

## ARTICLE



# EGF-SNX3-EGFR axis drives tumor progression and metastasis in triple-negative breast cancers

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Epidermal growth factor receptor (EGFR) has critical roles in epithelial cell physiology. Over-expression and over-activation of EGFR have been implicated in diverse cancers, including triple-negative breast cancers (TNBCs), prompting anti-EGFR therapies. Therefore, developing potent therapies and addressing the inevitable drug resistance mechanisms necessitates deciphering of EGFR related networks. Here, we describe Sorting Nexin 3 (SNX3), a member of the recycling retromer complex, as a critical player in the epidermal growth factor (EGF) stimulated EGFR network in TNBCs. We show that SNX3 is an immediate and sustained target of EGF stimulation initially at the protein level and later at the transcriptional level, causing increased SNX3 abundance. Using a proximity labeling approach, we observed increased interaction of SNX3 and EGFR upon EGF stimulation. We also detected colocalization of SNX3 with early endosomes and endocytosed EGF. Moreover, we show that EGFR protein levels are sensitive to SNX3 loss. Transient RNAi models of *SNX3* downregulation have a temporary reduction in EGFR levels. In contrast, long-term silencing forces cells to recover and overexpress *EGFR* mRNA and protein, resulting in increased proliferation, colony formation, migration, invasion in TNBC cells, and increased tumor growth and metastasis in syngeneic models. Consistent with these results, low *SNX3* and high *EGFR* mRNA levels correlate with poor relapse-free survival in breast cancer patients. Overall, our results suggest that SNX3 is a critical player in the EGFR network in TNBCs with implications for other cancers dependent on EGFR activity.

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## INTRODUCTION

EGFR (Epidermal Growth Factor Receptor) regulates signal transduction pathways that control diverse cell phenotypes, including proliferation, migration, and survival [1]. Overexpression and over-activation of EGFR in breast tumors are associated with poor clinical outcomes, including early recurrence, increased risk of metastasis, and decreased survival [2–6]. Therefore, it is essential to understand mechanisms regulating EGFR levels and activity. Among other critical processes, internalization and recycling steps regulate surface EGFR levels and activity in normal and in cancer cells.

SNX3 belongs to sorting nexins and is a critical player in the recycling pathway. SNX3 is involved in an early step of endocytosis by binding to phosphatidylinositol-3-phosphate (PI3P) of early endosomes. Multiple lines of evidence established that SNX3 is an essential component of the VPS35:VPS26:VPS29 retromer complex. Retromer complexes recycle the endocytosed receptors such as Wnt receptor Wntless (Wls) and EGFR back to the *trans*-Golgi network or plasma membrane, respectively [7–16]. If not recycled, endosomes mature into late endosomes, which later fuse with the lysosomes, and the cargo is degraded [17, 18].

SNX proteins, SNX1, SNX2, SNX5, SNX6, SNX12, and SNX16, have been implicated in lysosomal targeting of internalized receptors,

including EGFR [19–24]. SNX3, on the other hand, plays a direct role in multivesicular body formation and has a cargo recognition function within the retromer [25, 26]. Hence, SNX3 is involved in the recycling of receptors rather than directly targeting them to lysosomes. Only in the absence of SNX3 function, endocytosed receptors are degraded in the lysosomes. For example, genetic loss of *Snx3* leads to lysosomal degradation of internalized Wls receptors in *C. elegans* and *Drosophila* [27, 28]. Likewise, lysosomal degradation of EGFR increases upon siRNA silencing of SNX3 in epidermoid carcinoma cells [14].

Despite the role of SNX3 in recycling, knowledge on its contribution to cancer phenotypes is limited [29]. The objective of this study was to investigate SNX3 regulation with EGF stimulation in EGFR positive mammary cells and determine whether SNX3 has a role in breast tumorigenesis.

## RESULTS

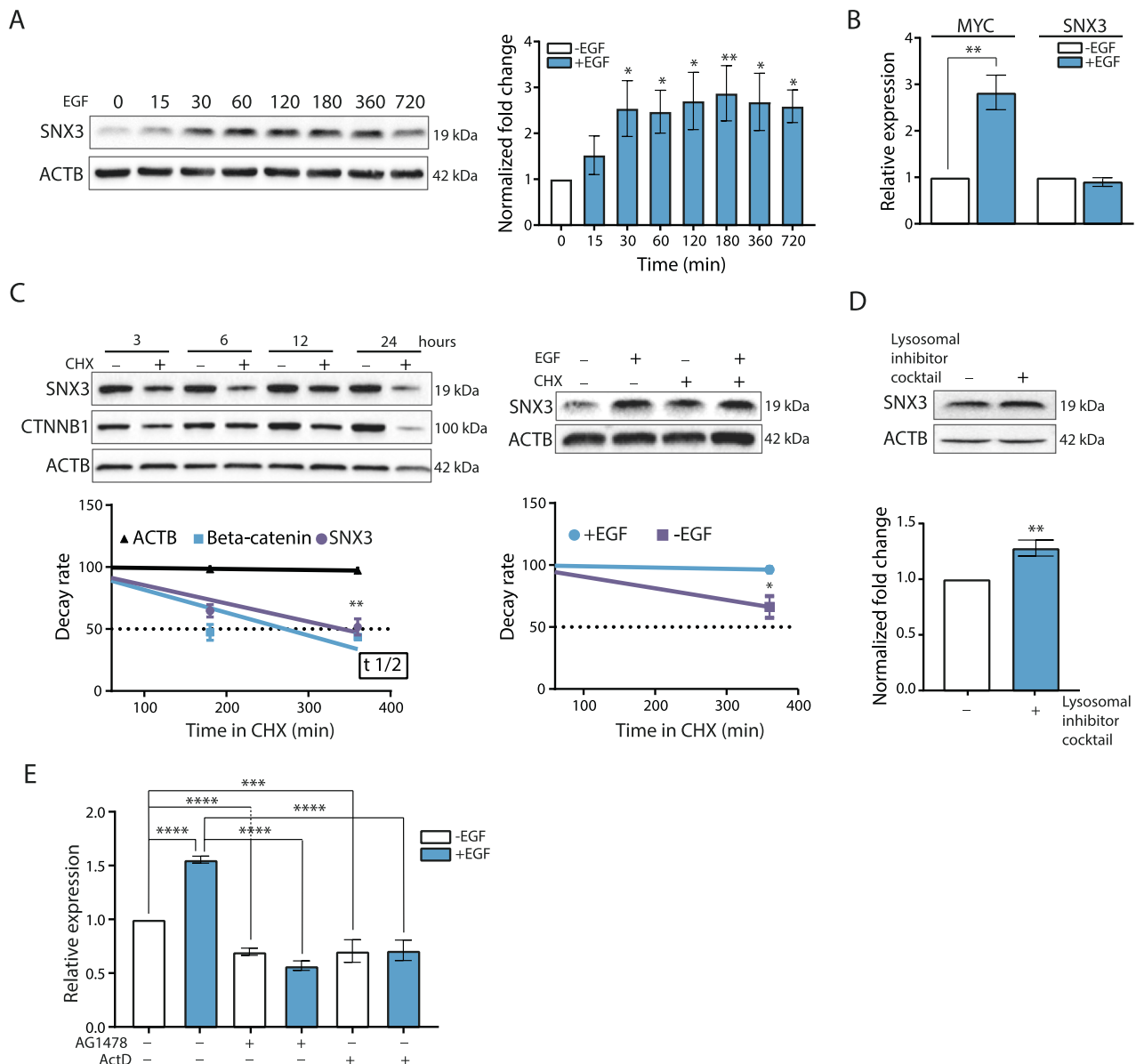
### SNX3 is a target of EGF stimulation in EGFR positive cells, first at the protein level and later at transcript level

To begin investigating the role of SNX3 concerning EGFR in breast cancers, we first asked whether SNX3 is an EGF/EGFR induced

