

Active G*α*/o mutants accelerate breast tumor metastasis via the c-Src pathway

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Conflicts of Interest Statement

The authors have declared that no conflict of interest exists.

Abstract

Constitutively active mutations in the $G\alpha_{i2}$ and $G\alpha_{oA}$ subunits of heterotrimeric G proteins have been identified in several human cancers including breast cancer, but their functional significance in tumorigenesis and metastasis has not been well characterized. In this study, we show that expression of the constitutively active $G\alpha_{oA}R243H$ and $G\alpha_{i2}R179C$ mutants alone was insufficient to induce mammary tumor formation in mice. However, in transgenic mouse models of breast cancer induced by Neu expression or PTEN loss, we found that the $G\alpha_{i2}R179C$ mutant enhanced spontaneous lung metastasis while having no effect on primary tumor initiation and growth. Additionally, we observed that ectopic expression of the $G\alpha_{oA}R243H$ and $G\alpha_{i2}R179C$ mutants in tumor cells promote cell migration *in vitro* as well as dissemination into multiple organs *in vivo* by activating c-Src signaling. Thus, our study uncovers a critical function of $G\alpha_{i/o}$ signaling in accelerating breast cancer metastasis via the c-Src pathway. This work is clinically significant, as it can potentially pave the way to personalized therapies for patients who present with active $G\alpha_{i/o}$ mutations or elevated $G\alpha_{i/o}$ signaling by targeting c-Src to inhibit breast cancer metastasis.

Introduction

Breast cancer is the most commonly diagnosed cancer in women (1). As of 2020, breast cancer surpassed lung cancer to become the most prevalent form of cancer worldwide (1). In the United States, approximately 1 in 8 women is expected to be diagnosed with breast cancer in her lifetime (1). Despite major and rapid advances in therapeutic approaches, breast cancer remains the leading cause of death from malignant tumors among women. Metastasis is the major cause of tumor mortality. While the average 5-year survival rate for localized breast cancer is 99%, it falls to only 29% for advanced breast cancer diagnoses with distant metastasis (www.cancer.org). Since currently available therapies have very limited efficacy in treating metastatic disease, further research into the mechanisms of tumor metastasis is crucial. Understanding these mechanisms on a molecular level is critical for developing novel therapies that will ultimately improve the survival rates of advanced breast cancer patients.

G protein coupled receptors (GPCRs) are the largest cell surface receptor family that regulates diverse cellular processes and functions (2). GPCRs transmit signals through heterotrimeric G proteins that consist of $G\alpha$, $G\beta$ and $G\gamma$ subunits. Based on sequence and functional similarity of $G\alpha$ subunits, heterotrimeric G proteins are divided into four major subclasses: $G_{i/o}$, G_s , $G_{q/11}$ and $G_{12/13}$ (3, 4). GPCRs regulate diverse cellular processes including cell recognition, proliferation and motility. Not surprisingly, dysregulated GPCR and G protein signaling has been implicated in cancer initiation, progression, and metastasis (5).

Current data suggest that more than 20% of tumors contain mutations in GPCRs and G proteins (6). Activating mutations in GNAQ, GNA11 and GNAS have been shown to act as oncogenic drivers to promote tumorigenesis in several tumor types including uveal melanomas, intestinal cancers, and pancreatic adenocarcinomas (7-12). Previous work has also identified several activating mutations in the $G\alpha$ subunits of the $G_{i/o}$ subclass, specifically GNAI2 (R179C and R179H) and GNAO1 (R209C and R243H), in multiple tumor types including ovarian and adrenal tumors (13), as well as breast cancer and leukemia (14-16). A recent study by Song et al. has established the role of GNAO1 R209C in tumorigenesis of acute lymphoblastic leukemia (16). However, the functional significance of the other $G\alpha_{i/o}$ mutants remains to be determined in clinically relevant tissues and models *in vivo*.

Our previous work found that many $G_{i/o}$ -coupled GPCRs are upregulated in breast cancer, and the upregulated $G_{i/o}$ -coupled GPCRs play a significant role in HER2-induced breast tumor initiation, growth, and metastasis (17, 18). $G_{i/o}$ -coupled GPCRs can transmit signals through both the dissociated $G\alpha_{i/o}$ and $G\beta\gamma$ subunits. We and others previously showed that $G\beta\gamma$ serves as a point of convergence for many $G_{i/o}$ -coupled GPCR signals mediating breast cancer growth and metastasis (19, 20), but the role of $G\alpha_{i/o}$ signaling in breast cancer progression remains to be characterized. In this study, we investigate the function of activating $G\alpha_{i/o}$ mutations in breast cancer development, using cell lines and transgenic models. Our experiments revealed that the active $G\alpha_{i2}$ R179C and $G\alpha_{oA}$ R243H mutants alone were insufficient to induce breast cancer formation. In the transgenic mouse models of breast cancer induced by Neu expression or PTEN loss, the $G\alpha_{i2}$ R179C mutant enhanced spontaneous lung metastasis but did not affect mammary tumor formation and growth. Additional data showed that the active $G\alpha_{i2}$ R179C and $G\alpha_{oA}$ R243H mutants promote breast cancer metastasis through activation of the c-Src pathway. Together, our study not only elucidates the function of active GNAI2 and GNAO1 mutations in breast cancer metastasis but also provides compelling evidence to suggest that targeting the c-Src pathway may be an effective way to block tumor metastasis in breast cancer patients with active $G\alpha_{i/o}$ mutants or elevated $G\alpha_{i/o}$ signaling.

Results

The active $G\alpha_{i2}$ R179C and $G\alpha_{oA}$ R243H mutants alone are insufficient to induce mammary tumor formation.

To determine the role of the $G\alpha_{i2}$ R179C and $G\alpha_{oA}$ R243H mutants in breast cancer formation, we expressed each of these mutants in a mammary epithelial cell line stably expressing firefly luminescence,

MCF10A-FLU, using a doxycycline-inducible system (Figure 1A). As observed in Figure 1B, inducing expression of either $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ in MCF10A cells activated c-Src and STAT3 as previously reported for $G\alpha_{oA}R243H$ in NIH3T3 cells (15). Activation of both c-Src and STAT3 was successfully blocked using the specific inhibitors, saracatinib (Sara) and STAT3-IN-1 (STAT3i), respectively. Interestingly, inhibition of c-Src by saracatinib did not affect STAT3 activation by $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$. Similarly, inhibition of STAT3 by STAT3-IN-1 had no effect on c-Src activation (Figure 1B). These observations suggest that $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ activate c-Src and STAT3 through independent mechanisms.

In 2D cell culture, MCF10A cells expressing $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ grew faster than control cells expressing GFP (Figure 1C). However, when grown in Matrigel, these cells did not show a significant difference in growth (Figure 1D). We then implanted MCF10A cells expressing either GFP, $G\alpha_{i2}R179C$, or $G\alpha_{oA}R243H$ into the mammary gland of nude mice. After monitoring the mice for tumor growth over a period of four months, using mammary palpation and highly sensitive bioluminescence imaging, we found that none of the cells formed detectable tumors in any of the mice (Figure 1E). These results suggest that activation of $G\alpha_{i/o}$ signaling alone is insufficient to drive the formation of breast tumors.

