

ELF3 suppresses gallbladder cancer development through downregulation of the EREG/EGFR/mTOR complex 1 signalling pathway

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Abstract

The prognosis of gallbladder cancer (GBC) remains poor, and a better understanding of GBC molecular mechanisms is important. Genome sequencing of human GBC has demonstrated that loss-of-function mutations of E74-like ETS transcription factor 3 (*ELF3*) are frequently observed, with *ELF3* considered to be a tumour suppressor in GBC. To clarify the underlying molecular mechanisms by which *ELF3* suppresses GBC development, we performed *in vivo* analysis using a combination of autochthonous and allograft mouse models. We first evaluated the clinical significance of *ELF3* expression in human GBC tissues and found that low *ELF3* expression was associated with advanced clinical stage and deep tumour invasion. For *in vivo* analysis, we generated *Pdx1-Cre; Kras^{G12D}; Trp53^{R172H}; Elf3^{fl/fl}* (KPCE) mice and *Pdx1-Cre; Kras^{G12D}; Trp53^{R172H}; Elf3^{wt/wt}* (KPC) mice as a control and analysed their gallbladders histologically. KPCE mice developed larger papillary lesions in the gallbladder than those developed by KPC mice. Organoids established from the gallbladders of KPCE and KPC mice were analysed *in vitro*. RNA sequencing showed upregulated expression of epiregulin (*Ereg*) in KPCE organoids, and western blotting revealed that EGFR/mechanical targets of rapamycin complex 1 (mTORC1) were upregulated in KPCE organoids. In addition, ChIP assays on *Elf3*-overexpressing KPCE organoids showed that *ELF3* directly regulated *Ereg*. *Ereg* deletion in KPCE organoids (using CRISPR/Cas9) induced EGFR/mTORC1 downregulation, indicating that *ELF3* controlled EGFR/mTORC1 activity through regulation of *Ereg* expression. We also generated allograft mouse models using KPCE and KPC organoids and found that KPCE organoid allograft tumours exhibited poorly differentiated structures with mTORC1 upregulation and mesenchymal phenotype, which were suppressed by *Ereg* deletion. Furthermore, EGFR/mTORC1 inhibition suppressed cell proliferation and epithelial–mesenchymal transition in KPCE organoids. Our results suggest that *ELF3* suppresses GBC development via downregulation of EREG/EGFR/mTORC1 signalling. EGFR/mTORC1 inhibition is a potential therapeutic option for GBC with *ELF3* mutation.

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Introduction

Gallbladder cancer (GBC) is the most common type of biliary tract cancer (BTC) with high lethality [1]. Its poor prognosis is partially attributed to the anatomical characteristics of the gallbladder. Lack of both submucosal and serosal layers on the hepatic side of gallbladder facilitates tumour invasion and metastasis, which is associated with diagnosis at an advanced

stage [2]. Lack of adequately effective therapies remains a major cause of poor prognosis of GBC. Although several efforts have been made to establish effective therapies, the 5-year survival rate of GBC remains under 30% [3]. To improve the prognosis, understanding the molecular mechanisms underlying GBC formation and identifying new therapeutic targets are important.

Several studies on genome sequencing of BTC, including GBC, recently revealed that many significantly altered

genes were included in the erb-b2 receptor tyrosine kinase (ERBB) pathway [4–6]. The ERBB family comprises four receptor tyrosine kinases: EGFR/ERBB1, ERBB2, ERBB3, and ERBB4 [7,8]. Activated ERBB stimulates various downstream pathways, such as the RAS/MAPK and PI3K/AKT pathways, which are involved in the development of many types of human cancers [7–10]. GBCs that harbour gene mutations in the ERBB pathway are associated with worse prognosis; therefore, the ERBB pathway is an important factor in GBC development [4].

Integrated genomic analysis of human GBC revealed that E74-like ETS transcription factor 3 (*ELF3*) is the second most frequently mutated gene and is altered in up to 20% of samples [6]. Other genome sequencing analyses also demonstrated that *ELF3* was highly mutated in extrahepatic cholangiocarcinoma and ampullary carcinoma [5,11]. *ELF3* is an epithelium-specific transcription factor belonging to the ETS transcription family and is involved in various physiological functions, such as cell differentiation/proliferation, cell cycle adjustment, apoptosis, and angiogenesis, in normal tissues [12]. In cancers, various malignancies harbour *ELF3* alterations [13]. *ELF3* plays an important role in cancer development; however, whether *ELF3* functions as a tumour suppressor or promoter depends on the primary organ [13–20]. In BTCs, including GBC, most *ELF3* mutations are the loss-of-function type, such as frameshifts and stop gains [5,6,11]. Moreover, xenografts of *ELF3*-overexpressing human BTC cells showed a better-differentiated tubular structure than that seen in control cells [21]. Thus, *ELF3* is considered a suppressor of tumour development and epithelial–mesenchymal transition (EMT) in BTC; however, the molecular mechanism by which *ELF3* suppress BTC development and EMT has not been fully elucidated.

In this study, we show that *ELF3* plays a tumour-suppressive role in GBC development in mice. *ELF3* negatively regulates mRNA expression of *Ereg*, the gene-encoding ligand of ERBB receptors, and consequently suppresses EGFR/mTOR complex 1 (mTORC1) activity.

Materials and methods

Study approval

This study was conducted in compliance with the Helsinki Declaration and the Ethical Guidelines for Medical and Health Research Involving Human Subjects. The protocol was approved by the Human Ethics Review Committee of Kyoto University Hospital (Approval No. G616). Informed consent was obtained from all patients with an opt-out method. All experiments with mice were approved by the Ethics Committee for Animal Experiments and performed in accordance with the guidelines of the Animal Experiments of Kyoto University (Approval No. 22581).

Patients

GBC specimens were obtained from surgically resected or liver biopsy tissues of 85 consecutive patients admitted to Kyoto University Hospital (Kyoto, Japan) between November 2004 and October 2020. Patient medical records were reviewed from their first visit to our institute until November 2021, loss to follow-up, or death. The detailed clinical information of patients with GBC is shown in supplementary material, Table S1.

Mice

Pdx1-Cre [22], *LSL-Kras*^{G12D} [23], and *LSL-Trp53*^{R172H} mice [24] were established as described previously. *Elf3* flox mice (C57BL/6N-*Elf3*^{tm1a(KOMP)Wtsi}, RRID:MMRRC_047524-UCD) were purchased from UC Davis KOMP Repository (Davis, CA, USA) and crossed with *B6-Tg (CAG-FLPe)*³⁶ mice to delete the inserted cassette floxed by the FRT site. NOD.CB17-*Prkdc*^{scid}/Jcl mice were purchased from CLEA Japan (Tokyo, Japan). All mice were maintained in a specific-pathogen-free facility at the Kyoto University Faculty of Medicine (Kyoto, Japan) with food and water provided *ad libitum*. Mice were housed at two to six per cage irrespective of their genotype and used without distinction of sex. For the gallbladder histological analysis, 9 or 10 mice per group were sacrificed after fasting for 2 h. For 5-bromo-4-chloro-3-indolyl- β -D-galactoside (X-gal, Wako, Osaka, Japan) staining of the biliary tract, three *Pdx1-Cre; Rosa26*^{LSL-LacZ} mice were used. For organoid establishment, three or four mice were used in each organoid group.

Semiquantification of immunostaining intensity in human GBC samples

For the semiquantification of immunostaining intensity in human GBC samples, the dominant immunostaining intensity was determined on a scale of 0 to 3: 0 = negative, 1 = weak, 2 = moderate, 3 = strong. For *ELF3* semiquantification, moderate intensity was defined as one similar to that of normal gallbladder epithelial cells, and low and high expression of *ELF3* were defined as staining intensities of 0–1 and 2–3, respectively. Scoring was performed by two investigators.

Statistical analysis

Data are presented as the mean \pm SEM. Statistical comparisons between groups were performed using Student's *t*-test, paired *t*-test, or one-way ANOVA test. $P < 0.05$ was considered statistically significant. All statistical analyses were performed using either GraphPad Prism 9 (GraphPad, San Diego, CA, USA, RRID:SCR_002798) or R version 4.0.3. Overall survival (OS) was defined as the interval from the date of diagnosis to the date of death or censoring for any cause and estimated using the Kaplan–Meier method. The difference between the two groups was calculated using the Gehan–Breslow–Wilcoxon test. Hazard ratios were

calculated using Cox proportional hazards regression analysis (Mantel–Haenszel method).

Further descriptions of allograft mouse models, histology, organoids and cell lines, lentivirus transfection, RNA isolation and RNA sequencing (RNA-seq), western blot analyses, RT-qPCR, CHIP, cell proliferation assays, scratch assays, invasion assays, and analysis of human GBC RNA-seq data can be found in Supplementary materials and methods. The antibodies and primer sets used in this study are listed in supplementary material, Tables S2 and S3, respectively.

