ORIGINAL ARTICLE

Elevation of adenylate energy charge by angiopoietin-like 4 enhances epithelial–mesenchymal transition by inducing 14-3-3γ expression

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Metastatic cancer cells acquire energy-intensive processes including increased invasiveness and chemoresistance. However, how the energy demand is met and the molecular drivers that coordinate an increase in cellular metabolic activity to drive epithelial–mesenchymal transition (EMT), the first step of metastasis, remain unclear. Using different in vitro and in vivo EMT models with clinical patient’s samples, we showed that EMT is an energy-demanding process fueled by glucose metabolism–derived adenosine triphosphate (ATP). We identified angiopoietin-like 4 (ANGPTL4) as a key player that coordinates an increase in cellular energy flux crucial for EMT via an ANGPTL4/14-3-3γ signaling axis. This augmented cellular metabolic activity enhanced metastasis. ANGPTL4 knockdown suppresses an adenylate energy charge elevation, delaying EMT. Using an in vivo dual-inducible EMT model, we found that ANGPTL4 deficiency reduces cancer metastasis to the lung and liver. Unbiased kinase inhibitor screens and Ingenuity Pathway Analysis revealed that ANGPTL4 regulates the expression of 14-3-3γ adaptor protein via the phosphatidylinositol-3-kinase/AKT and mitogen-activated protein kinase signaling pathways that culminate to activation of transcription factors, CREB, cFOS and STAT3. Using a different mode of action, as compared with protein kinases, the ANGPTL4/14-3-3γ signaling axis consolidated cellular bioenergetics and stabilized critical EMT proteins to coordinate energy demand and enhanced EMT competency and metastasis, through interaction with specific phosphorylation signals on target proteins.

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INTRODUCTION

The exponential growth of the primary tumor exposes cancer cells to various microenvironmental stresses, including hypoxic, acidic and inflammatory milieus.1,2 Interestingly, though these constraints can kill cancer cells, they also exert selective pressures on cells to exploit adverse microenvironments by modifying their cellular behavior and selecting for a highly invasive phenotype that facilitates metastasis.3 Metastasis, the spread of cancer cells to distant tissues and/or organs, is a complex and multistep process, making it a challenging therapeutic target.4 Metastasis accounts for >90% of cancer mortality.5 Initiation of the metastatic process is considered a rate-limiting event and resembles the epithelial–mesenchymal transition (EMT) that occurs during embryogenesis and wound healing.1,5 Microenvironmental stimuli can initiate the EMT process by controlling the expression and function of specific transcription factors, such as Snai1, Slug, Twist and ZEB-1.6 During EMT in cancer, epithelial cells lose their cell polarity and cell–cell adhesions are weakened as they adopt a mesenchymal phenotype. This change is accompanied by energy-demanding processes such as cytoskeletal remodeling and increased chemoresistance, ultimately allowing the cells to gain motility and the ability to invade distant organs.7 However, the mechanism that secures the energy supply for EMT competency remains unknown.

Our current understanding of cancer cell metabolism is based primarily on a comparison of metabolic status between primary tumors and normal healthy cells. Cancer cells derive a substantial amount of adenosine triphosphate (ATP) from aerobic glycolysis (that is, the Warburg effect) instead of oxidative phosphorylation to support their anabolic growth and proliferation.8 Cancer cells also exploit glutaminolytic flux, amino acid and lipid metabolism, mitochondrial biogenesis, the pentose phosphate pathway and macromolecular biosynthesis to progress malignancy.9,10 Recent studies have suggested that oncogenes and tumor suppressors function as critical modulators of metabolic reprogramming to support tumor progression.11,12 Hypoxia and transforming growth factor-β (TGF-β), which are known initiators of EMT, can also modulate cancer cell metabolism during tumorigenesis.3,13 Thus, these microenvironmental signals may also alter cancer cell bioenergetics to increase their motility and enhance their invasive capabilities. Despite the significance of EMT in metastasis, little is known about the changes in cellular bioenergetics that occur during this process and whether a demand for metabolic energy is a functional prerequisite. Furthermore, the molecular drivers of metabolic reprogramming in EMT remain unknown.

Using three different in vitro and in vivo EMT models to compliment human cancer biopsies, we identified angiopoietin-like 4 (ANGPTL4) as a key player that adenylates energy charge and coordinates the energy demands required for EMT. We further revealed an ANGPTL4/14-3-3 signaling influences biological processes by protein–protein interactions.
RESULTS
ANGPTL4 increases cellular bioenergetics needed for EMT competency

ANGPTL4 is best known for its role as an adipokine required for systemic glucose and lipid metabolism. ANGPTL4 has been identified as a prometastatic gene that is involved in tumor growth, anoikis resistance, angiogenesis and tumor invasion. However, its involvement in EMT, particularly its function in energy homeostasis in the cellular level, is unknown. Herein, to examine changes in energy demand during metastasis, we first measured the energy charge and examined the expression of ANGPTL4 in human tumor biopsies. Clinical samples showed significant correlations between the energy charge status and tumor grades, with the higher-grade metastatic tumors exhibiting between 26.7% and 106% increases in energy charge compared with cognate stage I tumors (Figure 1a; Supplementary Table S1). The energy charge status of the cells is indicative of the cellular metabolic activity. Importantly, our analysis also revealed that human metastatic cancers expressed higher ANGPTL4 expression compared with their lower-grade counterparts, suggesting a role for ANGPTL4 in regulating cellular bioenergetics for metastasis (Figure 1b).