To further confirm these findings, we generated mice to selectively express $G\alpha_{i2}R179C$ in the mammary gland using the tetO promoter system. We crossed transgenic mice carrying $G\alpha_{i2}R179C$ under the tetO promoter with transgenic mice expressing the transactivator, tTA, from the mammary gland-specific MMTV promoter, MMTV-tTA. Western blot comparison of mammary glands from 5-month-old mice expressing tTA alone and tTA/ $G\alpha_{i2}R179C$ ($G\alpha_{i2}R179C$) confirmed $G\alpha_{i2}R179C$ expression in the tTA/ $G\alpha_{i2}R179C$ mice in the absence of doxycycline treatment (Figure 2A). Neither the littermate control (tTA) mice, nor $G\alpha_{i2}R179C$ mice formed palpable tumors for up to 20 months. Additionally, whole-mount in situ staining of mammary glands extracted from 3- and 13-month-old tTA and $G\alpha_{i2}R179C$ 2 mice did not reveal any differences in morphology (Figure 2B). This indicates that $G\alpha_{i2}R179C$ does not affect normal mammary development and does not induce mammary gland hyperplasia. Together, these results indicate that activating mutations in GNAI2 or GNAO1 alone are insufficient to induce breast cancer formation.

$G\alpha_{i/o}$ signaling promotes breast cancer metastasis

Since $G_{i/o}$ -GPCR signaling is upregulated in HER2⁺ breast cancer, we further explored the role of elevated $G\alpha_{i/o}$ signaling in Neu tumor formation. We generated trigenic mice, tTA/ $G\alpha_{i2}R179C$ /Neu (Neu/ $G\alpha_{i2}R179C$), by crossing tTA/ $G\alpha_{i2}R179C$ mice with MMTV-Neu (Neu) transgenic mice, which selectively express an activated rat ErbB2/HER2 homologue in their mammary glands. The Neu mice spontaneously formed mammary tumors, which then metastasized to the lungs. The median time for tumor formation in the Neu mice (167 days) was not significantly different from the Neu/ $G\alpha_{i2}R179C$ mice (174 days) (Figure 2A). There was also no significant difference between tumor growth in Neu mice compared to Neu/ $G\alpha_{i2}R179C$ mice (Figure 2C). However, $G\alpha_{i2}R179C$ significantly increased the number of Neu-induced lung metastases (Figure 2D).

To test whether the $G\alpha_{i2}R179C$ mutant might regulate the progression of breast cancer that has already been induced by other existing genetic abnormalities, such as deletion of the tumor suppressor gene, PTEN (Phosphatase And Tensin Homolog), we crossed tTA/ $G\alpha_{i2}R179C$ mice with transgenic mice exhibiting mammary gland-specific deletion of one PTEN allele (MMTV-Cre/PTEN^{fl/+}; PTEN^{+/-}). As seen in Figure 3A, PTEN^{+/-} mice generated mammary tumors after an extended latency (407 days) compared to Neu mice (167 days). Following a similar trend as we saw previously with Neu/ $G\alpha_{i2}R179C$ mice, $G\alpha_{i2}R179C$ expression in PTEN^{+/-} mice (PTEN^{+/-}/ $G\alpha_{i2}R179C$) did not affect PTEN loss-induced tumor onset and growth (Figure 3A and 3B), but it did significantly increase the number of lung metastases (Figures 3C and 3D). These results suggest that although $G\alpha_i$ signaling is not required for the primary growth of mammary tumors, it is critical in the metastasis of tumors to distant organs.

We then question whether $G\alpha_{oA}$ signaling may play a similar role as $G\alpha_i$ signaling in promoting breast tumor metastasis. To this end, we injected Neu cells expressing inducible GFP or $G\alpha_{oA}R243H$ into the left ventricle of nude mice to mimic the metastatic spread of tumor cells into multiple organs. Beginning immediately after injection, the mice were continuously fed with doxycycline-containing chow to induce GFP and $G\alpha_{oA}R243H$ expression. Bioluminescence imaging showed that the $G\alpha_{oA}R243H$ -injected mice exhibited enhanced tumor growth compared to the GFP-injected mice (Figure 4A). While all the injected mice formed tumors in multiple organs, in particular the lung, there were significantly more metastatic tumors found in the mice injected with $G\alpha_{oA}R243H$ -expressing Neu cells (Figure 4B). These results therefore confirm that $G\alpha_{oA}$ signaling is also important in tumor metastasis.

To understand the mechanisms by which $G\alpha_i$ and $G\alpha_{oA}$ signaling promote tumor metastasis, we determined their impact on cell proliferation and migration. Neu cells expressing GFP, $G\alpha_{i2}R179C$, and $G\alpha_{oA}R243H$ grew at a similar rate, showing no significant difference in proliferation capacity (Figure 4C). Notably, however, the Neu cells expressing $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ exhibited enhanced trans-well migration ability compared to cells expressing GFP (Figure 4D). This difference is evident in the basal state but is amplified upon treatment with LPA. Data from wound-healing assays also demonstrated increased migration capacity of $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ expressing cells compared to cells expressing GFP (Figure 4E). These results suggest that $G\alpha_{i/o}$ signaling may promote tumor metastasis by enhancing tumor cell motility.

$G\alpha_{i/o}$ signaling promotes tumor metastasis via the c-Src pathway

We showed previously that $G_{i/o}$ -coupled GPCRs transactivate EGFR and HER2, and that their signaling converges at the AKT and c-Src pathways to promote Neu-induced breast cancer growth and metastasis (17). Western blotting analysis shows that, compared with Neu tumors, $G\alpha_{i2}R179C$ tumors showed significantly increased phosphorylation of EGFR^{Y1068} and c-Src^{Y416} but no significant changes in phosphorylation of HER2^{Y1221/1222}, STAT3^{Y705} and AKT^{S473} (Figure 5A and 5B). Similarly, phosphorylation of c-Src^{Y416}, but not AKT^{S472}, STAT3^{Y705} and EGFR^{Y1068}, was significantly increased in PTEN^{+/-}/ $G\alpha_{i2}R179C$ tumors as compared to PTEN^{+/-} tumors (supplemental Figure 1A and 1B). These findings suggest that $G\alpha_{i/o}$ signaling might be involved in $G_{i/o}$ -GPCR-stimulated c-Src activation, and in a context-dependent manner, EGFR transactivation.

Additional immunoblotting data showed that phosphorylation of EGFR^{Y1068} and c-Src^{Y416} was increased in Neu cells expressing $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$, as compared to Neu cells expressing GFP (Figure 5C). In contrast, there appears to be no significant difference in AKT^{S473}, STAT3^{Y705} and HER2^{Y1221/1222} phosphorylation (Figure 5C). Furthermore, treatment of Neu cells with the Src-specific inhibitor, saracatinib, abolished the observed increase in both c-Src^{Y416} and EGFR^{Y1068} phosphorylation in Neu cells expressing $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$. This finding suggests that EGFR is likely a downstream target for transactivation by c-Src. This proposition was corroborated by the finding that EGFR^{Y1068} phosphorylation was increased after inducing expression of a constitutively active c-Src mutant, c-Src^{Y527F} in Neu cells (Figure 5D).

We then sought to identify the mechanisms by which $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ may activate c-Src. We first considered the possibility of the $G\beta\gamma$ subunit being involved. Theoretically, the active $G\alpha_{i/o}$ mutants could release free $G\beta\gamma$ subunits, which could activate c-Src independent of the $G\alpha_{i/o}$ subunits (21). To test this, we inhibited $G\beta\gamma$ signaling by overexpressing $G\alpha_t$, a scavenger of $G\beta\gamma$, in Neu cells as we previously reported (20). As expected, overexpression of $G\alpha_t$ suppressed LPA-stimulated ERK phosphorylation, which is known to be activated by $G\beta\gamma$ (20), in Neu cells expressing GFP or $G\alpha_{i2}R179C$ (supplemental Figure 2A). However, the enhanced c-Src activation in Neu cells expressing $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ was unaffected by $G\alpha_t$ overexpression (supplemental Figure 2B). These findings indicate that the active $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ mutants activate c-Src independent of the dissociated $G\beta\gamma$ subunits.