Results

Low ELF3 expression in GBC is associated with advanced clinical stage and deep invasion of tumours

To elucidate ELF3 expression in epithelial cells of gallbladder and other parts of the extrahepatic biliary tract (EHBT) in humans, we performed immunohistochemistry (IHC) on normal human tissues. ELF3 was expressed in the epithelial cells of the EHBT, including the gallbladder, extrahepatic bile duct (EHBD), and papilla of Vater (Figure 1A). We also examined tissues from patients with chronic cholecystitis and precancerous lesions of the EHBT and revealed that ELF3 was also expressed in the epithelial cells of these tissues (Figure 1A).

Next, to evaluate ELF3 expression in GBC tissues, we performed IHC on GBC tissues obtained from GBC patients and semiquantitatively evaluated staining intensity (supplementary material, Table S1). Low ELF3 expression was observed in more than half of all samples (Figure 1B). We further investigated the clinical significance of ELF3 expression by comparing the clinical characteristics of the low- and high-ELF3-expression groups. Notably, low ELF3 expression was associated with advanced clinical stage and deep tumour invasion (Table 1). The Kaplan–Meier method and Gehan–Breslow–Wilcoxon test demonstrated that the OS of the low-ELF3-expression group (median: 672 days) tended to be shorter than that of the high-ELF3-expression group (median: 1,713 days; Figure 1C). These results suggest that ELF3 plays a tumour-suppressive role in human GBC, especially in terms of tumour invasiveness.

ELF3 is not essential for the development and maintenance of the EHBT in mice

Pdx1 is expressed in embryonic extrahepatic biliary epithelial cells [25], and Cre recombinase expression driven by the *Pdx1* promoter in extrahepatic biliary epithelial cells was confirmed using X-gal staining on the EHBT of adult *Pdx1-Cre; Rosa26^{LSL-LacZ}* mice (supplementary material, Figure S1). Therefore, we generated *Pdx1-Cre; Elf3^{fl/fl}* mice to investigate the functional role of ELF3 in EHBT development (supplementary material, Figure S2A). The EHBT

of *Pdx1-Cre; Elf3^{fl/fl}* mice showed no obvious changes in gross appearance or histological abnormalities compared with that of control *Pdx1-Cre; Elf3^{wt/wt}* mice (supplementary material, Figure S2B,C). IHC for ELF3 revealed that, similar to that in human tissues, ELF3 was expressed in extrahepatic biliary epithelial cells of control mice and confirmed that *Elf3* was efficiently deleted in the extrahepatic biliary epithelial cells in *Pdx1-Cre; Elf3^{fl/fl}* mice (supplementary material, Figure S2C).

To investigate the role of ELF3 in EHBT homeostasis, we evaluated these mice at 1 year of age. The EHBT of *Pdx1-Cre; Elf3^{fl/fl}* mice showed no changes in gross appearance or histological abnormalities compared with that of control mice, and IHC for ELF3 showed that *Elf3*-deleted extrahepatic biliary cells were maintained in *Pdx1-Cre; Elf3^{fl/fl}* mice (supplementary material, Figure S2D). These data suggest that ELF3 is not essential for EHBT development and homeostasis in mice.

Elf3 deletion accelerates tumourigenesis in the gallbladder in the context of *Kras* and *Trp53* mutation in mice

TP53 is the most frequently mutated gene and is found altered in half of human GBC cases [4,6]. Therefore, we generated *Pdx1-Cre; Trp53^{R172H}; Elf3^{fl/fl}* (PCE) mice and *Pdx1-Cre; Trp53^{R172H}; Elf3^{wt/wt}* (PC) mice as controls to evaluate whether *Elf3* deletion and *Trp53* mutation cause tumourigenesis in the gallbladder (supplementary material, Figure S3A). Although the supranuclear mucinous change in gallbladder epithelial cells in PCE mice at 40–45 weeks of age was slightly more prominent than that in PC mice, neither PCE nor PC mice showed obvious tumour formation in the EHBT (supplementary material, Figure S3B,C). This indicates that *Trp53* inactivation is not sufficient to cause tumour formation in the gallbladder of mice, regardless of the presence or absence of *Elf3*, and additional oncogenic driver mutations are essential.

In human GBC, several substantially altered genes have been identified in the ERBB/RAS/PI3K pathway, including *KRAS* [4,6], and oncogenic *Kras^{G12D}* mutations have been reported to induce gallbladder adenomas in mice [26]. Therefore, we generated *Pdx1-Cre; Kras^{G12D}; Trp53^{R172H}; Elf3^{fl/fl}* (KPCE) mice and *Pdx1-Cre; Kras^{G12D}; Trp53^{R172H}; Elf3^{wt/wt}* (KPC) control mice (Figure 2A). While KPC mice are widely accepted models of pancreatic ductal cancer [27], we used them to investigate the impact of *Elf3* deletion on *Kras* activation and *Trp53* inactivation in the gallbladder epithelium. At 16–18 weeks of age, the gallbladder wall of KPCE mice was thicker than that of KPC mice (Figure 2B). Histologically, massive papillary lesions were observed in KPCE mice, whereas papillary lesions in KPC mice were relatively thin (Figure 2C). The occupancy rate of papillary lesions in the gallbladder lumen was significantly higher in KPCE mice than in KPC mice (Figure 2D). Furthermore, the epithelial cells in KPCE mice showed a higher grade of cellular atypia

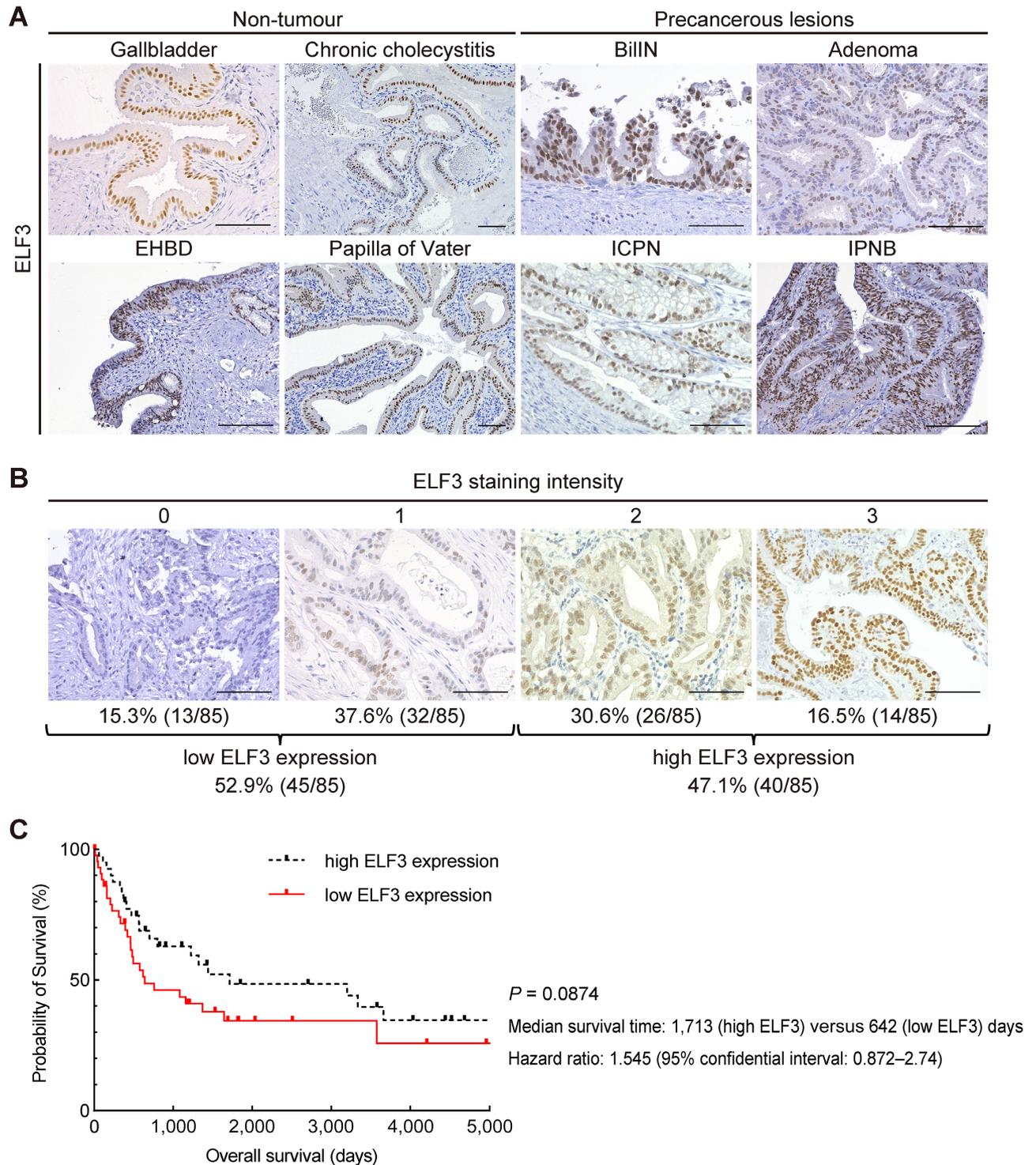


Figure 1. ELF3 expression of human EHB tissues. (A) Representative immunostaining for ELF3 in non-tumour tissues (left) and precancerous lesion tissues (right) of human EHB. Non-tumour tissues include normal gallbladder, EHBD, papilla of Vater, and chronic cholecystitis. Precancerous lesions include biliary intraepithelial neoplasia (BiIIN), adenoma, intracystic papillary neoplasm (ICPN), and intraductal papillary neoplasm of bile duct (IPNB). Scale bars, 100 μ m. (B) Representative immunostaining for ELF3 in human gallbladder cancer and scaling of dominant staining intensity. Scale bars, 100 μ m. (C) Kaplan–Meier curves of overall survival of low and high ELF3 expression groups ($n = 45$ and 40 , respectively). $P = 0.0874$, Gehan–Breslow–Wilcoxon test. See also Table 1 and supplementary material, Table S1, for clinical information of GBC patients and differences between the two groups.