Next, we carried out a series of molecular and biochemical analyses to examine whether change in cellular bioenergetics was observed in in vitro EMT of cancer cells. First, we established three in vitro EMT models using the polarized gastric carcinoma cell line MKN74. In two models, EMT was initiated by either hypoxia (1% O2, Figures 1c–e) or TGF-β (Supplementary Figures S1a–c) to mimic the microenvironment of stress-induced EMT. The third model utilized the transgenic MKN74 cell line harboring a Snai1-ER transgene (MKN74Snai1ER) that allows direct initiation of EMT by 4-hydroxytamoxifen (4-OHT) (Supplementary Figures S1d–f). Upon exposure to either microenvironmental stress or 4-OHT, the colonies of MKN74 underwent EMT, with diminished E-cadherin staining bordering the epithelial cells after 48–96 h (Figure 1c and Supplementary Figures S1c and f). Real-time PCR and immunoblot analysis showed the expression of critical epithelial genes (E-cadherin (CDH1); discoidin domain receptor 1 (DDR1); receptor-tyrosine kinase (ErBb3)) was downregulated, with a concomitant increase in mesenchymal gene expression (Snai1; zinc finger E-box-binding homeobox 1 (ZEB-1); Figures 1d and e and Supplementary Figures S1a, b, d, and e). Regardless of the EMT stimuli, we detected an increase in glucose uptake and energy charge in cancer cells, indicative of increased cellular metabolic activity during EMT (Figures 1f and g). Importantly, ANGPTL4 mRNA and protein levels, specifically the C-terminal fibronogen-like form (cANGPTL4), were elevated during EMT (Figure 1h). An increase in cANGPTL4 expression (Supplementary Figure S1g) and augmented cellular metabolic activity were also observed in hypoxia-induced EMT of MCF7, II4, HSC and HepG2 cancer cell lines (Figure 1i and Supplementary Figure S1h). Our observations indicate that an increase in cellular bioenergetics in cancer cells undergoing EMT is widespread.

To confirm a role for cANGPTL4 in metabolic activity of cancer cells during EMT, we investigated the effects of ANGPTL4 deficiency (through knockdown or immunoneutralization) and treatment of recombinant human cANGPTL4 (rh-cANGPTL4) on energy charge and EMT. The increases in glucose uptake and energy charge were suppressed by either neutralizing cANGPTL4 antibody (α-cANGPTL4) or small interfering RNA against ANGPTL4 (ΔANGPTL4) in MKN74 (Figures 2a and b and Supplementary Figures S2a and b). MCF7, II4, HSC and HepG2 cancer lines during hypoxia-induced EMT (Supplementary Figure S1h). Exogenous rh-cANGPTL4 alone elevated 2-deoxyglucose uptake associated with increased GLUT1 protein expression and augmented the energy charge status during EMT (Figures 2a–c and Supplementary Figure S2c). Importantly, an impaired ANGPTL4-mediated increase in energy charge delayed EMT. At 48 and 96 h after EMT induction, the ANGPTL4-deficient cells still retained their epithelial-like morphology characterized by clear E-cadherin staining (Figure 2d and Supplementary Figures S2d and f). Quantitative PCR (qPCR) and immunoblot analysis revealed a delay in the expression of critical EMT genes (Figure 2e and Supplementary Figures S2e and g). Conversely, MKN74 cells treated with rh-cANGPTL4 resulted in an incomplete disruption of E-cadherin staining (Figure 2f), repressed expression of epithelial markers and significant elevation of mesenchymal markers (Figure 2g). Taken together, these observations suggest that ANGPTL4 modulates cellular bioenergetics needed for EMT competency.

