogenesis(35) and preliminary analyses of specific macrophage and endothelial cell markers in our *in vivo* models (Supplementary Figure 16 C-D), suggest that ZEB1 expression in tumor cells may have a more important role in the reprogramming of entire stromal compartment through CCN2/CTGF in late stages of cancer to support tumor growth and dissemination. These evidences together with the fact that CTGF is an independent prognostic factor indicator of both tumor recurrence and overall survival for intrahepatic CCA(36) ensures its potential as a target for the future of this disease. In this sense, Pamrevlumab, a CTGF neutralizing antibody, recently reached the clinics and results from trials in pancreatic cancer are encouraging,(37) showing that the use of Pamrevlumab in combination with gemcitabine and nab-paclitaxel has potential to enhance tumor response and increase resection rates. Looking at the similarities of pancreatic cancer and CCA, both exhibiting a high degree of desmoplasia due to extensive extracellular matrix deposition, once more the evidence points to the potential use of the anti-CTGF antibody in CCA.

We also found that ZEB1-overexpressing tumor cells displayed an elevated expression of HGF, in both cells and murine preclinical models of CCA, and direct binding of ZEB1 to HGF promoter was demonstrated by CHIP analysis for the first time to our knowledge. Furthermore, ZEB1 expression in CCA tumor cells induced an upregulation of HGF in stromal cells from xenografts and CM from ZEB1-overexpressing cells increased HGF expression in myofibroblasts *in vitro*. A recent work describes a similar paracrine regulatory mechanism by which ZEB1 expression in tumor cells is able to stimulate stromal macrophage production of signaling molecules (*i.e.* ROS and IL-1 β) to promote inflammatory colorectal carcinoma.(38) Nonetheless, the exact paracrine

signal that favors HGF production by CAF in CCA is yet to be identified. Interestingly, we also observed that manipulation of ZEB1 expression in CCA cells and fibroblasts correlated with the ability of these cells to respond to exogenous HGF *in vitro*. Since expression of the main HGF receptor (MET) did not explain this ZEB1-dependent response, we hypothesized that ZEB1 may be modulating the expression of other mediators that could influence HGF signaling. MET has been described as a promiscuous receptor able to interact with multiple partners that act as co-receptors, modulating its signaling.(39) Among them, integrins are membrane receptors described to play important roles in the establishment of EMT.(4) Previous results from our group(16) led us to hypothesize a potential implication of Integrin Beta 3 (*ITGB3*, also known as CD61), which has also been described as a CTGF receptor in fibroblasts.(40) Indeed, expression analysis indicate that mRNA and protein expression of *ITGB3* perfectly correlated with that of *ZEB1* in HuCCT1 and hTERT-HSC modified cells (Supplementary Figure 17A-B). Although very preliminary data, further functional experiments are guaranteed to unravel the exact functions of *ITGB3* downstream of ZEB1.

The tumor microenvironment plays a role in cancer progression from initiation to metastasis and therapeutic response.(41) Stromal CAF from CCA are known to produce a wide variety of growth-stimulatory, pro-angiogenic, immune-modulatory and pro-invasive factors, as well as different ECM components.(42) IL6 binds to specific receptors on tumor and stromal cells leading to cancer progression.(43) It has been demonstrated that IL6 and IL8 are directly regulated by ZEB1 in breast cancer cells,(44) but we did not find any relationship between ZEB1 and IL6 expression in CCA tumor cells. Here we found that ZEB1 invalidation in fibroblasts reduces the expression of IL6 to the surrounding stroma. Recently it has been described that IL6 can be indirectly regulated by ZEB1 through a mechanism that involves p53 acetylation and recruitment to *IL6* promoter in stromal fibroblast from tumor of mammary origin.(9) Intriguingly, we observed a similar regulation of both *IL6* and *CXCL8* (IL8) in hTERT-HSC invalidated for ZEB1 and exposed to TGF- β 1 (Figure 8D and Supplementary Figure 18), although the absence of a murine IL8 ortholog made impossible its analysis in the stroma of xenografts. The *in vitro* experiments showed a modest effect of IL6 on CCA cell lines viability. However, effects of IL6 at other levels of the tumorigenic process(43) remain to be explored in our experimental models.

In conclusion, our results highlight ZEB1 as a key player in the progression of CCA, not only by allowing tumor cells to become more dedifferentiated and aggressive, by participating in the activation of CAFs and the production of extracellular matrix, but also through regulation of crosstalk signals that boost CCA growth (summarized in Figure 8G).

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“Author names in bold designate shared co-first authorship”

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Author contributions:

Conceptualization and design: JV, LF.

Data generation: CL, MV, AA, EGS, LI, AP, NG, CL, CC, JV.

Data analysis and interpretation: CL, AA, EGS, NG, CL, VP, CC, JV, LF.

Manuscript preparation: CC, NG, JV, LF.

Review and editing: CL, MV, AA, EGS, LI, AP, NG, CL, VP, JMB, CC, CH, JV, LF.

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Figure legends

Figure 1. ZEB1 expression is related to tumor dedifferentiation in CCA. **A.** Representative IHC staining of ZEB1 in human non-tumoral liver (left panel), Grade 1 (middle panel) and Grade 3 (right panel) intrahepatic CCA. **B-C.** Quantification of CCA with ZEB1 positive signal in tumor epithelial cells according to tumor grade (B) and vascular invasion (C). Statistics were performed by applying the Chi square test. **D.** ZEB1, E-cadherin and vimentin protein expression in a panel of 8 human CCA cell lines, as determined by western blot. **E.** Correlation between *CDH1*, *ESRP1* and *VIM* and *ZEB1* expression in Rennes and TCGA CCA cohorts, determined using Spearman's correlation analysis.

Figure 2. ZEB1 gain/loss of function in *in vitro* models derived from CCA cell lines. **A.** Schematic representation of the generation of gain/loss of function cellular models derived from HuCCT1 (which express negligible levels of ZEB1), SK-ChA-1 cells (which express high levels of ZEB1) and HuCCT1 Resistant (R) cells to erlotinib (which express high levels of ZEB1 as cause of the resistance). **B.** Representative phase contrast pictures showing the mesenchymal or epithelial morphology. **C.** Changes in ZEB1 mRNA and protein expression, as determined by RT-QPCR and western blot. Values are expressed as means $\pm$ SEM from 5 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, ****, $p < 0.0001$, as compared with control cells or shControl clone 1. #####, $p < 0.0001$, as compared with shControl clone 2.

Figure 3. ZEB1 expression promotes the acquisition of an EMT/CSC phenotype in CCA cells. **A-C.** Changes in *CDH1* (E-cadherin), *ESRP1* and *VIM* (vimentin) mRNA and protein expression and localization in HuCCT1 control cells or cells overexpressing Zeb1, as determined by RT-QPCR, western blot and immunofluorescence, respectively. Scale: 50 μ m. **D-E.** Changes in CD44 and CD24 mRNA and protein expression in the same cells, as determined by RT-QPCR (D) and by flow cytometry (E). Quantification in the right panel shows the % of cells in the lower right quadrant corresponding to a CD44⁺/CD24⁻ population. **F.** Cell tracking analyzed by time-lapse microscopy during 24 h. Tracks of 8 representative cells for each condition are shown. Bar graphs show quantification of the distance travelled by the cells. **G-H.** Colony (G) and sphere (H) formation assay after 7 days in culture. Representative pictures of colonies and spheres are shown. Scale: 100 μ m. Values are expressed as means $\pm$ SEM from at least 3 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, ****, $p < 0.0001$, as compared with Control cells.

Figure 4. ZEB1 expression promotes tumorigenicity and stroma expansion in CCA xenografts. **A.** Tumor volume of mice bearing HuCCT1 Control (white) or Zeb1 (black) cells. **B-D.** Representative images of tumors (B), HE (C) and Picro-Sirius Red stainings (D) from HuCCT1

control and Zeb1 xenograft tumors. Scale: 50 μ m. Quantification of collagen positive area was analyzed with ImageJ. **E.** mRNA expression of mouse *Acta2* (α -SMA), *Col1a1* and *Col4a1* in xenograft tumors derived from HuCCT1 Control and Zeb1 cells. **F-G.** mRNA expression of *TGFB1*, *CCN2* and *PDGFB* in tumor cells (corresponding to human genes) in xenograft tumors derived from HuCCT1 Control and Zeb1 cells (F), or HuCCT1 WT and Resistant (R) cells (G). Values are expressed as means $\pm$ SEM from 5 animals. *, $p < 0.05$; **, $p < 0.01$; comparing with control or WT tumors. **H-I.** Changes in CTGF protein expression in xenograft tumors derived from HuCCT1 Control and Zeb1 cells (H), or HuCCT1 WT and R cells (I), as determined by western blot.

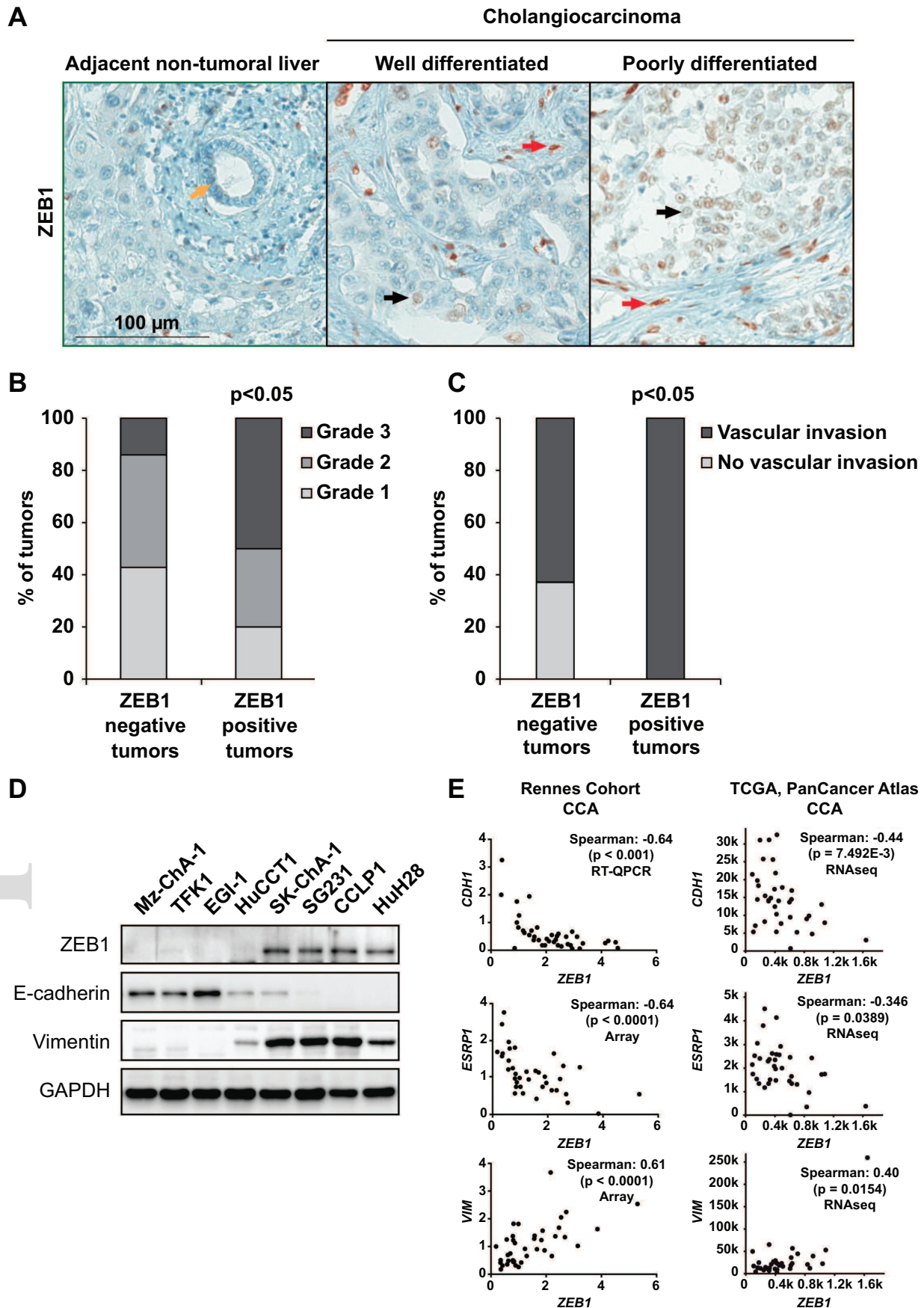
Figure 5. ZEB1 regulates the expression of CCN2/CTGF in CCA. **A.** *CCN2* (CTGF) mRNA and protein expression in gain/loss of function models derived from SK-ChA-1, HuCCT1 and HuCCT1 Resistant (R) cells. Results are expressed as means $\pm$ SEM from 5 independent cultures. **B.** Correlation between *CCN2* and *ZEB1* expression in the Rennes and the TCGA CCA cohorts. **C.** Effect of recombinant CTGF 100 ng/ml on the viability of hTERT-HSC cells. **D.** Effect on the viability of hTERT-HSC cells of conditioned media (CM) from HuCCT1 Control (white bar) and CM from HuCCT1-Zeb1 cells in absence (black) or presence of an unspecific IgG (dark grey) or a CTGF neutralizing antibody (light grey). **E.** Chromatin immunoprecipitation PCR (ChIP-PCR) experiment using primers targeting *CCN2* promoter in HuCCT1 Control (white bars) and Zeb1 (black bars) cells. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, as compared with Control cells, shRNA-Control clone 1 or vehicle. ##, $p < 0.01$; ###, $p < 0.001$, as compared with shRNA-Control clone 2.

Figure 6. ZEB1 expression in cancer-associated fibroblasts impacts their activation status. **A.** Representative IHC co-immunostaining of ZEB1 (brown) and α -SMA (pink) in human iCCA showing ZEB1 expressing CAF. **B.** Correlation between *ACTA2* (α -SMA), *COL4A1*, *TGFB1* and *CCN2* (CTGF) with *ZEB1* expression in microdissected stroma from 10 iCCA samples. **C-D.** mRNA expression of *Zeb1* in the stromal compartment (corresponding to murine genes) in xenograft tumors derived from HuCCT1 Control and Zeb1 cells (C), or from HuCCT1 WT and Resistant (R) cells (D). **E.** Changes in mRNA and protein expression of ZEB1, determined by RT-QPCR and WB in hTERT-HSC shControl or shZEB1 cells. **F.** mRNA expression of *TGFB1*, *CCN2* (CTGF), *PDGFB*, *ACTA2* (α -SMA) and *COL4A1*, in hTERT-HSC shControl or shZEB1 cells. Results are expressed as means $\pm$ SEM from 5 animals or 5 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, as compared with Control tumors, WT tumors or shControl cells.

Figure 7. Depletion of ZEB1 in myofibroblasts reduces CCA progressiveness. **A.** Tumor volume of mice bearing EGI-1+hTERT-HSC-shControl or EGI-1+hTERT-HSC-shZEB1 cells. Representative images of tumors are shown. **B.** mRNA expression of *Zeb1*, *Acta2* (α -Sma),

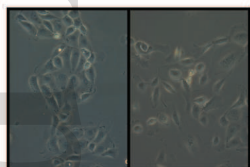
Col4a1, *Tgfb1*, *Ccn2* (Ctgf) and *Pdgfb* in the stromal compartment (corresponding to murine genes). **C.** Effect of conditioned media (CM) from hTERT-HSC shControl or shZEB1 cells on the viability of EGI-1 and HuCCT1 cells. **D.** Representative image of western blot analysis of the phosphorylated and total forms of STAT3, AKT and ERK after 30 minutes of exposure to the indicated CM in EGI-1 and HuCCT1 cells. Results are expressed as means $\pm$ SEM from 5 animals or 5 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, as compared with shControl tumors or vehicle. ##, $p < 0.01$; ###, as compared with CM from hTERT-HSC shControl.

Figure 8. ZEB1 expression in tumor cells and CAF impacts their crosstalk. **A** and **C.** mRNA expression of *HGF* (A) and *IL6* (C) in HuCCT1 Control and Zeb1, HuCCT1 WT and Resistant (R) CCA cell lines, and hTERT-HSC-shRNA-Control and shRNA-ZEB1 cell lines, and their respective derived CCA xenografts. For each model, gene expression in cultured cells (*in vitro*), tumor cells (corresponding to human genes) and stromal cells (corresponding to murine genes) of the xenografts (*in vivo*) is shown. **B.** mRNA expression of *HGF* in hTERT-HSC cells exposed to conditioned media (CM) from HuCCT1 Control or Zeb1 cells. **D.** mRNA expression of *ZEB1* and *IL6* in hTERT-HSC-shRNA-Control and shRNA-ZEB1 cells after 24 h after exposure to TGF- β 1 10 ng/ml or the vehicle. **E-F.** Effect of HGF (F) and IL6 (G) on the viability of CCA and fibroblast cell lines. **G.** Schematic diagram depicting the major findings of the study. Overexpression of ZEB1 leads to induction of EMT and stemness in tumor cells and to fibroblast activation. Furthermore, ZEB1 at both sides of the tumor-stroma interface, regulates the production of soluble signaling molecules (such as CTGF, HGF and IL6) to boost tumor progression. Values are expressed as means $\pm$ SEM from 5 independent cultures/animals. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$; ****, $p < 0.0001$, as compared with Control cells, Control tumors or the vehicle.