than those in KPC mice and exhibited pleomorphic nuclei with anisonucleosis and distinct nucleoli, whereas those in KPC mice showed almost no atypical change

(Figure 2C). These results indicate that *Elf3* deletion accelerates tumourigenesis in the gallbladder in the context of *Kras* and *Trp53* mutations in mice.

Table 1. Association between histological ELF3 expression and clinicopathological characteristics in GBC patients.

	Low ELF3 expression (n = 45)	High ELF3 expression (n = 40)	P value
Sex, male/female, n	25/20	15/25	0.128
Median age (range), years	70 (44–88)	73 (49–85)	0.142
Clinical stage (UICC 8th), n			
0/I/II/III/IVA/IVB	6/2/7/14/2/14	4/6/14/5/0/11	
0, I, II versus III, IVA, IVB	15/30	24/16	0.0172*
T stage, (UICC 8th), n			
Tis/T1/T2/T3/T4	6/2/9/23/5	4/6/17/8/5	
Tis, T1, T2 versus T3, T4	17/28	27/13	0.00895**
N stage (UICC 8th), n			
N0/N1/N2	23/21/1	28/11/1	0.125
M stage (UICC 8th), n			
M0/M1	31/14	29/11	0.813
Main histological type, n			
pap, tub1/tub2-3, por, undiff/others	17/19/6	22/14/4	0.498
Initial treatment, n			
Surgery/Chemotherapy/BSC	40/4/1	33/7/0	0.335
Recurrence after resection, n			
Yes/no	20/20	10/23	0.101

BSC, best supportive care; UICC, Union for International Cancer Control.

* $p < 0.05$;

** $p < 0.01$, Student's *t*-test.

ERBB signalling pathway is upregulated in KPCE organoids and ELF3 directly downregulates *Ereg* and *Areg* expression

We further investigated the mechanism underlying tumour formation in the gallbladders of KPCE mice before the development of papillary lesions using organoids generated from the gallbladder epithelium of KPC and KPCE mice at 10–11 weeks of age (Figure 3A). A previous report revealed that EGF and FGF removal from the culture medium could efficiently select tumour organoids harbouring *KRAS* mutation [28], and thus we used serum-free culture medium without EGF or FGF for organoid culture and confirmed the recombination of *Kras*^{G12D} and *Trp53*^{R172H} alleles and efficient selection of organoids (supplementary material, Figure S4A). In addition, to verify that these organoids were derived from gallbladder epithelial cells, we also established organoids from the common bile duct of the same mice and evaluated mRNA expression of the gallbladder and common bile duct organoids using RT-qPCR (supplementary material, Figure S4B). RT-qPCR revealed that the gallbladder-derived organoids had higher levels of the gallbladder-specific marker *Sox17* and lower levels of the common bile duct-specific markers *Slc28a3* and *Kcne3* compared to those in the common bile duct organoids (supplementary material, Figure S4C) [29], indicating that the gallbladder organoids could be established without contamination by epithelial cells from other regions of the bile duct.

To investigate the underlying transcript changes in the gallbladders of KPCE mice before the development of papillary lesions, RNA-seq was performed on the

gallbladder organoids (Figure 3B). Gene set enrichment analysis (GSEA) demonstrated that the ERBB signalling pathway was significantly enriched in KPCE organoids compared to that in KPC organoids (Figure 3C). In addition, EMT-related genes tended to be enriched in KPCE organoids (Figure 3C). Kyoto Encyclopaedia of Genes and Genomes pathway analysis also revealed that the ERBB signalling pathway and the downstream PI3K/AKT and MAPK signalling pathways were significantly enriched in KPCE organoids (Figure 3D). Among the differentially expressed genes (DEGs) in the ERBB signalling pathway in KPCE organoids, we focused on *Ereg* and *Areg* (Figure 3E). *Ereg* and *Areg* encode epiregulin (EREG) and amphiregulin (AREG), respectively, which are members of the EGF family of proteins and ligands of the ERBB receptors. EREG and AREG can upregulate ERBB downstream PI3K/AKT/mTORC1 and RAS/MAPK pathways and are involved in cancer progression [8,30,31]. Based on these findings, we hypothesised that ELF3 downregulates *Ereg* and *Areg* expression and consequently suppresses the downstream signalling pathway of ERBB receptors.

To test this hypothesis, we generated *Elf3*-overexpressing KPCE (KPCE-*Elf3* OE) organoids. RT-qPCR analysis confirmed that *Elf3* mRNA levels were increased and *Ereg* mRNA levels were significantly decreased in KPCE-*Elf3* OE organoids compared with those in KPCE organoids (Figure 3F). *Areg* mRNA levels tended to be lower in KPCE-*Elf3* OE organoids than in KPCE organoids, though not significantly (Figure 3F). Furthermore, ChIP assays performed on KPCE-*Elf3* OE organoids clarified that ELF3 directly bound to *Ereg* and *Areg* promoter regions (Figure 3G). These results indicate that ELF3 directly downregulates *Ereg* and *Areg* mRNA expression.

EGFR/mTORC1 signal and EMT are upregulated in KPCE gallbladder tumours

To further study the ERBB downstream signalling pathways, we generated organoids from the gallbladder of KPC and KPCE mice at 16–18 weeks of age and performed several analyses on these organoids (Figure 4A). RT-qPCR analysis revealed decreased *Cdh1* expression and increased expression of *Snail* and *Cd133* in KPCE organoids, indicating that dedifferentiation and EMT were induced in KPCE organoids (Figure 4B). We next evaluated their proliferation abilities. Consistent with the result of histological analysis on mouse models, KPCE organoids showed a tendency toward higher proliferation than KPC organoids in proliferation assays (Figure 4C). We also evaluated the protein expression levels of the pathways using western blotting of proteins from these organoids. Interestingly, the phosphorylated EGFR (p-EGFR) level was increased in KPCE organoids (Figure 4D). Ribosomal protein S6 (RPS6) kinase (RPS6K) and eukaryotic translation initiation factor 4E-binding protein 1 (4E-BP1), the main downstream targets of mTORC1, were also highly phosphorylated in KPCE organoids (Figure 4D). Additionally, the expression of the