ANGPTL4 modulates energy charge in metastasis

EMT is an integral step of metastasis. An increase in cellular metabolic activity could fuel energy-demanding processes required for enhanced invasiveness to the cancer cells. To further strengthen the link between ANGPTL4 and energy-demanding EMT, we generated a dual-inducible EMT xenograft mouse model. The MKN74Snai1ER-rhANGPTL4 cell line for tumor xenografts has a doxycycline (Dox)-inducible shANGPTL4 expression plasmid introduced into MKN74ShiER cells. This cell line undergoes EMT when treated with 4-OHT, whereas endogenous ANGPTL4 will be suppressed when treated with Dox (Supplementary Figure S3a). Mice on a normal diet and injected with 4-OHT harbored significantly more lung metastases than the vehicle control mice or mice treated with 4-OHT and fed a Dox diet (Figure 3a). Next, we used human-specific TBP primers to identify and quantify human MKN74ShiER cells that have metastasized to the recipient mouse lung. The 18S primers were used to detect both human and mouse cells for normalization. In line with the above observation, the qPCR result showed the presence of human cancer cells in mouse lung sections after 4-OHT induction (Figure 3a). The cANGPTL4 protein was elevated in the xenograft tumors of 4-OHT-treated mice, whereas mice fed simultaneously with a Dox diet exhibited in vivo suppression of cANGPTL4 in MKN74ShiER-rhANGPTL4-derived tumors (Figure 3b). Histological analysis of tumor sections revealed that vehicle-treated control tumors showed strong defined E-cadherin and laminin 332 staining (Figure 3c). The 4-OHT-treated tumors lost E-cadherin staining at the cell–cell borders, with clear disruption of basement membrane laminin 332 compared with vehicle-treated tumors. After in vivo suppression of ANGPTL4 by the Dox diet, the punctuated E-cadherin staining was diminished (Figure 3c). Gelatin zymography revealed elevated activities for proinvasive matrix metalloproteinases 2 and 9 in 4-OHT-treated tumors, indicating an aggressive tumor phenotype (Supplementary Figure S3b). The in vivo 2-deoxyglucose imaging showed that 4-OHT-treated tumors had significantly elevated glucose uptake compared with vehicle-treated tumors (Figure 3d). The suppression of ANGPTL4 in 4-OHT-treated tumors reduced the 2-deoxyglucose uptake to levels comparable to vehicle-treated tumors (Figure 3d). Consistent with our findings, 4-OHT-treated tumors expressed higher levels of GLUT1 than vehicle- and Dox-treated tumors. ANGPTL4 suppression in 4-OHT-treated tumors resulted in decreased GLUT1 expression (Figure 3b). Importantly, in vivo xenograft tumors exhibited a similar elevation in the energy charge status during EMT, and this elevation was diminished by Dox diet-induced ANGPTL4 depletion (Figure 3e). Taken together, these observations indicate a role for cANGPTL4 in cellular metabolic changes during EMT in vivo.

ANGPTL4 is a molecular driver of metabolic changes during EMT

The phosphatidylinositol-3-kinase (PI3K)/AKT and AMPK (adenosine monophosphate-activated protein kinase) pathways were identified as two major regulatory conduits for cellular
bioenergetics in cancer cells. To gain further insights into how cellular bioenergetics affects EMT, dominant-negative (dn) and constitutive active forms (ca) of AKT and AMPK were introduced into MKN74 cells (Supplementary Figure S3c). As expected, MKN74control cells underwent EMT after 48 h of hypoxia as evidenced by diminished immunofluorescence signal for E-cadherin at the cell–cell junction (Supplementary Figure S3d), reduced E-cadherin and concomitant increased Snail1 protein...
Figure 2. ANGPTL4 increases cellular bioenergetics for EMT competency. (a) Fluorescence-activated cell sorting (FACS) analysis of the fluorescent glucose analog 2-NBDG uptake and (b) percentage increase in energy charge in MKN74 and MKN74Snai1ER cells after the indicated treatments. (c) Immunoblot analysis of GLUT1 expression in MKN74 cells after the indicated treatments. (d, f) Immunofluorescence staining of E-cadherin in hypoxia-treated MKN74 cells in the presence of neutralizing human ANGPTL4 antibodies (α-cANGPTL4) (d) and recombinant human ANGPTL4 (rh-cANGPTL4) -treated MKN74 (f) at the indicated time intervals. Cells were counterstained with DAPI (blue) for nuclei and phalloidin (red) for actin cytoskeleton. Scale bar = 40 μm. (e, g) Relative mRNA expression (left panel) and immunodetection (middle and right panels) of EMT markers in hypoxia-treated MKN74 cells exposed to α-cANGPTL4 (e) and rh-cANGPTL4 (g) at the indicated time intervals. For immunoblot analyses, representative immunoblot pictures and densitometric quantification plots are shown. Loading controls for the immunoblot analyses were from the same sample. For qPCR, TBP was used as reference gene. Data are represented as mean ± s.d. from at least three independent experiments. *P < 0.05, **P < 0.01, ***P < 0.001; n.s., not significant.
levels (Figure 4a). In normoxic condition, the colony morphology of MKN74 cells harboring dnAKT (MKN74 dnAKT), caAMPK (MKN74 caAMPK) and dnAMPK (MKN74 dnAMPK) resembled that of MKN74 CTRL (empty vector). The MKN74 caAKT cells revealed a reduced E-cadherin signal compared with MKN74 CTRL cells (Supplementary Figure S3d). Upon exposure to hypoxia-induced EMT, MKN74caAKT and MKN74caAMPK cells exhibited an earlier (24 h) and more pronounced EMT phenotype than corresponding MKN74CTRL cells (48 h) (Supplementary Figure S3d). Indeed, immunoblots for E-cadherin and Snai1 indicated that dnAKT or dnAMPK was attenuated (Figure 4a), whereas caAKT and caAMPK facilitated hypoxia-induced EMT (Figure 4a). Importantly, the enhanced EMT observed in MKN74 caAMPK cells was associated with an increase in energy charge compared with MKN74 CTRL (Figure 4b). Conversely, the delayed EMT in MKN74dnAKT and MKN74dnAMPK was accompanied by the abolishment of the rise in energy charge for EMT of these cells (Figure 4b). Similar observations were also obtained using 4-OHT-induced EMT of MKNSnai1ER cells (Figure 4c).