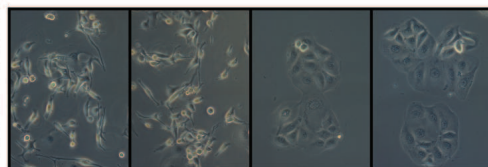


A

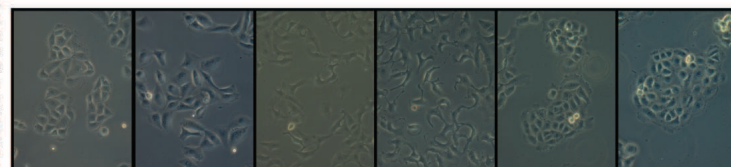
Overexpression model

HuCCT1
(Low ZEB1)Retroviral
Infection
(murine
Zeb1)HuCCT1
Control
(Low ZEB1)HuCCT1
Zeb1
(High Zeb1)

Downregulation model

SK-ChA-1
(High ZEB1)Lentiviral
Infection
(shRNA)Sk-ChA-1 shControl
Clones 1&2
(High ZEB1)Sk-ChA-1 shZEB1
Clones 1&2
(Low ZEB1)

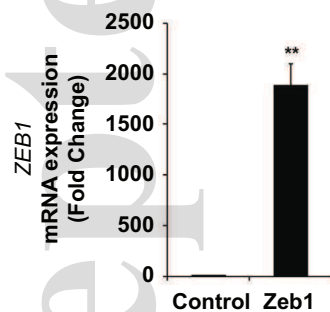
Overexpression & downregulation model

HuCCT1
(Low ZEB1)Vaquero J et al. 2018
(Natural ZEB1
overexpression)Lentiviral Infection
(shRNA)HuCCT1
WT
(Low ZEB1)HuCCT1
Resistant
(High Zeb1)HuCCT1 Resistant
shControl Clones 1&2
(High ZEB1)HuCCT1 Resistant
shZEB1 Clones 1&2
(Low ZEB1)

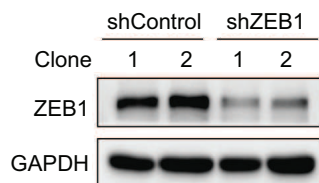
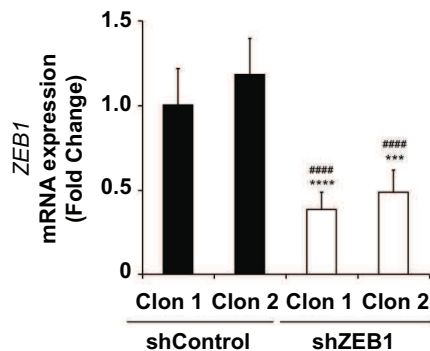
B

C

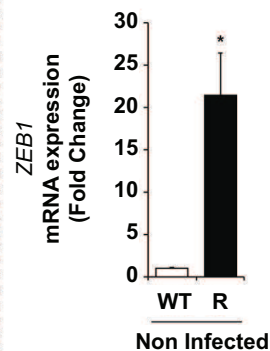
HuCCT1



SK-ChA-1



HuCCT1



HuCCT1 Resistant

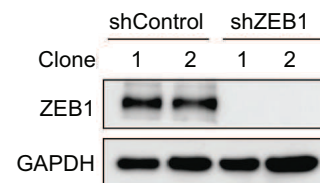
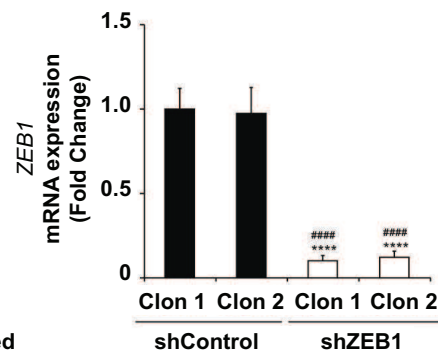


Figure 2

hep_32069_f2.eps

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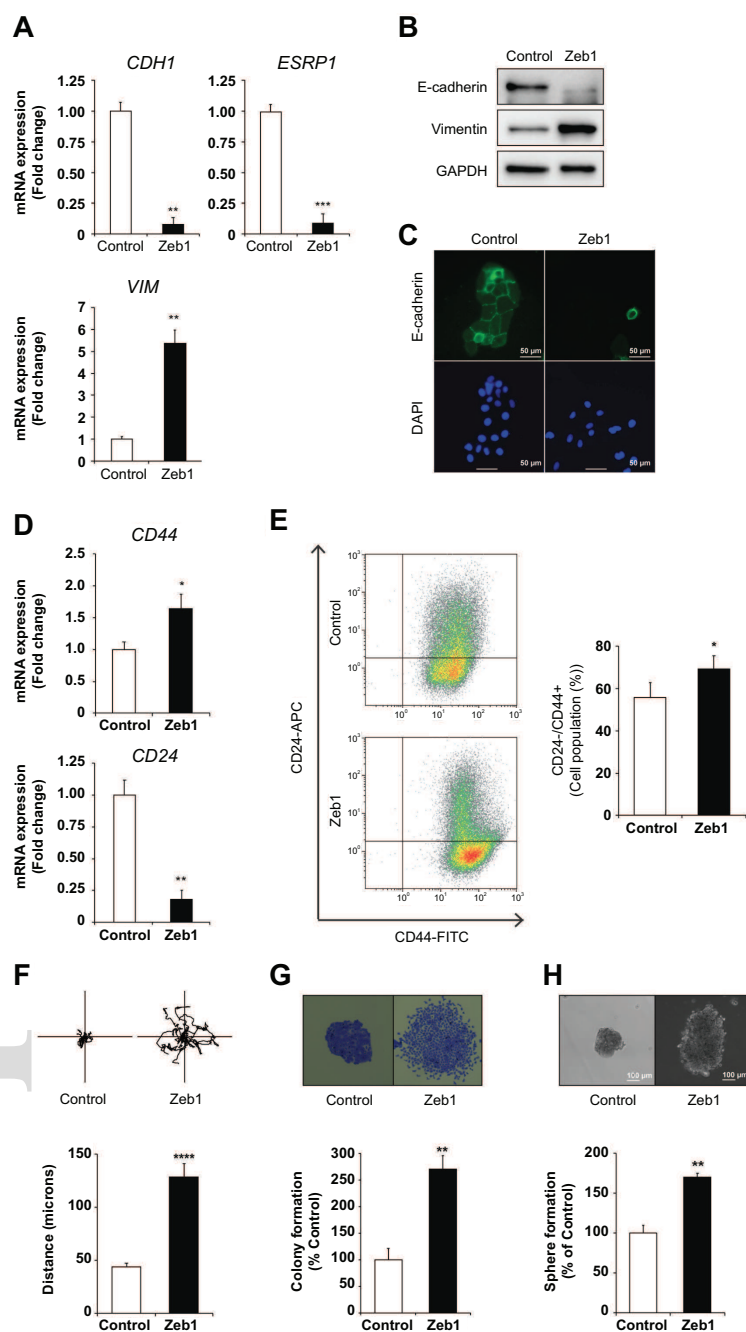