We then pursued the idea that $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ may activate c-Src by direct interaction as previously reported for $G\alpha_i$ (22). We performed co-immunoprecipitation assays with Neu cell lysates and found that $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ interacted with c-Src, but not $G\beta\gamma$ (Figure 5E). Transwell migration assays also showed that overexpression of $G\alpha_t$ or G protein coupled receptor kinase 2 (GRK2-ct), another

G $\beta\gamma$ scavenger, did not affect the enhanced Neu cell migration induced by G α_{oA} R243H (supplemental Figure 2C). Suppression of c-Src by saracatinib, however, abolished the enhanced cell migration by G α_{i2} R179C and G α_{oA} R243H while not significantly altering the migration of GFP-expressing cells (Figure 6A). These results indicate that G α_{i2} R179C and G α_{oA} R243H promote tumor cell migration via the c-Src pathway.

To validate our suppositions *in vivo*, we established tumor metastases in multiple organs by injecting Neu cells expressing GFP or G α_{oA} R243H into the left ventricle of nude mice and feeding them doxycycline-containing chow to induce GFP and G α_{oA} R243H expression. Eight days post-injection, the mice were treated with a vehicle control or saracatinib and tumor growth was monitored by bioluminescence imaging. As measured by whole-body bioluminescence intensity, saracatinib treatment did not affect the growth of GFP-expressing Neu tumors but diminished the enhanced growth of G α_{oA} R243H-expressing Neu tumors (Figure 6B). Moreover, the overall survival of mice injected with either GFP- or G α_{oA} R243H-expressing Neu cells was significantly increased by saracatinib (Figure 6C; 20 and 20 days for vehicle control vs 28 and 25 days for saracatinib, respectively). At the end of the treatment for two weeks, *ex vivo* bioluminescence imaging of key tissues from the mice harvesting GFP tumors showed that GFP tumors primarily grew in the lung and heart and saracatinib treatment significantly reduced tumor growth in these tissues (Figure 6D). Together, these results indicate that G $\alpha_{\text{i/o}}$ signaling can accelerate Neu tumor metastasis via the c-Src pathway.

Discussion

In this study, we demonstrated that overexpression of the constitutively active G α_{i2} R179C or G α_{oA} R243H mutant alone in mammary epithelial cells is insufficient to induce mammary tumor formation but can cooperate with Neu/HER2 hyperactivation or PTEN deficiency to drive tumor metastasis. Somatic GNAI2 R179C and GNAO1 R243H mutations have been found in human cancers, including ovarian and adrenal tumors as well as breast cancer (13-15). Although these mutants were shown to promote oncogenic transformation and anchorage-independent growth of fibroblasts or mammary epithelia cells *in vitro*, their function as oncogenes has never been characterized in relevant tissues *in vivo*. Thus, our study provides the first evidence for the significant role of these mutants in breast cancer progression.

Notably, the active G α_{i2} R179C and G α_{oA} R243H mutants selectively affect breast tumor cell migration and tumor metastasis but have no effect on tumor cell proliferation *in vitro*. The G α_{i2} R179C mutant also shows no significant effect on tumor initiation and growth in the two different genetic mouse models of breast cancer, Neu overexpression and PTEN deficiency. The mechanisms by which G α_{i2} R179C and G α_{oA} R243H selectively enhance tumor cell migration and tumor metastasis are unclear but may be related to their ability to activate only a limited number of pathways. Indeed, in Neu cells and tumor samples, G α_{i2} R179C and G α_{oA} R243H primarily enhance c-Src and EGFR transactivation. This is in stark contrast to our previous data showing that G i/o -coupled receptor signaling contributes to the activation of multiple pathways, including c-Src and AKT, and transactivation of both EGFR and HER2 (17). The activation of additional pathways by G i/o -GPCRs likely involve G $\beta\gamma$ subunits liberated from heterotrimeric G i/o proteins since activation of heterotrimeric G i/o proteins by GPCRs generates both functional G $\alpha_{\text{i/o}}$ and G $\beta\gamma$ subunits and G $\beta\gamma$ is known to be involved in PI3K/AKT activation (3, 4, 23-25). Moreover, we and others have established that G $\beta\gamma$ signaling is critical for breast cancer cell growth and migration induced by multiple G i/o -GPCRs (20, 24). Given that blocking G i/o -coupled receptor signaling not only delays Neu-induced tumor initiation, but also suppresses tumor growth and metastasis (17), G i/o -coupled GPCRs likely signal through both G $\alpha_{\text{i/o}}$ and G $\beta\gamma$ to accelerate breast tumor progression. While G $\alpha_{\text{i/o}}$ signaling primarily drives tumor metastasis, G $\beta\gamma$ signaling may promote both primary tumor growth and metastasis.

Our data indicate that G α_{i2} R179C and G α_{oA} R243H primarily activate the c-Src pathway to promote tumor metastasis. In breast tumor cells and tissues, overexpression of G α_{i2} R179C and G α_{oA} R243H enhanced c-Src activation. These findings are consistent with several reports that active G α_{oA} mutants can activate c-Src in fibroblasts (15, 26). The activation of c-Src by G α_{i2} R179C and G α_{oA} R243H does not seem to involve G $\beta\gamma$ since

co-expression with a G $\beta\gamma$ scavenger has no effect on c-Src activation. Rather, G α_{i2} R179C and G α_{oA} R243H may activate c-Src by direct interaction as previously reported for active G α_i since they are co-immunoprecipitated with c-Src in breast tumor cells (22). We cannot, however, exclude the possibility that there are other proteins mediating or regulating this interaction.

The activation of c-Src by G α_{i2} R179C and G α_{oA} R243H is functionally important since blocking c-Src activation with a specific inhibitor, saracatinib, suppresses the mutant-promoted tumor cell migration *in vitro* and metastatic spread of tumor cells to multiple organs *in vivo*. Saracatinib, however, has no significant effect on Neu-driven tumor cell migration and tumor metastasis, although it significantly improves the survival of Neu mice. The reason for the differential effect of saracatinib on GFP tumor growth and overall survival is unclear but may be related to the inhibition of metastatic tumor growth in critical organs such as the lung and heart, as indicated by *ex vivo* bioluminescence imaging of the tissues from the surviving mice at the end of the experiments. These results suggest that Neu-driven tumor metastasis may involve multiple redundant signaling pathways and combining Src inhibition with other therapeutic reagents may be required for effectively blocking Neu tumor metastasis.

We also found that activation of c-Src by G α_{i2} R179C and G α_{oA} R243H causes EGFR transactivation in Neu cells and Neu tumors, supporting a functional interaction between c-Src and EGFR as previously reported (27). However, although G α_{i2} R179C also enhances c-Src activation in PTEN tumors, it does not cause EGFR transactivation, suggesting the role of c-Src in EGFR transactivation is context dependent. Since G α_{i2} R179C promotes tumor metastasis in both Neu and PTEN tumor models, these results also suggest that the transactivation of EGFR by c-Src is not essential for G α_{i2} R179C- and G α_{oA} R243H-promoted tumor metastasis.