The above observations prompted us to examine whether ANGPTL4 could modulate these two pathways. MKN74 CTRL cells exposed to hypoxia and 4-OHT-treated MKNSnai1ER displayed increased phospho-activation of AKT and AMPK (Figures 4d and e). We examined the temporal phosphorylation profiles of AMPK and AKT during rh-ANGPTL4-induced EMT. The expression of phospho-AMPK peaked by 48 h and returned to basal level by 96 h. Notably, the expression of phospho-AKT peaked at 72 h and remained elevated even at 96 h (Figure 4f). This suggests that ANGPTL4-mediated activation of AMPK and AKT may play nonredundant roles to metabolically prime cancer cells for EMT. The analysis of MKN74Snai1ER:shANGPTL4-derived xenograft tumors and MCF7, II4, HSC and HepG2 cancer cell lines also revealed similar phospho-activation of AKT (Figure 4g and Supplementary Figures S3c and e). Higher AKT activity was also detected in patients’ metastatic tumors compared with lower-grade tumors (Figure 4h). Taken together, these observations suggest that ANGPTL4 modules the AKT and AMPK pathways and that the consolidation of cellular bioenergetics is important for EMT competency.

The ANGPTL4/14-3-3γ axis coordinates energy demand for EMT

The 14-3-3 protein family consists of adaptor proteins that play critical roles in numerous important cellular functions via binding
to specific phosphorylation signals on target proteins. Previous studies showed that ANGPTL4 influenced the expression of 14-3-3 adaptor protein family during wound healing, reminiscent of EMT. Thus, we hypothesize that ANGPTL4 consolidates cellular bioenergetics to enhance EMT by regulating several key regulatory networks.

In response to EMT-inducing signals, only the expression of 14-3-3 was consistently elevated by ANGPTL4 in MKN74CTRL and

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**Figure 4.** ANGPTL4 is a molecular driver of metabolic changes during EMT. (a) Immunoblot analysis of E-cadherin and Snai1 and (b) percentage increase in energy charge in MKN74 cells (MKN74CTRL, MKN74dnAKT, MKN74caAMPK and MKN74dnAMPK) under normoxic or hypoxic conditions. (c) Percentage increase in energy charge in 4-OHT-induced EMT in MKN74Snai1ER cells under similar modifications in (b). (d–g) Immunoblot analysis of pAKT (S473), total AKT, pAMPK (T172) and AMPKα in MKN74ctrl, MKN74Snai1ER (e), MKN74ΔANGPTL4 (f) cells and MKN74Snai1ER:shANGPTL4-derived xenograft tumors (g) after indicated treatments. (h) Immunoblot analysis of pGSK3α/β (S21/9) and total GSK3α/β in breast, colorectal, gastric and head and neck tissues at various tumor stages. For immunoblot analyses, representative immunoblot pictures and densitometric quantification plots are shown. Loading controls for the immunoblot analyses were from the same sample. Data are represented as mean ± s.d. from at least three independent experiments. *P < 0.05, **P < 0.01, ***P < 0.001; n.s., not significant.
MKN74_CTRBL (Figures 5a and b and Supplementary Figures S4a and b). The remaining 14-3-3 isotypes were unchanged. Consistently, the suppression of ANGPTL4 resulted in a significant decrease in the expression of 14-3-3γ, whereas the exposure of MKN74_CTRL cells to rh-cANGPTL4 enhanced 14-3-3γ expression (Figure 5c and Supplementary Figure S4c). Similar changes in 14-3-3γ expression were also observed in four other cancer lines during EMT as well as in metastatic human tumor biopsies (Figure 5d and Supplementary Figures S4d–g). Notably, the knockdown of 14-3-3γ in MKN74_CTRL attenuated the expression...
of mesenchymal markers (Supplementary Figures S4h and i). These observations suggest that ANGPTL4 may exert its effect on cancer cell bioenergetics and EMT competency by modulating the expression of 14-3-3γ.