Figure 3

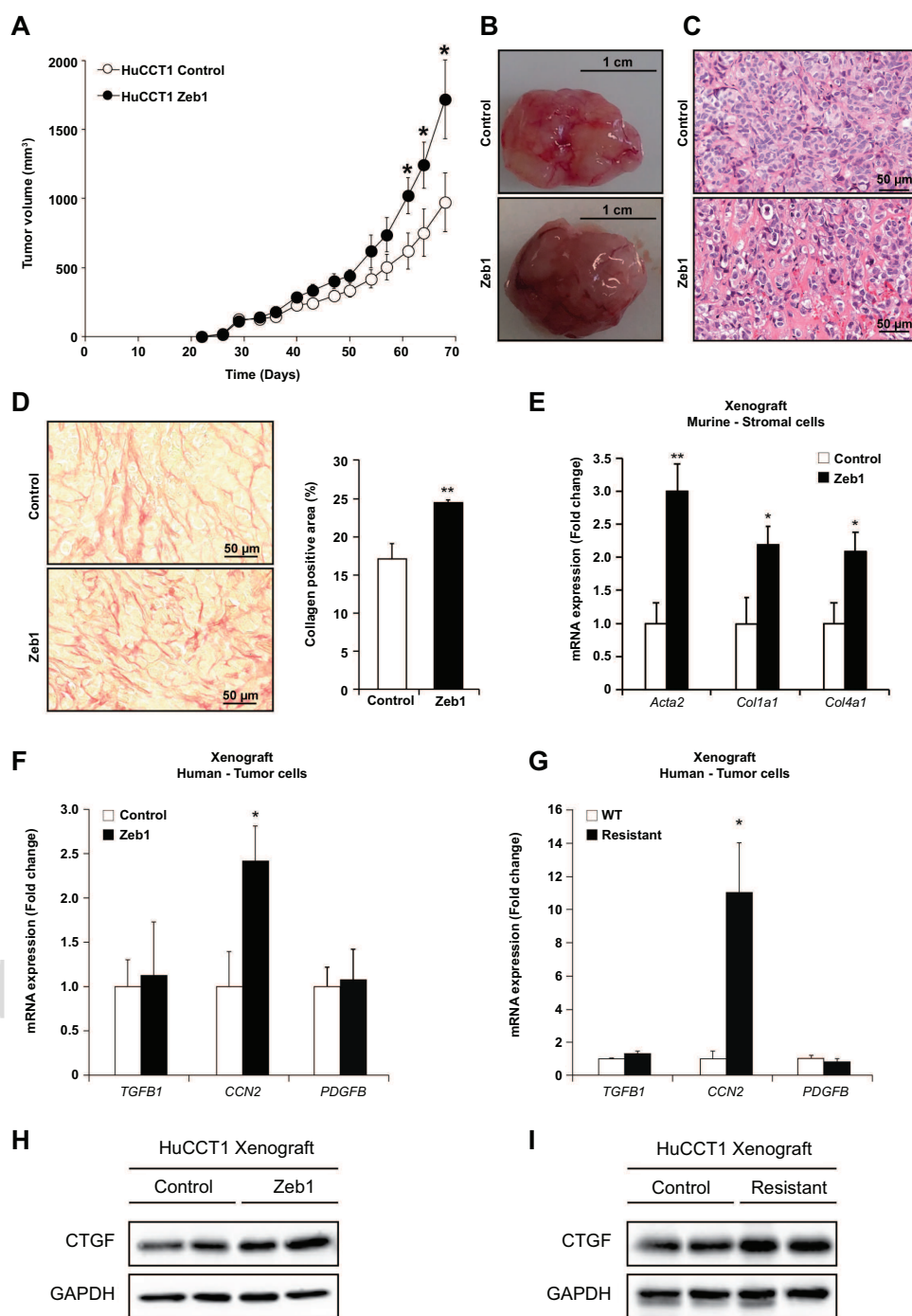


Figure 4

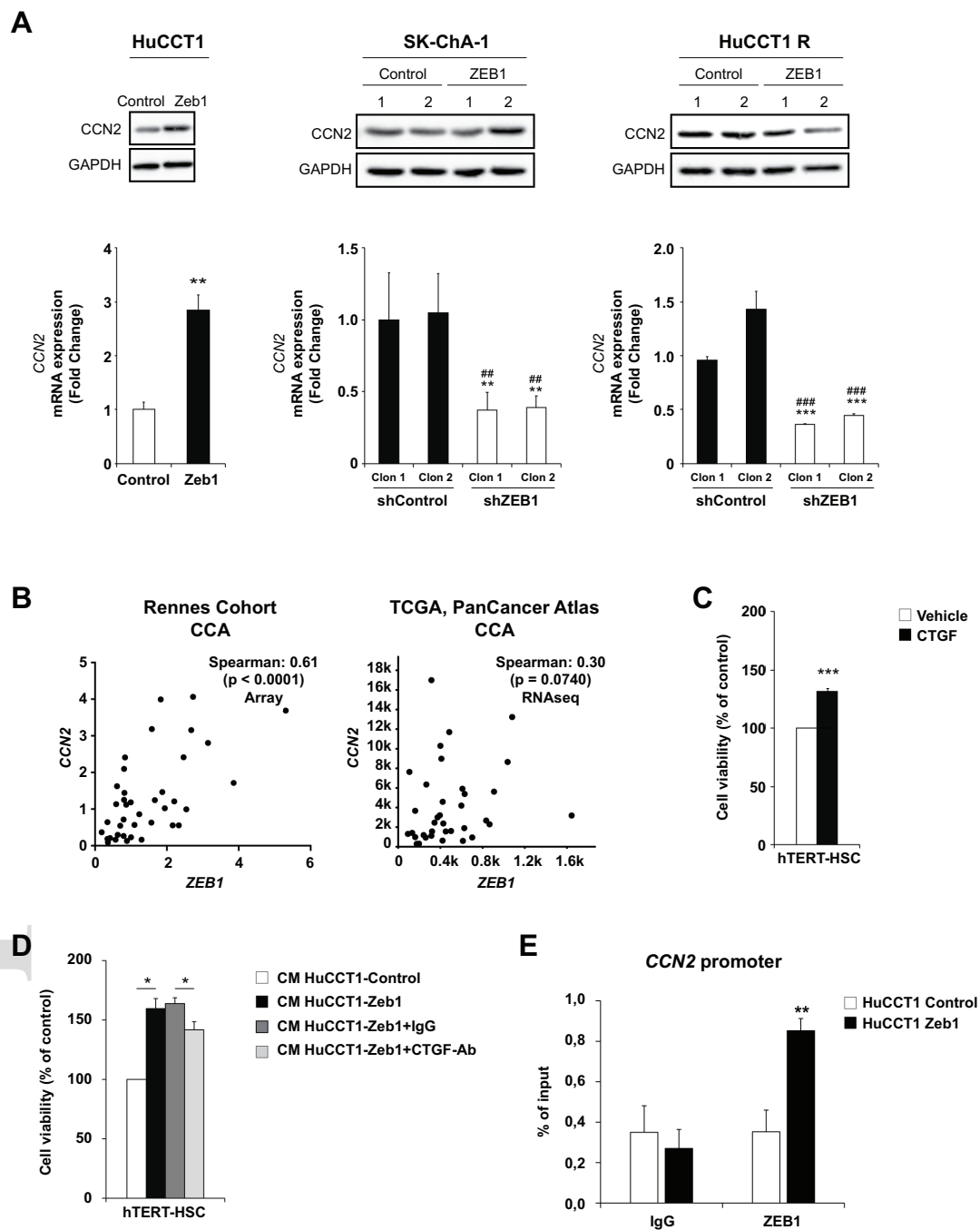


Figure 5

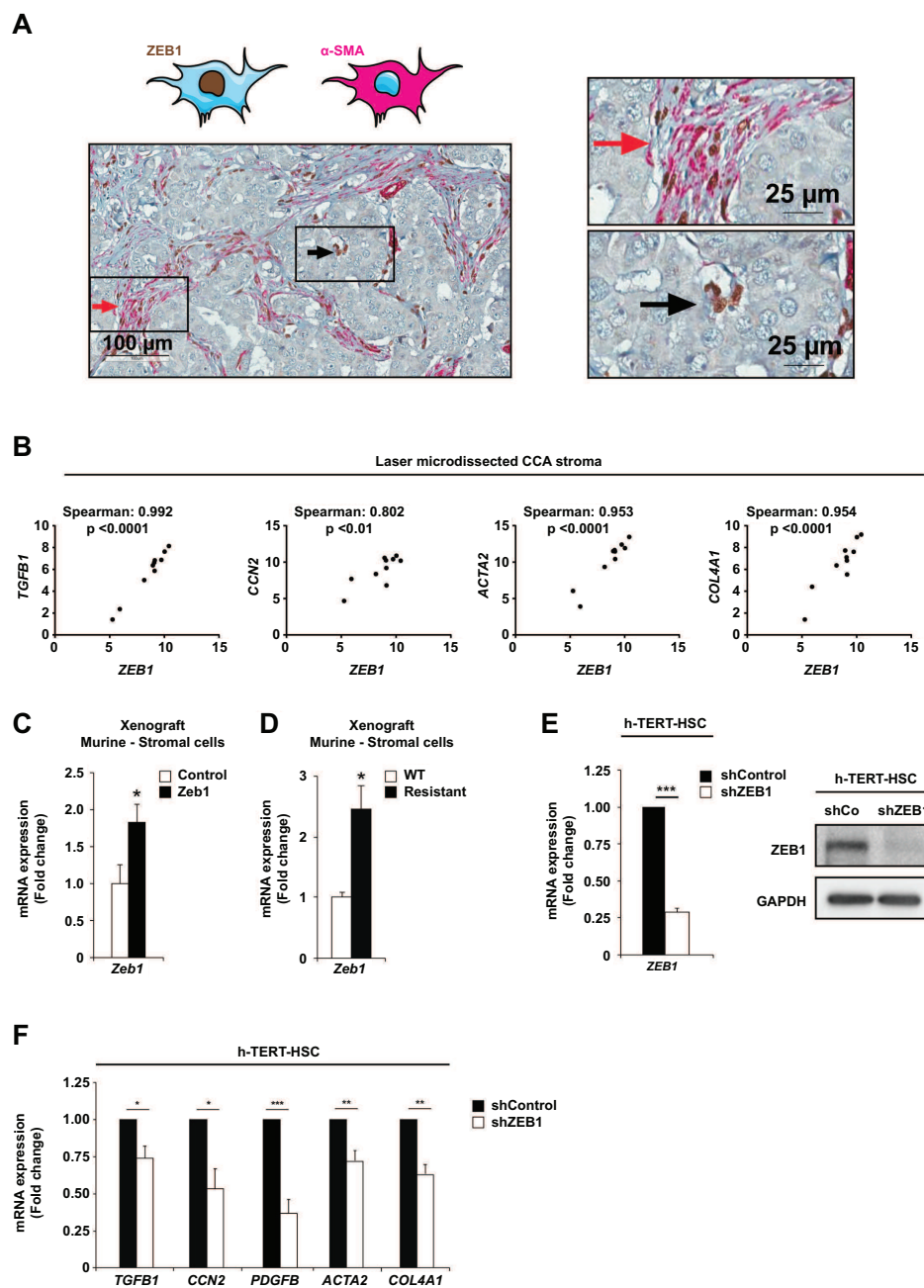


Figure 6

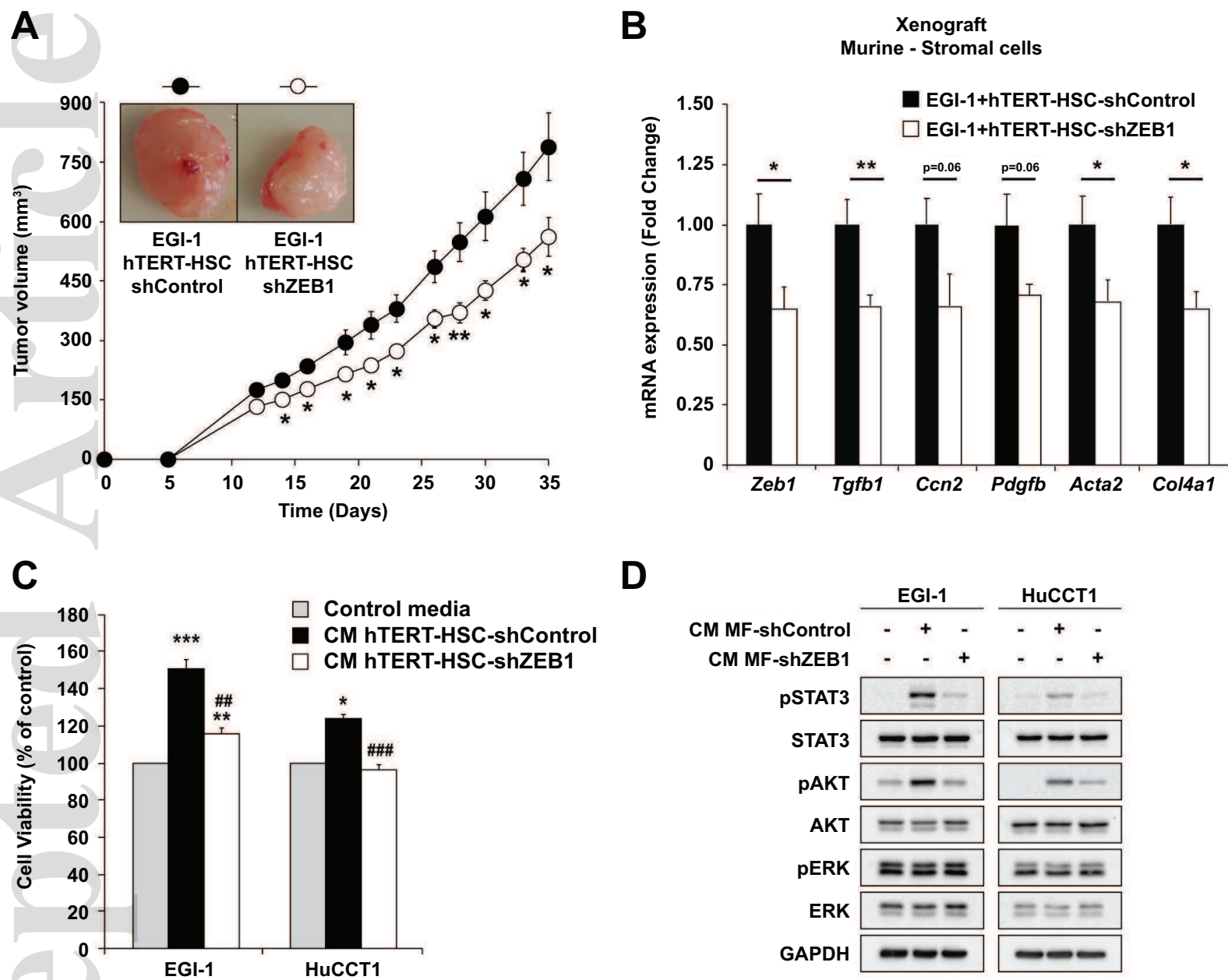


Figure 7

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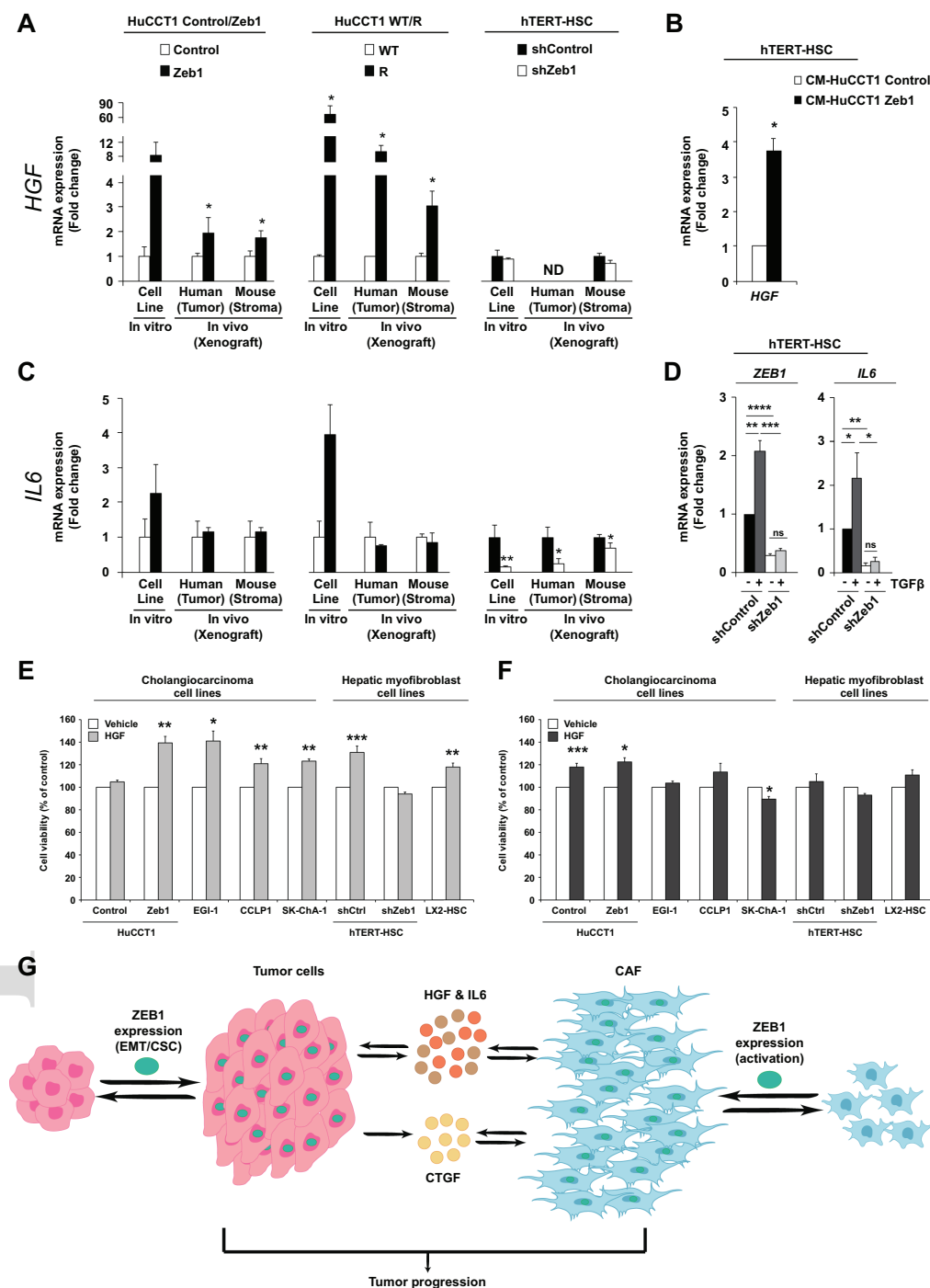


Figure 8

Supplementary information

ZEB1 promotes cholangiocarcinoma progression through tumor dedifferentiation and tumor-stroma paracrine signaling

Cindy Lobe^{1,*}, Marie Vallette^{1,*}, Ander Arbelaiz¹, Ester Gonzalez-Sanchez^{1,2,3}, Laura Izquierdo^{3,4}, Anna Pellat¹, Nathalie Guedj⁶, Corentin Louis⁷, Valérie Paradis⁶, Jesus M Banales^{3,4,5}, Cédric Coulouarn⁷, Chantal Housset^{1,8}, Javier Vaquero^{1,2,3,9,#,§} & Laura Fouassier^{1,#,§}

1-Sorbonne Université, Inserm, Centre de Recherche Saint-Antoine (CRSA), Paris, France.

2-TGF- β and Cancer Group, Oncobell Program, Bellvitge Biomedical Research Institute (IDIBELL), Barcelona, Spain.

3-National Biomedical Research Institute on Liver and Gastrointestinal Diseases (CIBERehd), Instituto de Salud Carlos III, Madrid, Spain.

4-Department of Liver and Gastrointestinal Diseases, Biodonostia Health Research Institute - Donostia University Hospital, University of the Basque Country (UPV/EHU), San Sebastián, Spain.

5-IKERBASQUE, Basque Foundation for Science, Bilbao, Spain.

6-Service d'Anatomie Pathologique Hôpital Beaujon, F-92110 Clichy, France; INSERM, UMR 1149, Centre de Recherche sur l'Inflammation, F-75018 Paris, France.

7-Inserm UMR 1241, Univ Rennes 1, Rennes, France.

8- Assistance Publique-Hôpitaux de Paris (AP-HP), Saint-Antoine Hospital, Reference center for inflammatory biliary diseases and autoimmune hepatitis, ERN Rare-Liver, Paris, France.

9- LPP, Sorbonne Université, CNRS, Ecole Polytech., Univ. Paris-Sud, Observatoire de Paris, Université Paris-Saclay, PSL Research University, 4 Place Jussieu, 75252 Paris, France.

* Share co-first authorship

Share co-senior authorship

Supplementary Materials and Methods

Reagents

CTGF, EGF, HGF, IGF2 and IL6 were purchased from PreproTech.

Cell culture and treatment

HuCCT1 cells, derived from intrahepatic CCA, were kindly provided by Dr. G. Gores (Mayo Clinic, MN). SG231, CCLP1 and HuH28 cells, derived from intrahepatic CCA were kindly provided by Dr. Cédric Coulouarn (NuMeCan, Rennes, France). EGI-1 and TFK1 cells, derived from extrahepatic CCA, were obtained from the German Collection of Microorganisms and Cell Cultures (DSMZ, Germany). SK-ChA-1 and Mz-ChA-1 cells, derived from extrahepatic CCA, were obtained from Dr. A. Knuth (Zurich University, Switzerland). HuCCT1, CCLP1, EGI-1, Mz-ChA-1 and SK-ChA-1 cells were cultured in DMEM supplemented with 1 g/L glucose, 10 mmol/L HEPES. TFK-1 cells were cultured in RPMI supplemented with HCO_3^- 0.75%. SG231 cells were cultured in MEM alpha supplemented with 10 mmol/L HEPES. HuH28 were cultured in MEM. All culture medias were supplemented with glutamax, 10% fetal bovine serum (FBS), antibiotics (100 UI/mL penicillin and 100 mg/mL streptomycin), and antimycotic (0.25 mg/mL amphotericin B; Invitrogen). The human hepatic stellate cell (HSC) lines hTERT-HSC and LX2-HSC, were kindly provided by Dr. L. Aoudjehane (ICAN, Paris, France). hTERT-HSC and LX2-HSC were cultured in DMEM supplemented with 4.5 g/L glucose, antibiotics (100 UI/mL penicillin and 100 $\mu\text{g/mL}$ streptomycin), antimycotic (0.25 mg/mL amphotericin B), and 10% FBS and 2% FBS, respectively. Cell lines were routinely screened for the presence of mycoplasma and authenticated for polymorphic markers to prevent cross-contamination.

CCN2 RNA silencing

CCN2 expression was silenced by using a pool of four small interfering RNAs (siRNA) (On-Targetplus SMARTpool, Dharmacon). Control siRNA was a pool from Dharmacon (siGENOME nontargeting siRNA Pool 2). Transient transfections were performed with 100 nM siRNA using DharmaFECT 4 (Thermo Scientific). Cells were plated in 6-well plates and after 24 h they were

transfected with the corresponding siRNA. 24 h after transfection cell culture media was changed to remove transfection complexes. Then, the appropriate protocol for each experiment was followed.

Conditioned media

HuCCT1-Control, HuCCT1-Zeb1, hTERT-HSC-shControl and hTERT-HSC-shZEB1 cells were grown to confluence, washed twice with phosphate-buffered saline (PBS), and 0% FBS DMEM (supplemented as above) was added and collected after 48h. Conditioned media were centrifuged to remove cell debris and stored at -80°C until use. 0% FBS DMEM was used as control.

Gene expression profiling

Gene expression datasets established from whole(1) or laser microdissected(2) CCA tumors were previously described. Clinical and pathological features of a cohort of 39 patients with iCCA (thereafter referred to as the Rennes cohort) were previously reported.(1). Freshly frozen tumor samples were obtained through the French liver cancer biobanks network – INCa (BB-0033-00085). Of note, because the probe designed for *CDH1* (A_23_P206359) was unreliable providing poorly reproducible data, we performed a RT-qPCR analysis using validated pairs of primers for *CDH1*, *ZEB1* and *TBP* (housekeeping gene) (Supplementary Table 1) to evaluate the correlation between these genes in the Rennes cohort.

Comparison of mRNA expression from public databases

The mRNA expression levels of *ZEB1* and selected markers (*CDH1*, *VIM*, *CCN2*, *ESRP1*, *IGF2*, *EGF*, *HGF* and *IL6*) in human cancer cell lines derived from the Broad-Novartis Cancer Cell Line Encyclopedia (CCLE) and in tumors (breast invasive carcinoma, hepatocellular carcinoma, head and neck squamous cell carcinoma, and, colorectal, lung, pancreatic and prostate adenocarcinomas) from the repository of The Cancer Genome Atlas (TCGA) were analyzed through the cBioPortal database (<https://www.cbioportal.org/>) and represented as Spearman correlation coefficients (r) and P-values to measure the significance.(3-5)

RNA and reverse transcription-PCR

Total RNA extraction and RT-qPCR were performed as previously described.⁽⁶⁾ Primer sequences are provided in Supplementary Tables 1, 2 and 3. Gene expression was normalized to *GAPDH* mRNA content for human genes and to *Hprt* mRNA content for mouse genes and was expressed relatively to the control condition of each experiment. The relative expression of each target gene was determined from replicate samples using the formula $2^{-\Delta\Delta C_t}$.

Chromatin immunoprecipitation PCR (ChIP-PCR) assays

ChIP assays were performed using the Chromatin Immunoprecipitation Assay Kit from Milipore (#17-295). 4×10^6 cells were plated in a T75 for 24h and sonicated with Bioruptor sonicator (Diagenode) for 20 cycles of 30 seconds sonication and 30 seconds cooling. ChIP was done with 2 μ g of ZEB1 Antibody from NovusBiologicals (NBP1-05987). Internal control has been done with a normal rabbit IgG (Cell Signaling Technologies) (CST2729). ChIP DNA product and input DNA, extracted by phenol chloroform method, were suspended in 20 μ L and Input were diluted 1 to 100, to perform the qPCR (2 μ L of sample). Primer sequences are provided in Supplementary Table 4.

Immunoblot analysis

For obtaining whole-cell lysates for immunoblotting, cell cultures were lysed in RIPA buffer (Sigma) supplemented with 1 mmol/L orthovanadate and a cocktail of protease inhibitors. Proteins were quantified using a BCA kit (Pierce). Western blot analysis was performed as previously described.⁽⁶⁾ Primary antibodies are provided in Supplementary Table 5.

Immunofluorescence

Immunofluorescence assays were performed as previously described.⁽⁶⁾ Primary antibodies are provided in Supplementary Table 5. Cells were observed with an Olympus Bx 61 microscope (Olympus).

Viability assays

Cells were plated in 96-well plates. Twenty-four hours later, the medium was replaced by fresh culture medium in the absence (vehicle) or presence of the corresponding growth factors or conditioned media (CM) prepared from HuCCT1 or hTERT-HSC cells. For neutralization experiments unspecific IgG or a neutralizing antibody against CTGF (10 µg/ml) (PreproTech 500-P252) were added to the CM of HuCCT1-Zeb1 cells. Cells were then incubated for 72 h before determining the viability by the crystal violet method. Absorbance was quantified with a spectrophotometer (Tecan) at 595 nm.