Once activated, c-Src can interact with various substrates and effectors, including STAT3, to promote tumor cell proliferation, migration, and invasion (28, 29). Previous studies showed that the active G α_{i2} and G α_{oA} mutants can induce oncogenic transformation of NIH3T3 fibroblasts via the c-Src-stimulated STAT3 pathway (26, 30). We found that G α_{i2} R179C and G α_{oA} R243H can activate STAT3 in MCF10A cells but have no effects in Neu cells. Moreover, the expression of G α_{i2} R179C enhances c-Src but not STAT3 activation in Neu and PTEN^{+/-} tumors. The activation of STAT3 by G α_{i2} R179C and G α_{oA} R243H in MCF10A cells also does not seem to involve c-Src. These findings suggest that the activation of STAT3 by the G α_{i2} and G α_{oA} mutants is likely to be cell type dependent. Further studies are required to identify the downstream effectors of c-Src that mediate G α_{i2} R179C- and G α_{oA} R243H-promoted tumor cell migration and metastasis.

In conclusion, we demonstrate through both *in vitro* and *in vivo* studies that active G $\alpha_{i/o}$ mutants accelerate breast cancer metastasis via the c-Src pathway. In addition to mutations in G $\alpha_{i/o}$ genes, multiple G $\alpha_{i/o}$ -coupled receptors and G $\alpha_{i/o}$ proteins are frequently upregulated in many cancers including breast cancer (6, 17, 31). Moreover, the activity of c-Src is increased in a variety of neoplastic tissues and targeting c-Src has been proposed as a promising strategy for blocking breast cancer metastasis (29, 32-34). Given that tumor metastasis is a major cause of cancer mortality, our findings that c-Src is a key mediator of G $\alpha_{i/o}$ signaling in promoting tumor metastasis have important ramifications in the development of personalized therapies to improve the survival of breast cancer patients with active G $\alpha_{i/o}$ mutants or enhanced G $\alpha_{i/o}$ signaling.

Methods

Reagents

Saracatinib was obtained from LC Laboratories. Antibodies for EGFR (no. 2232), phospho-EGFR^{Y1068} (no. 3777), EGFR (no. 4267), HER2 (no. 2165), phospho-HER2^{Y1221/1222} (no. 2243), AKT (no. 4685), phospho-AKT^{S473} (no. 4060), Src (no. 2109), phospho-Src^{Y416} (no. 6943), ERK1/2 (no. 4696) and phospho-ERK1/2^{T202/Y204} (no. 4370) were obtained from Cell Signaling Technology; GADPH (sc-47724) from Santa Cruz Biotechnology; mouse anti- α -tubulin antibody (no. 901801) from Biolegend.

Cell lines

Neu cells were generated from tumors formed by the transgenic mice, MMTV-c-Neu, and cultured in DMEM media containing 10% FCS. Cell lines were tested for *Mycoplasma* using a Mycoplasma Detection kit (ATCC). MCF10A cells were purchased from the ATCC, and cultured in DMEM/F12 media containing 5% horse serum supplemented with EGF at 20ng/ml, hydrocortisone at 500ng/ml, cholera toxin at 100ng/ml, and insulin at 10µg/ml. Each cell line was cryopreserved at low passage numbers (less than six passages after receipt) and used in experiments for a maximum of 18 passages.

Plasmid

The lentiviral vector (Tet-CA-Src-GFP) for tetracycline-inducible expression of the GFP-tagged, constitutively active Src mutant, Src/Y527F, was obtained from Addgene (item #83469). $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ mutants were constructed using pcDNA3.1 wild-type human ee-tagged- $G\alpha_{i2}$ - and ee-tagged- $G\alpha_{oA}$ plasmids as template (cDNA Resource Center) and Q5 Site-directed mutagenesis kit (New England BioLabs). The lentiviral vector for tetracycline-inducible expression of the $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ mutants was constructed by first cloning $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ from pcDNA3.1 to pENTR vector (ThermoFisher Scientific) and then into the destination vector pLIX_402 (Addgene, item #41394) using the Gateway cloning system.

The adenovirus vectors for $G\alpha_t$ and GRK2ct were constructed by first cloning $G\alpha_t$ and GRK2ct into the pENTR vector and then into the destination vector, pAd/CMV/DEST (ThermoFisher Scientific), using the Gateway cloning system.

Lentiviral production

Lentiviruses were generated in HEK293FT cells as described previously (23). Lentiviruses collected from the cell culture supernatants were concentrated using the Lenti-X-concentrator (Takara Bio).

Adenoviral production and transduction

Adenoviruses were generated in HEK293A cells using the ViraPower Adenoviral Expression System (ThermoFisher Scientific) and purified with a $CsCl_2$ density gradient combined with ultracentrifugation. Transduction of Neu cells was conducted by incubation of cells with purified adenoviruses overnight.

Establishment of stable cell lines

The MCF10A and Neu cells were transduced with lentiviruses encoding GFP-tagged Src^{Y527F}, GFP, $G\alpha_{i2}R179C$, or $G\alpha_{oA}R243H$, and selected with puromycin (2 µg/ml) for at least 1 week to establish stable lines.

Cell proliferation and viability assays

Cell proliferation in two-dimensional culture or in Matrigel was analyzed as we described previously (17, 23). Cell viability was quantified using AlamarBlue (Thermo Fisher Scientific) assays as per the manufacturer's instructions.

Cell migration and wound healing assays

Transwell migration and wound healing assays were performed as we described previously (17, 23). To exclude the influence of cell proliferation, cells were treated with 5 µg/ml mitomycin.

Western blotting analysis

Protein lysates were prepared from cells and tumor tissues and analyzed by Western blotting as we described. Blots were imaged using the iBright 1500 (Thermo Fisher Scientific) or Odyssey (LI-COR Biotechnology) imaging system (17, 23).

Co-immunoprecipitation assay

Neu cells were lysed in modified RIPA buffer containing 50mM Tris-HCl, 150mM NaCl, 1mM EDTA and 1% Triton, pH 7.4 as we reported previously. Cell lysates were incubated overnight at 4 °C with mouse anti-ee antibodies pre-bound to Dynabeads protein G (Thermo Fisher Scientific). The immunoprecipitated protein complex was analyzed by Western blotting.

Mouse studies

MMTV-c-Neu (no. 005038), TetO- $G\alpha_{i2}$ R179C (no. 017333), MMTV-Cre (no. 003553) and PTEN^{fl/fl} (no. 034621) mice were purchased from the Jackson Laboratory and MMTV-tTA mice were obtained from Dr. Wagner's laboratory (50). All mice were in the FVB/N genetic background. Mice were genotyped by PCR as reported previously (24, 50, 51). Female mice were kept as virgins throughout the experiments. To determine tumor onset, mice were checked twice per week by palpation beginning four months after birth. To assess tumor progression, the largest tumor was measured weekly by caliper. To evaluate lung metastasis, the lung was harvested once the largest tumor reached a size of 2 cm in diameter and was perfused and fixed with 4% paraformaldehyde before paraffin embedding. The number of metastases in the lung was analyzed by serial sectioning followed by HE staining.

To generate metastatic mouse models, Neu cells ($\sim 5 \times 10^5$ in 100 μ l of phosphate-buffered saline) expressing luciferase and doxycycline-inducible GFP or $G\alpha_{oA}$ R243H were injected into 6–8-week-old female nude mice (Charles River) via left ventricle. Immediately post-injection, bioluminescence imaging was performed to confirm successful injection. Mice were then continuously fed with a doxycycline-containing diet (TD.01306; ENVIGO) to induce target gene expression. Tumor growth was monitored by bioluminescence imaging of the mice at both dorsal and ventral positions as we reported previously. To determine the effect of c-Src inhibition, mice were administered with saracatinib (25mg/kg, daily gavage) dissolved in 2% DMSO, 30% PEG300 and H₂O, eight days post-injection.