Many studies showed that 14-3-3γ can associate with diverse proteins, modulating complex regulatory networks, but how 14-3-3γ is transcriptionally regulated remains unclear. We observed that rh-cANGPTL4 increased the expression of 14-3-3γ mRNA in MKN74 cells in 6 h (Supplementary Figure S4j). To obtain further insight into the signaling pathways, we performed an unbiased kinase inhibitor screen to identify key signaling mediators whose inhibition would suppress ANGPTL4-mediated upregulation of 14-3-3γ mRNA (Supplementary Figure S4k). Coincidently, qPCR followed by Ingenuity Pathway Analysis revealed a convergence to the PI3K/AKT and mitogen-activated protein kinase signaling cascades and implicated several transcription factors known to be involved in cancer progression (Figure 5e). Immunoblot analysis showed that three of these transcription factors, namely CREB, cFOS and Stat3, were phospho-activated during EMT (Figure 5f and Supplementary Figure S4l). Next, we performed transient small interfering RNA knockdown of TSC2 (Figures 6a and Supplementary Figure S5e) and patients’ benign counterparts, highlighting the importance of 14-3-3γ/Snai1 complexes in 4-OHT-treated tumors than in vehicle-treated tumors from mice fed a normal diet (Figure 6h and Supplementary Figure S6b). PLA analysis of tumor biopsies from patients further confirmed that higher-grade, more aggressive tumors harbored significantly more 14-3-3γ/Snai1 complexes than their benign counterparts, highlighting the importance of 14-3-3γ/Snai1 complexes during malignancy (Figure 6l and Supplementary Figures S6c–e). These observations underscore the ANGPTL4/14-3-3γ signaling axis as an important driver of the observed changes in bioenergetics during EMT.

The interaction of 14-3-3γ with Snai1 was proposed to stabilize Snai1 binding on the E-cadherin promoter to facilitate transcriptional repression. We showed that reduced 14-3-3γ/Snai1 complex in MKN74/ANGPTL4 decreased Snai1 occupancy of the E-cadherin promoter during hypoxia-induced EMT, indicating that the repression of E-cadherin gene expression was impaired (Figure 6j). Furthermore, EMT was delayed in the absence of ANGPTL4, highlighting the importance of 14-3-3γ/Snai1 complexes during malignancy. Altogether, our data revealed a previously unknown mechanism where the ANGPTL4/14-3-3γ signaling axis coordinates energy demand and stability of critical EMT proteins necessary to drive EMT (Figure 6k).

**DISCUSSION**

Metastatic cancer cells acquire energy-intensive processes including increased invasiveness and chemoresistance. However, how the energy demand is met and the mechanisms that coordinate cellular metabolic activity with EMT remain unclear. Using experimental and clinical metastatic tumors, we showed that an increase in adenylate energy charge during cancer cell EMT is widespread and is required for metastasis competency. We identified ANGPTL4, a matricellular protein with no intrinsic enzymatic activity, as a novel extracellular driver of cellular metabolic activity during EMT. ANGPTL4 increased energy charge for EMT through 14-3-3γ-dependent pathways.

**Figure 5.** ANGPTL4 regulates 14-3-3γ expression to facilitate EMT. (a–c) Relative fold change of the various 14-3-3 isoforms (protein level) in hypoxia-induced MKN74Ctrl and MKN74/ANGPTL4 (a), 4-OHT-induced MKN74/Snai1ER (b) and rh-cANGPTL4-induced MKN74Ctrl conditions (c). (d) Immunoblot analysis of 14-3-3γ in breast, colorectal, gastric and head and neck tissues at various tumor stages. (e) Analysis of map of kinases (solid line) whose inhibition attenuates ANGPTL4-induced 14-3-3γ expression and possible downstream transcriptional regulators (dotted line). (f) Relative fold changes of the various transcription factors (phospho- and total-proteins) in hypoxic-induced (left panel) and rh-cANGPTL4-treated (right panel) MKN74Ctrl cells. The plots were derived based on western blots shown on Supplementary Figure S4l. (g) Representative ChIP assay performed using preimmune IgG (p.i.) or antibodies against pCREB, pSTAT3 or p-cFOS in rh-cANGPTL4-treated MKN74 cells. Putative specific binding sites for respective transcription factors spanning the promoter region of 14-3-3γ are shown on the left panel, along with their cognate primers. A control region 2 kb upstream of the promoter served as a negative control. For the immunoblot analyses, densitometric quantification plots are shown. Loading controls were from the same sample. Data are represented as mean ± s.d. from at least three independent experiments. *P < 0.05, **P < 0.01.
phosphorylation signals on target proteins, consolidated cellular bioenergetics and stabilized critical EMT proteins to coordinate energy demand and enhanced EMT.

Many studies have linked ANGPTL4 to metastasis, from cell motility, angiogenesis to vascular dysfunction to facilitate distal dissemination. Indeed, ANGPTL4 has been recently implicated in EMT of non-small-cell lung cancer; however, a role for ANGPTL4 in cellular metabolic changes is still unknown. We identified ANGPTL4 as a molecular driver of EMT-enriched metabolic changes as supported by several lines of experimental evidence. First, in vitro and in vivo EMT models showed that cancer cells deficient in ANGPTL4 were unable to increase the cellular energy charge during EMT, resulting in reduced metastasis. Molecular and biochemical analyses confirmed the down- and