Migration assay

Cell motility was evaluated by time-lapse microscopy. Cells were plated in 24-well plates (TPP) and cultured for 24h. Then, medium was replaced for fresh culture medium and images were collected every 20 min for 24h with an Olympus Bx 61 microscope (Olympus). During the entire experiment, cells were kept at 37°C under 5% CO₂. 8 cells were randomly selected in the first frame of the videos and then, their movement was followed in all the frames from 0 to 24h. Cell trajectories were traced using the center of the nucleus as a reference point and were used to calculate the mean speed of the cells with ImageJ software.

Flow cytometry analysis

For the determination of the CD44/CD24 phenotype, cells were washed with PBS, detached with trypsin incubation, and resuspended in PBS supplemented with 0.5% BSA. Combinations of fluorochrome-conjugated monoclonal antibodies against human CD44 (FITC) and CD24 allophycocyanin (APC) were obtained from Beckman Coulter. Specific antibodies or the respective isotype controls were added to the cell suspension, as recommended by the manufacturer, and incubated at 4°C for 20 minutes. Cells were washed with PBS containing 0.5% BSA, centrifuged, and resuspended in PBS with 2% paraformaldehyde. The labelled cells were analyzed with a Gallios flow cytometer (Beckman-Coulter).

Colony formation assay

Cells were plated in 6-well plates at a density of 500 cells/well and then they were cultured for 7 days. Colonies were fixed with 4% paraformaldehyde, stained with 1% crystal violet and counted. Only colonies with more than 50 cells were counted.

Sphere formation assay

Cells were plated in 6-well ultralow attachment plates (Greiner Bio One) in serum-free DMEM/F12 medium, supplemented with 100 mg/mL gentamycin (Sigma-Aldrich), B27 (Life Technologies), 20 ng/mL human epidermal growth factor (EGF, Life Technologies), 20 ng/mL human basic fibroblast growth factor (bFGF, Life Technologies), and 1% antibiotic–antimycotic solution (Life Technologies) at a density of 4×10^4 cells/well. Fresh sphere medium was added to plates after 4 days. After 7 days, spheres were counted on EVOS FL Cell Imaging System (Thermo Fisher Scientific).

Xenograft tumor model

Animal experiments were performed in accordance with the French Animal Research Committee guidelines and all procedures were approved by a local ethic committee (APAFIS#14678-2018041510038914). In a first set of experiments, HuCCT1-Control and HuCCT1-Zeb1 cells (5×10^6) suspended in 60 mL of PBS were mixed with 60 mL of Matrigel growth factor reduced (Corning) and implanted subcutaneously into the flank of 5-week-old female NMRI-nu (nu/nu) mice (Envigo). In a second set of experiments EGI-1+hTERT-HSC-shControl or EGI-1+hTERT-HSC-shZEB1 (2×10^6 in a ratio 1:1) were injected into the animals following the same protocol. Mice were housed under standard conditions in individually ventilated cages enriched with a nesting material and kept at 22°C on a 12-h light/ 12-h dark cycle with ad libitum access to food and tap water. Tumor growth was followed with a caliper, and tumor volume (V) was calculated as follows: xenograft volume = $xy^2/2$, where x is the longest and y, the shortest of two perpendicular diameters. During the experiments, animals showed no sign of toxicity, such as body weight loss (>15%), decreased food intake, or

diarrhea. The day of the sacrifice, a slice of the tumor was fixed in 10% formalin and embedded in paraffin, and the rest was cut in pieces and frozen in liquid nitrogen.

CCA specimens

We retrospectively retrieved from the files of surgical and pathology department, cases of patients who had undergone liver resection for intrahepatic CCA between 2002 and 2014 at Beaujon Hospital, Clichy, France. All clinicopathological and follow up data of patients were collected and registered in data base (Supplementary Table 6). Informed consent was obtained in all cases.

Immunohistochemical analysis on Tissue MicroArray (TMA)

Paraffin embedded tissue blocks were used for tissue microarray construction. The slides were reviewed to identify and mark representative areas of viable tumor tissue. Taking tumor heterogeneity into account, three tissue cores of 1mm each were punched from selected from selected tumor areas of any donor tissue block and brought into a recipient paraffin block. We used a tissue arraying instrument (MTA-1, Beecher Instrument, Inc., Sun Prairie, WI, USA). A total of 5 TMA were built.

Immunohistochemistry against ZEB1 and smooth muscle actin (α -SMA) (Supplementary Table 5) was realized. Immunohistochemical study was performed on TMA using an automated immunohistochemical stainer according to the manufacturer's guidelines (streptavidin-peroxidase protocol; Benchmark, Ventana, Tucson, AZ). All sections were evaluated by pathologists (NG and VP) using a semi-quantitative analysis. ZEB1 was considered positive when nuclear staining in tumor cells was observed whatever the intensity. α -SMA was considered positive when CAF were immunostained.

We also performed a co-immunostaining between ZEB1 and α -SMA according the ultraview double immunostained protocol (optiview DAB/ultraview RED).

Statistical analysis

Results were analyzed using the GraphPad Prism 5.0 statistical software. Data are shown as means $\pm$ standard error of the mean (SEM). For comparisons between two groups, parametric Student t test or nonparametric Mann–Whitney test were used. For comparisons between more than two groups, parametric one-way ANOVA test followed by a posteriori Bonferroni test was used. Histopathological parameters were statistically analyzed Chi-square test.

Supplementary References.

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Supplementary Table 1. Human primers used for determination of *CDH1* and *ZEB1* correlation in the Rennes cohort by quantitative real-time PCR.

Gene	Protein	Forward (5'→3')	Reverse (5'→3')	Accession Number
<i>CDH1</i>	E-cadherin	CGAGAGCTACACGTTACGG	GGGTGTCGAGGGAAAAATAGG	NM_004360.4
<i>TBP</i>	TBP	GAGCTGTGATGTGAAGTTTCC	TCTGGGTTTGATCATTCTGTAG	NM_003194.5
<i>ZEB1</i>	ZEB1	CCCACACGACCACAGATACG	TGAGGAGAACTGGTTGCCTG	NM_001128128.2

Supplementary Table 2. Mouse primers used for quantitative real-time PCR.

Gene	Protein	Forward (5'→3')	Reverse (5'→3')	Accession Number
<i>Acta2</i>	α -SMA	CTGTCAGGAACCCTGAGACGCT	TACTCCCTGATGTCTGGGAC	NM_007392.3
<i>Adgre1</i>	F4/80	CTTTGGCTATGGGCTTCCAGTC	GCAAGGAGGACAGAGTTTATCGTG	NM_010130.4
<i>Ccl2</i>	Mcp1	GCCTGCTGTTACAGTTGC	CAGGTGAGTGGGGCGTTA	NM_011333
<i>Ccn2</i>	Ctgf	GGGCCTCTTCTGCGATTTC	ATCCAGGCAAGTGCATTGGTA	NM_010217.2
<i>Col1a1</i>	Collagen I	GAAACCCGAGGTATGCTTGA	GACCAGGAGGACCAGGAAGT	NM_007742.4
<i>Col4a1</i>	Collagen IV	ATTTCCAGGCGTGTCATTGC	AGCTCTCTCCTTTCTGACCTT	NM_009931.2
<i>Egf</i>	Egf	TGCCTCAGAAGGAGTGGGTTA	GTGTTCCAAGCGTTCCTGAGA	NM_010113.4
<i>Hgf</i>	Hgf	ATGTGGGGGACCAAACCTTCTG	GGATGGCGACATGAAGCAG	NM_001289458.1
<i>Hprt</i>	Hprt	TCAGTCAACGGGGGACATAA	TGCTTAACCAGGGAAAGCAAA	NM_013556.2
<i>Igf2</i>	Igf2	GCATCGTGGAAGAGTGCTG	GGGTATCTGGGGAAGTCGTC	NM_001315489.1
<i>Il6</i>	Il6	TAGTCCTTCCTACCCCAATTTCC	TTGGTCCTTAGCCACTCCTTC	NM_031168.2
<i>Pdgfb</i>	Pdgfb	CATCCGCTCCTTTGATGATCTT	GTGCTCGGGTCATGTTCAAGT	NM_011057.4
<i>Pecam1</i>	Cd31	AGCCTCCAGGCTGAGGAAAA	GATGTCCACAAGGCACTCCA	NM_001305158.1
<i>Tgfb1</i>	Tgfb1	CTCCCGTGGCTTCTAGTGC	GCCTTAGTTTGGACAGGATCTG	NM_011577.2
<i>Vwf</i>	Vwf	GCTGCAGTTATCTCCTGGCT	ACAGGCTCATTCTCTTGCCA	NM_011708.4
<i>Zeb1</i>	Zeb1	CAGATTCCGATGATGAAGACAAAC	TCTTTCACGTTGTCTTGCCAGC	NM_011546.3

Supplementary Table 3. Human primers used for quantitative real-time PCR.

Gene	Protein	Forward (5'→3')	Reverse (5'→3')	Accession Number
<i>ACTA2</i>	α -SMA	GACAATGGCTCTGGGCTCTGTAA	CTGTGCTTCGTCACCCACGTA	NM_001141945.1
<i>CCN2</i>	CTGF	CAGCATGGACGTTCTGCTG	AACCACGGTTTGGTCCTTGG	NM_001901.3
<i>CD24</i>	CD24	CTCCTACCCACGCAGATTTATTC	AGAGTGAGACCACGAAGAGAC	NM_013230.3
<i>CD44</i>	CD44	TGCAGTCAACAGTCGAAGAAG	CTGTCCTCCACAGCTCCATT	NM_000610.4
<i>CDH1</i>	E-Cadherin	AAGAAGCTGGCTGACATGTACGGA	CCACCAGCAACGTGATTCTGTCAT	NM_004360.4
<i>COL1A1</i>	Collagen I	AGTTCGAGTATGGCGG	CAGTGACGCTGTAGGT	NM_000088.3
<i>COL4A1</i>	Collagen IV	CCGCTGCGAAGGGTGG	CCCTTTTGTCTGGTGGTCCC	NM_001845.5
<i>CXCL8</i>	IL8	ACTGAGAGTGATTGAGAGTGGAC	AACCCTCTGCACCCAGTTTTTC	NM_000584
<i>EGF</i>	EGF	AGTGCATCCACTTGACAAC	CAATTTGCAAAGTTTCTGCTCA	NM_001963.4
<i>EPCAM</i>	EpCAM	CCATGTGCTGGTGTGTGAA	TGTGTTTTAGTTCAATGATGATCCA	NM_002354.2
<i>ESRP1</i>	ESRP1	CCCACCGCCATGTAAGTT	GCAGGAGCTGGAAATGTGTAG	NM_017697.4
<i>GAPDH</i>	GAPDH	AGCCACATCGCTCAGACAC	GCCCAATACGACCAAATCC	NM_002046.5
<i>HGF</i>	HGF	AAAGGACTTCCATTCACTTGC	CCACTCTTAGTGATAGATACTG	NM_000601.4
<i>IGF2</i>	IGF2	CGTCGCAGCCGTGGCATCGTTGA	GCCCACGGGGTATCTGGGGAAG	NM_000612.5
<i>IL6</i>	IL6	TACATCCTCGACGGCATCTC	TCCAGATTGGAAGCATCCATCCATC	NM_000600.5
<i>IL6R</i>	IL6R	CCCCTCAGCAATGTTGTTTGT	CTCCGGGACTGCTAACTGG	NM_001206866.2
<i>IL6ST</i>	GP130	CGGACAGCTTGAACAGAATGT	ACCATCCCACTCACACCTCA	NM_002184.4
<i>ITGB3</i>	CD61	CAAGATTGGAGACACGGTGA	CGGGCTTTATGGTAAAGGACT	NM_000212.3
<i>MET</i>	MET	AGCAATGGGGAGTGTAAGAGG	CCCAGTCTTGTACTCAGCAAC	NM_000245.4
<i>SNAI1</i>	SNAIL	CCAGAGTTTACCTTCCAGCAGCC	CAGGACAGAGTCCCAGATGAGCAT	NM_005985.3
<i>PDGFB</i>	PDGFB	GGGCATCGGCAGGAGAGT	AGACAGACGGACGAGGGAAA	NM_002608.2
<i>SNAI2</i>	SLUG	ATCTCCGGAGGTGGGATG	GAGAGACATTCTGGAGAAGG	NM_003068.4
<i>TGFB1</i>	TGFB1	GCCCTGGACACCAACTATTGC	GGCTCCAAATGTAGGGGCAGG	NM_000660.5
<i>TWIST1</i>	TWIST1	AGCTACGCCTTCTCGGTCT	CCTTCTCTGGAACAATGACATC	NM_000474.3
<i>TWIST2</i>	TWIST2	GCAAGAAGTCGAGCGAAGAT	GCTCTGCAGCTCCTCGAA	NM_001271893.4
<i>VIM</i>	Vimentin	CCCTGCAATCTTTCAGACAG	ATCTGAGCCTGCAGCTCCT	NM_003380.3
<i>ZEB1</i>	ZEB1	ACAATCGTGGCCATTGCTGA	TGGTTCTTGGACTGCAGGGCT	NM_001128128.2
<i>ZEB2</i>	ZEB2	AGGGACAGATCAGCACCAAATGCTA	GGCACTCGTAAGGTTTTTCACCACTG	NM_014795.3

Supplementary Table 4. Human primers used for chromatin immunoprecipitation PCR (CHIP-PCR) experiments.

Gene	Forward (5'→3')	Reverse (5'→3')	Accession Number
<i>CCN2</i>	GCTGGAGTGTGCCAGCTTT	CGAGGCTTTTATACGCTCCG	NG_016131.1
<i>HGF</i>	GGGTAAGGGAGCTGGAGGACAA	AGAACAGTGCCCCTGTCCA	NG_016274.2

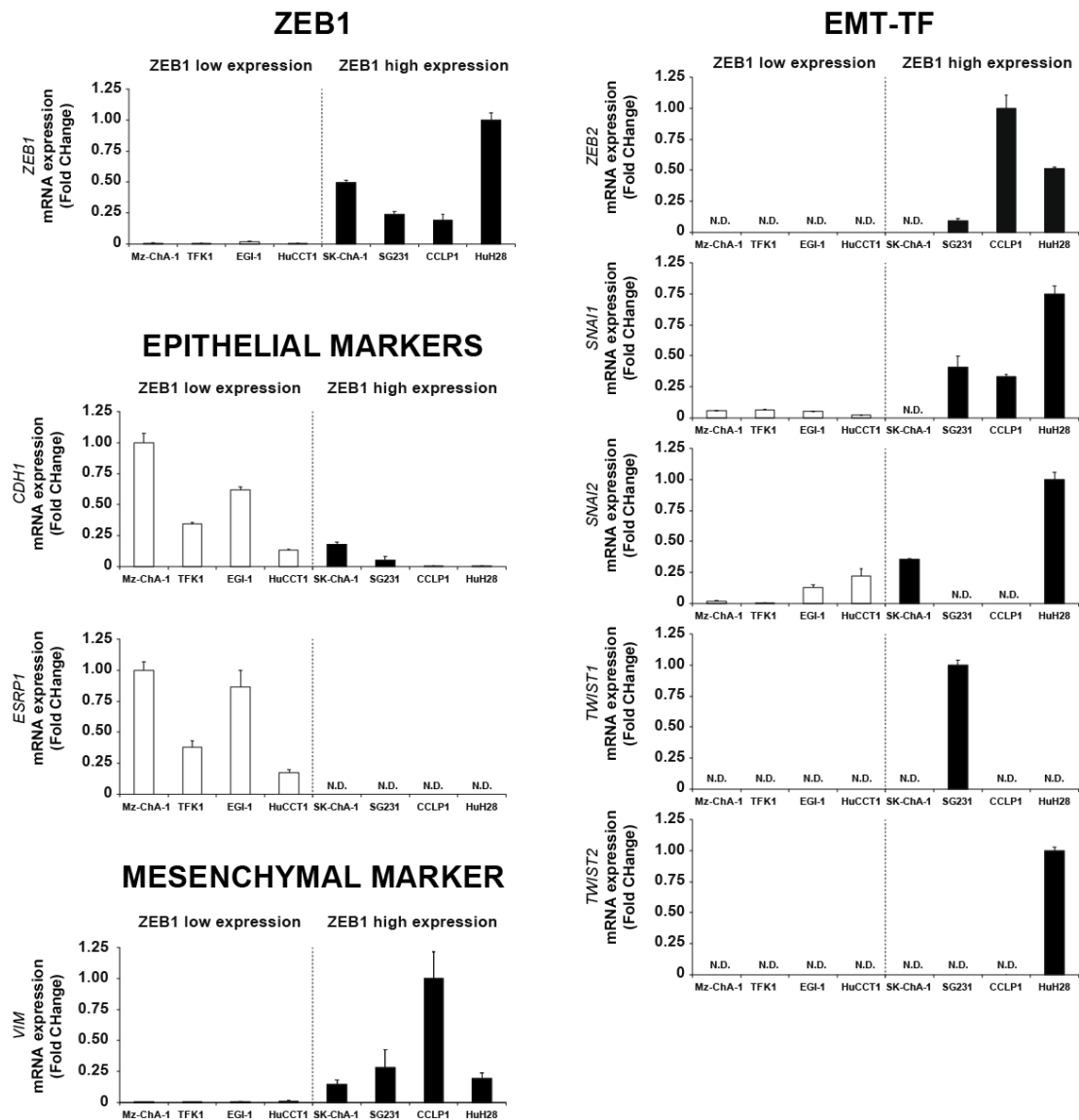
Supplementary Table 5. Primary antibodies used for immunodetection.