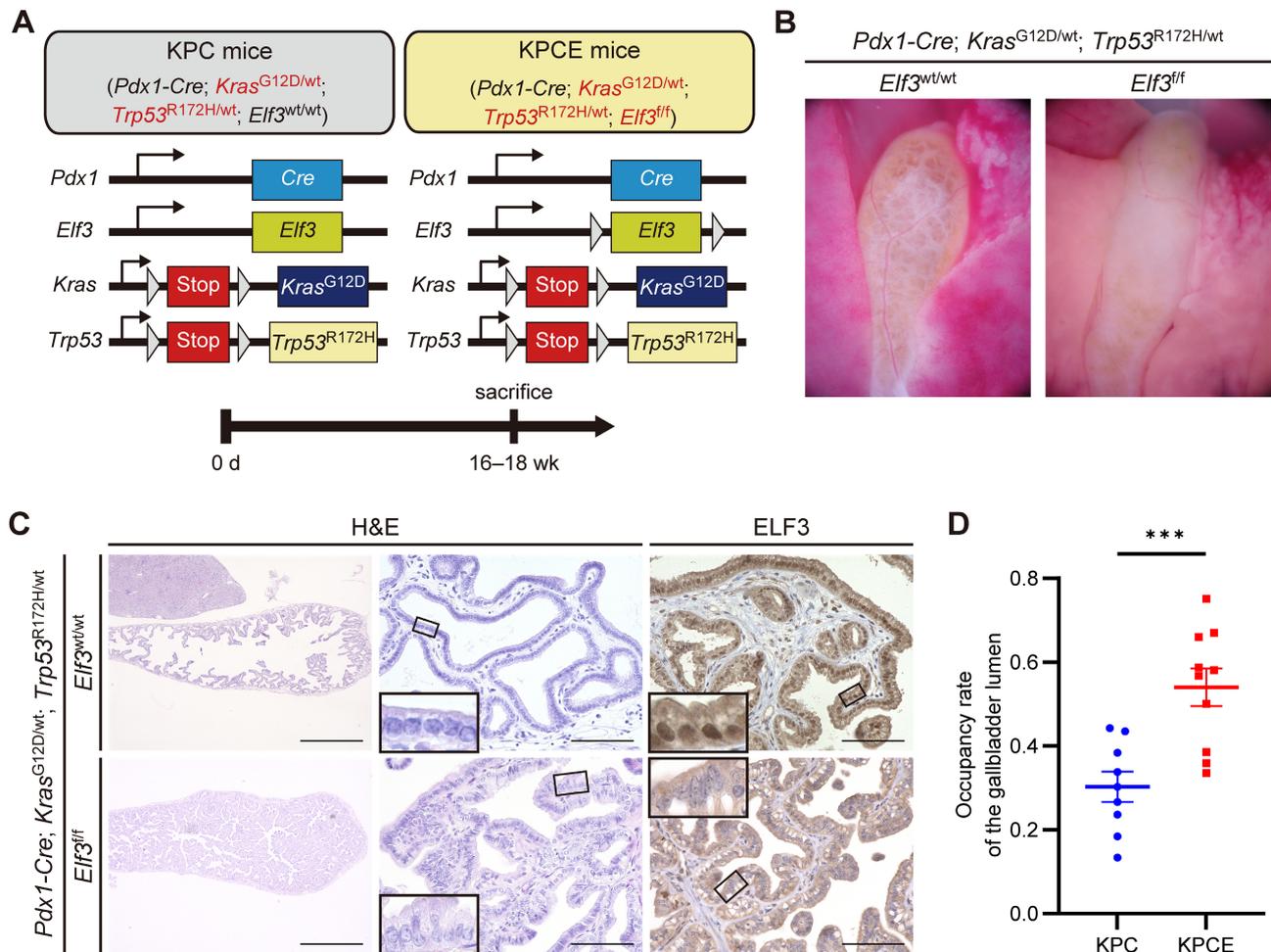


Figure 2. *Elf3* deletion accelerates tumorigenesis in gallbladder in context of *Kras* and *Trp53* mutation in mice. (A) Genetic designs of KPC (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{wt/wt}*) and KPCE (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{fl/fl}*) mice. Arrowheads indicate loxP sites. Cre-mediated deletion induces monoallelic mutation of *Kras* and *Trp53* and biallelic deletion of *Elf3*. (B) Gross appearance of gallbladder of KPC and KPCE mice at 16–18 weeks of age. (C) H&E staining and immunostaining for ELF3 in gallbladder tissues of KPC and KPCE mice at 16–18 weeks of age. Insets show higher-magnification images. Scale bars (left pictures), 1 mm; (middle and right pictures), 100 μm. (D) Quantification of papillary lesion occupancy rates in gallbladder lumen of KPC and KPCE mice (*n* = 9 and 10, respectively). ****P* < 0.01, Student's *t*-test. Occupancy rate was calculated as follows: 1 – inner-gallbladder cavity area/area enclosed by outermost layer of gallbladder wall.

mesenchymal marker proteins, N-cadherin, ZEB1, and SNAI1, was higher in the KPCE than KPC organoids (Figure 4D). To validate these results in mouse models, we performed IHC on KPC and KPCE mice. Notably, EREG, p-EGFR, phosphorylated RPS6 (p-RPS6), and phosphorylated 4E-BP1 (p-4E-BP1) were highly expressed in KPCE gallbladder (Figure 4E). These findings suggest that EREG and p-EGFR expression are increased by *Elf3* deletion, which leads to mTORC1 upregulation and EMT in murine gallbladder organoids.

We further analysed the signalling pathways between EGFR and mTORC1. The PI3K/AKT and RAS/MAPK signalling pathways, which are main targets of ERBB downstream signalling, activate mTORC1 [32]. In addition, AMP-activated protein kinase (AMPK) also modulates mTORC1 activity. AMPK is activated by intracellular ATP/AMP level, low glucose level, hypoxia, ischemia, and heat shock, and then suppresses mTORC1 [32,33]. Western blotting revealed that phosphorylated PI3K and AKT levels were increased, while

the phosphorylated AMPK level was decreased in KPCE organoids (Figure 4D). These findings indicate that mTORC1 in KPCE organoids is modulated by several signalling cascades, including PI3K/AKT and AMPK. To validate this suggestion, we evaluated PI3K/AKT, AMPK, and mTORC1 activity in KPCE organoids following administration of an EGFR inhibitor, afatinib. Thus, EGFR inhibition induced PI3K/AKT suppression and AMPK activation, both of which can suppress mTORC1 activity (supplementary material, Figure S5). This finding validates the idea that the EGFR signal upregulated by *Elf3* deletion activates mTORC1 activity through PI3K/AKT and AMPK.

Elf3 deletion induces the dedifferentiation and mesenchymal phenotype with EREG/EGFR/mTORC1 upregulation in allograft tumour models

Since KPC and KPCE mice died from pancreatic cancers before gallbladder tumours progressed to invasive