Figure 6. The ANGPTL4/14-3-3γ axis coordinates energy demand during EMT. (a, e) Immunoblot analysis of 14-3-3γ expression after TSC2$_{939}$ (a) or Snai1 (e) immunoprecipitation in MKN74$_{CTRL}$ and MKN74$_{ANGPTL4}$ cells. (b–d and f–i) Proximity ligation assay (PLA) quantification of 14-3-3γ/TSC2$_{939}$ and 14-3-3γ/Snai1 interactions in MKN74$_{CTRL}$ and MKN74$_{ANGPTL4}$ cells (b, f), xenograft tumors (c, h), and tumor biopsies from patients (d, i) after the indicated treatments or with indicated disease stages. (g) Immunodetection of the indicated proteins in hypoxic-induced MKN74$_{CTRL}$ and MKN74$_{ANGPTL4}$ cells at 0 and 48 h. (j) Representative ChIP results using anti-Snai1 (upper panel) followed by re-ChIP with anti-14-3-3γ (lower panel) in MKN74$_{CTRL}$ and MKN74$_{ANGPTL4}$ cells under normoxic or hypoxic conditions. Preimmune serum (IgG) served as control. (k) Schematic diagram depicting the coordination of metabolic reprogramming by the ANGPTL4/14-3-3γ axis during EMT. Data are represented as mean ± s.d. from at least three independent experiments. *P < 0.05, **P < 0.01. Loading controls for the immunoblot analyses were from the same sample.
up-regulation of epithelial and mesenchymal markers, respectively. Histological analysis of xenograft tumors also showed increased matrix metalloproteinase activities and a disrupted laminin 332 phenotype. Second, ANGPTL4 activated the proglycolysis AKT and AMPK pathways to enhance the glycolytic phenotype and increase the cellular energy charge necessary to fuel biological processes of EMT. The active remodeling of the actin cytoskeleton requires ATP binding and hydrolysis to confer invasive abilities to the cancer cells.31-34 AMPK has been proposed to play both ‘conditional’ tumour suppressor and ‘contextual’ oncopgene roles.35,36 In cancer cells, AMPK functions as an energy sensor that inhibits the main anabolic processes that cancer cells needed for proliferation and growth, similarly observed in cells at 0 h. During EMT, AMPK was rapidly phospho-activated to carry out its ‘contextual’ oncogene role to increase glucose uptake and ATP production. Upon completion of EMT, AMPK returned to basal level. Independently, phospho-activation of AKT modulates the mTOR machinery regulating energy flux for EMT progression. After EMT, phospho-activated AKT remained elevated for subsequent metastatic events, such as resistance to anoikis.29 Finally, ANGPTL4 regulates the expression of specific 14-3-3 proteins that coordinates the energy supply during EMT. Notably, these changes were also detected in clinical tumor biopsies and strongly correlated with the stage and aggressiveness of the tumors, further underscoring the clinical relevance of our findings. TGF-β and hypoxia are well-established stimuli of EMT. The human metastasis-related ANGPTL4 gene is transcriptional upregulated by TGF-β29 and hypoxia.14,15 Hypoxia stabilizes the hypoxia-inducible factor-1α that upregulates several genes to promote metastasis.35,36 The hypoxia-inducible factor-1α has been shown to directly upregulate ANGPTL4 expression that facilitates transendothelial migration and increases angiogenesis.37-40 We showed that the expression of ANGPTL4 was increased during hypoxia-induced EMT. The ANGPTL4-mediated upregulation of 14-3-3-γy expression stabilizes Snai1-mediated repression of E-cadherin and elevated adenylate charge to fuel the various biological activities required for EMT.

Metastasis necessitates the cooperative effects of multiple signaling events and biological progresses.1,13,41 A key mechanistic insight is that ANGPTL4 regulates 14-3-3-γy expression and contributes to an important new signaling axis that coordinates multiple biological processes for metastasis. Unlike protein kinases, 14-3-3 adaptor protein associates with many mediators of key signaling hubs to either augment or attenuate the activities of specific pathways. We showed that ANGPTL4/14-3-3γy adaptor protein interactome with TCS and Snai1, respectively. The TSC2 pathway coordinates catabolic processes, such as glycolysis, to provide the necessary substrates for the PI3K/AKT signaling pathways to signal anabolic metabolism. The phospho-inhibition of TSC2 by ANGPTL4-mediated AKT activation altered cellular energy charge and primed cancer cells for EMT. ANGPTL4 also modulated Snai1 complex stabilization through the 14-3-3 y/Snai1 interaction to stably suppress E-cadherin gene expression and augment the EMT progress. Interestingly, the loss of E-cadherin has been reported to trigger a glycolytic switch that favors EMT.42,43 Previous studies showed that ANGPTL4 triggers the 14-3-3 adaptor protein to sequester the pro-apoptotic Bad protein from mitochondria, conferring resistance to anoikis and favoring tumor survival and growth that has been suggested to be a prerequisite for EMT.29 In addition, ANGPTL4/14-3-3 is involved in rapid integrin recycling during cell migration, an important biological process for metastasis.21

Our study shows that an increase in adenylate energy charge by angiotensin-like 4 protein enhances EMT by inducing 14-3-3γy expression. Our findings of the ANGPTL4/14-3-3γy signaling axis revealed the integration of protein–protein interactome with kinase network to synchronize cancer cell metabolic activities with EMT.