Statistics

Data were expressed as mean $\pm$ SEM. Statistical comparisons between groups were analyzed by two tail Student's *t* test or ANOVA ($P < 0.05$ was considered significant). The survival curves were analyzed according to the Kaplan–Meier method.

Study approval

All animal studies were conducted in accordance with an IACUC-approved protocol at the University of Iowa.

Authors' Contributions

Data curation, C. Lyu, A. Bhimani and W. Draus. Formal analysis, C. Lyu and S. Chen; Methodology, C. Lyu; Project administration, R. Weigel and S. Chen; Writing-original draft, S. Chen.; Writing-review & editing, A. Bhimani, R. Weigel and S. Chen.

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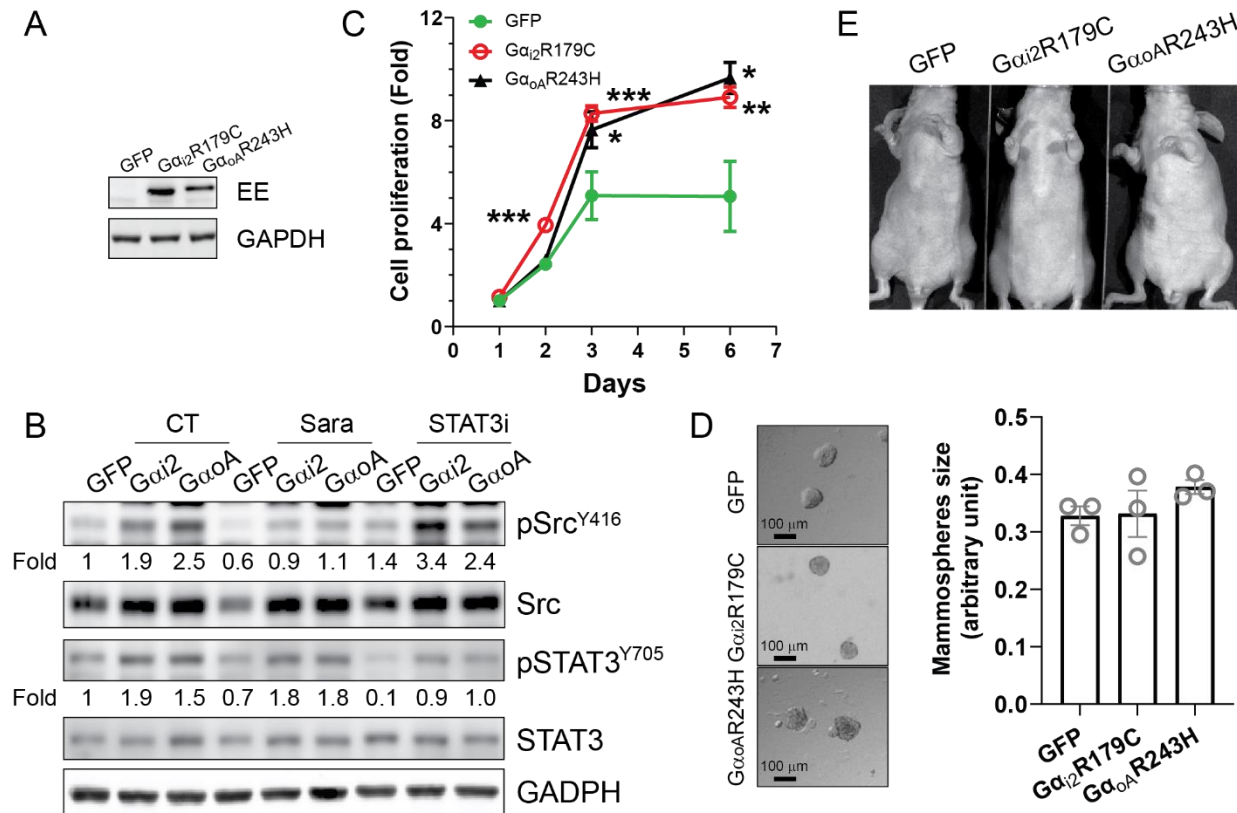


Figure 1

Figure 1. The active $G\alpha_{i/o}$ mutants alone are insufficient to cause neoplastic transformation of mammary epithelial cells. **A-B**, representative immunoblots showing the expression of $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ in MCF10A cells (**A**) and activation of c-Src and STAT3 by phosphorylation of Src^{Y416} and STAT3^{Y705} in MCF10A cells treated with vehicle control (CT), 2 μ M of saracatinib (sara) and STAT3-IN-1 (STAT3i) (**B**). **C-D**, the effect of $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ expression on the growth of MCF10A cells in 2D culture (**C**) and in Matrigel (**D**). **E**, representative images showing the absence of tumor formation in nude mice orthotopically implanted with MCF10A cells expressing GFP, $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$. *, **, *** $p < 0.05$, 0.01 and 0.001 vs. GFP, $n = 4-6$. One-way ANOVA was used for statistical analysis in this Figure.