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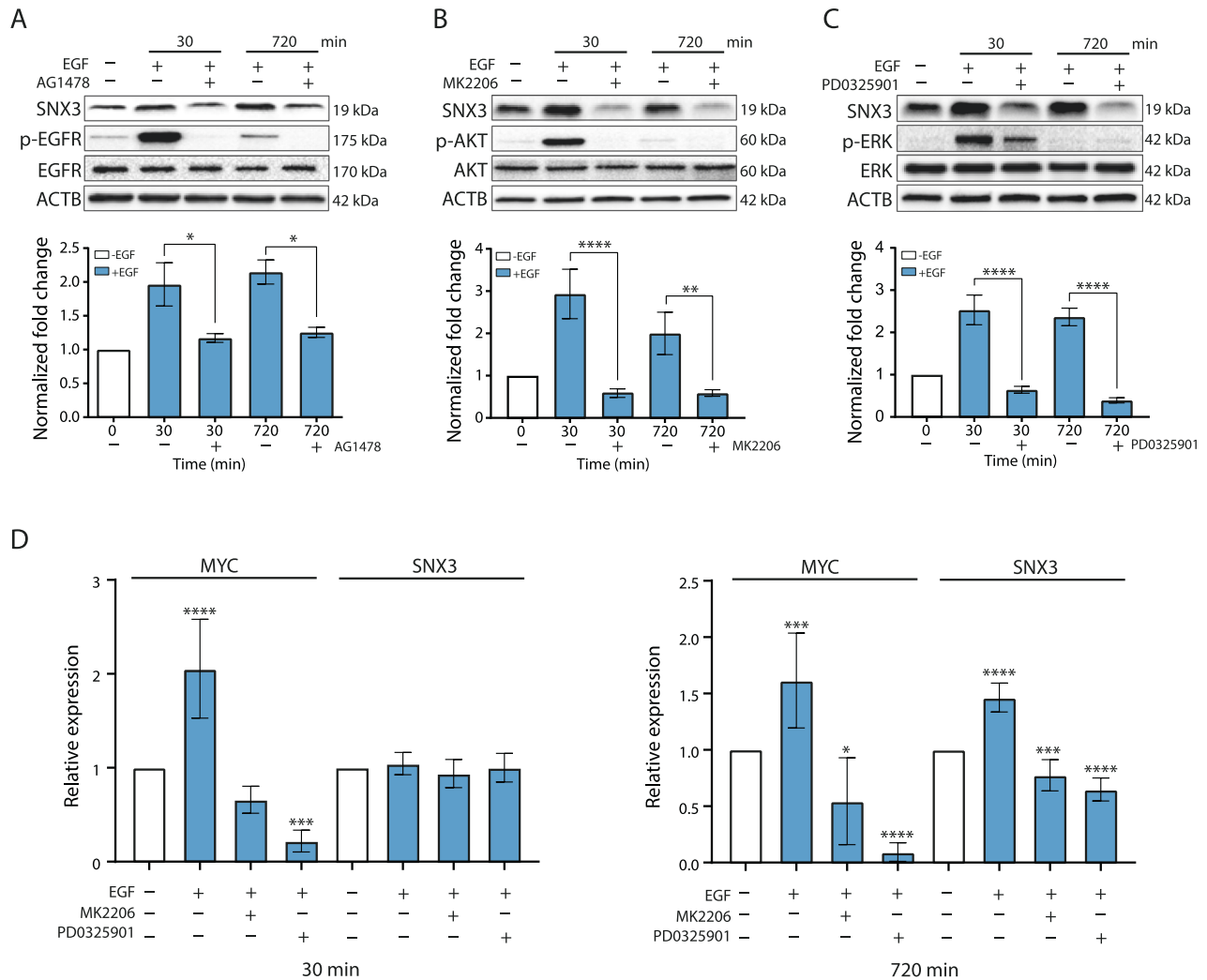
**Fig. 1** **EGF induction of SNX3.** **A** Western blotting analysis for SNX3 in MCF10A cells treated with EGF (20 ng/ml) for the indicated time points (minutes). ACTB was used as a loading control. Blots are representative of 3 independent treatments. Graphs show densitometric quantification of bands. The data represent the mean (SD) of 3 experiments. One-way ANOVA with Tukey's multiple comparison test was used, \* $p < 0.05$ , \*\* $p < 0.01$ . **B** RT-qPCR for the expression of MYC and SNX3 upon EGF stimulation (20 ng/ml, for 30 min) relative to untreated MCF10A cells. The data represent the mean (SD) of 3 independent treatments. Unpaired  $t$ -test was performed for statistical significance, \*\* $p < 0.01$ . **C** Western blotting analysis for the decay rate of SNX3 protein in MCF10A cells treated with CHX (30  $\mu$ g/ml) with or without EGF (20 ng/ml) for the indicated time points. Cell lysates were immunoblotted for SNX3, for ACTB as a loading control, and for CTNNB1 (Beta-catenin) as a positive control for CHX treatments. The data represent the mean (SD) of 3 independent treatments. Unpaired  $t$ -test with Welch's correction was performed for statistical significance, \* $p < 0.05$ , \*\* $p < 0.01$ . **D** MCF10A cells were treated with a lysosomal inhibitor cocktail (10  $\mu$ M Leupeptin, 1.5  $\mu$ M Pepstatin A, 1  $\mu$ M E-64) for 24 h, and lysates were collected. Blots are representative of 3 independent treatments. Graphs show densitometric quantification of bands. Unpaired  $t$ -test was performed for statistical significance, \*\* indicates  $p < 0.01$ . **E** RT-qPCR for SNX3 mRNA. MCF10A cells were pre-treated with actinomycin D (10  $\mu$ g/ml) for 1 h and with (EGF 20 ng/ml) for 12 h. The data represent the mean (SD) of 3 independent treatments. One-way ANOVA with Tukey's multiple comparison test was used \*\*\*\* $p < 0.0001$ , \*\*\* $p < 0.001$ .

gene. We used the MCF10A cell line, an EGFR positive, non-tumorigenic, and hormone receptor-negative mammary cell line. MCF10A is also a well-established model for studying gene expression patterns in response to EGF stimulation [30–32]. We stimulated cells with EGF in a time-course experiment to detect dynamic changes in gene expression and collected lysates and RNA.

Shortly after EGF (15–30 min) stimulation, the abundance of SNX3 protein was increased and remained high throughout the

EGF treatment (i.e., 12 h) (Fig. 1A). This increase in SNX3 following EGF treatment was also seen in HEK293 (EGFR positive human embryonic kidney cells) (Supplementary Fig. S1). These results showed that the EGF responsiveness of SNX3 was not MCF10A specific.

To understand how SNX3 protein abundance increased in response to EGF, we first tested whether regulation was transcriptional. Transcriptional upregulation of MYC mRNA was evident as an EGF responsive gene [32]. However, we did not



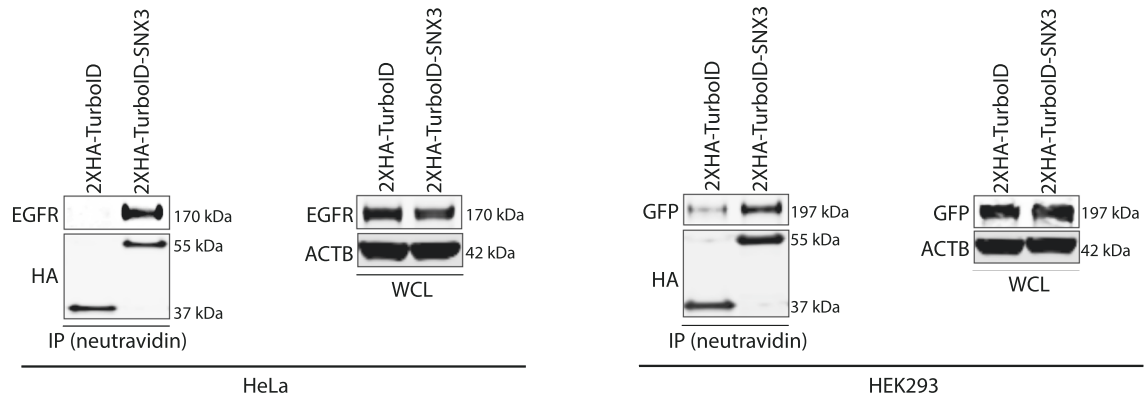
**Fig. 2 Effect of blocking of EGFR and downstream pathways on SNX3 protein levels.** Western blotting for SNX3, p-EGFR, p-AKT, p-ERK, EGFR, AKT, and ERK in MCF10A cells pre-treated with (A) AG1478 (25  $\mu$ M) for 12 h, (B) with MK2206 (1  $\mu$ M), (C) with PD0325901 (1  $\mu$ M) for 4 h. Pre-treated cells were stimulated with EGF (20 ng/ml) for the indicated time points. Blots are representative of 3 independent experiments. Bars show densitometric quantification of SNX3 levels (mean (SD) of 3 experiments). One-way ANOVA with Tukey's multiple comparison test was used, \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\*\* indicates  $p < 0.0001$ . (D) RT-qPCR for the mRNA levels of SNX3 in MCF10A cells upon EGF stimulation (30 min or 720 min) and inhibitor treatments compared with untreated cells. MYC expression was used as a positive control. The fold change for SNX3 mRNA was calculated as described. The data represent the mean (SD) of 3 independent treatments. One-way ANOVA with Dunnett's multiple comparison test was used, \* $p < 0.05$ , \*\*\* $p < 0.001$ , \*\*\*\* $p < 0.0001$ .

observe transcriptional upregulation of SNX3 mRNA upon EGF stimulation at early time points (Fig. 1B, Supplementary Fig. S2). Next, we turned to protein stability as a possible mechanism to explain increased SNX3 protein levels at early time points of EGF stimulation. In a time-course experiment, we measured SNX3 protein half-life after blocking translation with cycloheximide (CHX) treatment alone or with EGF stimulation and collected cell lysates for western blotting. CTNNB1 (Beta-catenin) was used as a control for the CHX treatment [33]. We also confirmed that CHX treatment did not impair the lysosomal degradation of EGF-induced EGFR (Supplementary Fig. S3). Inhibition of translation by CHX caused a reduction in SNX3 protein levels (Fig. 1C, left panel). Unlike SNX3, EGFR protein decay was not rapid (Supplementary Fig. S3). Interestingly, EGF treatment enhanced the abundance of SNX3 protein levels in CHX treated cells (Fig. 1C, right panel). EGF inflicted increase in protein levels suggested post-translational regulation of SNX3 protein, possibly through degradation. Accordingly, blocking proteasomal degradation did not recover SNX3 protein levels (Supplementary Fig. S4), whereas lysosomal inhibition enhanced SNX3 abundance (Fig. 1D).