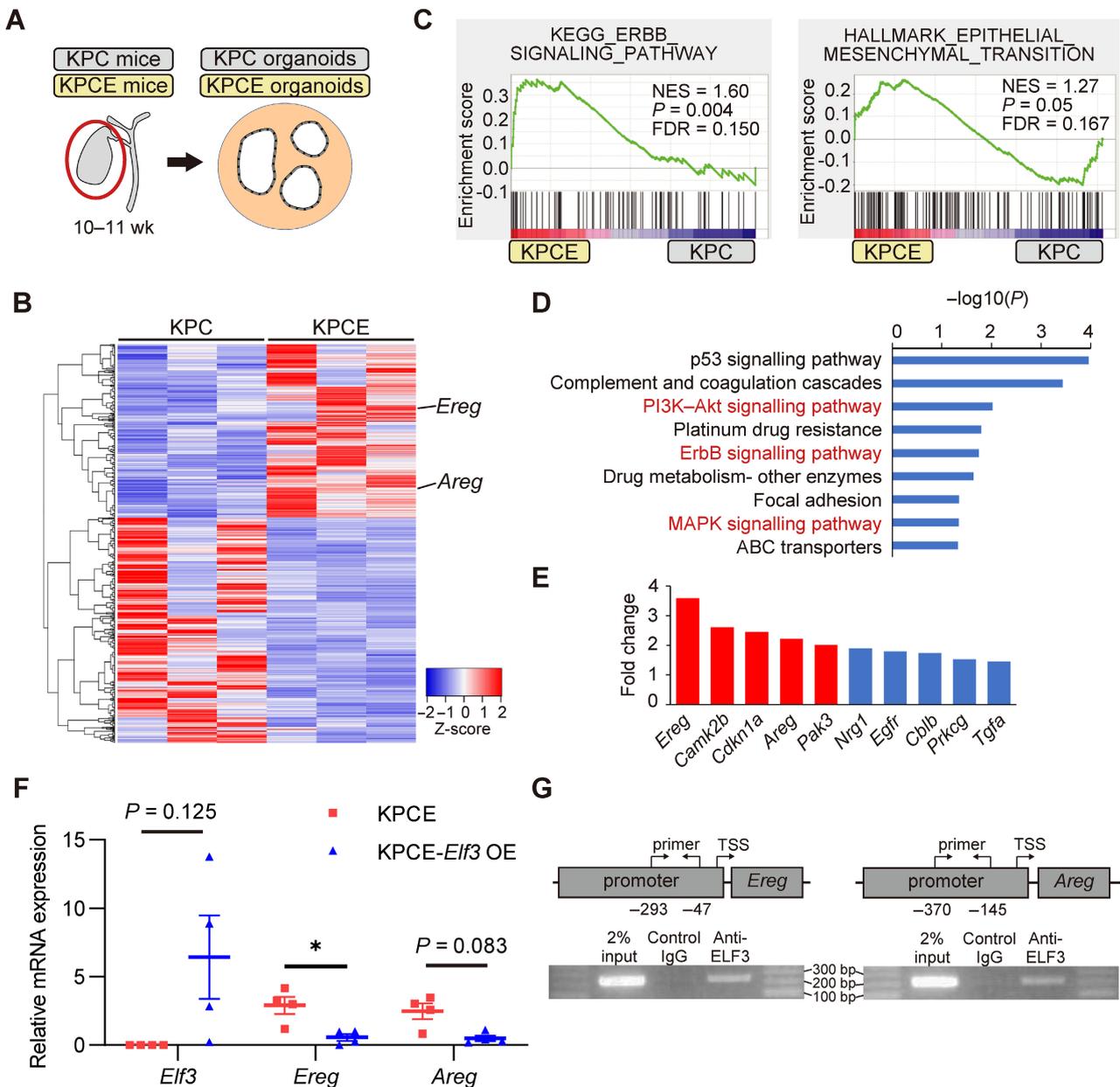


Figure 3. The ERBB signalling pathway is upregulated in KPCE organoids, and ELF3 directly downregulates *Ereg* and *Areg* expression. (A) Organoids were generated from gallbladder of KPC (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{wt/wt}*) and KPCE (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{fl/fl}*) mice at 10–11 weeks of age. (B) Heat map showing DEGs calculated from RNA-seq data. (C) Enrichment plots from gene set enrichment analysis showing that ERBB and EMT signalling pathways were upregulated in KPCE organoids. The normalised enrichment score (NES), nominal *p* value, and false discovery rate (FDR) are shown in each plot. (D) Upregulated signalling pathways in KPCE organoids analysed through Kyoto Encyclopaedia of Genes and Genomes pathway analysis using DEGs calculated from RNA-seq data. (E) Upregulated genes related to ERBB signalling pathway in KPCE organoids. Red and blue bars show the genes satisfied fold change ≥ 2 and < 2 , respectively. (F) RT-qPCR analysis of *Elf3*, *Ereg*, and *Areg* in KPCE and KPCE-*Elf3* OE organoids ($n = 4$ in each group). The expression level of each gene in KPCE and KPCE-*Elf3* OE organoids is shown as a relative ratio to that of KPC organoids ($n = 4$). Means \pm SEM. * $P < 0.05$, paired *t*-test. (G) Schematic illustrations showing promoter regions of *Ereg* and *Areg* and PCR primer sets used in ChIP assay, demonstrating that DNA fragments co-precipitated with anti-ELF3 antibody were amplified using PCR using primer sets specific for respective promoter region. TSS, transcription start site.

cancers, we could not evaluate the GBC phenotype under the *Elf3* deletion condition. To overcome this, we generated a subcutaneous allograft mouse model with KPC and KPCE organoids at 16–18 weeks of age (Figure 5A). Although no significant difference in volume was observed between the allograft tumours derived from KPC and KPCE organoids (Figure 5B), histological analysis revealed that allograft KPC-derived

tumours had more cystic areas than those in KPCE-derived tumours and the ratio of non-cystic to whole sectional area tended to be greater in KPC than in KPCE allograft tumours, suggesting that the accurate volume of KPCE allograft tumours was larger than that of KPC allograft tumours (Figure 5C). Furthermore, allograft tumours of KPCE organoids showed poorly differentiated structures, whereas allograft tumours of

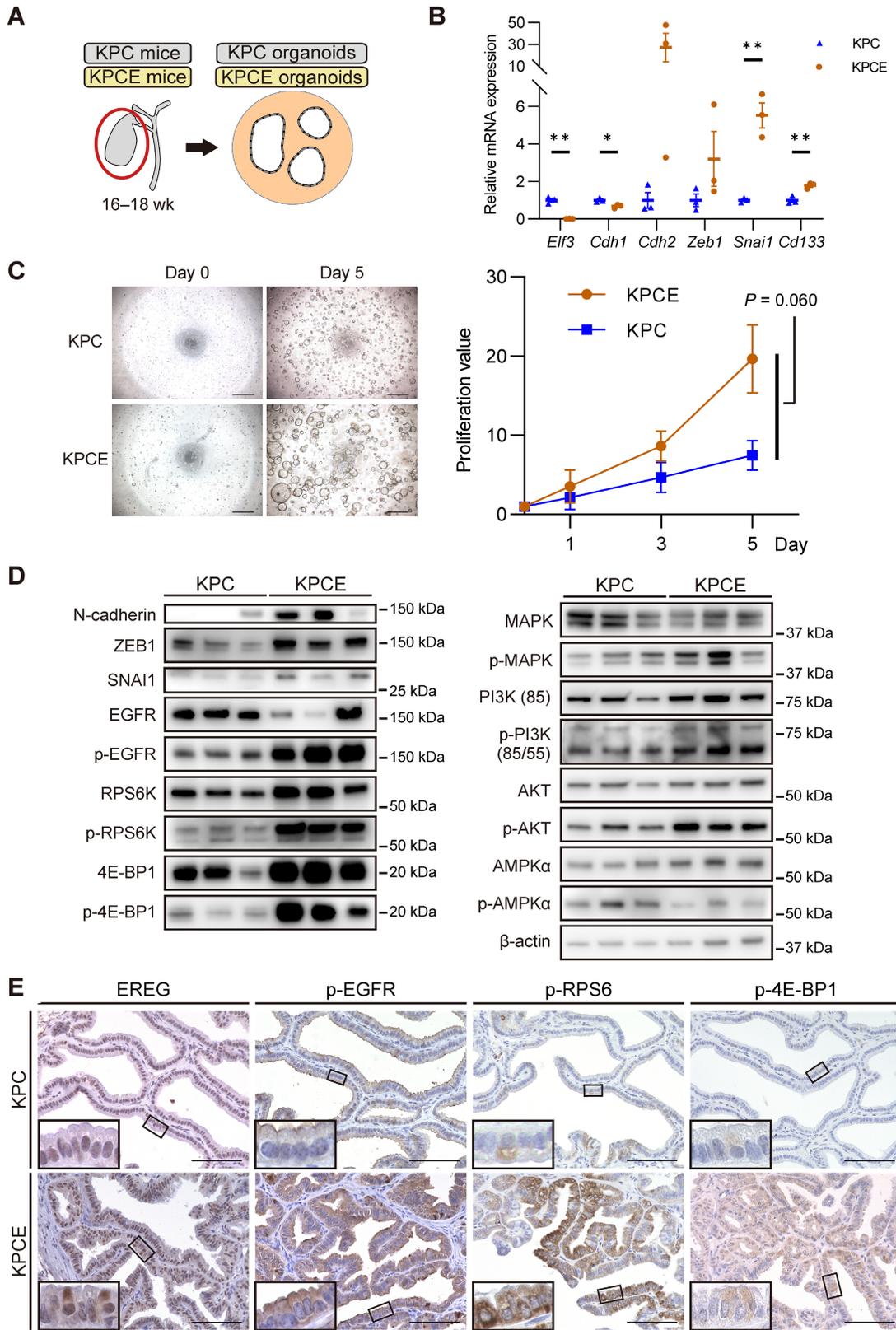


Figure 4. EGFR/mTORC1 signal and EMT are upregulated in KPCE gallbladder tumours. (A) Organoids were generated from gallbladder tissues in KPC (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{wt/wt}*) and KPCE (*Pdx1-Cre; Kras^{G12D/wt}; Trp53^{R172H/wt}; Elf3^{fl/fl}*) mice at 16–18 weeks of age. (B) RT-qPCR analysis on *Elf3*, an epithelial cell marker gene *Cdh1*, mesenchymal cell marker genes (*Cdh2*, *Zeb1*, and *Snai1*), and a cancer stem cell marker gene *Cd133* in KPC and KPCE organoids ($n = 3$ in each group). Means \pm SEM. * $P < 0.05$, ** $P < 0.01$, Student's *t*-test. (C) Cell proliferation assay of KPC and KPCE organoids ($n = 3$ in each group). Means \pm SEM. Student's *t*-test. (D) Western blot analysis using proteins extracted from KPC and KPCE organoids. (E) Representative immunostaining for EREG, p-EGFR, p-RPS6, and p-4E-BP1 in gallbladder tissues of KPC and KPCE mice. Insets show higher-magnification images. Scale bars, 100 μ m.