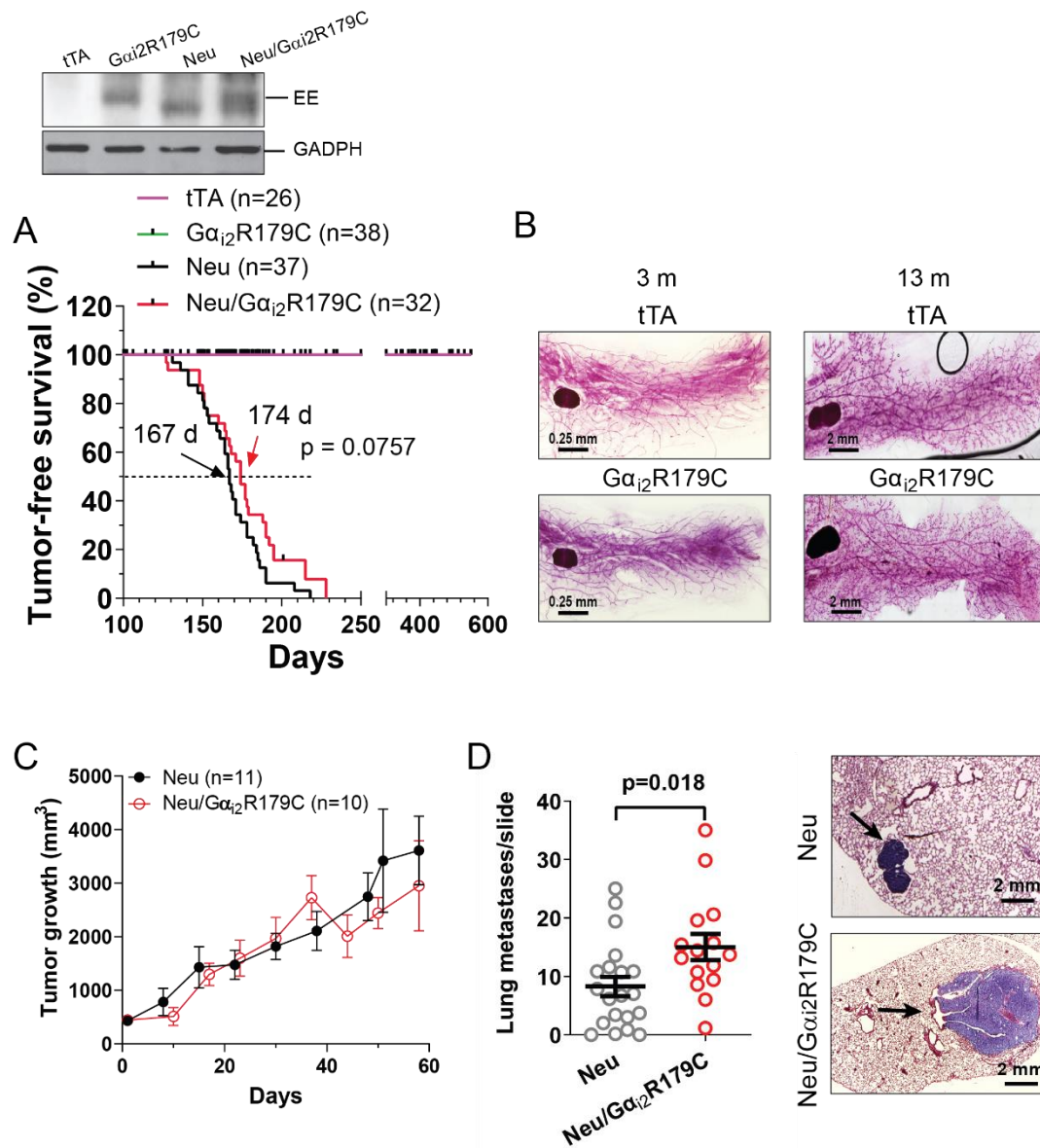


Figure 2

Figure 2. Expression of Gα_{i2}R179C neither induces mammary tumor formation nor affects HER2-driven primary tumor development, but accelerates lung metastasis. **A**, tumor-free survival curves of mice expressing tTA, tTA/Gα_{i2}R179C (Gα_{i2}R179C), tTA/Neu (Neu), and tTA/Neu/Gα_{i2}R179C (Neu/ Gα_{i2}R179C), and representative immunoblots showing ee-tagged Gα_{i2}R179C expression in the mammary gland and Neu tumors. **B**, whole-mount *in situ* staining showing the morphology of the mammary glands from 3 and 13-month-old tTA and Gα_{i2}R179C mice. **C**, Neu and Neu/Gα_{i2}R179C tumor growth curves. **D**, a graph to show the number of lung metastases and images representative of HE-stained lung metastasis (indicated by arrows) from the transgenic Neu and Neu/Gα_{i2}R179C mice. Two tail unpaired Student's *t* test was used for statistical analysis in this Figure.

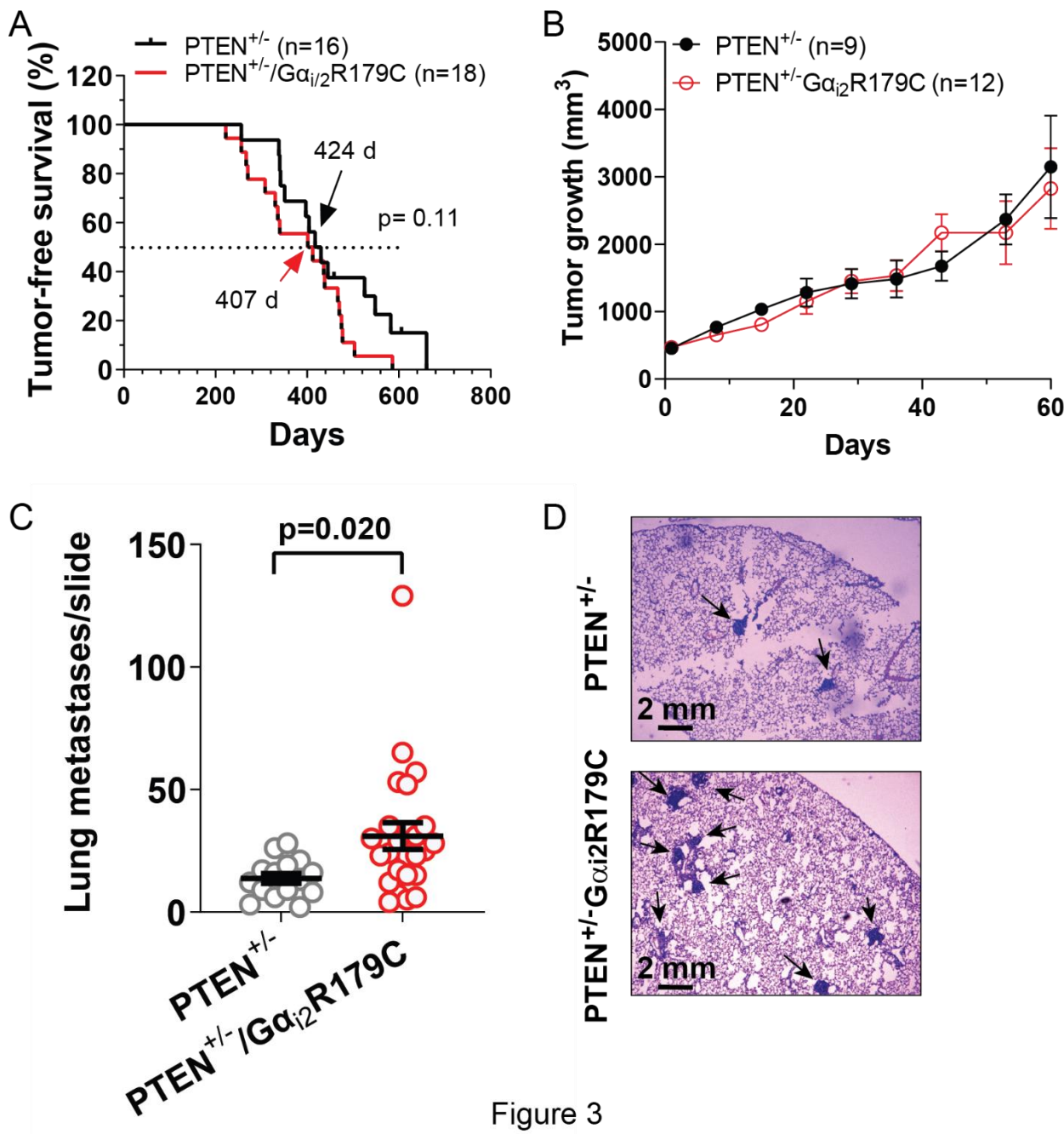


Figure 3

Figure 3. Expression of Gα_{i2}R179C enhanced PTEN deficiency-driven lung metastasis. **A**, tumor-free survival curves of MMTV-Cre/PTEN^{fl/+} (PTEN^{+/-}) and MMTV-Cre/PTEN^{fl/+}/MMTV-tTA/Gα_{i2}R179C (PTEN^{+/-}/Gα_{i2}R179C) mice. **B**, PTEN^{+/-} and PTEN^{+/-}/Gα_{i2}R179C tumor growth curves. **C-D**, a graph to show the number of lung metastases (**C**) and images representative of HE-stained lung metastases (indicated by arrows) (**D**) from PTEN^{+/-} and PTEN^{+/-}/Gα_{i2}R179C mice. Two tail unpaired Student's *t* test was used for statistical analysis in this Figure.