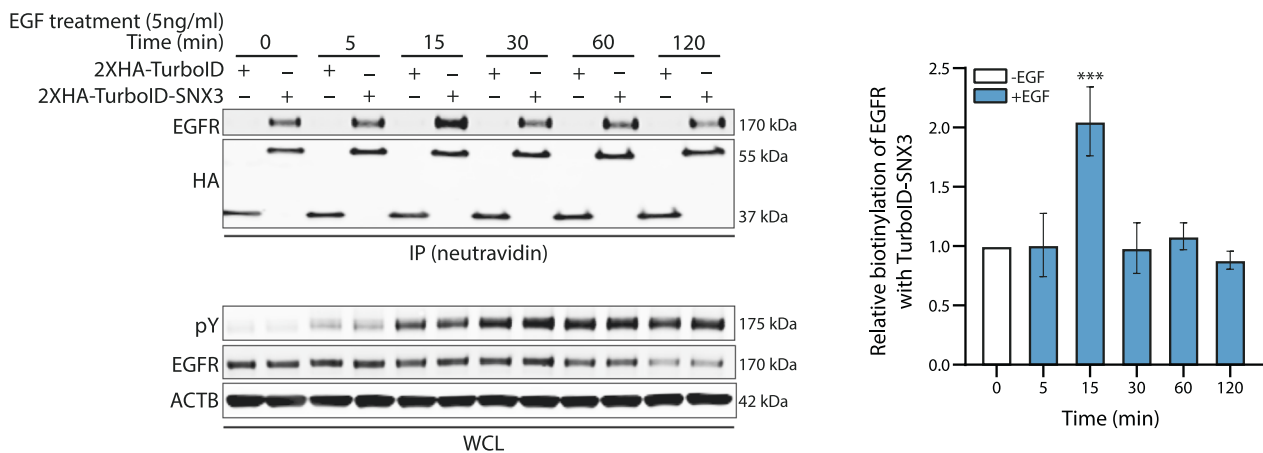
On the other hand, EGF stimulation for 12 h resulted in increased levels of SNX3 mRNA (Fig. 1E). To test whether the increase in SNX3 mRNA levels was transcriptional or due to enhanced mRNA stability, we treated cells with actinomycin D, a transcription inhibitor, or AG1478, a selective inhibitor of EGFR tyrosine kinase activity [34]. Both actinomycin D and AG1478 treatments reversed the EGF-induced increase of the SNX3 mRNA (Fig. 1E). In addition, we measured the decay rate of SNX3 mRNA by blocking transcription for up to 12 h. We showed that the SNX3 transcript was not rapidly degraded (Supplementary Fig. S5). Therefore, it is unlikely that SNX3 mRNA stability is enhanced further by EGF stimulation.

Furthermore, upregulation of SNX3 protein upon EGF stimulation was reversible by AG1478 [34] (Fig. 2A), by MK2206, an Akt inhibitor [35] (Fig. 2B), and by PD0325901, a MEK inhibitor [36] (Fig. 2C). These results showed that inhibiting EGFR and downstream pathways blocked the upregulation of SNX3. Treating EGF stimulated cells with MK2206 and PD0325901 further decreased SNX3 protein levels (Supplementary Fig. S6). As part of the late response, transcriptional regulation of SNX3 was also inhibited

A



B



**Fig. 3 Proximity-based labeling by SNX3-TurboID.** **A** Western blotting analysis on lysates from cells transfected with 2XHA-TurboID or 2XHA-TurboID-SNX3 constructs for proximity-based labeling assay. For HeLa cells, immune-precipitated (IP) samples were immunoblotted with EGFR and HA antibodies. In parallel, the 2XHA-TurboID or 2XHA-TurboID-SNX3 and EGFR-GFP constructs were transfected to HEK293 cells. IP samples were immunoblotted with HA and GFP antibodies. GFP-antibody was used to detect the EGFR-GFP in the IP samples. HA-antibody was used to verify the expression of TurboID or TurboID-SNX3 fusion protein for both cells. Whole-cell lysates (WCL) were used to show the equal loading of proteins. Blots are representative of 3 independent experiments. **B** Western blotting analysis on lysates from HeLa cells transfected with 2XHA-TurboID or 2XHA-TurboID-SNX3 constructs and treated with EGF as indicated. Biotinylated EGFR levels were detected. HA-antibody was used to verify the expression of TurboID or TurboID-SNX3 fusion protein. WCL shows EGFR, phosphotyrosine (pY), and ACTIN levels. Blots and the graph for densitometry analysis represent 3 independent experiments. One-way ANOVA and Dunnett's multiple comparison tests were used to measure statistical significance, \*\*\* $p < 0.001$ .

after blocking EGFR and downstream signaling pathways, preventing the increase of SNX3 protein (Fig. 2D). In support of these results, using the same inhibitors prevented upregulation of SNX3 in HEK293 cells (Supplementary Fig. S7).

In addition, we tested whether any other proliferative signal would induce SNX3 mRNA or protein in a similar pattern. However, estradiol stimulation of estrogen receptor (ER) positive MCF7 breast cells did not induce SNX3 at the mRNA or protein level (Supplementary Fig. S8).

Taken altogether, SNX3 is a specific target of EGF stimulation in EGFR positive cells, first at the protein level and later at the transcriptional level.

#### SNX3 interacts with EGFR, co-localizes with early endosomes and endocytosed EGF

After establishing EGF/EGFR specific upregulation of SNX3, we began examining how SNX3 might, in turn, regulate EGFR. First, we took the proximity ligation approach to confirm the dynamic interaction between SNX3 and EGFR [37, 38]. We fused SNX3 protein to the C-terminus of 2XHA-tagged TurboID (2X HA-TurboID-SNX3).

We transfected this construct to HeLa cells. In parallel, we transfected HEK293 cells with the same TurboID constructs and an EGFR-GFP fusion construct. The reasons for choosing these cells were their high transfection rates, EGFR positivity, and previous literature that established the functionality of the retromer and the endocytic pathways [25, 39, 40].

Following biotin incubation and neutravidin-mediated pull-down, immuno-blotting showed TurboID-SNX3 biotinylated endogenous EGFR and EGFR-GFP protein in HeLa and HEK293 cells, respectively (Fig. 3A). Immuno-blotting with whole-cell lysates (WCL) showed equal EGFR levels in TurboID vector control and 2XHA-TurboID-SNX3 transfected cells.

Next, we stimulated HeLa cells with EGF (5 ng/ml) for a period of 5, 15, 30, 60, and 120 min. EGFR is known to be recycled back to the plasma membrane upon low-dose EGF treatment [41]. Accordingly, biotinylation of EGFR by SNX3-TurboID was highest at 15 min of low-dose EGF stimulation (Fig. 3B). In contrast, following long-term EGF stimulation, EGFR is degraded in lysosomes to circumvent the over-activation of downstream pathways [41]. In agreement, total EGFR levels did not change

until EGF stimulation was extended to 60 min when EGFR levels were downregulated (Fig. 3B). Biotinylation of EGFR by SNX3-TurboID was also decreased after 15 min of EGF stimulation. Of note, EGFR was also biotinylated in unstimulated cells, albeit at a reduced level. We may explain this by the low-level internalization of EGFR upon non-EGF ligand interactions [42]. It is also plausible that overexpressing SNX3 may have enhanced the interaction with EGFR.

Next, we further examined the dynamic interaction of SNX3 and EGFR with confocal microscopy. We used Alexa Fluor™ labeled-EGF (EGF647) to follow endocytosis of EGF-induced EGFR. SNX3 colocalization with early endosomes (EEA1) and endocytosed EGF was highest at 15 min of EGF stimulation, consistent with the biotinylation results (Fig. 4A, B).

### Effects of SNX3 silencing on EGFR levels in TNBC cells

Next, we wanted to ascertain further the functional relationship between SNX3 and EGFR in TNBCs. For this, we examined whether SNX3 silencing would affect EGFR protein levels. We tested MDA-MB-231, MDA-MB-468 TNBC cell lines [43], and additional cell lines that are EGFR positive (SK-BR-3, HeLa, and HEK293). All of the cell lines transfected with a siRNA pool targeting SNX3 had decreased levels of EGFR protein compared to non-targeting siRNA transfected controls within 48–72 h of silencing (Fig. 5A). SNX3-siRNA transfected MCF10A cells also had decreased total EGFR levels and decreased p-EGFR (Fig. 5B).