Name	Species	Manufacturer	Reference	Dilution	Antigen unmasking
α-SMA	M	Dako	M851	1/500 (IHC)	
pAKT	R	Cell Signaling	#4060	1/1000 (WB)	
AKT	R	Cell Signaling	#4691	1/1000 (WB)	
CTGF	M	Santa Cruz	Sc-365970	1/1000 (WB)	
E-cadherin	M	Abcam	ab1416	1:5000 (WB) / 1:250 (IF)	
pERK	M	Santa Cruz	sc-7383	1:250 (WB)	
ERK 1	R	Santa Cruz	sc-93	1:500 (WB)	
GAPDH	M	Santa Cruz	sc-32233	1:5000 (WB)	
SLUG	R	Cell Signaling	#9585	1/1000 (WB)	
SNAIL	R	Cell Signaling	#3879	1/1000 (WB)	
pSTAT3	R	Cell Signaling	#9145	1:500 (WB)	
STAT3	R	Cell Signaling	#4904	1:500 (WB)	
TWIST1	M	Santa Cruz	sc-81417	1:500 (WB)	
TWIST2	R	Abcam	ab66031	1:1000 (WB)	
Vimentin	M	Sigma	V6630	1:15000 (WB)	
ZEB1	R	Sigma	HPA027524	1/200 (IHC)	yes
ZEB1	R	Santa Cruz	sc-25388	1:500 (WB)	
ZEB2	R	Santa Cruz	sc-48789	1:500 (WB)	

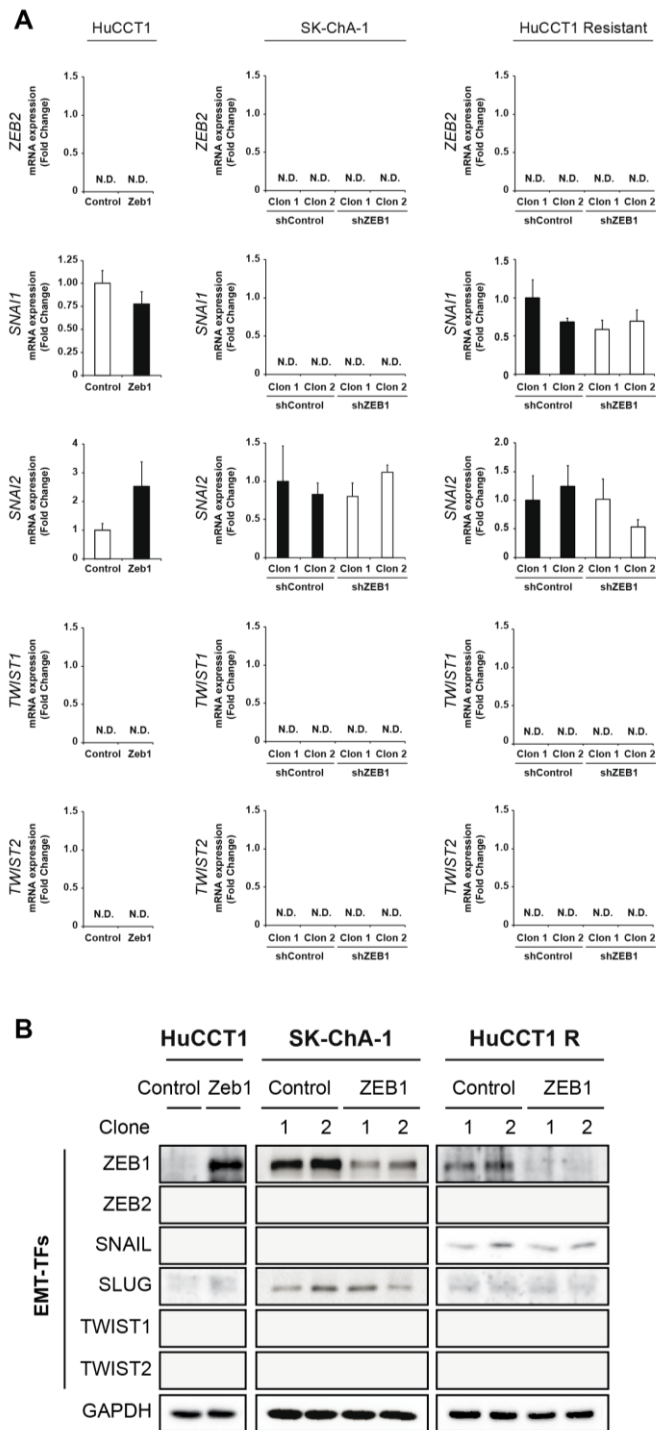
M, mouse; R, rabbit; WB, western blot; IF, immunofluorescence; IHC, immunohistochemistry.

Supplementary Table 6. Clinical and pathological characteristics of patients with CCA (n=45)

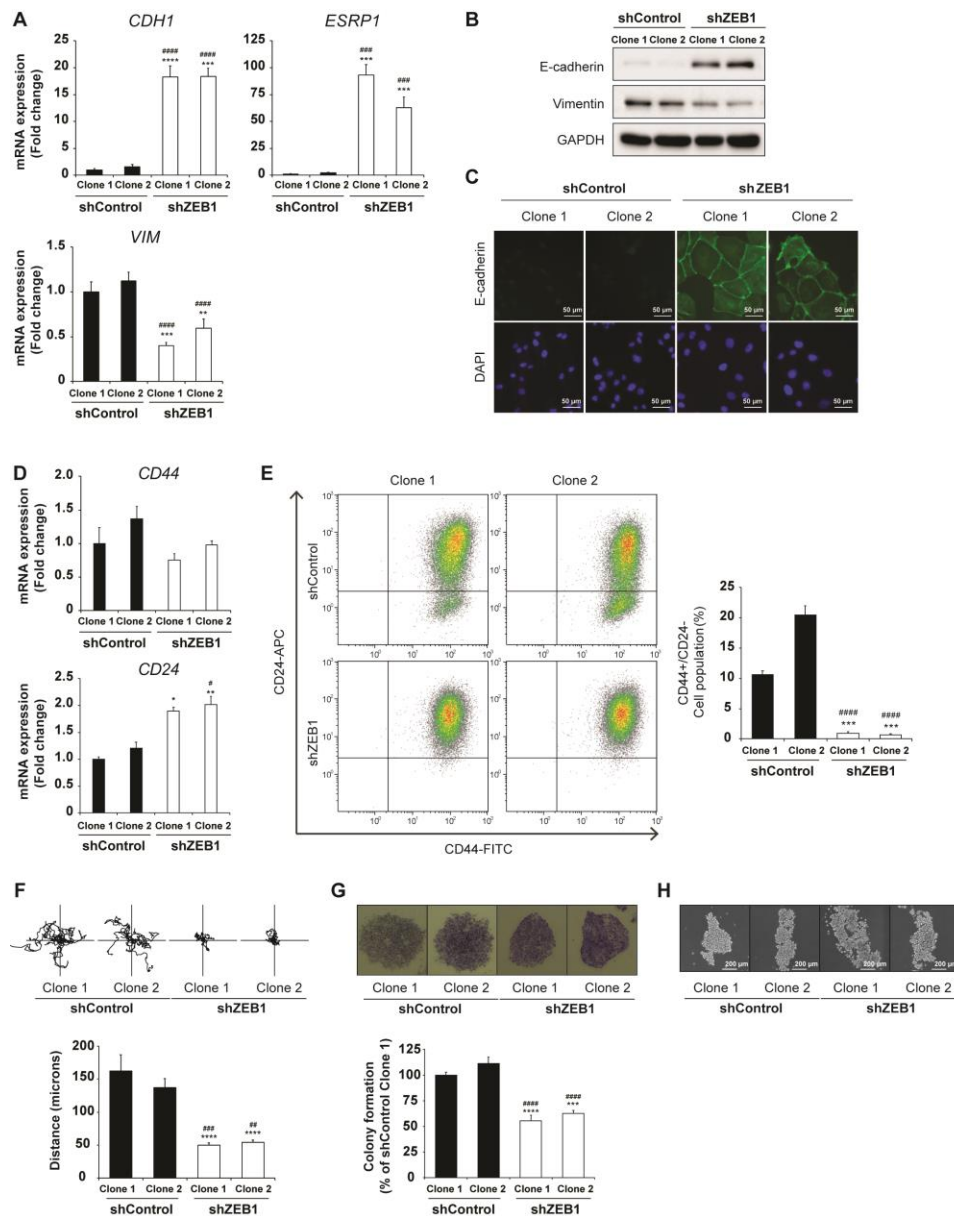
Age (years)	62 +12
Mean ($\pm$ SD)	
Sex ratio (M/F)	0.9
Tumor size (mm)	82 (15-220)
Mean ($\pm$ SD)	
Tumor grade	
Well differentiated	17/45 (38%)
Moderately differentiated	18/45 (40%)
Poorly differentiated	10/45 (22%)
pTNM (7 th edition)	
T1a	13%
T1b	9%
T2	78%
Vascular invasion	71%
Perineural invasion	31%
Lymph node metastasis	37%



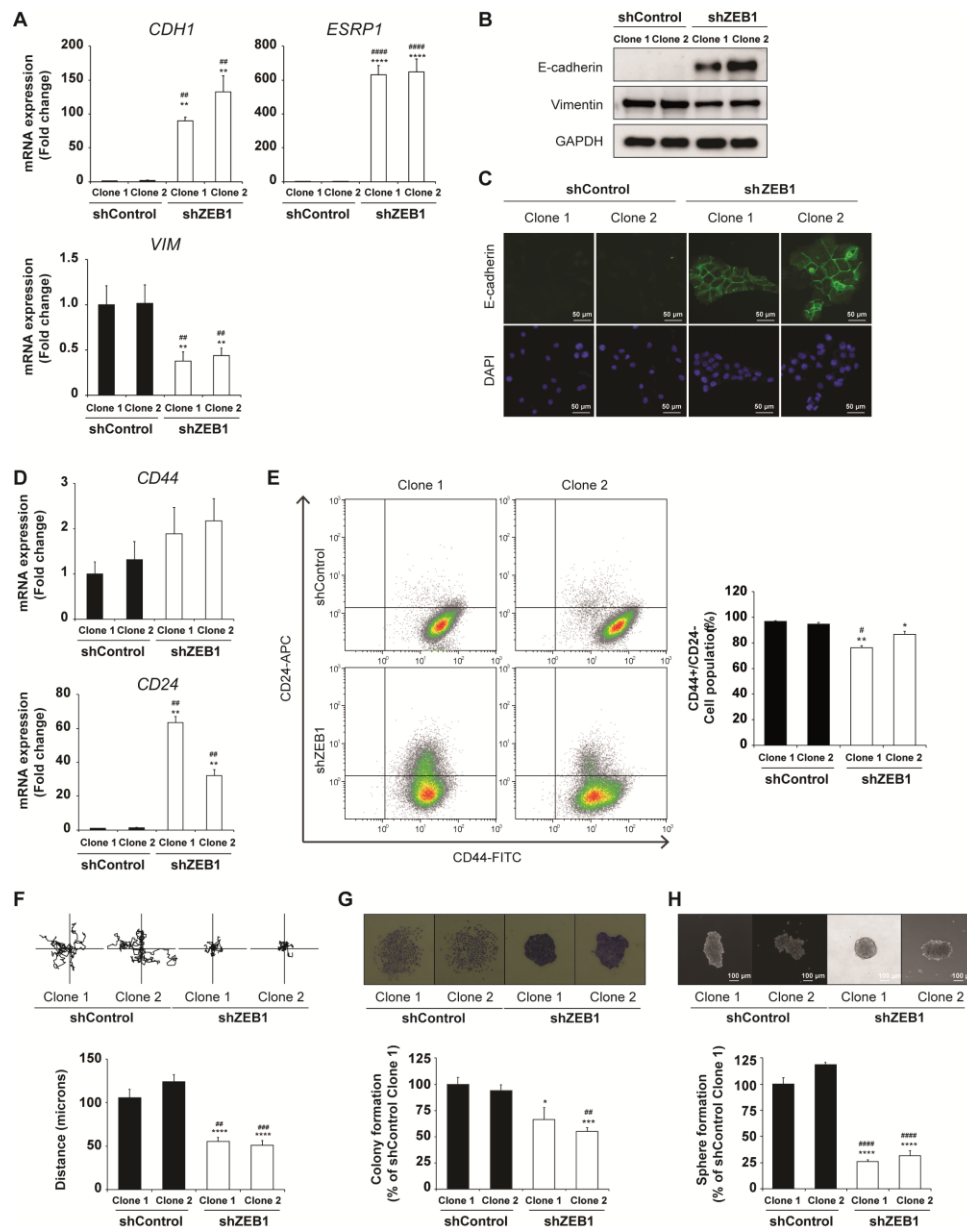
Supplementary Figure 1. ZEB1 expression correlates with that of EMT markers. EMT markers *CDH1* (E-cadherin), *ESRP1* and *VIM* (vimentin) and EMT-TFs *ZEB2*, *SNAI1* (SNAIL), *SNAI2* (SLUG), *TWIST1* and *TWIST2* mRNA expression in a panel of 8 human CCA cell lines with epithelial morphology and negligible ZEB1 expression (white bars) or mesenchymal morphology and high ZEB1 expression (black bars). Values are expressed as means $\pm$ SEM from 5 independent cultures.



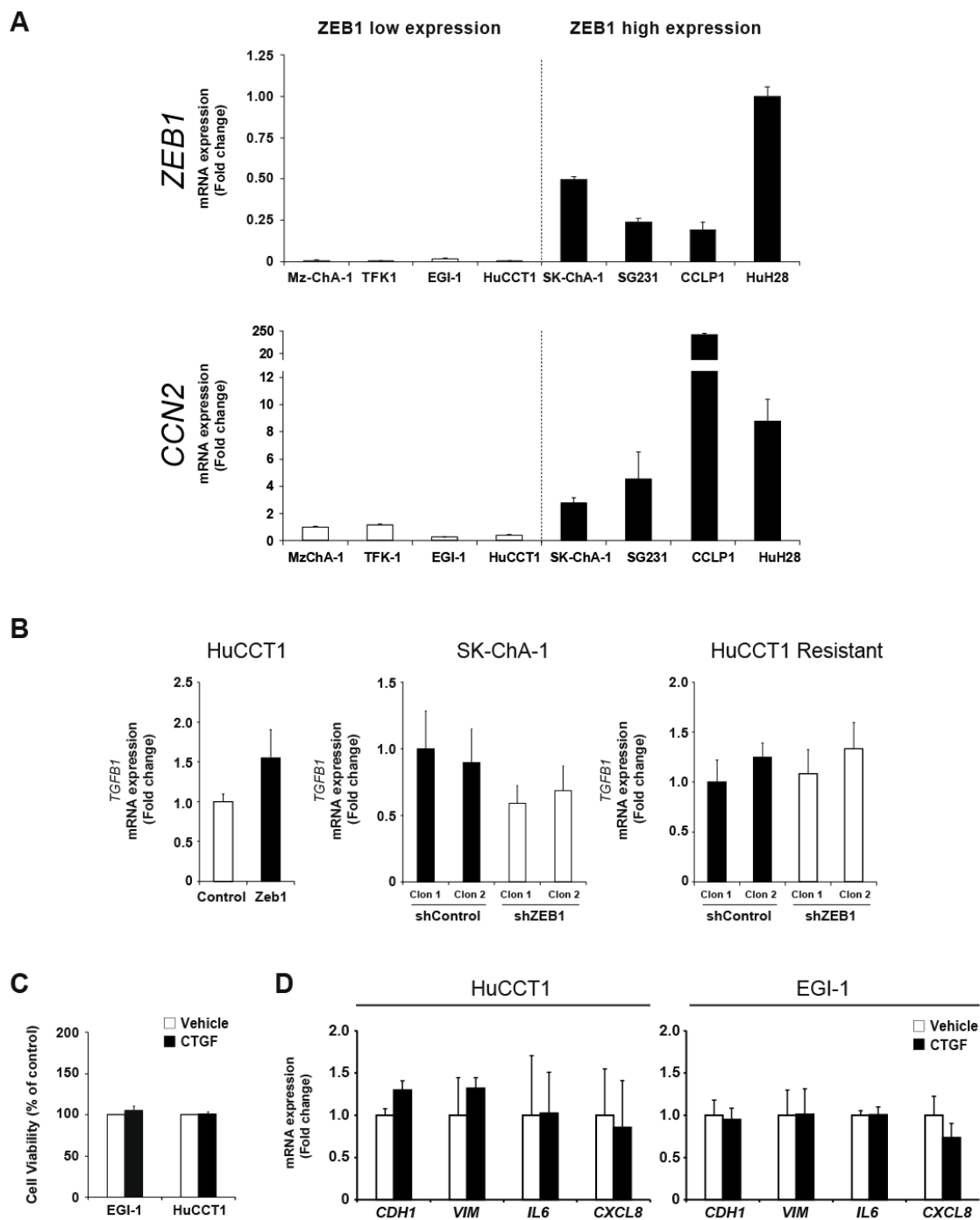
Supplementary Figure 2. ZEB1 gain/loss of function in *in vitro* models do not modify the expression of other EMT-TFs as a compensatory mechanism. A-B. Changes in EMT-TFs ZEB2, SNAI1 (SNAIL), SNAI2 (SLUG), TWIST1 and TWIST2 mRNA (A) and protein (B) expression in gain/loss of function models derived from HuCCT1, SK-ChA-1 and HuCCT1 R cells, as determined by RT-QPCR and western blot. Values are expressed as means $\pm$ SEM from 5 independent cultures. ND, Not Detected.