**MATERIALS AND METHODS**

**Antibodies**

Otherwise indicated, all antibodies were from Cell Signaling (Boston, MA, USA); N-cadherin and Snai2 from Millipore (Billerica, MA, USA); β-tubulin, Snai1, hypoxia-inducible factor-1α and 14-3-3-α from Santa Cruz (Dallas, TX, USA); Vimentin, laminin 332 and mouse monoclonal anti-human cANGPTL4 mAb11F6C4 from Abcam (Cambridge, MA, USA); pan-cytokeratin and IRdye 680-conjugated secondary antibodies from Thermo Scientific (Waltham, MA, USA); and Alexa Fluor 488-conjugated secondary antibodies and Alexa Fluor 594-conjugated phalloidin from Molecular Probes (Waltham, MA, USA).

**Human tumor biopsies**

Fresh frozen breast, colon, gastric and head and neck tumor biopsies (Proteogenex, Culver City, CA, USA) were used for protein, energy charge and immunofluorescence analysis (Supplementary Table S1).

**Expression and purification of recombinant cANGPTL4 proteins**

The expression and purification of recombinant cANGPTL4 were performed as previously described.12,22

**Cell culture**

MKN74 human gastric carcinoma (JCRB0255, Osaka, Japan) was cultured in RPMI-1640 supplemented with 10% fetal bovine serum in a humidified atmosphere of 5% CO2 at 37 °C. All other cell lines were cultured in Dulbecco’s modified Eagle’s medium with 10% fetal bovine serum. HSC and I4 were from German Cancer Research Center (DKFZ, Heidelberg, Germany), and MCF7 (HTB-22) and HepG2 (HB-8065) were from American Type Culture Collection (ATCC, Manassas, VA, USA). Smartpool small interfering RNAs (Thermo Scientific) against human ANGPTL4 (L-007807-00-0050) and TSC2 (L-003029-00-0005) were used to knockdown respective genes in cells of interest. AKT activity was measured using the AKT activity kit (Cell Signaling).

**Retrovirus transduction**

MKN74Snai1ER cells were generated using retroviral transduction as previously described.44 Briefly, HEK 293T cells were transfected with pCL-10A1 retrovirus packaging vector (ImageX, Littleton, CO, USA) and pWZL-Snai1-ER (Addgene, Cambridge, MA, USA). Harvested pseudoviruses were used to infect MKN74 cells. Following transduction, cells were selected with 5 μg/ml of blasticidin hydrochloride (Sigma Aldrich, St Louis, CO, USA).

**Generation of dual-inducible MKN74Snai1ER/shANGPTL4 cell line**

Dox-inducible pSingle-TS-shRNA vector carrying either shANGPTL4 or scrambled shRNA was introduced into MKN74Snai1ER cells using Fugene HD reagent as per the manufacturer protocol (Promega, Madison, WI, USA). MKN74Snai1ER/shANGPTL4 Cells were selected with 1 mg/ml of G418 (Sigma Aldrich). The efficiency of knockdown was verified by qPCR and immunoblotting after treatment with various concentrations of Dox (up to 10 μg/ml).

**Glucose uptake assay**

Briefly, Cells were incubated with 10 μM of 2-NBDG for 12 h. The cells were then harvested and subjected to fluorescein-activated cell sorting analysis using BD Accuri C6 flow cytometer (BD Biosciences, San Jose, CA, USA). Data were analyzed using FlowJo (Ashland, OR, USA) and plotted based on mean intensity.

**In vitro induction of EMT**

Cells were seeded at a density of 5.25 × 10^4 cell per cm². Hypoxia treatment (1% O2 concentration) was conducted in hypoxic chamber (Stem Cell Technology, Cambridge, MA, USA). For TGF-β-induced and Snai1-mediated EMT, cells were treated with 10 ng/ml of TGF-β for 2 days and with 20 ng/ml of 4-OHT for 4 days, respectively.
**In vivo induction of EMT**

Six-week-old male NSG mice (20–22 g; NOD.Cg-Prkdcscid Il2rgtm1Wjl/SzJ) were injected subcutaneously with 1 × 10^6 cancer cells resuspended in matrigel. A week after the xenograft injection, each xenograft group was randomly divided into four subgroups. Two subgroups received intra-uterine injection of 4-OHT (4 mg/kg) and were fed on either normal chow diet or Dox diet (625 mg/kg, Harlan Laboratories, Indianapolis, IN, USA).

The other two subgroups received vehicle (sunflower seed oil) and similarly fed on the two different diets. After 8 weeks, tumors and lungs were harvested for analysis. Power analysis was used to determine the required number of animal necessary to ensure adequate power to detect a prespecified effect size. All the experimental groups were allocated with double-blinded randomization. Animal experiments were carried out in accordance to the guidelines of the institutional animal care and use committee (ARF-SBS/NIE-A0250A, -A0324 and -A0321) of Nanyang Technological University, Singapore.

**ChIP and Re-ChIP**

ChIP experiments were carried out as previously described. Sonicated chromatin complexes were immunoprecipitated using antibodies against SnaI, pCRB, pSTAT3 or p-p38 and Re-ChIP using anti-14-3-3y. Immunocomplexes were affinity precipitated by Protein A/G Sepharose (Santa Cruz). The ChIP primer sequences are listed in Supplementary Table S2.