Moreover, a targeted transcriptomic analysis in SNX3-siRNA transfected MCF10A cells using NanoString technology showed decreased expression of cell cycle and survival-related genes including *MYC*, *CCDN1*, *CCDN2*, and *BIRC5* (Survivin) (Fig. 5C). Downregulation of these genes was consistent with the downregulation of EGFR in SNX3 silenced cells. However, while transient SNX3 silencing caused decreased EGFR availability and activity, to our surprise, *EGFR* mRNA was upregulated in SNX3-targeted siRNA transfected cells (Fig. 5C). RT-qPCR also confirmed upregulation of the *EGFR* mRNA in SNX3 silenced MCF10A and MDA-MB-231 cells (Fig. 5D). Upregulation of *EGFR* mRNA suggested a compensatory response to rescue downregulated EGFR levels upon SNX3 silencing. To test this in another cancer cell line and investigate this further in vivo, we first generated shRNA silenced SNX3 (Snx3sh) models in the 4T1 cell line, a well-established syngeneic model for TNBCs, mimicking tumor growth and metastatic spread of human breast cancers [44]. We observed upregulated levels of EGFR protein (Fig. 6A) upon long-term silencing of SNX3. *EGFR* mRNA was also upregulated in these Snx3sh cells (Fig. 6B).

### Long-term silencing of SNX3 enhances neoplastic phenotypes in vitro and promotes tumor progression and metastasis in vivo

Next, we wanted to determine whether cellular phenotypes were altered in Snx3sh 4T1 cells due to increased EGFR protein levels. SNX3sh cells had increased cell proliferation, measured by xCELLigence live proliferation system (Fig. 6C), increased number of large colonies (Fig. 6D), increased migration, and invasion detected by transwell assays compared to control cells (Fig. 6E). Because SNX3 may recycle other receptors, we tested whether EGFR overexpression was responsible for these phenotypes. Inhibition of EGFR kinase activity with AG1478 treatment in Snx3sh 4T1 cells reversed increased proliferative rate and colony-forming potential, confirming the role of EGFR activity in these in vitro phenotypes (Fig. 6F, G).

When we moved to an orthotopic mouse model and injected Snx3sh-4T1 cells subcutaneously near mammary fat pads (MFP) of BALB/c mice without surgery, tumors were larger in Snx3sh 4T1 injected mice compared with control animals. There were no significant changes to body weight (Fig. 7A, B). Although we did subcutaneous injection near the MFP (instead of injection to MFPs

with surgery), there was gross metastasis in the lungs of mice bearing Snx3sh tumors compared with controls ( $p < 0.0001$ ) (Fig. 7C–E). Later, tumors isolated from animals were lysed and examined for Snx3 and Egfr status. Snx3 silencing was maintained in the primary tumors, and Egfr protein was significantly overexpressed (Fig. 7F).

Of note, SNX3sh-2 tumors with higher EGFR expression had higher numbers of metastatic nodules.

These results showed that 4T1 cells became further aggressive due to overexpression of EGFR, inflicted by SNX3 loss. To investigate the clinical relevance of these observations, we analyzed mRNA expressions of *SNX3* and *EGFR* in TNBC patients in the METABRIC (Molecular Taxonomy of Breast Cancer International Consortium) dataset [45]. We found that low *SNX3* and high *EGFR* mRNA levels correlated with poor relapse-free survival in patients (Fig. 7G). Similarly, low *SNX3* and high *EGFR* mRNA levels correlated with poor relapse-free survival in TNBC patients in an independent dataset (GSE31519) (Fig. 7G, right panel).

Collectively, our results showed SNX3 to be a critical player in the EGFR network as an EGF responsive target and a regulator of EGFR protein levels. Based on these results, we speculate that SNX3 loss may be linked to EGFR overexpression in not only TNBCs but possibly other cancers where gene-level amplification and/or mutations do not exist.

### DISCUSSION

We report multiple lines of evidence suggesting that the EGF-SNX3-EGFR axis drives tumor progression and metastasis in triple-negative breast cancers (Fig. 8).

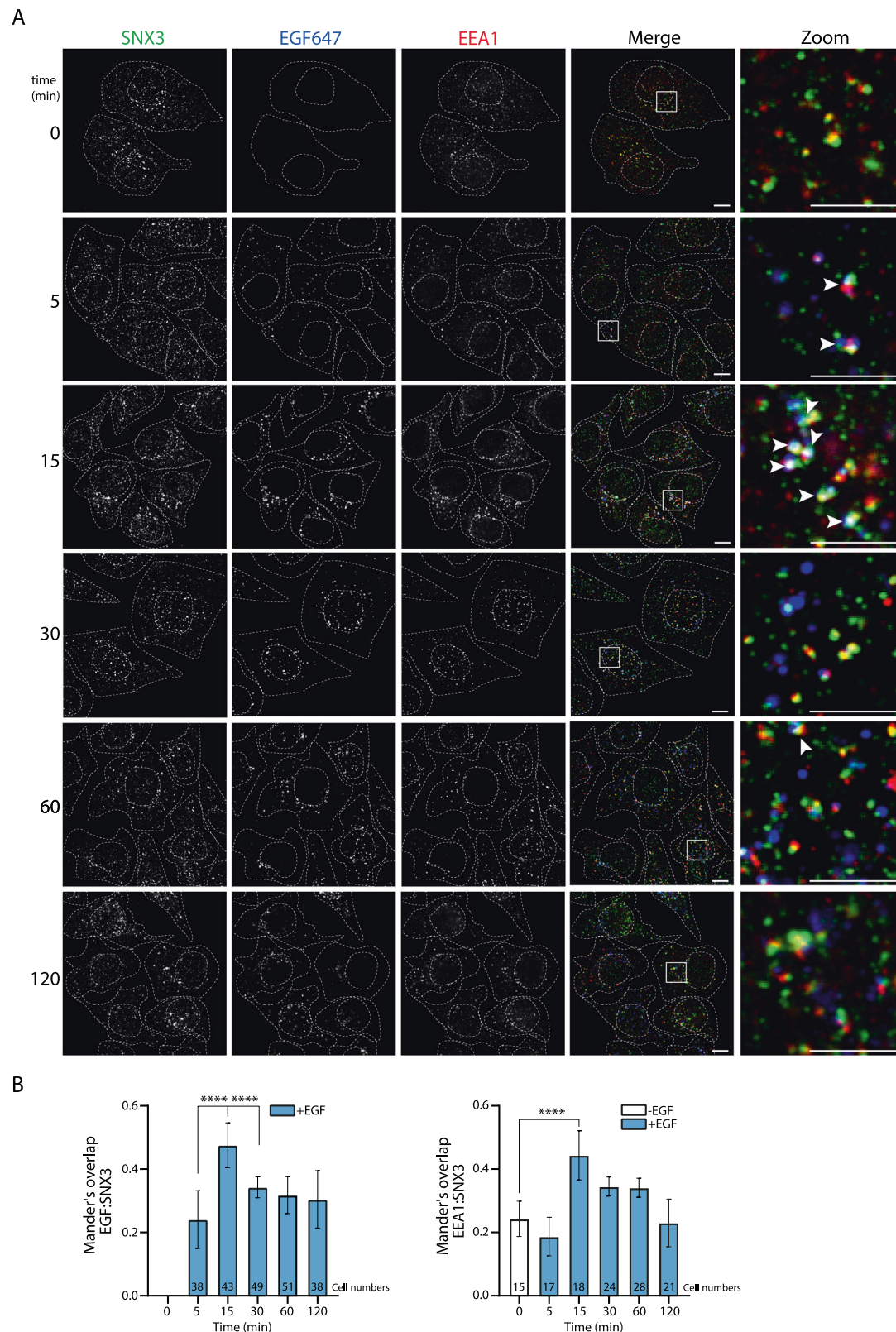
First, we identified SNX3 as an immediate and sustained target of EGF stimulation. In EGFR positive mammary cells, SNX3 protein levels were upregulated quickly after EGF stimulation. The rapid upregulation of SNX3 protein was possibly due to the delayed turnover of SNX3 in lysosomes. SNX3 protein levels were high throughout EGF stimulation, with the contribution of transcriptional upregulation only at much later time points (i.e., 12 h). These results, showing multiple mechanisms to regulate SNX3 mRNA/protein abundance in response to EGF, highlight the role of SNX3 in EGFR biology.

Next, to provide insight into SNX3 function, we used a proximity-based labeling approach in live cells. We showed close proximity between SNX3 and EGFR, mainly when ligand-activated and endocytosed EGFR is recycled most [41]. These observations were consistent with increased colocalization of SNX3, early endosomes, and endocytosed EGF at 15 min of EGF stimulation.