Supplementary Figure 3. ZEB1 expression promotes the acquisition of an EMT/CSC phenotype in CCA cells. **A-C.** Changes in *CDH1* (E-cadherin), *ESRP1*, and *VIM* (vimentin) mRNA (A), and protein expression (B) and localization (C) of E-cadherin and vimentin in SK-ChA-1 cells downregulated or not for ZEB1 by shRNA by shRNA, as determined by RT-QPCR, western blot and immunofluorescence. Scale: 50 μ m. **D-E.** Changes in *CD44* and *CD24* mRNA and protein expression in the same cells, as determined by RT-QPCR (D) and by flow cytometry (E) using APC-conjugated anti-CD24 and FITC-conjugated anti-CD44 antibodies. Gates are based on the isotype controls. Quantification in the right panel shows the % of cells in the lower right quadrant corresponding to a CD44⁺/CD24⁻ cell population. **F.** Cell tracking analyzed by time-lapse microscopy during a 24 h long time-lapse experiment. Tracks of 8 representative cells for each condition are shown. Bar graphs show quantification of the distance travelled by the cells. **G.** Colony formation assay after 7 days in culture. Representative pictures of colonies from each cell line. **H.** Sphere formation assay after 7 days in culture. Representative pictures of spheres from each cell line. SK-ChA-1 sphere counting was impossible due to aggregate formation. Scale: 200 μ m. Values are expressed as means $\pm$ SEM from at least 3 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, ****, $p < 0.0001$, as compared with shControl clone 1. #, $p < 0.05$; ##, $p < 0.01$; ###, $p < 0.001$; ####, $p < 0.0001$, as compared with shControl clone 2.

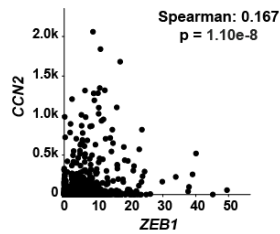


Supplementary Figure 4. ZEB1 expression promotes the acquisition of an EMT/CSC phenotype in CCA cells. **A-C.** Changes in *CDH1* (E-cadherin), *ESRP1*, and *VIM* (vimentin) mRNA (A), and protein expression (B) and localization (C) of E-cadherin and vimentin in HuCCT1 R cells downregulated or not for ZEB1 by shRNA, as determined by RT-QPCR, western blot and immunofluorescence. Scale: 50 μ m. **D-E.** Changes in *CD44* and *CD24* mRNA and protein expression in the same cells, as determined by RT-QPCR (D) and by flow cytometry (E) using APC-conjugated anti-CD24 and FITC-conjugated anti-CD44 antibodies. Gates are based on the isotype controls. Quantification in the right panel shows the % of cells in the lower right quadrant corresponding to a CD44⁺/CD24⁻ cell population. **F.** Cell tracking analyzed by time-lapse microscopy during a 24 h long time-lapse experiment. Tracks of 8 representative cells for each condition are shown. Bar graphs show quantification of the distance travelled by the cells. **G.** Colony formation assay after 7 days in culture. Representative pictures of colonies from each cell line. **H.** Sphere formation assay after 7 days in culture. Representative pictures of spheres from each cell line. Scale: 200 μ m. Values are expressed as means $\pm$ SEM from at least 3 independent cultures. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$; ****, $p < 0.0001$, as compared with shControl clone 1. #, $p < 0.05$; ##, $p < 0.01$; ###, $p < 0.001$; ####, $p < 0.0001$, as compared with shControl clone 2.

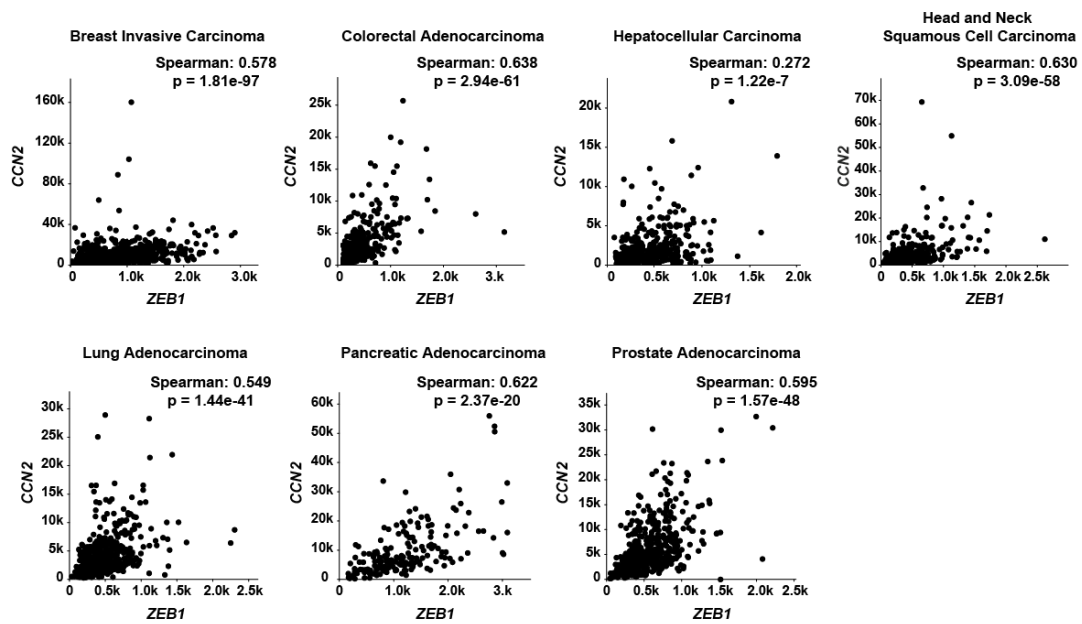


Supplementary Figure 5. A. Changes in CCN2 (encoding CTGF) mRNA expression in a panel of 8 CCA cell lines with epithelial morphology and negligible ZEB1 expression (white bars) or mesenchymal morphology and high ZEB1 expression (black bars). Values are expressed as means $\pm$ SEM from 5 independent cultures. **B.** Changes in *TGFB1* mRNA expression in gain/loss of ZEB1 function models derived from HuCCT1, SK-ChA-1 and HuCCT1 R cells, as determined by RT-QPCR. **C-D.** Effect of exogenous CTGF 100 ng/mL on cell viability (C) and mRNA expression of *CDH1* (E-cadherin), *VIM* (vimentin), *IL6* and *CXCL8* (D) in HuCCT1 and EGI-1 cells. Cell viability and mRNA level were measured after 72 h and 24 h of incubation with CTGF, respectively. Values are expressed as means $\pm$ SEM from 5 independent cultures.

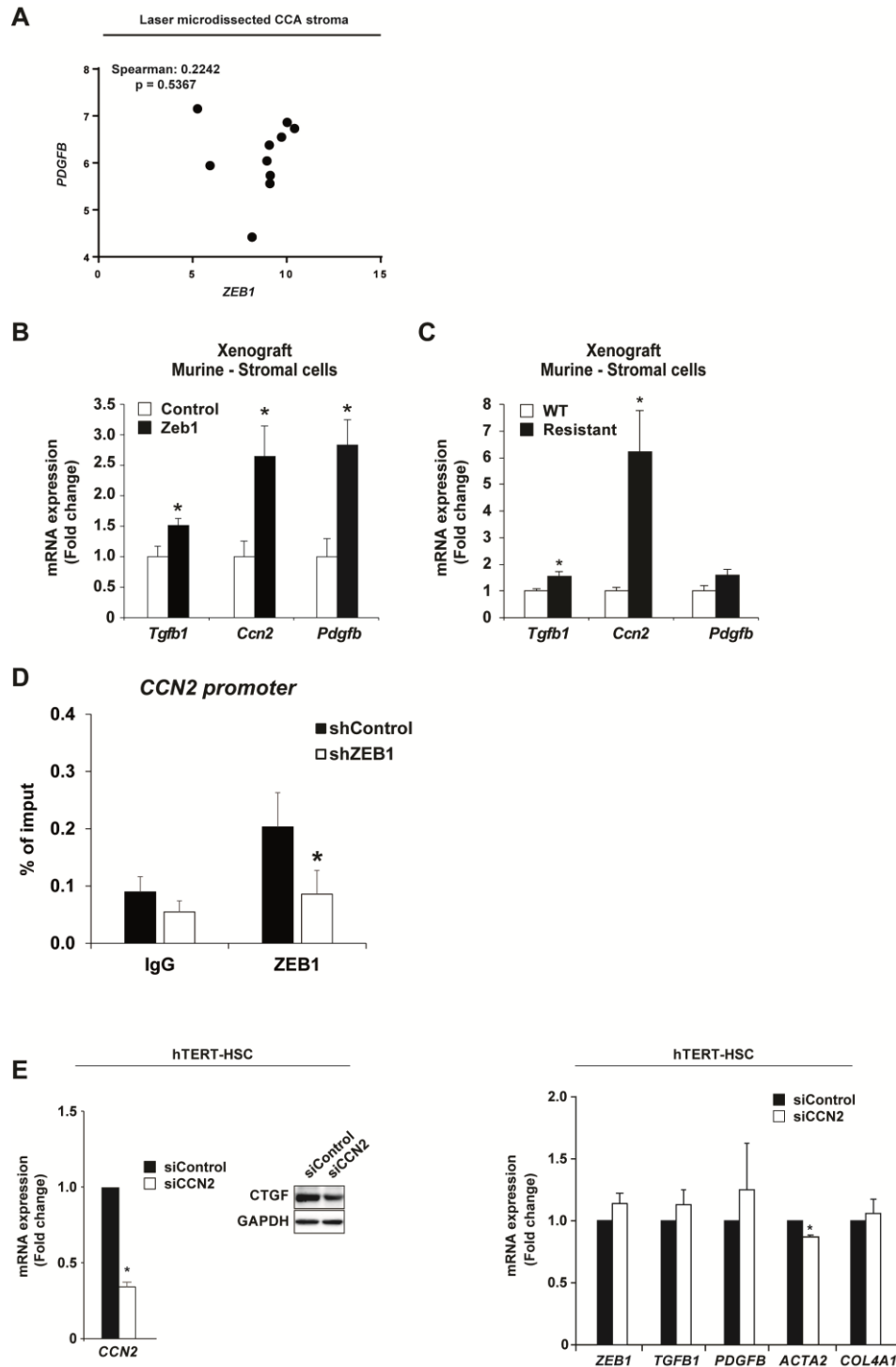
A (Cancer Cell Line Encyclopedia)



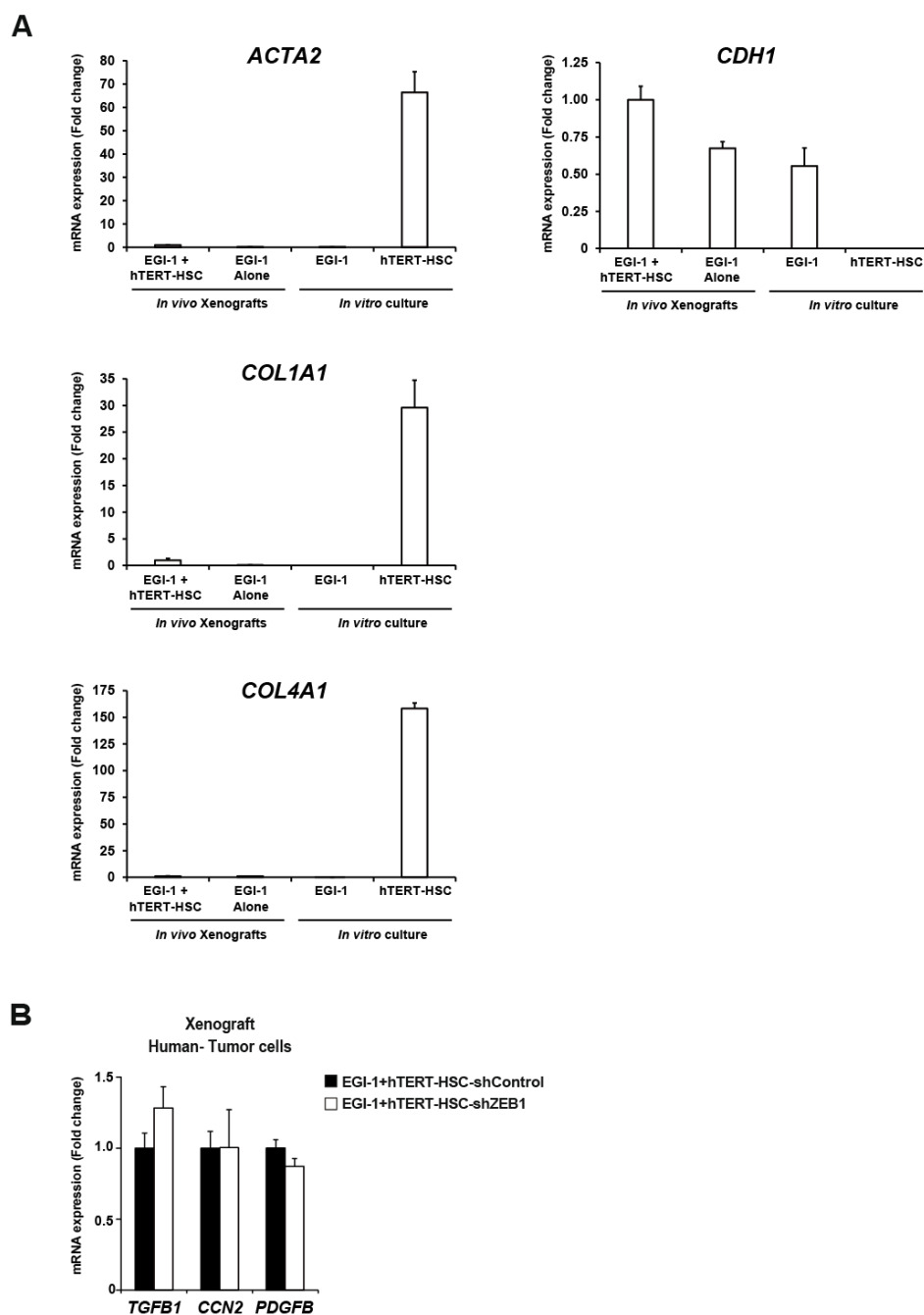
B (TCGA, PanCancer Atlas)



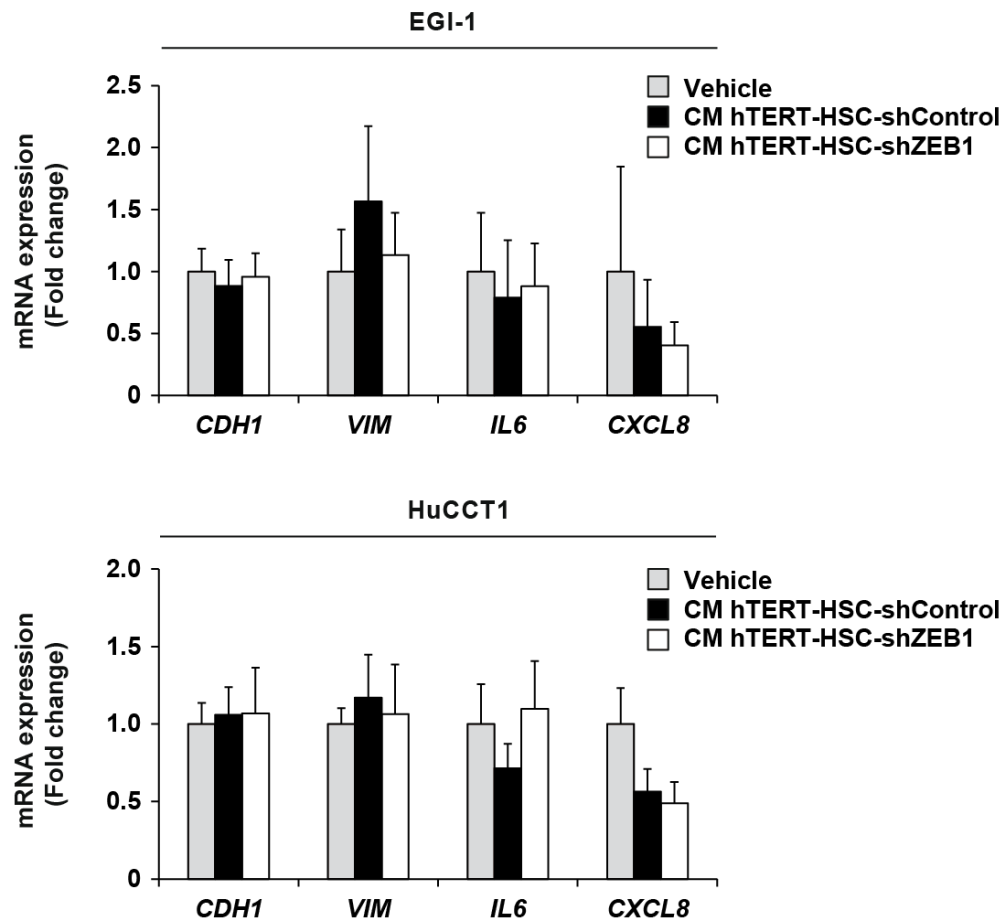
Supplementary Figure 6. CCN2 expression correlates with ZEB1 expression in multiple human cancers and cell lines. A-B. Correlation of CCN2 gene expression with ZEB1 in cell lines from the Cancer Cell Line Encyclopedia (A) and multiple cancers from the TCGA study (B), determined using Spearman's correlation analysis.