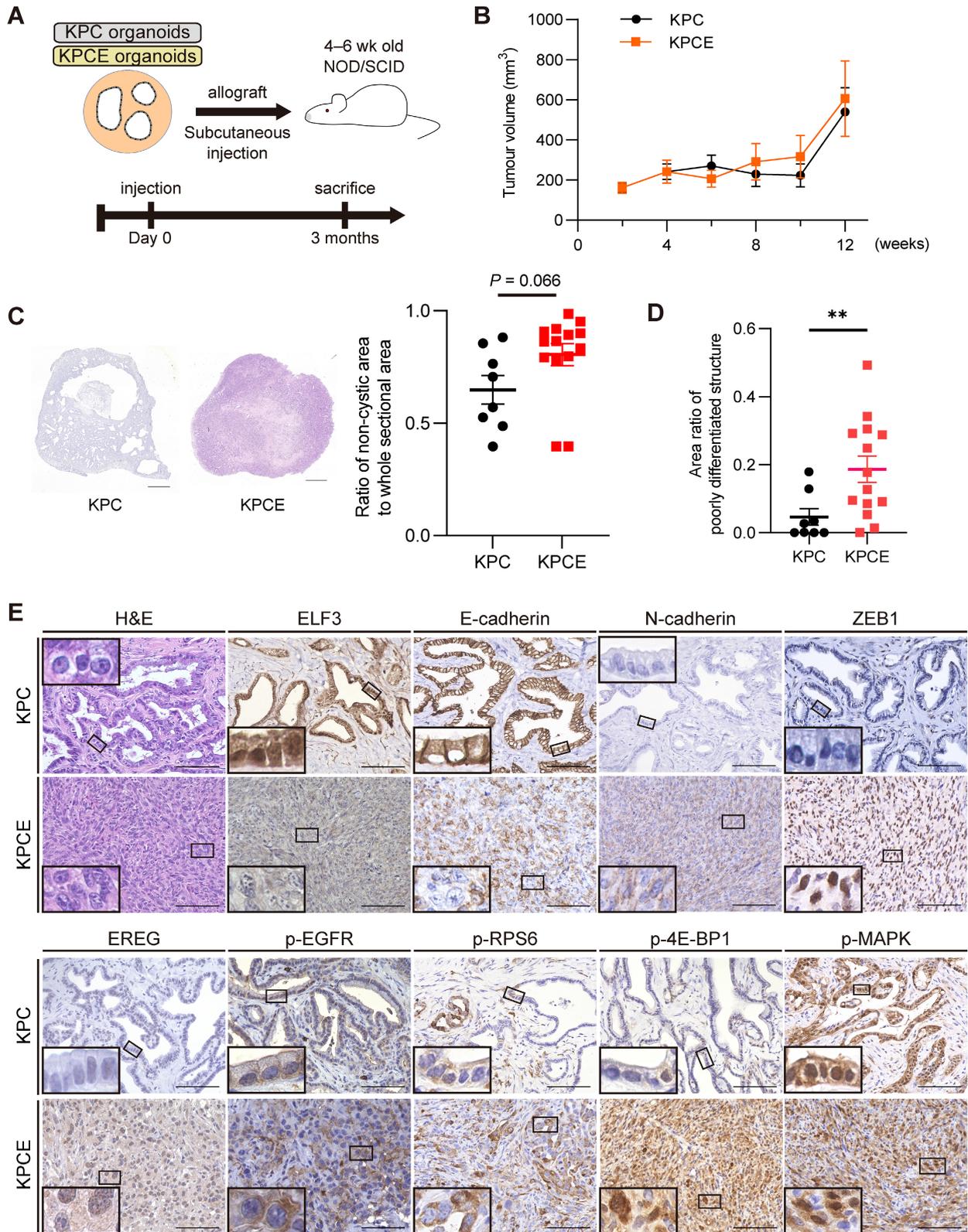


Figure 5. *Elf3* deletion induces dedifferentiation and mesenchymal phenotype with EREG/EGFR/mTORC1 upregulation in allograft tumour models. (A) Schematic illustration of generation method and analysis schedule of allograft mouse model using KPC and KPCE organoids. (B) Graph showing allograft tumour volumes of KPC and KPCE organoids ($n = 6$ and 9 in the KPC and KPCE group, respectively). Means \pm SEM are shown. (C) Representative H&E staining of allograft tumour cross sections for KPC and KPCE organoids and quantification of non-cystic to sectional area ratio ($n = 8$ and 14 in KPC and KPCE groups, respectively). Scale bars, 1 mm. Means \pm SEM. Student's *t*-test. (D) Representative H&E staining and immunostaining for ELF3, E-cadherin, N-cadherin, ZEB1, EREG, p-EGFR, p-RPS6, p-4E-BP1, and p-MAPK in allograft tumour tissues of KPC and KPCE organoids. Insets show higher-magnification images. Scale bars, 100 μ m. (E) Quantification of poorly differentiated structure to sectional area ratio ($n = 8$ and 14 in KPC and KPCE group, respectively). Means \pm SEM. ****** $P < 0.01$, Student's *t*-test.

KPC organoids mainly exhibited tubular structures (Figure 5D,E). IHC on KPCE allograft tumours revealed decreased expression of epithelial cell markers such as E-cadherin and increased expression of mesenchymal cell markers such as N-cadherin and ZEB1 (Figure 5D). IHC also confirmed that EREG, p-EGFR, p-RPS6, and p-4E-BP1 were highly expressed in KPCE allograft tissues compared with those in KPC tissues (Figure 5D). Meanwhile, IHC for phosphorylated MAPK (p-MAPK) expression showed no difference between KPCE and KPC allograft tumours (Figure 5D), although the MAPK signalling pathway was significantly upregulated in KPCE organoids in the pathway analysis, and p-MAPK expression by western blotting tended to be higher in KPCE organoids than in KPC organoids (Figures 3D and 4D).

Additionally, we generated tumour cell lines from both KPC and KPCE allograft tumours and evaluated their cellular properties *in vitro*. KPCE cells showed remarkably higher migration and invasion rates than KPC cells (supplementary material, Figure S6A,B). Furthermore, KPCE cells resulted in more extensive metastases in the livers of allograft mice than KPC cells (supplementary material, Figure S6C). These results indicate that KPCE cells have greater migration, invasion, and metastatic ability than KPC cells. Taken together, these findings suggest that *Elf3* deletion induces dedifferentiation and mesenchymal phenotype in gallbladder tumours in mice.

EREG upregulates the EGFR/mTORC1, and the EREG/EGFR/mTORC1 axis is an important factor in the formation of the mesenchymal structure of KPCE allograft tumours

A previous report revealed that, compared with other EGFR ligands such as EGF and AREG, EREG induces sustained EGFR signalling upregulation [34]. In addition, EREG showed more notable change in its mRNA expression than AREG in our analysis (Figure 3E,F); therefore, we focused on EREG. To elucidate the functional role of EREG in KPCE organoids, we established *Ereg*-knockout KPCE organoids (KPCE-*Ereg* KO) and non-targeting KPCE organoids (KPCE-NT) as controls (Figure 6A). RT-qPCR confirmed that *Ereg* mRNA expression was significantly decreased in KPCE-*Ereg* KO organoids (Figure 6B). Western blotting showed that p-EGFR level was decreased in KPCE-*Ereg* KO organoids, indicating that the ERBB downstream signal was attenuated (Figure 6C). Consistent with this finding, p-4E-BP1 level was decreased in KPCE-*Ereg* KO organoids (Figure 6C). On the other hand, the p-MAPK level was rather increased in KPCE-*Ereg* KO organoids (Figure 6C). In addition, the expression of mesenchymal markers (N-cadherin and SNAIL) decreased in KPCE-*Ereg* KO organoids (Figure 6C). These findings suggest that EREG upregulates the EGFR/mTORC1 pathway, not MAPK, and participates in the induction of EMT in murine gallbladder tumour organoids.

To evaluate the impact of EREG/EGFR/mTORC1 activity on the histological phenotype, we generated subcutaneous allograft mouse models with KPCE-*Ereg* KO and KPCE-NT organoids (Figure 6A). Interestingly, KPCE-*Ereg* KO tumours were significantly smaller than KPCE-NT tumours (Figure 6D). Histological analysis revealed that KPCE-*Ereg* KO tumours mainly showed tubular structures, whereas KPCE-NT tumours showed poorly differentiated structures (Figure 6E). IHC revealed that p-EGFR, p-4E-BP1, and p-RPS6 expression were lower in KPCE-*Ereg* KO tumour tissues than in KPCE-NT tumour tissues (Figure 6E). In addition, IHC showed decreased expression of N-cadherin and ZEB1 in KPCE-*Ereg* KO tumour tissues, indicating that EMT was suppressed in KPCE-*Ereg* KO tumours (Figure 6E). On the other hand, p-MAPK expression in KPCE-*Ereg* KO tumours did not clearly differ from that in KPCE-NT tumours (Figure 6E). These results indicate that the EREG/EGFR/mTORC1 pathway is an important factor in the formation of papillary lesions in the KPCE gallbladder and the mesenchymal structure of KPCE allograft tumours and that the mesenchymal phenotype of KPCE allograft tumours can be at least partially suppressed by *Ereg* deletion.

EGFR/mTORC1 inhibition suppresses cell proliferation and EMT in KPCE organoids

To investigate the pathophysiological role of EGFR/mTORC1 on *Elf3*-deleted organoids, we administered afatinib (an EGFR inhibitor) or rapamycin (mTORC1 inhibitor) to KPCE organoids. Western blotting confirmed suppression of p-EGFR in KPCE organoids treated with afatinib and suppression of phosphorylated RPS6K (p-RPS6K) and p-4E-BP1 in KPCE organoids treated with either drug (supplementary material, Figure S7A). Moreover, western blotting revealed the increased expression of E-cadherin in KPCE organoids treated with afatinib and decreased expression of N-cadherin, ZEB1, and SNAIL in KPCE organoids treated with either drug, suggesting that EGFR/mTORC1 inhibition suppressed EMT (supplementary material, Figure S7A). In cell proliferation assay, KPCE organoids treated with either drug showed significantly lower proliferation activity than those treated with DMSO (supplementary material, Figure S7B). These results suggest that EGFR/mTORC1 signalling plays an important role in tumour progression and EMT induction in the gallbladder of KPCE mice, suggesting that EGFR/mTORC1 inhibition could be a therapeutic option for GBC with an *Elf3* mutation.