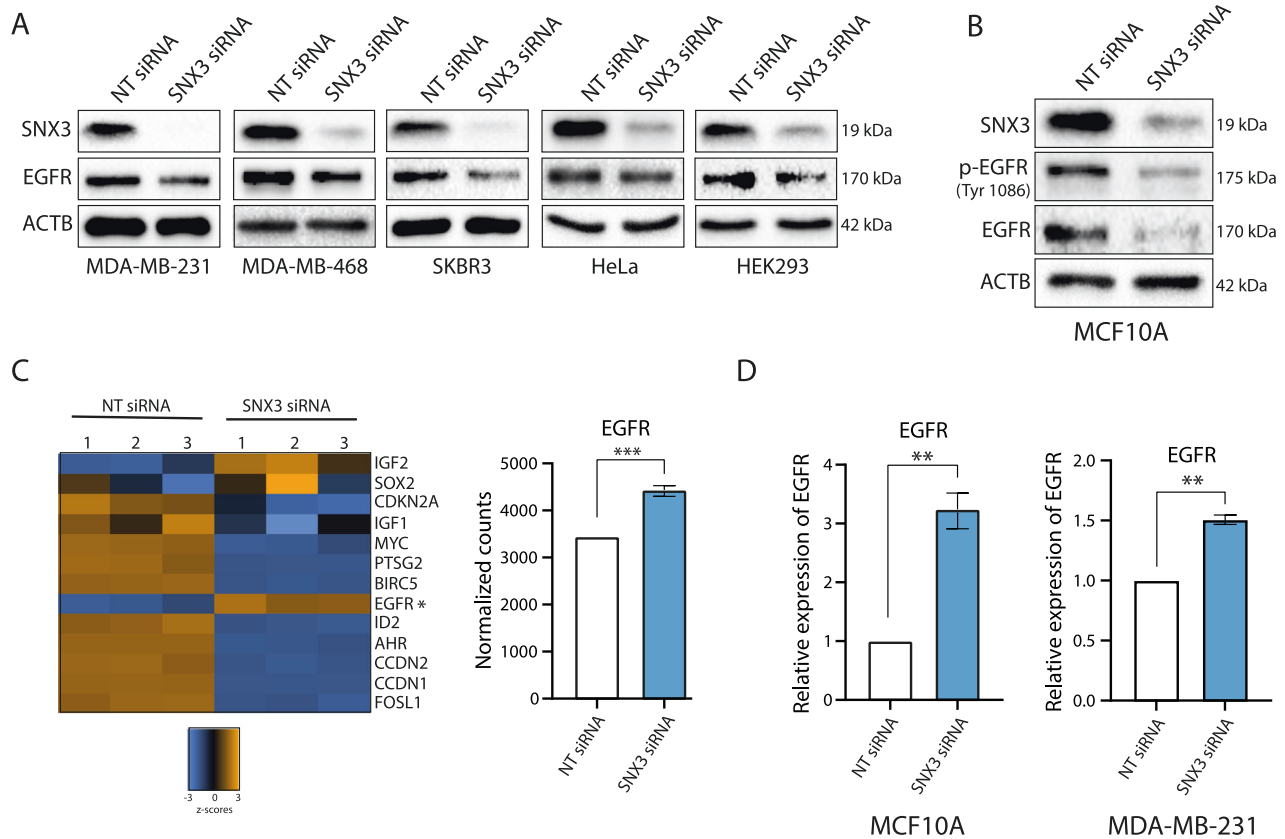
After establishing SNX3 upregulation upon EGF stimulation and confirming its association with EGFR, we investigated the role of SNX3 in TNBCs. To this end, we performed short and long-term silencing of SNX3 and conducted a targeted gene expression analysis. This approach allowed us to capture dynamic changes of the *EGFR* mRNA and protein levels. As expected, short-term silencing of SNX3 in a panel of breast and non-breast cancer cell lines showed decreased EGFR protein levels. However, transcriptome level analysis showed increased levels of *EGFR* mRNA in SNX3 silenced cells, suggesting activation of a compensatory mechanism to restore EGFR protein levels. Indeed, when we investigated the long-term effects of SNX3 silencing, we observed significant upregulation of EGFR protein in TNBC cells. The same gene expression pattern (low *SNX3*, high *EGFR*) correlates with the shorter survival of TNBC patients. In agreement with the clinical data, in vivo studies showed increased tumor growth and metastasis upon SNX3 loss and EGFR overexpression. Importantly, oncogenic phenotypes in vitro were reversible by pharmacological inhibition of EGFR.

Overexpression of EGFR and its association with poor prognosis is known in many cancer types, including breast cancer [46–48]. However, *EGFR* gene amplifications and/or mutations can only





**Fig. 4 Time-dependent interactions of SNX3.** **A** Representative images of confocal microscopy show SNX3, EEA1, and EGF in HeLa cells. Boxed regions in the merge images indicate zoomed areas. The colocalization of SNX3 (green), EGF (blue), and EEA1 (red) are highlighted with white arrows in the zoom images. Dashed white lines indicate cell and nuclear boundaries. Scale bars are 10  $\mu$ m. **B** Graphs of colocalization analyses were plotted using Mander's overlap values, which were quantified from multi-cell images. Cell numbers are shown in the bar graphs. Data are from 3 independent experiments. Statistical significance was calculated using one-way ANOVA with Tukey's multiple comparison test. \*\*\*\* $p < 0.0001$ .



**Fig. 5** Effects of transient SNX3 silencing. **A** Western blotting analysis for SNX3 and EGFR on lysates from indicated cell lines treated with SNX3 siRNA pool or non-targeting siRNAs (NT) for 48–72 h. Blots are representative of at least two independent experiments. **B** Western blotting analysis for SNX3, p-EGFR (Tyr1086), and EGFR on lysates from MCF10A cells transfected with NT siRNA and SNX3 siRNA for 72 h. Blots are representative of 2 independent experiments. **C** Transcription level changes in NT siRNA and SNX3 siRNA transfected MCF10A cells using the NanoString Vantage 3D RNA cartridge. Blue and orange colors on the heat map represent z scores. The asterisk shows *EGFR* mRNA. The graph shows normalized mRNA counts for *EGFR*. Unpaired *t*-test was performed for statistical significance, \*\*\**p* < 0.001. **D** RT-qPCR for the expression of *EGFR* relative to *RPLP0* in MCF10A and MDA-MB-231 cells transfected with SNX3 siRNA compared to NT siRNA transfected cells. Unpaired *t*-test was performed for statistical significance, \*\**p* < 0.01.

partly explain cases of EGFR overexpression. Other mechanisms have been proposed to explain the upregulation of EGFR. For example, increased translation or decreased protein degradation leads to EGFR overexpression in the hypoxic microenvironment [49, 50]. Hypoxia also induces Early Growth Response Factor 1 (EGR1) and activates the basal transcriptional activity of the EGFR promoter [51]. In addition, changes in the local epigenetic landscape of the *EGFR* promoter modulate the accessibility of other transcription factors in different cellular contexts [52]. Despite these findings, mechanisms of EGFR overexpression and activation in cancers are not entirely understood. Our results suggest that deregulated retromer function may also drive EGFR overexpression. The molecular network behind compensatory upregulation of EGFR remains to be investigated.

It is also worth noting that SNX3 plays a role in recycling other receptors (e.g., transferrin receptor, WIs, and possibly other members of the ErbB family). We observed increased transferrin receptor levels in SNX3 silenced 4T1 cells but not in primary tumors (Supplementary Fig. S9). This result suggests that deregulated recycling activates compensatory mechanisms to repopulate the receptor levels. However, these mechanisms are possibly dynamic and context-dependent. As in the case of the transferrin receptor, protein levels were back to normal in SNX3 silenced tumors, whereas EGFR was still overexpressed. The differential upregulation of transferrin receptor and EGFR in tumors could depend on the biological context and the inherent complexity of signaling networks in