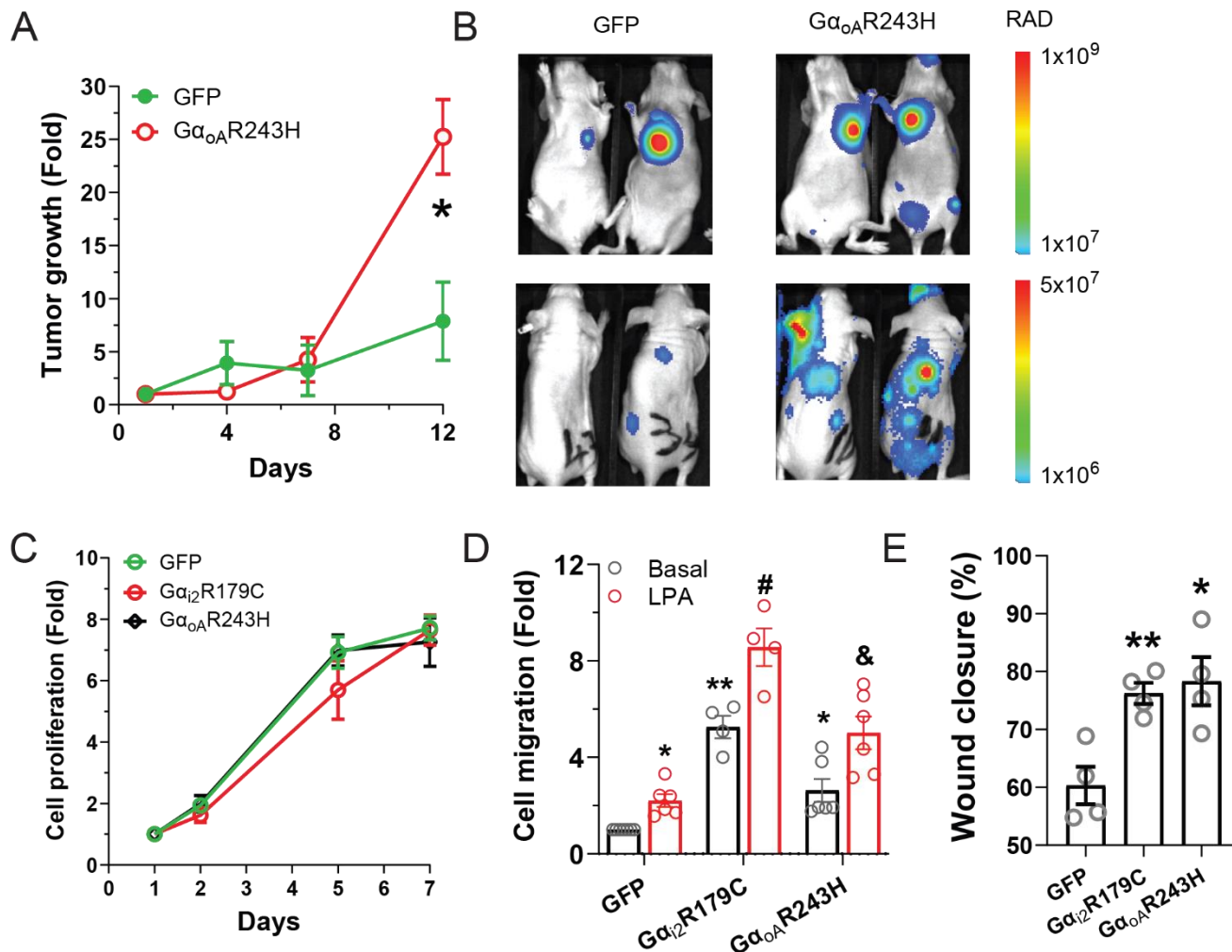


Figure 4

Figure 4. Effects of $G\alpha_{i2}R179C$ and $G\alpha_{oA}R243H$ on Neu cell growth and migration and tumor metastasis. **A-B**, the tumor growth curves of Neu cells expressing GFP or $G\alpha_{oA}R243H$ injected into nude mice via left ventricle (**A**) and representative bioluminescence images of mice at the dorsal (top) and ventral (bottom) positions (**B**). **C**, the *in vitro* growth curves of Neu cells expressing GFP, $G\alpha_{i2}R179C$, or $G\alpha_{oA}R243H$. **D**, basal and LPA (10 μ M)-stimulated transwell migration of Neu cells expressing GFP, $G\alpha_{i2}R179C$, or $G\alpha_{oA}R243H$. **E**, wound healing assay data of Neu cells expressing GFP, $G\alpha_{i2}R179C$, or $G\alpha_{oA}R243H$. *, ** $p < 0.05$ and 0.01 vs GFP basal; #, & $p < 0.05$ and 0.01 vs GFP LPA. Two-way and one-way ANOVA were used for statistical analysis in Figure D and E, respectively.