ELF3 regulates EREG expression and EGFR/mTOR signalling in human GBC tissues

To determine whether *ELF3* expression affected mTORC1 activity in human GBC tissue, we analysed the RNA-seq data of human GBC published by Pandey *et al* [6]. Most *ELF3* mutations were loss-of-function type; therefore, we divided the samples into wild-type

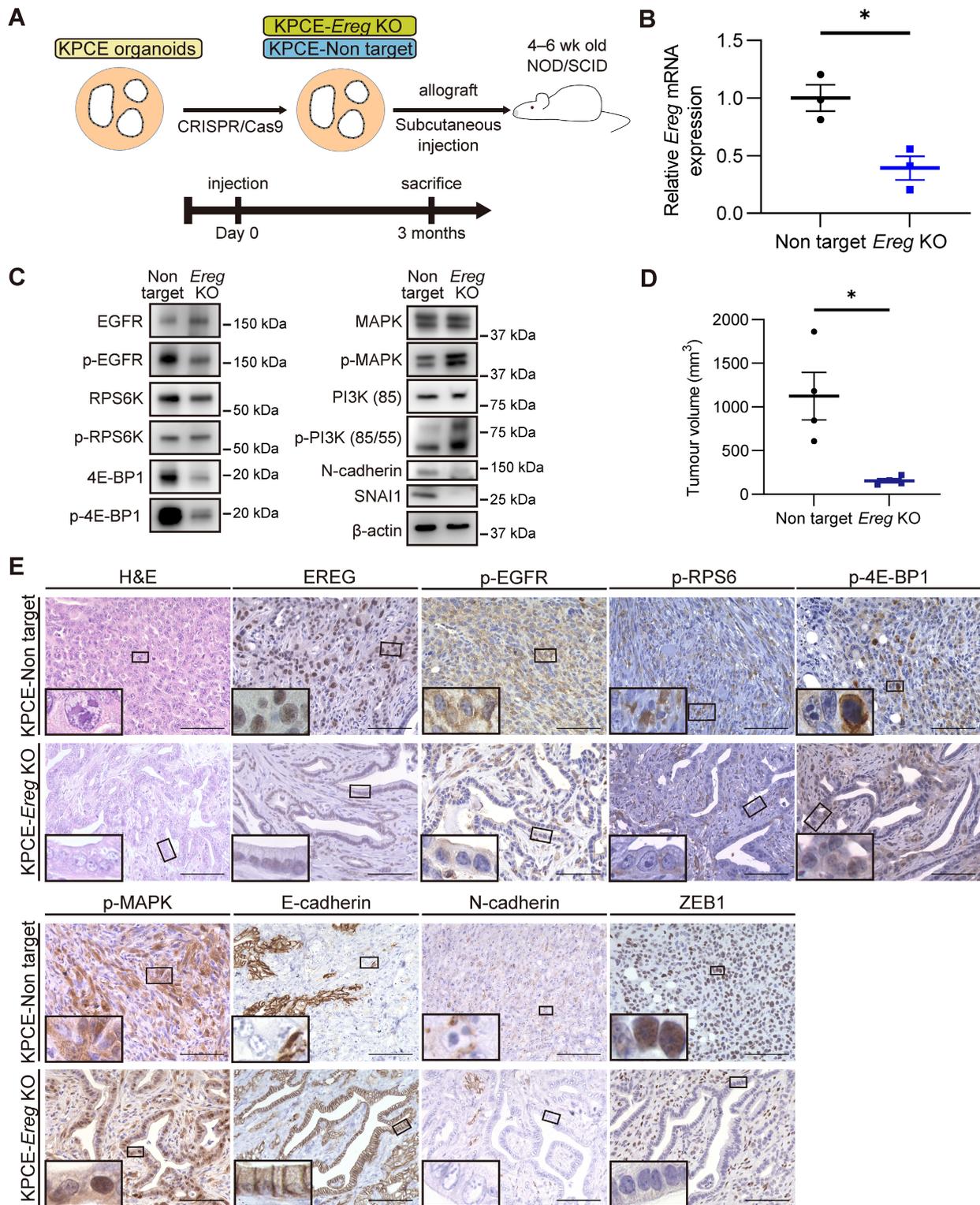


Figure 6. EREG/EGFR/mTORC1 axis is an important factor in the formation of the mesenchymal structure of KPCE allograft tumours. (A) *Ereg*-knockout organoids (KPCE-*Ereg* KO) and negative control organoids (KPCE-NT) were generated from KPCE (*Pdx1-Cre*; *Kras*^{G12D/wt}; *Trp53*^{R172H/wt}; *Elf3*^{fl/fl}) organoids using CRISPR/Cas9 lentivirus system. An allograft mouse model was generated using the organoids. (B) RT-qPCR analysis of *Ereg* expression in KPCE-*Ereg* KO and KPCE-NT organoids ($n = 3$ each). The expression level of KPCE-*Ereg* KO organoids is shown as a ratio relative to that of KPCE-NT organoids. Means \pm SEM. * $P < 0.05$, Student's *t*-test. (C) Western blot analysis of KPCE-*Ereg* KO and KPCE-NT organoids. (D) Volumes of allograft tumours using KPCE-*Ereg* KO and KPCE-NT organoids ($n = 4$ in each group). Means \pm SEM. * $P < 0.05$, Student's *t*-test. (E) Representative H&E staining and immunostaining for EREG, p-EGFR, p-RPS6, p-4E-BP1, p-MAPK, E-cadherin, N-cadherin, and ZEB1 in allograft tumour tissues of KPCE-*Ereg* KO and KPCE-NT organoids. Insets show higher-magnification images. Scale bars, 100 μ m.

ELF3 (*ELF3*-WT) and mutated *ELF3* (*ELF3*-MT) groups ($n = 92$ and 23, respectively) and analysed the transcript differences between the two groups using

GSEA. Consistent with our mouse model data, gene sets related to the EGFR, mTOR, and EMT signalling pathway were significantly enriched in the *ELF3*-MT group

(supplementary material, Figure S8A). Furthermore, there was a mild negative correlation between *ELF3* and *EREG* mRNA expression in the *ELF3*-WT group (supplementary material, Figure S8B). In addition, high *EREG* expression also significantly upregulated the gene sets related to EGFR, mTORC1, and EMT (supplementary material, Figure S8C). These findings suggest that ELF3 regulates *EREG* expression and the EGFR/mTOR signalling pathway in human GBC.

Lastly, we evaluated ELF3/EREG/EGFR/mTORC1 activity in human GBC tissues using IHC. Out of 85 GBC cases, we selected the 47 cases with a T stage of T2 or greater, for which the tissues included both intraepithelial/submucosal portions and invasive portions. We semi-quantitatively evaluated the staining intensities of ELF3, EREG, p-EGFR, and p-4E-BP1 by IHC (supplementary material, Figure S8D) and found that ELF3 expression was significantly reduced in invasive portions compared with that in intraepithelial portions of the tumour. Furthermore, p-4E-BP1 expression was significantly increased, and p-EGFR expression tended to be higher in invasive portions than in intraepithelial/submucosal portions, consistent with our observations with the mouse models. Meanwhile, EREG staining intensity did not differ between intraepithelial/submucosal and invasive portions. Collectively, these results also support the idea that ELF3 regulates EGFR/mTORC1 activity.

Discussion

In this study, we demonstrated that *Elf3* deletion promoted tumour progression and EMT in mice via direct *Ereg* upregulation and subsequent EGFR/mTORC1 signalling pathway activation. Although previous reports showed that ELF3 acted as a tumour suppressor in GBC [6], the underlying molecular mechanism has not been fully elucidated. With the use of an *in vivo* mouse model, this study provides insights into the functional roles of ELF3 in GBC.

We investigated the clinical significance of ELF3 expression in GBC using human samples and showed that low ELF3 expression in GBC was associated with advanced clinical stage and deep tumour invasion. Additionally, the OS of the low ELF3 expression group tended to be shorter than that of the high-ELF3-expression group in human GBC. Consistent with previous reports, these results suggest that ELF3 functions as a tumour suppressor in human GBC development [6,21].