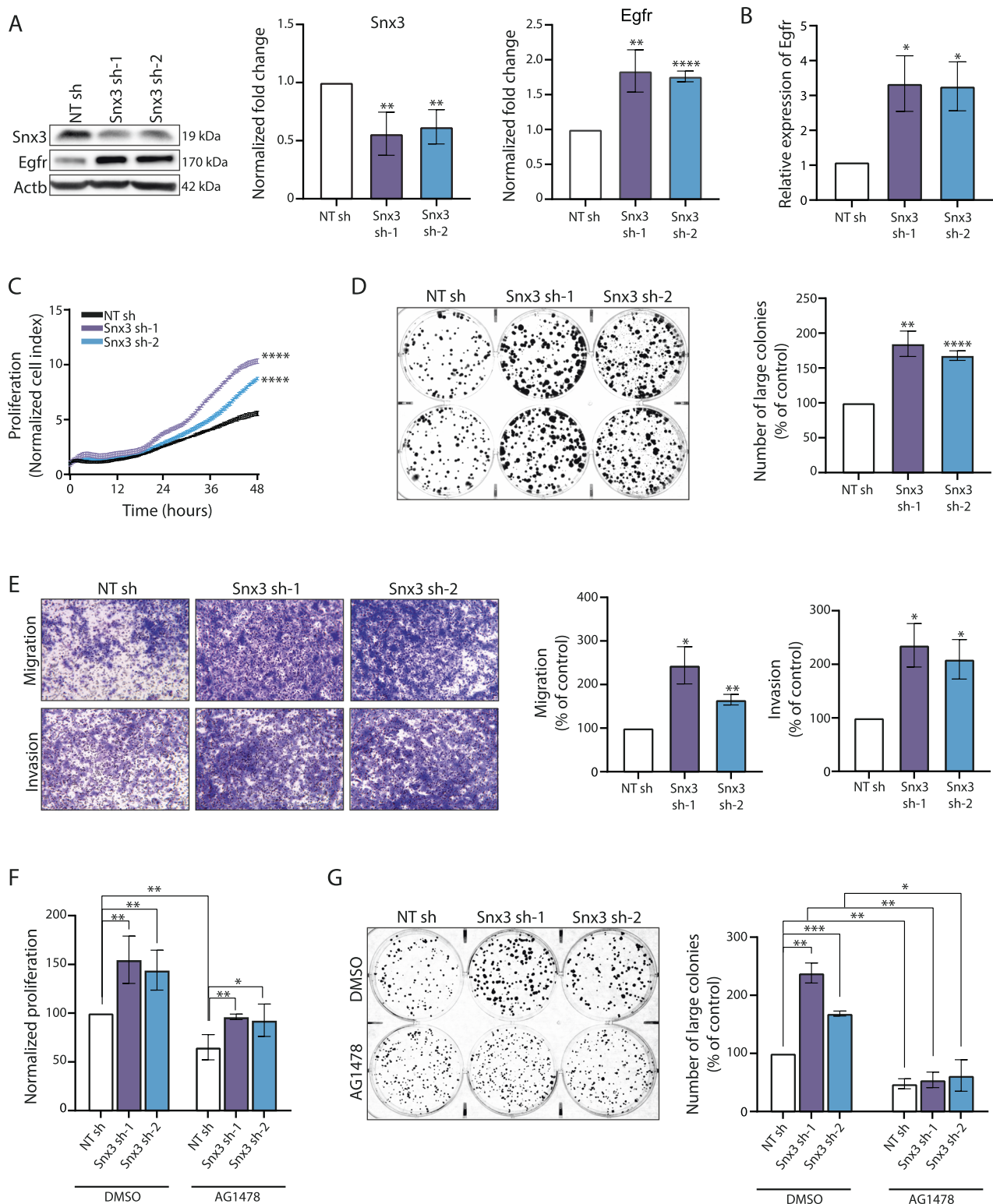
TNBC cells. Hence, it is plausible that deregulated recycling causes diverse phenotypic outcomes in a cell context-dependent manner. This view may also clarify why SNX3 is linked to various diseases, including cancer, cardiac hypertrophy [53], and Alzheimer's disease [54].

In closing, our results provide insight into SNX3 gene/protein regulation, with important implications for the overexpression of EGFR protein in a group of breast and possibly other cancers. This observation may partly explain EGFR overexpression cases with no genomic amplification or mutations. Thus, the EGF-SNX3-EGFR axis presented here may have future implications for developing potent therapies targeting the EGFR network and possibly understanding resistance mechanisms to anti-EGFR therapies.

## METHODS

### Cell lines and treatments

MCF10A, MDA-MB-231, HEK293, HeLa cells were grown as described [55]. 4T1 cells were grown in DMEM high glucose medium with 10% FBS (Biowest, France, 181T-500), 1% P/S, and 1% nonessential amino acids (Biological Industries, Israel, 03-031-1B). All cell lines were grown as monolayers and were incubated at 37 °C with 95% humidified air and 5% CO<sub>2</sub>. Cells were routinely monitored for mycoplasma contamination. 4T1 cells expressing stable shRNAs in pSUPER vector were maintained in G418 (175 µg/ml, Roche, Germany, 04727894001) containing medium. For EGF (Thermo Fisher Scientific, MA, USA, PHG0313) treatments, cells were serum-starved and then stimulated with EGF for the indicated time points. Cycloheximide, actinomycin D, and AG1478 were purchased from Tocris,



UK (0970, 1299, 1276). Akt inhibitor MK2206 (Selleckchem, USA, 1078) and MEK inhibitor PD0325901 (Selleckchem, USA, 1036) were gifts from Dr. Onur Cizmecioglu (Bilkent University). E2 ( $\beta$ -Estradiol) was purchased from Sigma-Aldrich (E2257).

#### Plasmids and siRNAs

The EGFR-GFP vector was purchased from Addgene (Plasmid 32751) [56]. The coding sequence of SNX3 was cloned into 2X HA-TurboID (pEGFP-C1)

vector (a gift from Dr. S. Neeffjes, Leiden University) and sequenced. Mouse SNX3 short-hairpin RNA (shRNA) (5'-AACTTCCTCGAGATCGACG-3') and non-targeting (NT) shRNA (5'-CGTACGCGGAATACTTCGATT-3') were cloned into pSUPER retro.neo-GFP (a gift from Dr. U. Tazebay, Gebze Teknik University) and sequenced. SNX3 siRNA pool (Dharmacon, USA, SMARTpool siGENOME M-011521-01,) and NT siRNAs (Qiagen, Germany, 1027310) were used for transient silencing of SNX3. DharmaFect (Dharmacon, T-2004) was used to transfect siRNAs.



**Fig. 6 Effects of long-term silencing of SNX3.** **A** 4T1 cells were stably transfected with shRNA vectors (NTsh and Snx3sh). Immuno-blotting shows sustained silencing of SNX3 and overexpression of EGFR in two monoclonal populations (Snx3sh-1 and Snx3sh-2) compared with 4T1 control (NTsh) cells. Bars show densitometry analysis of 3 independent experiments ( $^{**}p < 0.001$ ,  $^{****}p < 0.0001$ , unpaired *t*-test). **B** *Egfr* mRNA levels in NTsh and Snx3sh cells, detected by RT-qPCR. ( $^{*}p < 0.05$ , unpaired *t*-test,  $n = 3$ ). **C** Real-time proliferation analysis of Snx3sh 4T1 cells with xCELLigence RTCA. NTsh and Snx3sh-1 and Snx3sh-2 cells were monitored for 48 h with three technical replicates ( $^{****}p < 0.0001$ , two-way ANOVA). **D** Colony formation of NTsh and Snx3sh cells in seven days of seeding. Large colonies ( $\geq 1000$  pixels) were counted using CountPHICS software ( $n = 3$  independent experiments,  $^{*}p < 0.05$ ,  $^{**}p < 0.001$ ,  $^{***}p < 0.0001$ , unpaired *t*-test). **E** Transwell migration and invasion of Snx3sh cells. Cells were allowed to migrate for 8 h. For invasion assay, cells were allowed to invade Matrigel-coated chambers for 12 h. Pictures were taken at  $\times 4$  magnification. Three independent experiments were analyzed with unpaired *t*-test ( $^{*}p < 0.05$ ,  $^{**}p < 0.001$ ). **F** Effect of EGFR blockage on the proliferation of Snx3sh 4T1 cells by AG1478 treatment for 72 h, detected by MTT ( $n = 3$  independent experiments,  $^{*}p < 0.05$ ,  $^{**}p < 0.001$ , unpaired *t*-test). **G** Colony-forming potential upon AG1478 treatment. NTsh and Snx3sh cells were treated with AG1478 or DMSO for seven days. Bars show the number of large colonies ( $\geq 1000$  pixels) counted using the CountPHICS software ( $n = 3$  independent experiments, unpaired *t*-test,  $^{*}p < 0.05$ ,  $^{**}p < 0.001$ ,  $^{***}p < 0.0001$ ).

### Proximity-based labeling assay

Cells were transfected with either pEGFP\_2XHA\_TurboID or pEGFP\_2XHA\_TurboID\_SNX3. After 24 h, cells were starved with serum-free media for 4 h. 5 ng/ml EGF was given to cells for the indicated time points. 50  $\mu$ M biotin (Sigma-Aldrich, MO, USA, B4639) was given to cells for 5 min during the last 5 min of EGF treatment. Cell lysis and IP with high capacity neutravidin beads (Thermo Scientific, 29202) were performed as described [38]. Lysates were loaded to 4–12% SDS-PAGE for western blotting.

### Confocal microscopy

$4 \times 10^5$  HeLa cells were seeded onto 24 well plates with glass coverslips (Menzel Gläser, Germany MENZCB00130RAC). Cells were kept in serum-free media for 4 h and induced with 100 ng/ml Alexa Fluor™ 647-tagged EGF for indicated times. Cells were stained as described [57]. For permeabilization and blocking, 0.1% Triton X-100 (Sigma-Aldrich, T8787) and 1% BSA (ChemCruz, TX, USA, sc-2323A) in PBS were added to cells and incubated for 30 min. 3  $\mu$ g/ml primary antibodies in blocking buffer was given to cells for 1 h. Alexa Fluor™ tagged secondary antibodies were used (1:300). Following PBS washes, 4% paraformaldehyde was used for 30 min for post fixation. Cells were then mounted using ProLong Gold antifade Mounting medium with DAPI (Life Technologies, CA, USA, P36941). Imaging was performed using appropriate solid-state lasers on Leica SP8 microscope. HyD detectors and HCX PL 63X magnification oil emersion objectives were used. Images were taken with 2.5 digital zoom in 1024 $\times$ 1024 scanning format with line averaging. Mander's overlap was used to indicate colocalization.