**Cell energy charge determination**

Cellular levels of AMP, ADP and ATP were determined by high-performance liquid chromatography. A Shimadzu LC-20AD (Kyoto, Japan) series Quaternary Gradient HPLC system with PDA detector using a series Quaternary Gradient HPLC system with PDA detector was used. Nucleotides were extracted as previously described. Nucleotide standards were from Sigma Aldrich. Separation of the nucleotides was carried out using the mobile phase containing 60 mmol/l of KH₂PO₄, 0.45 mmol/l of tetrabutylammonium bromide and 1.26 mol/l of acetoni-trile. The pH of the mobile phase was adjusted to 3.20 using concentrated phosphoric acid. Retention times of the individual nucleotides were matched to that of the respective nucleotide standards. Areas under the peaks for the respective nucleotides were tabulated and energy charge was calculated using the formula: Energy Charge = [ATP]+½[ADP]/[ATP]+[ADP]+[AMP].

Kinase inhibitor array

MKN74 cells were treated with 95 different kinase inhibitors (SYN-2103; Synkinase, Victoria, Australia) in the absence and presence of rh-ANGPTL4 (10 μg/ml) for 6 h. RNA was isolated and reverse transcribed using the iScript Reverse Transcription Supermix (Bio-Rad, Hercules, CA, USA). qPCR was done using the KAPA SYBR qPCR Universal Master Mix (KAPA Biosystems). The qPCR primer sequences are listed in Supplementary Table S3.

**Immunofluorescence staining**

EMT was identified by immunofluorescence staining for E-cadherin and SnaI2. MKN74 cells were fixed with 4% paraformaldehyde for 10 min, permeabilized with 0.2% Triton X-100 for 10 min and blocked with 3% normal goat serum (NGS) containing 0.05% Triton X-100 in a humidified chamber for 1 h at room temperature. Following three washes in phosphate-buffered saline, cells were incubated overnight at 4 °C with either anti-human-E-cadherin (1:2000) or SnaI (1:100) antibodies in 3% NGS. After three washes, cells were incubated for 1 h at room temperature with Alexa488-secondary antibodies (1:250) and counterstained with Alexa594-phalloidin for F-actin and DAPI (4',6-diamidino-2-phenylindole) for nuclei. Immunostainings performed without primary antibodies served as negative controls. Images were acquired using Zeiss (Munich, Germany) LSM710 confocal microscope with a Plan-APochromat 40×/1.4 oil objective and ZEN2009 software.

**PLA and analysis**

MKN74CTRL or MKN74ANGPTLA (5 × 10^4 cells) were subjected to either normoxia or hypoxia for 2 days at 37 °C. At indicated time, MKN74 cells were fixed, permeabilized and blocked like immunofluorescence staining. Next, MKN74 cells were incubated with monoclonal mouse-anti human 14-3-3y antibody (1:200) and monoclonal rabbit anti-human SnaI antibody (1:200) in 3% NGS overnight at 4 °C. PLA was carried out as per the manufacturer’s protocol (Olink Bioscience, Watertown, MA, USA). Images were taken using Carl Zeiss confocal microscope LSM710 using a Plan-APOCHROMAT 63 ×/1.4 oil DIC objective, and ZEN2012 LE software with constant exposure and gain. Number of protein interaction was quantified using the BlobFinder software.

**Immunoprecipitation and immunoblot**

For immunoprecipitation, at the indicated time points after treatments, cells were lysed using IP lysis buffer (25 μl Tris-HCl pH7.4, 150 μl NaCl, 1% NP-40, 1 μl EDTA). The lysate was then incubated with indicated antibodies overnight at 4 °C with constant rotation. Following that, the antibodies were affinity precipitated using protein A/G beads (Santa Cruz). Proteins were released by boiling for 10 min in Laemmli’s buffer. Protein extracts were resolved using 12% SDS–polyacrylamide gel electrophoresis and electrotransferred (25 μl Tris, 192 μl glycine, 20% methanol) onto a low fluorescence polyvinylidene difluoride membrane for immunoblot analysis. Membranes were stripped and reprobed for other proteins as previously described. Protein bands were detected using Odyssey Clx Infrared Imaging system (LI-COR, Lincoln, NE, USA).

**Statistical analysis**

Statistical analyses were performed using two-tailed Mann–Whitney or one-way analysis of variance test with SPSS software (New York, NY, USA) where appropriate. A P-value of <0.05 is considered significant.

**CONFLICT OF INTEREST**

The authors declare no conflict of interest.

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**AUTHOR CONTRIBUTIONS**

ZT performed experiments, analyzed the results and wrote the article; MKS, JSKC, YL, MMKL, TP, JYHL, ZWT and PZ performed experiments and contributed to discussion. JSKC, YL, LL and PZ performed and analyzed the results from the animal experiments. NST analyzed the results, contributed to discussion, reviewed and edited the article.

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