We further analysed the role of ELF3 in GBC using mouse models. *Elf3* deletion resulted in tumour progression in the gallbladder under the condition of *Kras* oncogenic activation and *Trp53* inactivation. Specifically, organoids derived from *Elf3*-deleted gallbladder tumours demonstrated dedifferentiation and EMT. Moreover, transcriptome analysis showed

upregulation of the ERBB signalling pathway. In general, alterations in the ERBB family have been identified in various cancers and are involved in cancer progression [9]. Regarding GBC, somatic ERBB family mutations occurred at a rate of 36.8%, and GBC patients with ERBB pathway mutations had a worse outcome [4]. In a mouse model, constitutive ERBB2 expression in gallbladder epithelium led to gallbladder adenocarcinoma development [35]. Considering these reports, ERBB is one of the most important signalling pathways in GBC.

Among the ERBB signalling pathway-related genes, we focused on *Ereg* and found that ELF3 negatively regulated *Ereg* expression via direct interaction with its promoter region. Moreover, *Ereg* deletion suppressed tumour growth and EMT through EGFR/mTORC1 downregulation. EREG is upregulated in various cancers, and its expression correlates with tumour cell growth, invasiveness, and metastasis [36–39]. By contrast, the role of EREG in GBC has not yet been elucidated. Our results suggest that EREG is a key factor for GBC development through EGFR/mTORC1 upregulation, and ELF3 plays an important role by regulating its expression.

To verify the significance of EGFR/mTORC1 in *Elf3*-deleted gallbladder tumours, we administered an inhibitor of EGFR or mTORC1 to KPCE organoids *in vitro*, which suppressed proliferation and EMT in KPCE organoids, suggesting that EGFR/mTORC1 assume a crucial role in development and EMT induction in *Elf3*-deleted gallbladder tumours. Although the therapeutic effect of EGFR or mTORC1 inhibition on human GBC samples was not evaluated in this study, we found that gene sets related to EGFR, mTOR, and EMT were significantly enriched in human GBC with *ELF3* mutations. Moreover, IHC analyses showed that ELF3 deletion was correlated with EGFR/mTORC1 activity and tumour invasion in human GBC tissues. Considering that *Elf3* deletion leads to EGFR/mTORC1 upregulation, EGFR/mTORC1 inhibition can be an effective therapeutic option for human GBC with *ELF3* alterations. However, further investigation is essential to verify the therapeutic effects in human GBC.

Concerning the EREG/EGFR/mTORC1 axis, some issues remain unelucidated in this study. First, the mTOR signal is frequently activated in human GBC, and high mTOR activation is correlated with the invasive and migratory capacity of GBC cell lines and poor prognosis of human GBC patients [40,41]; however, the molecular mechanism by which mTORC1 induces EMT has not been fully clarified. A previous report revealed that loss of 4E-BP1 function induced EMT via translational activation of SNAI1, which repressed E-cadherin expression and was, thus, considered a key regulator of EMT [42]. In our study, *Snai1* mRNA and SNAI1 expression was higher in KPCE than in KPC organoids. In addition, *Ereg* deletion or EGFR/mTORC1 inhibition in KPCE organoids decreased SNAI1 expression. These results suggest that SNAI1 might play an important role in inducing EMT as the downstream molecule of the EREG/EGFR/mTORC1 axis in KPCE organoids.

Further studies on the relationship between mTORC1 and EMT are needed. Second, AMPK activity was downregulated in KPCE organoids; however, the molecular interaction between AMPK and the ELF3/EREG/EGFR axis has not yet been elucidated. In lung cancer, inhibition of the EGFR tyrosine kinase inhibits glucose uptake and reduces intracellular ATP levels through liver kinase B1 (LKB1), which induces AMPK activation [43], consistent with our results. Therefore, EGFR signalling might suppress AMPK activation through LKB1 downregulation. Because EGFR activity was upregulated in the KPCE organoids in our study, EGFR activation might also induce AMPK suppression in gallbladder tumours. However, the role of AMPK and its association with the ELF3/EREG/EGFR axis in GBC requires further investigation. Third, EREG is a ligand of EGFR, and EREG loss can induce downregulation of MAPK and mTORC1 signalling. Upon comparing KPC and KPCE organoids, we found that both MAPK and mTORC1 were upregulated in KPCE organoids; however, MAPK was upregulated in KPCE-*Ereg* KO organoids, in contrast to mTORC1. This contradiction could be explained by the MAPK-mTORC1 feedback loop. A previous report revealed that mTORC1 inhibition led to MAPK activation through a PI3K-dependent feedback loop [44]. In our study, phosphorylated PI3K was upregulated in KPCE-*Ereg* KO organoids (Figure 6C), indicating that MAPK was activated through the negative feedback loop. Furthermore, in the allograft mouse models using KPCE-*Ereg* KO and KPCE-NT organoids, mTORC1 rather than MAPK was suggested to play an important role in inducing EMT. On the other hand, EGFR inhibition, which led to the inhibition of both MAPK and mTORC1, induced E-cadherin upregulation and N-cadherin downregulation compared with mTORC1 inhibition alone in the *in vitro* inhibitor experiment of KPCE organoids, suggesting that EMT was suppressed more through dual inhibition of MAPK and mTORC1 than through mTORC1 inhibition alone. This indicates that MAPK may also play an important role in EMT. It is clear from our and previous findings that the MAPK and mTORC1 signalling pathways have an intimate and complicated relationship and are important factors in GBC, which warrants a more detailed investigation of these pathways. Fourth, unlike p-EGFR and p-4E-BP1, EREG staining intensity did not differ in comparison between intraepithelial/submucosal and invasive portions in human GBC tissues. While this result is contradictory to the result of human GBC RNA-seq data, it suggests that molecules other than EREG, such as AREG or EGF, might also play important roles between ELF3 and EGFR/mTORC1 in human GBC, and further study on human GBC is essential.

To develop tumours in the murine gallbladder, we induced oncogenic *Kras*^{G12D} and *Trp53*^{R172H} mutation. *TP53* is the most frequently mutated gene in GBC, and *KRAS* mutation is observed in 4–8% of cases, and several altered genes have been identified in the ERBB/RAS/PI3K pathway [4,6]. Therefore, we believe that this combination of gene mutations satisfyingly

imitates the mutation profile of human GBC. Whereas these mutations could lead to the development of pre-cancerous lesions in gallbladder, the mice harbouring these mutations died from pancreatic cancers before gallbladder lesions progressed into advanced cancers. As an alternative method, we used a subcutaneously implanted allograft mouse model. This model could generate invasive cancers derived from KPC and KPCE organoids. However, this approach has the demerit of changes in the tumour growth environment, which is also a limitation of our study. Therefore, establishment of a well-refined GBC mouse model is needed.

In conclusion, we have demonstrated that ELF3 functions as a tumour suppressor in a gallbladder tumour mouse model by directly downregulating *Ereg* expression and consequently downregulating the EGFR/mTORC1 signalling pathway. Further studies are warranted to elucidate the functional role of ELF3 in human GBC.

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Author contributions statement

TN, YN and MS conceived and designed the study. TN performed all the experiments and analysed the data with the assistance of YN, MS, MY, S Mat, YM, SO, HY, HO, TK, S Mar, T Mat and T Mar. TN analysed the clinical data with the assistance of YN. TN and HT performed the sequence analysis and interpreted the data. TN wrote the manuscript, and YN and MS revised it. NU, EH, YK and HS supervised the study. All authors had final approval of the submitted manuscript.

Data availability statement

The data of RNA-seq performed in this study are publicly available in DNA Data Bank of Japan Sequence Read Archive (<https://ddbj.nig.ac.jp/resource/sra-submission/DRA014895>). The other data generated in this study are available from the corresponding author upon reasonable request.

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SUPPLEMENTARY MATERIAL ONLINE

Supplementary materials and methods

Figure S1. Gene recombination in extrahepatic biliary epithelial cells confirmed using X-gal staining in biliary tract of *Pdx1-Cre; Rosa26^{LSL-LacZ}* mice

Figure S2. ELF3 is not essential for development and maintenance of extrahepatic biliary tract in mice

Figure S3. *Elf3* deletion does not cause tumour formation in gallbladder in context of *Trp53* mutation in mice

Figure S4. Genotyping and RT-qPCR analysis for verification of origin of organoids

Figure S5. EGFR signalling regulates mTORC1 activity through PI3K/AKT and AMPK

Figure S6. Dedifferentiation and epithelial–mesenchymal transition (EMT) are induced in KPCE allograft tumour cells *in vitro*

Figure S7. EGFR/mTORC1 inhibition suppresses cell proliferation and EMT in KPCE organoids

Figure S8. Gene sets related to EGFR, mTOR, and EMT are significantly enriched in human GBC tissues with ELF3 mutations

Table S1. Clinical information of 85 patients with GBC

Table S2. Antibodies for IHC, western blotting, and CHIP assay used in this study

Table S3. Primer sets for RT-qPCR, CHIP, and genotyping analyses used in this study