### Expression analysis

RNA isolation, quantification, cDNA synthesis and expression analysis were conducted according to MIQE Guidelines [58] and as described previously [59]. Ssoadvanced Universal SYBR Green Supermix (BioRad, CA, USA, 1725271) and CFX Connect Real-Time PCR Detection System (BioRad, 1855201) were used for RT-qPCR. Fold changes were normalized against *RPLPO* reference gene. For relative quantification, the reaction efficiency incorporated  $\Delta\Delta$ Ct formula was used [60].

### Western blotting

M-PER Mammalian Protein Extraction Reagent (Thermo Fisher Scientific, 78501) with protease and phosphatase inhibitor cocktails (Roche, 05056489001, 4906845001) was used. Lysate concentrations were measured using Pierce BCA Protein Assay Kit (Thermo Fisher Scientific, 23227), and western blotting was performed as described [55]. WesternBright ECL (Advanta, USA, K12045-D50) and Chemidoc MP Imaging System (BioRad) were used for visualization.

### Antibodies

SNX3 antibody (rabbit, 10772-1-AP) was from Proteintech (USA). P-EGFR (rabbit, 2220) and p-AKT (rabbit, 4060) antibodies were from Cell Signaling Technology (USA). EGFR (mouse, sc-373746), AKT (rabbit, sc-8312), p-ERK 1/2 (rabbit, sc-16982), ERK 1/2 (mouse, sc-514312), CTNNB1 (mouse, sc-133240), GFP (mouse, sc-9996), and HA (mouse, MMS-101R) antibodies were from Covance (US). EGFR (rabbit, 06-847) and phosphotyrosine (pY, 4G10, mouse, 05-321) antibodies were from Millipore (Germany). Anti-Transferrin receptor (TfR) was from Thermo Fisher Scientific (13-6890). ACTB (mouse, sc-47778, Santa Cruz Biotechnology) and ACTB (mouse, A5441, Sigma-Aldrich) antibodies were used. HRP-conjugated secondary

antibodies (mouse, R-05071; rabbit, R-05072) were purchased from Advanta. Fluorescent secondary antibodies (IRDye 800CW goat anti-rabbit IgG (H + L) 926-32211, IRDye 800CW goat anti-mouse IgG (H + L) 926-32210, IRDye 680LT goat anti-rabbit IgG (H + L) 926-68021, IRDye 680LT goat anti-mouse IgG (H + L) 926-68020) were purchased from Li-COR (USA). The following antibodies were used in confocal microscopy: SNX3 (Abcam, UK, ab56078), EEA1 (BD Transduction Laboratories, US 610457), Alexa 568 (Thermo Fisher Scientific, A10037), and Alexa 488 (Thermo Fisher Scientific, A21206).

### NanoString

Three independent siRNA transfections to MCF10A were performed, RNAs were pooled and loaded in three replicates. 100 ng of total RNA was loaded for each sample into the cartridge nCounter® Vantage 3D™ RNA Panel (VRXC-Wnt1-12). nSolver and advanced analysis softwares of NanoString Technologies were used to assess the quality of the run and process data. Two normalization analyses were done as recommended by the manufacturer to generate normalized counts. The first normalization included six internal positive controls whose geometric means were calculated, and a normalization factor was used to equilibrate counts. The same process was applied to each gene in the second normalization, using the geometric means of housekeeping genes (CC2D1B, COG7, EDC3, GPATCH3, HDAC3, MTMR14, NUBP1, PRPF38A, SAP130, SF3A3, TLK2, ZC3H14).

### Proliferation assays

pSUPER-NTshRNA and pSUPER-Snx3shRNA stably transfected 4T1 cells were seeded at a density of  $2 \times 10^4$  cells/well on E-plates placed to the RTCA plate station (xCELLigence Real-Time Cell Analyzer, ACEA Biosciences). Impedance was monitored live for 48 h with 15-min intervals. For MTT,  $2 \times 10^3$  cells were plated in 96-well flat-bottom plates and were treated with AG1478 (15  $\mu$ M) or vehicle control. MTT was performed as described [55].

### Colony formation assay

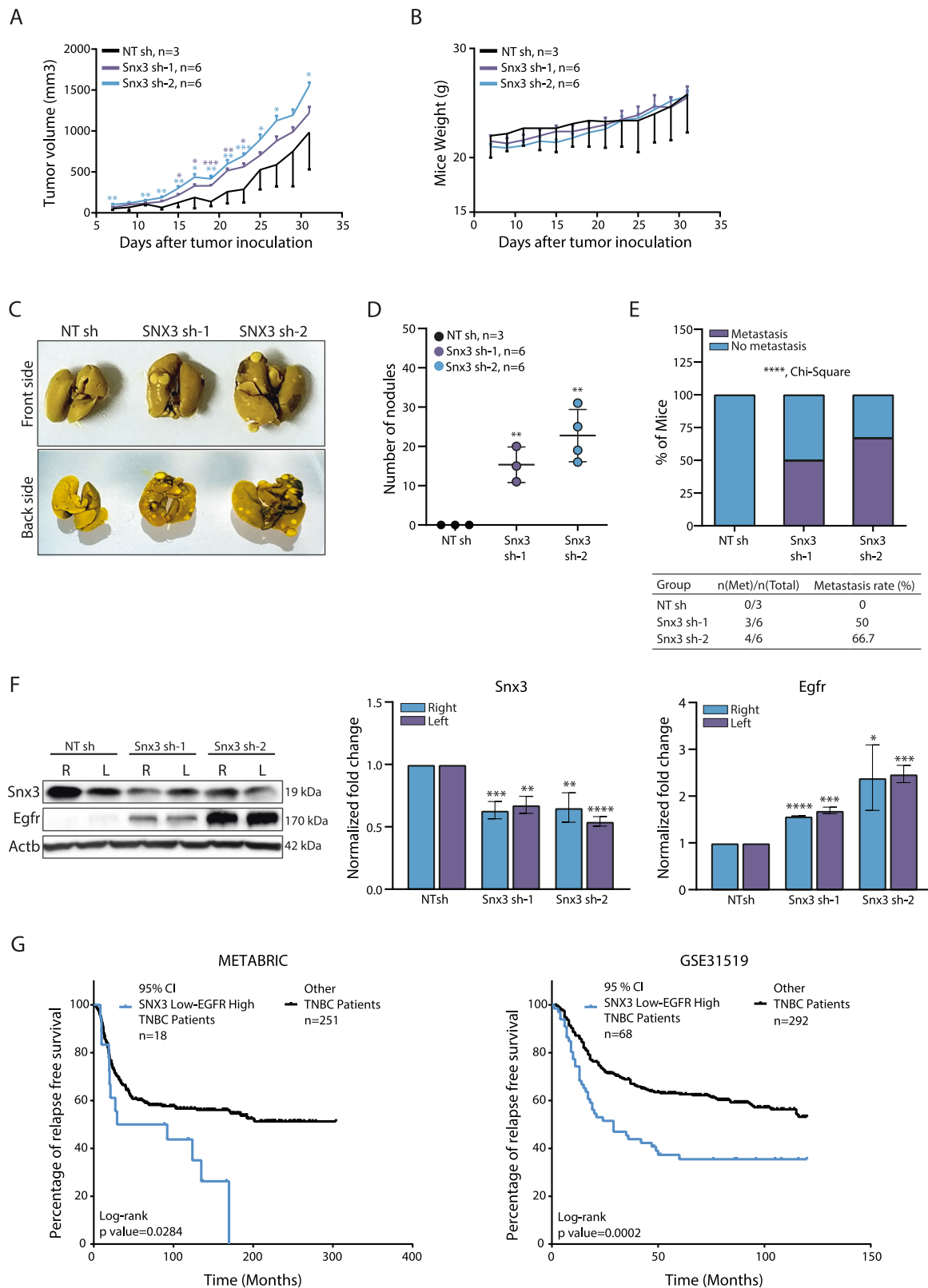
Stably transfected 4T1 cells (500 cells/well) were grown in 6-well plates for seven days. Following methanol fixation, colonies were stained with 0.5% crystal violet as described [61]. Cells were treated with AG1478 (IC50 = 15  $\mu$ M) or vehicle control DMSO for the same duration. Colonies ( $\geq 1000$  pixels) were determined and counted using CountPHICS software [62].