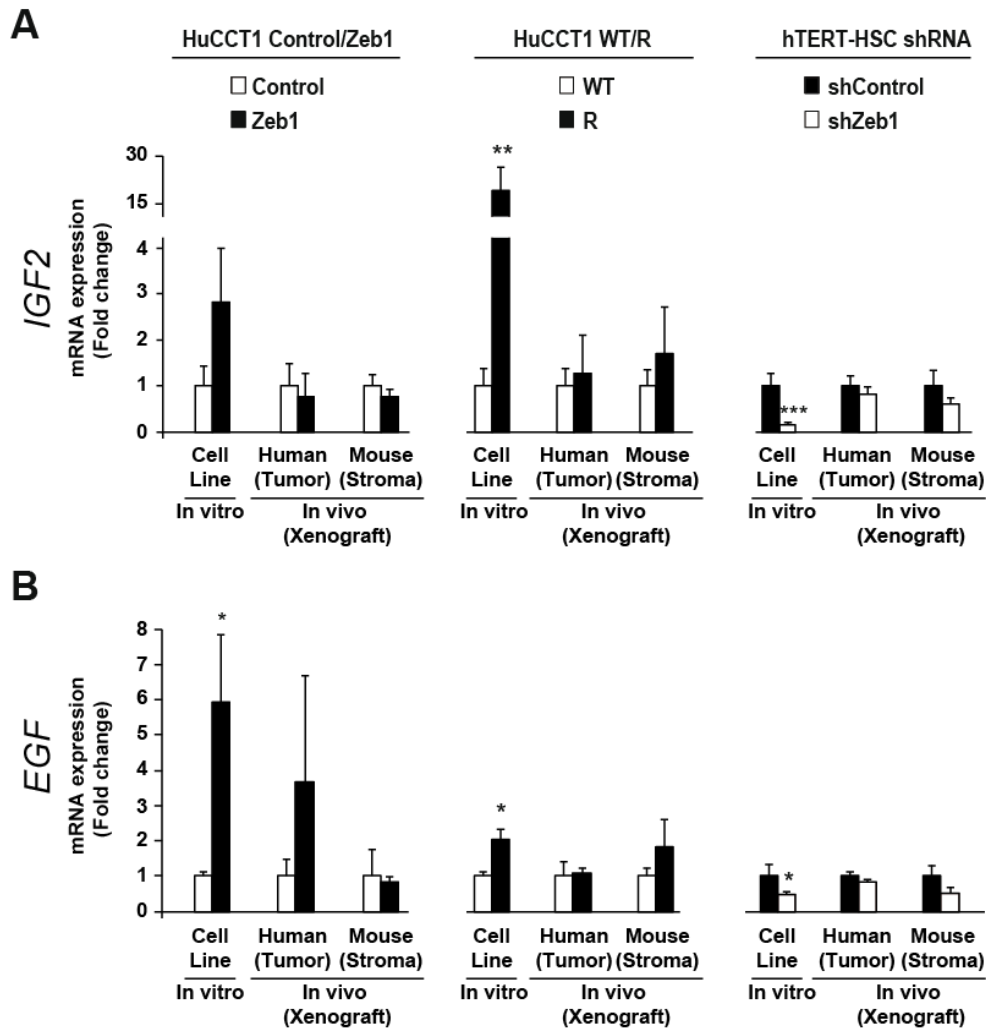
Supplementary Figure 7. A. Correlation between *PDGFB* with *ZEB1* expression in microdissected stroma from 10 iCCA samples determined using Spearman's correlation analysis. **B-C.** Changes in mRNA expression *Tgfb1*, *Ccn2* and *Pdgfb* in the stromal compartment (corresponding to murine genes) of xenograft tumors derived from HuCCT1 Control and Zeb1 cells (B), or HuCCT1 WT and Resistant (R) cells (C), analyzed by RT-QPCR. **D.** Chromatin immunoprecipitation PCR (ChIP-PCR) experiment using primers targeting *CCN2* promoter in hTERT-HSC-shControl (black bars) and shZEB1 (white bars) cells. **E.** Effect of *CCN2* downregulation by siRNA transfection in hTERT-HSC cells in the expression of *ZEB1*, *TGFB1*, *PDGFB*, *ACTA2* and *COL4A1*. Values are expressed as means $\pm$ SEM from 5 animals or 5 independent cultures. *, $p < 0.05$ comparing with Control tumors, WT tumors or sh/siControl cells.



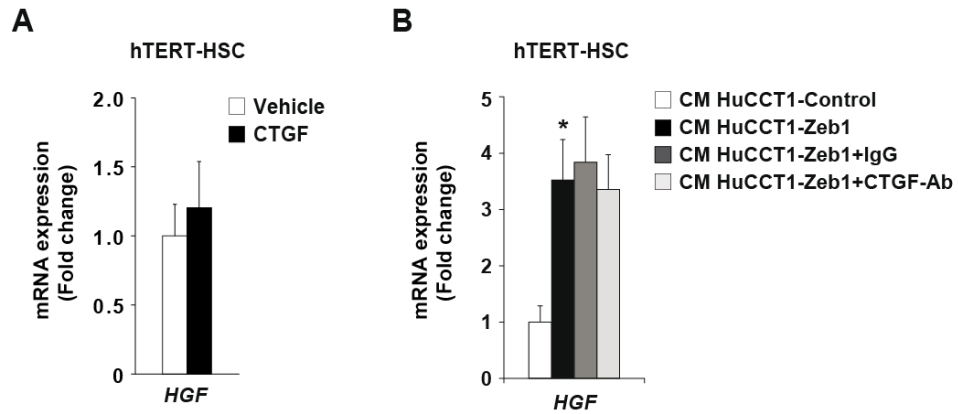
Supplementary Figure 8. A. mRNA expression of human *ACTA2* (encoding α -SMA), *COL1A1*, *COL4A1* and *CDH1* (encoding E-cadherin) determined by RT-QPCR in xenograft tumors derived from EGI-1+hTERT-HSC or EGI-1 alone, and in EGI-1 and hTERT-HSC cells *in vitro*. **B.** mRNA expression of *TGFB1*, *CCN2* and *PDGFB* in the tumor cells (corresponding to the human genes) of xenograft tumors from mice bearing EGI-1+hTERT-HSC-shControl or EGI-1+hTERT-HSC-shZEB1 cells. Results are expressed as means $\pm$ SEM from 5 animals or 5 independent cultures.



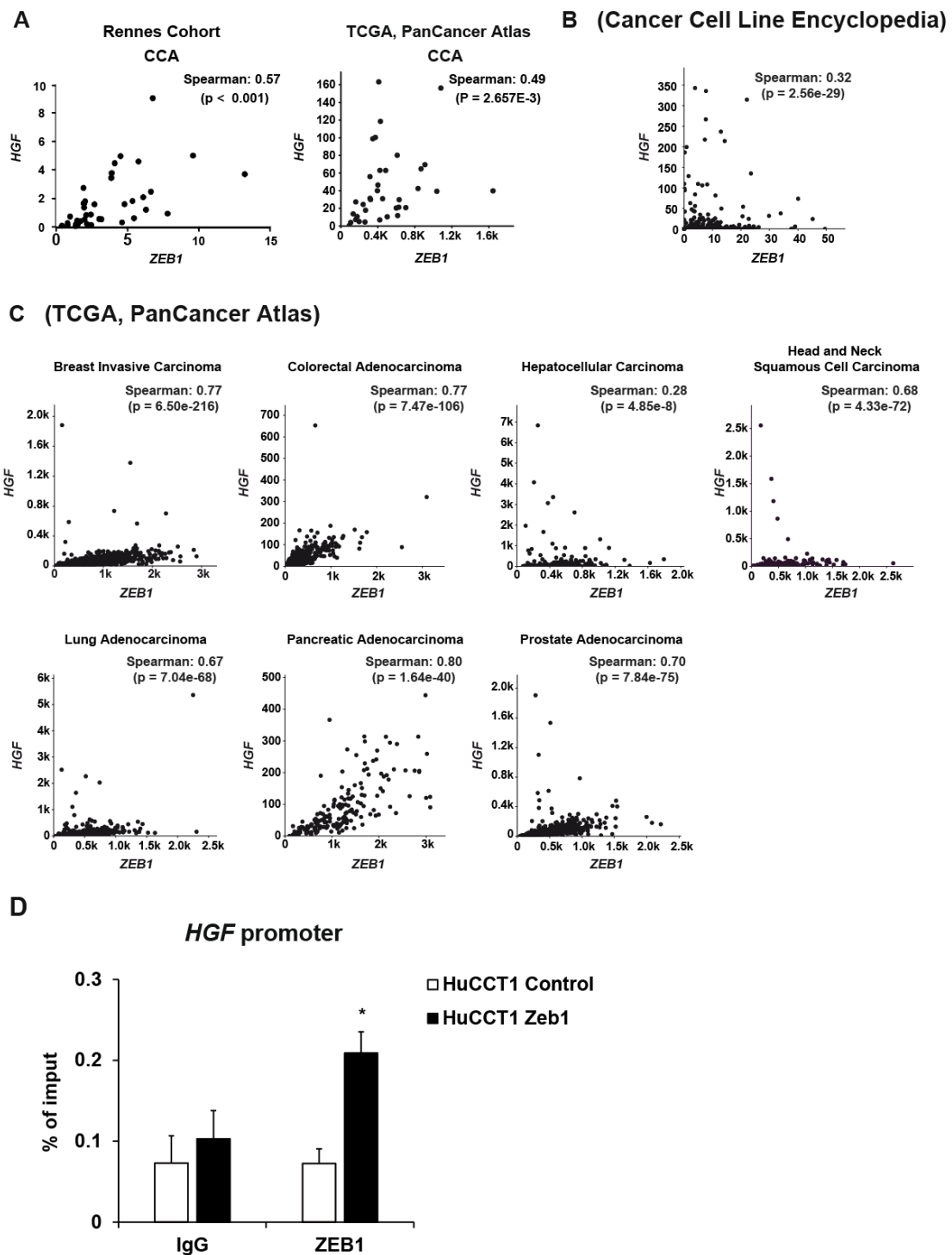
Supplementary Figure 9. Expression of *CDH1* (E-cadherin), *VIM* (vimentin), *IL6* and *CXCL8* mRNA analyzed by RT-QPCR in EGI-1 or HuCCT1 treated with conditioned media (CM) of hTERT-HSC shControl or shZEB1 cells. Values are expressed as means $\pm$ SEM from 5 independent cultures.



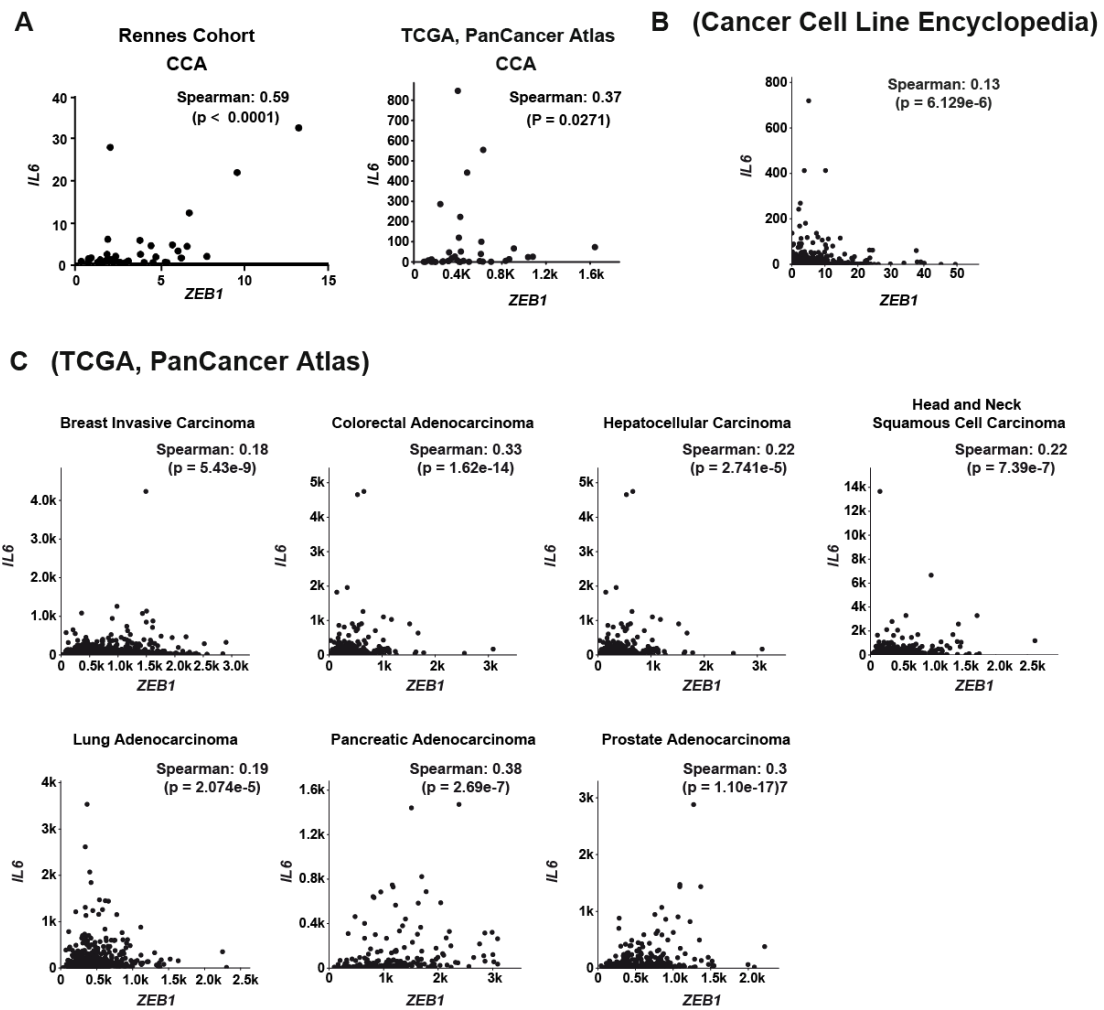
Supplementary Figure 10. A-B. Changes in mRNA expression of *IGF2* (A) and *EGF* (B) in HuCCT1 Control and Zeb1, HuCCT1 WT and Resistant (R) CCA cell lines, and in hTERT-HSC-shRNA-Control and shRNA-ZEB1 cell lines, and their respective derived CCA xenografts, determined by RT-QPCR. For each model gene expression in cultured cells (*in vitro*), tumor cells (corresponding to human genes) and stromal cells (corresponding to murine genes) of the xenografts (*in vivo*) is shown. Values are expressed as means $\pm$ SEM from 5 independent cultures/animals. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$, as compared with Control cells or Control tumors.



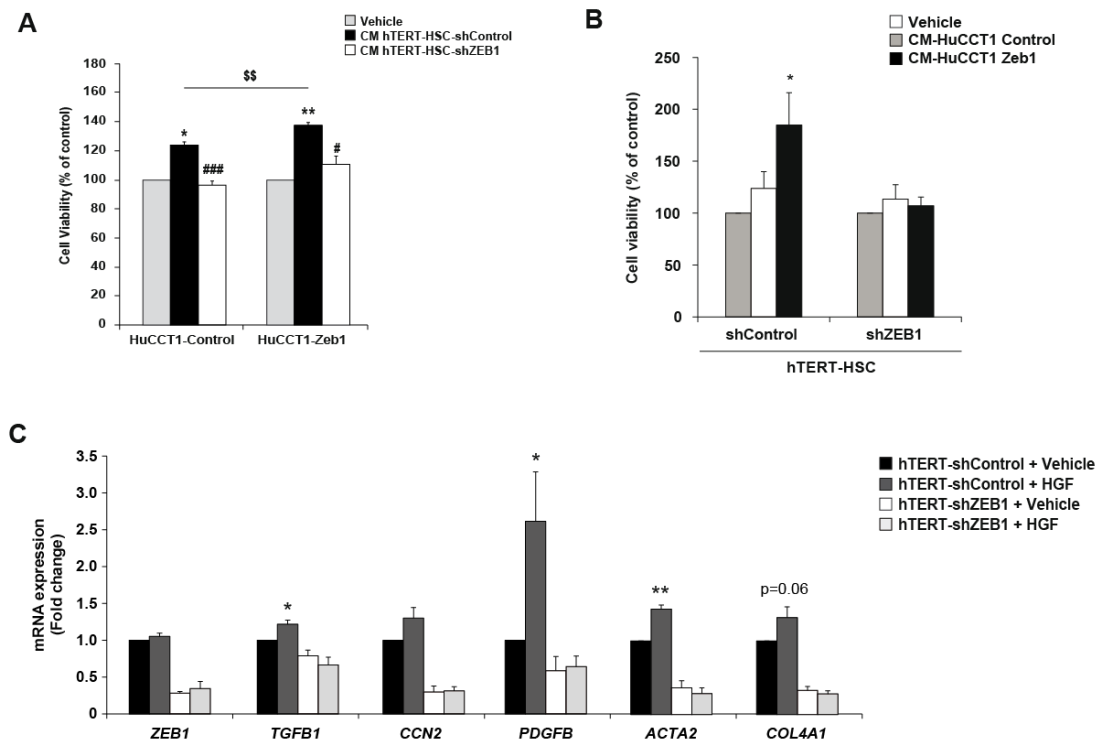
Supplementary Figure 11. Expression of *HGF* mRNA determined by RT-QPCR in hTERT-HSC cells incubated for 24 h with recombinant CTGF 100 ng/mL (A) and conditioned media (CM) (B) from HuCCT1 Control or Zeb1 cells treated with an unspecific IgG or a neutralizing antibody against CTGF. Values are expressed as means $\pm$ SEM from 5 independent cultures. *, $p < 0.05$.