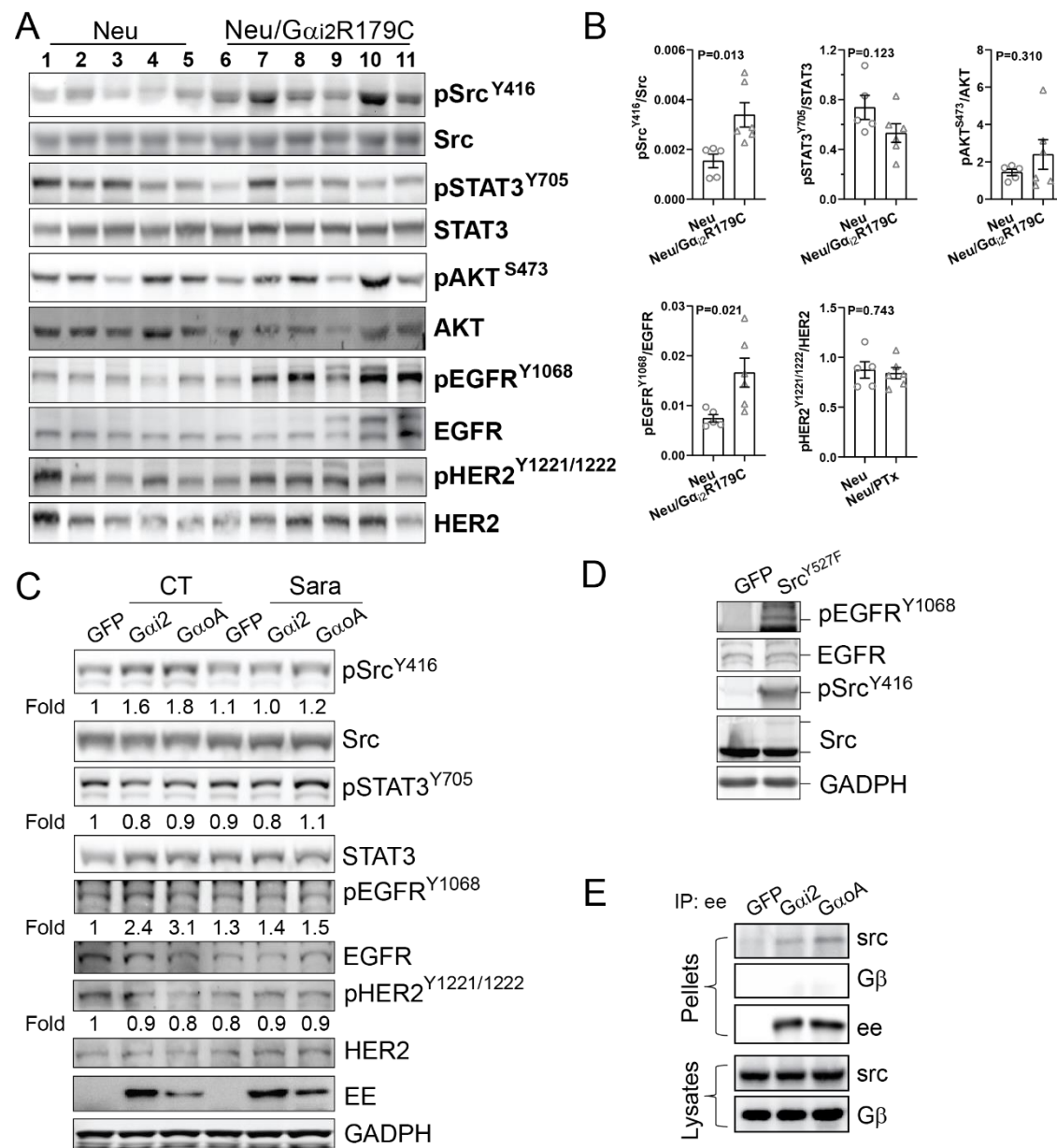


Figure 5

Figure 5. Gα_{i2}R179C and Gα_{0A}R243H activate c-Src and induce EGFR transactivation in tumor cells. A-B, Western blotting (A) showing increased phosphorylation of c-Src^{Y416} and EGFR^{Y1068} and no change in phosphorylation of AKT^{S473}, STAT3^{Y705} and HER2^{Y1221/1222} in Neu/Gα_{i2}R179C tumors as compared to Neu tumors. Each lane represents a sample from an individual tumor. The Western blotting data from A were quantified and expressed as the ratio of the phosphorylated to total proteins (B). Two tail unpaired Student's *t* test was used for statistical analysis of the data in B, and *p* values are shown. C-D, Western blotting showing phosphorylation of the indicated proteins in Neu cells expressing GFP, Gα_{i2}R179C (Gα_{i2}) or Gα_{0A}R243H (Gα_{0A}) and treated with vehicle control (CT) or saracatinib (sara; 2 μM) (C) and Neu cells overexpressing GFP or the constitutively active c-Src^{Y527F} mutant (D). E, co-immunoprecipitation of ee-tagged Gα_{i2}R179C and Gα_{0A}R243H with c-Src in Neu cells. Neu cells expressing GFP, Gα_{i2}R179C (Gα_{i2}) or Gα_{0A}R243H (Gα_{0A}) were immunoprecipitated with the anti-ee antibody and probed with the indicated antibodies.

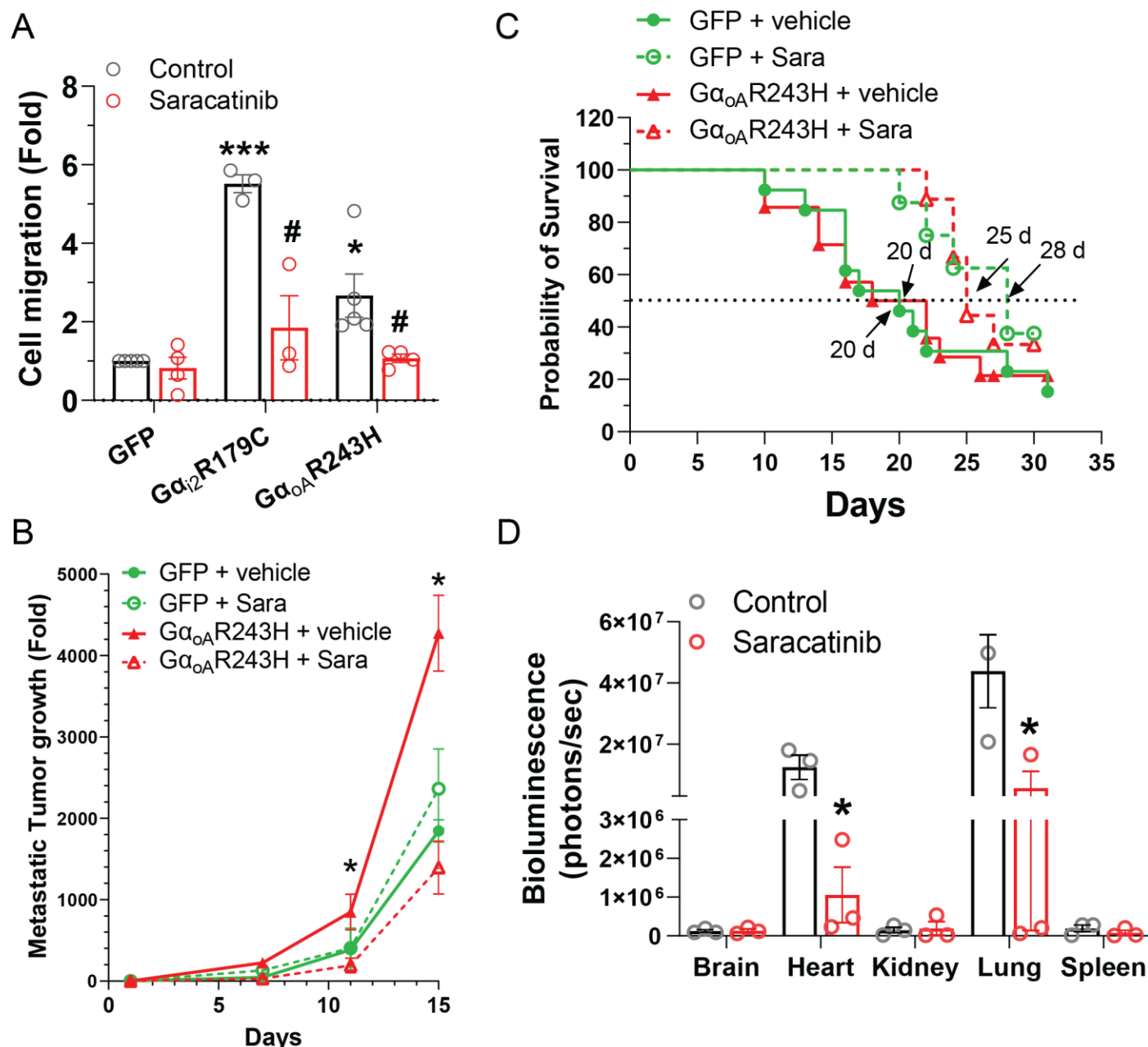


Figure 6

Figure 6. The effects of saracatinib on Neu cell migration and tumor metastasis. **A**, transwell migration of Neu cells expressing GFP, $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ and treated with vehicle (control) or saracatinib (2 μ M). *, *** $p < 0.05$ and 0.001 vs GFP control; # $p < 0.05$ vs $G\alpha_{i2}R179C$ or $G\alpha_{oA}R243H$ control. **B-C**, the growth curves (**B**) and survival curves (**C**) of mice harvesting GFP and $G\alpha_{oA}R243H$ metastatic tumors and treated with vehicle or saracatinib (25mg/kg initially for 7 days and then 50mg/kg for the rest of treatment, gavage daily; sara). Two-way ANOVA was used for statistical analysis in Figure B and C ($n = 9-13$). Log-rank test was used for statistical analysis in Figure D ($n = 9-13$). **D**, a graph showing the bioluminescence intensity of the indicated tissues measured via *ex vivo* bioluminescence imaging from the remaining mice harvesting GFP metastatic tumors and treated with vehicle (control) or saracatinib for two weeks. * $p < 0.05$ vs vehicle.