### Transwell migration and invasion assays

$5 \times 10^4$  cells in 100  $\mu$ l of 1%FBS growth medium were seeded onto upper chamber of transwell inserts (Greiner, The Netherlands). 10%FBS-medium was added to bottom chambers. Cells were allowed to migrate for 8 h at 37 °C. For invasion assay, transwell chambers were coated with 100  $\mu$ l Matrigel (300  $\mu$ g/ml, Corning, NY, USA), and  $8 \times 10^4$  cells were seeded per well for 12 h. Staining and counting of cells were performed as previously described [63].

### In vivo tumor growth and metastasis experiments

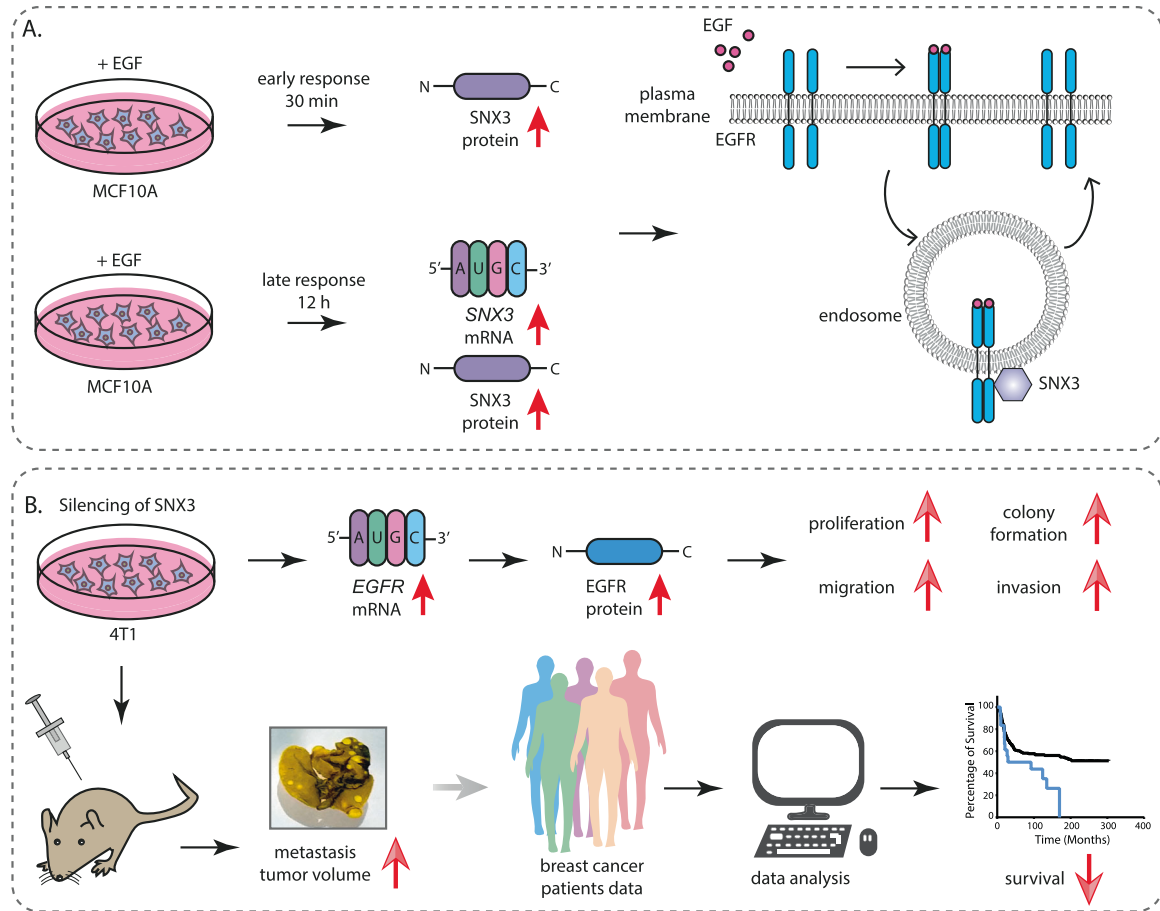
Animal experiments were conducted following institutional ethical guidelines and regulations for animal experimentations of Bilkent University. Mice were housed in a temperature-controlled room with 12-h light/dark cycles and received a standard diet with water ad libitum. Six to eight-week-old female Balb/c mice were injected with  $1 \times 10^6$  4T1



NTsh and two different Snx3sh monoclonal (Snx3sh-1, Snx3sh-2) cells into left and right mammary fat pads (MFP) subcutaneously. Cells were prepared in 1:1 PBS: Matrigel (Corning). Primary tumor growth was monitored by measuring tumor volume every other day with a caliper. Tumor volume was calculated using the formula  $(\text{length} \times \text{width}^2)/2$ . All mice were sacrificed when one of the groups reached the threshold of

1500 mm<sup>3</sup> volume. Tumors were collected for subsequent analysis. Lung metastasis was evaluated with Bouin's fixation [64]. Number of tumor nodules on lung surface was counted under a dissecting microscope, and metastasis rate was determined. One mouse from each injection group was sacrificed, and tumor samples were collected for western blot analysis.

**Fig. 7 Effect of Snx3 loss on tumor progression and metastasis in vivo.** **A** Changes in tumor volume of 4T1 syngeneic model upon Snx3 knockdown. Significance was calculated using multiple t-test for each data point for each day compared with control. **B** Mice body weight (g) of 4T1 syngeneic tumors. **C** Representative images of lungs collected from each group. Mice were sacrificed when one of the groups reached 1500 mm<sup>3</sup>. **D** The frequency of lung metastasis. Significance was determined with Chi-square analysis (\*\*\*\* $p < 0.0001$ ). **E** Number of nodules on the lung surface of each group (\* $p < 0.05$ , \*\* $p < 0.01$ ; \*\*\* $p < 0.001$ , Student's  $t$  test). **F** Primary tumors were collected, and lysates were used for Snx3 and Egfr protein expression analysis. Actb antibody was used as the loading control. Bars show the densitometry analysis of three independent experiments. Expression values of Snx3sh tumor samples were normalized to the average of NTsh tumor samples (\* $p < 0.05$ , \*\* $p < 0.001$ , \*\*\* $p < 0.0001$ , unpaired  $t$ -test) (R: right side, L: left side). **G** Expression of *SNX3* (ILMN\_1740180) and *EGFR* (ILMN\_1696521) mRNAs were determined using  $\pm 95\%$  ( $Z = 1.96$ ) confidence interval in the METABRIC dataset. Relapse-free survival (%) of low *SNX3* and high *EGFR* expressing TNBC breast cancer patients ( $n = 18$ ) was compared to other TNBC patients ( $n = 251$ ). For the GSE31519 dataset, expression values of *SNX3* (2160648\_x\_at) and *EGFR* (201983\_s\_at) mRNAs were determined using  $\pm 95\%$  ( $Z = 1.96$ ) confidence interval. Relapse-free survival of low *SNX3* and high *EGFR* expressing TNBC patients ( $n = 68$ ) was compared to other TNBC patients ( $n = 292$ ).



**Fig. 8 Schematic diagram showing the EGF-SNX3-EGFR axis.** **A** SNX3 protein levels are upregulated during the early response to EGF stimulation. Prolonged exposure to EGF causes transcriptional upregulation. In turn, SNX3, as a retromer protein, interacts with EGFR and endocytosed EGF in early endosomes. **B** Sustained silencing of SNX3 causes *EGFR* mRNA and protein upregulation, causing increased neoplastic phenotypes in vitro and in vivo. In support of these, TNBC patients with low *SNX3* and high *EGFR* mRNA have poor overall survival.

### Patient data analysis

The normalized mRNA expression data for *SNX3* (ILMN\_1740180) and *EGFR* (ILMN\_1696521) were downloaded from METABRIC [39] (European Genome-Phenome Archive with EGAS00000000083), and the clinical data of the study were downloaded from cBioPortal [65, 66]. TNBC patients were grouped based on negativity of ER, HER2, and PR expression ( $n = 318$ ). For disease-specific relapse-free survival (RFS) analyses, the patients who died of other causes were excluded from the analysis. The relapse-free survival was determined by comparing the low *SNX3*, high *EGFR* expressing tumors with other TNBC patients. Grouping was done by filtering the outlier normalized expression values of *SNX3* and *EGFR* according to  $\pm 95\%$  ( $Z = 1.96$ ) confidence interval. Patients in the GSE31519 dataset were also grouped based on low *SNX3* and high *EGFR* mRNA levels.

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## AUTHOR CONTRIBUTIONS

EC performed EGF induction experiments, AC developed and characterized RNAi models of SNX3, MO performed siRNA transfections, biotinylation, and microscopy experiments, OAC performed in vivo experiments, DND performed survival analysis, SGE and RCA performed and guided NanoString hybridizations, respectively, AS and HO oversaw protein interaction and localization experiments, OS managed mouse experiments, AEED designed and supervised experiments, and wrote the paper. All authors, except late HO read and commented on the paper.

## COMPETING INTERESTS

The authors declare no competing interests.

## ADDITIONAL INFORMATION

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