Supplementary Figure 12. *HGF* expression correlates with that of *ZEB1* in multiple human cancers and cell lines. A-C. Correlation of *HGF* gene expression with *ZEB1* in Rennes and TCGA CCA cohorts (A), in cell lines from the Cancer Cell Line Encyclopedia (B) and in multiple cancers from the TCGA study (C), determined using Spearman's correlation analysis. **C.** Chromatin immunoprecipitation PCR (ChIP-PCR) experiment using primers targeting *HGF* promoter in HuCCT1 Control (white bars) and Zeb1 (black bars) cells.

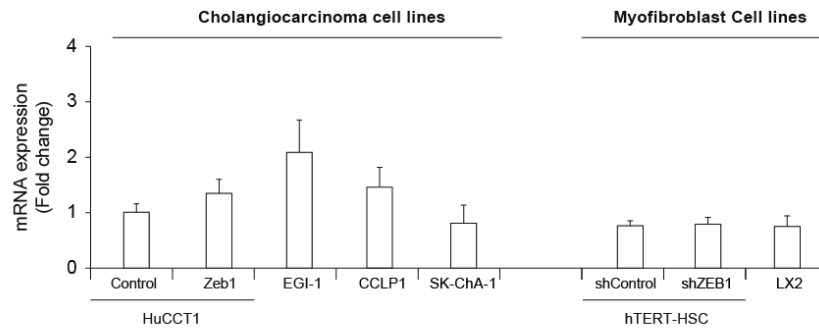


Supplementary Figure 13. *IL6* expression correlates with that of *ZEB1* in multiple human cancers and cell lines. A-C. Correlation of *IL6* gene expression with *ZEB1* in Rennes and TCGA CCA cohorts (A), in cell lines from the Cancer Cell Line Encyclopedia (B) and in multiple cancers from the TCGA study (C), determined using Spearman's correlation analysis.

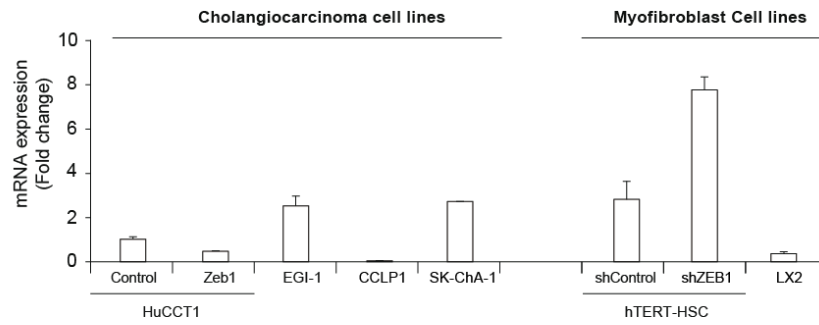


Supplementary Figure 14. A. Effect of conditioned media (CM) from hTERT-HSC shControl or shZEB1 cells on the viability of HuCCT1 Control and Zeb1 cells. **B.** Effect of CM from HuCCT1 Control and Zeb1 on the viability of hTERT-HSC shControl or shZEB1 cells. **C.** Effect of HGF on the expression of *ZEB1*, *TGFB1*, *CCN2*, *PDGFB*, *ACTA2* and *COL4A1* in hTERT-HSC shControl and shZEB1. mRNA expression was analyzed by RT-QPCR. Values are expressed as means $\pm$ SEM from at least 4 independent cultures. *, $p < 0.05$; **, $p < 0.01$, as compared with Control cells (hTERT-shControl + vehicle). *, $p < 0.05$; **, $p < 0.01$; as compared with vehicle. #, $p < 0.05$; ###, $p < 0.001$, as compared with CM from hTERT-HSC shControl. \$\$, $p < 0.01$ as compared with HuCCT1 Control cells.

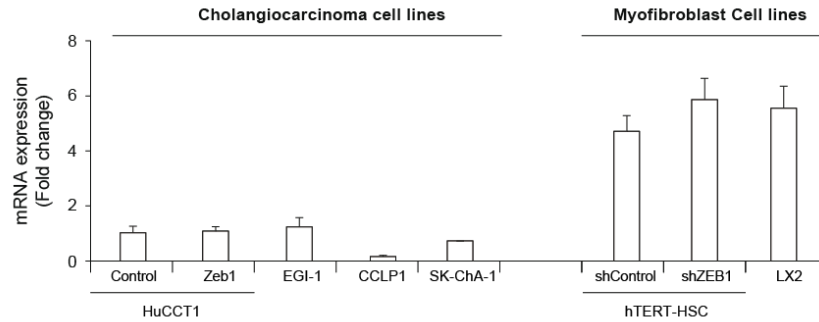
A-MET



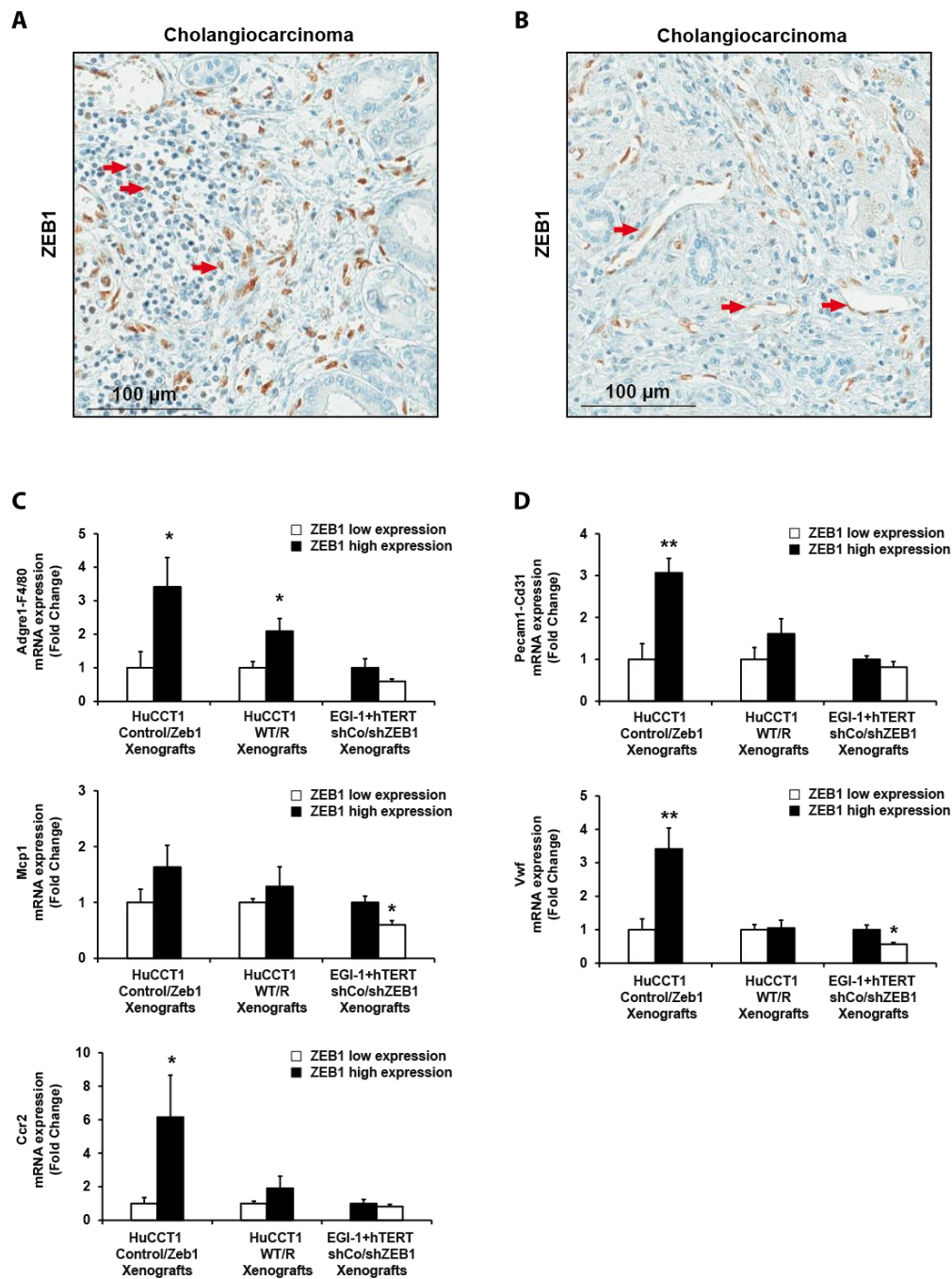
B-IL6R



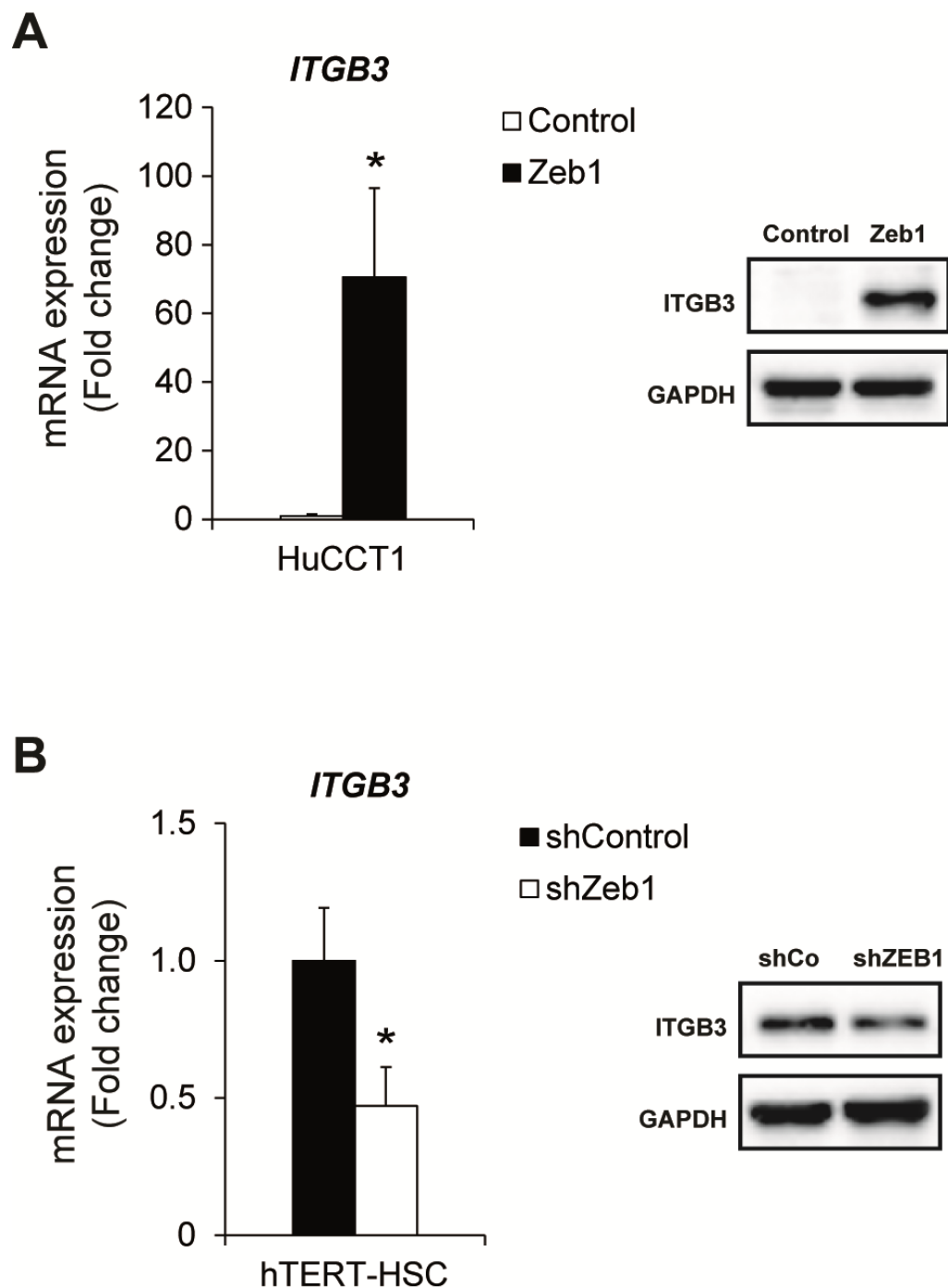
C-IL6ST



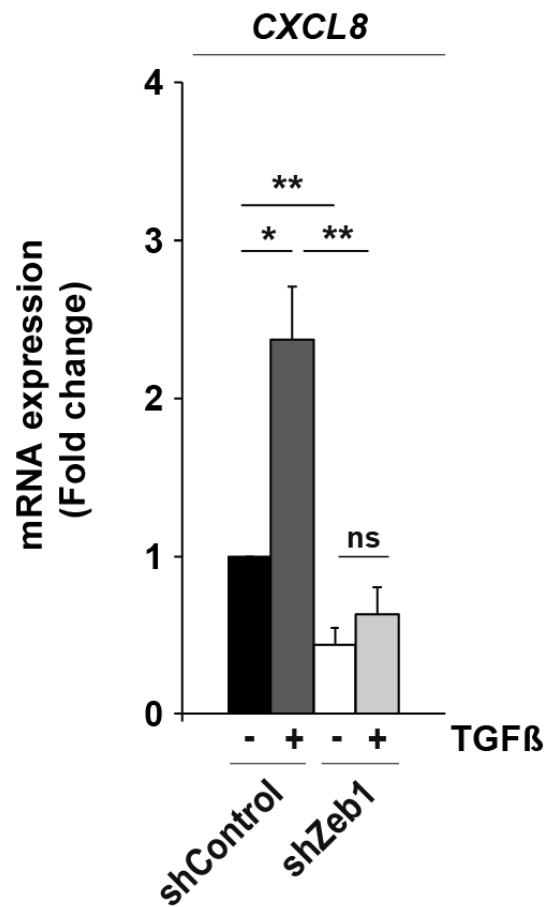
Supplementary Figure 15. mRNA expression of HGF receptor, *MET*, and of IL6 receptors, *IL6R* and *IL6ST* (GP130) in human CCA and hepatic myofibroblast cell lines. Expression is determined by RT-QPCR and values are expressed as means $\pm$ SEM from 4 independent cultures.



Supplementary Figure 16. A-B. Stromal cells other than CAF express ZEB1 in human intrahepatic CCA. A-B. Representative IHC staining of ZEB1, showing positive ZEB1 staining in immune cells (A) and endothelial cells (B). Red arrows point to either immune cells (A) or endothelial cells (B) expressing ZEB1. Scale: 100 μ m. C-D. ZEB1 expression in tumor cells modifies other stromal cells than CAF in murine model of CCA. mRNA expression of murine *Adgre* (F4/80), *Mcp1*, *Ccr2* (macrophage markers) (C), *Pecam-1* (CD31) and *Vwf* (endothelial markers) (D), in xenografts from mice bearing HuCCT1 Control and Zeb1, HuCCT1 WT and R and the combination of EGI-1 plus hTERT-HSC-shControl and shZeb1 cells, analyzed by RT-QPCR. Values are expressed as means $\pm$ SEM from 5 independent animals. *, $p < 0.05$; **, $p < 0.01$, as compared with ZEB1 low expression.



Supplementary Figure 17. A-B. Changes in *ITGB3* (Integrin beta 3, also known as CD61) mRNA and protein expression in HuCCT1 Control and Zeb1 (A), and in hTERT-HSC-shControl and shZEB1 (B), as determined by RT-QPCR and western blot. Results are expressed as means $\pm$ SEM from 5 independent cultures. *, $p < 0.05$; as compared with Control or shControl cells.



Supplementary Figure 18. ZEB1 in myofibroblasts impacts CXCL8 (IL8) expression. Changes in CXCL8 (IL8) in hTERT-HSC-shRNA-Control and shRNA-ZEB1 cells exposed to TGF- β 1 10 ng/ml or the vehicle for 24 h. Results are expressed as means $\pm$ SEM from 5 independent cultures. *, $p < 0.05$; **, $p < 0.01$, as indicated by lines.