SCIENTIFIC REPORTS

OPEN

The role of miR-26a and miR-30b in HER2+ breast cancer trastuzumab resistance and regulation of the *CCNE2* gene

Received: 12 August 2016
Accepted: 19 December 2016
Published: 25 January 2017

Eduardo Tormo^{1,*}, Anna Adam-Artigues^{1,*}, Sandra Ballester¹, Begoña Pineda¹, Sandra Zazo², Paula González-Alonso², Joan Albanell^{3,4,5}, Ana Rovira^{3,4}, Federico Rojo², Ana Lluch^{1,6} & Pilar Eroles¹

A subset of HER2+ breast cancer patients manifest clinical resistance to trastuzumab. Recently, miR-26a and miR-30b have been identified as trastuzumab response regulators, and their target gene *CCNE2* seems to play an important role in resistance to trastuzumab therapy. Cell viability was evaluated in trastuzumab treated HER2+ BT474 wt (sensitive), BT474r (acquired resistance), HCC1954 (innate resistance), and MDA-MB-231 (HER2−) cell lines, and the expression of miR-26a, miR-30b, and their target genes was measured. BT474 wt cell viability decreased by 60% and miR-26a and miR-30b were significantly overexpressed (~3-fold, $p = 0.003$ and $p = 0.002$, respectively) after trastuzumab treatment, but no differences were observed in resistant and control cell lines. Overexpression of miR-30b sensitized BT474r cells to trastuzumab ($p = 0.01$) and *CCNE2*, was significantly overexpressed after trastuzumab treatment in BT474r cells ($p = 0.032$), but no significant changes were observed in sensitive cell line. When *CCNE2* was silenced BT474r cell sensitivity to trastuzumab increased ($p = 0.03$). Thus, the molecular mechanism of trastuzumab action in BT474 cell line may be regulated by miR-26a and miR-30b and *CCNE2* overexpression might play an important role in acquired trastuzumab resistance in HER2+ breast cancer given that resistance was diminished when *CCNE2* was silenced.

Breast cancer is a complex and heterogeneous disease that accounts for 25% of all cancer cases and 15% of all female deaths from cancer worldwide¹. Approximately 20–30% of breast tumors overexpress HER2² and this subtype is characterized by high rates of cell proliferation and metastasis, a poor prognosis, low overall survival, and variable chemotherapy response³. However, the HER2 receptor is a target for personalized and directed therapy with monoclonal antibodies such as trastuzumab (Herceptin). This therapy blocks the function of the HER2 receptor (and related pathways) which promote migration, angiogenesis, and cell growth and survival, and thus prolongs patient survival and decreases the rate of tumor development⁴. Despite this, about 40% of metastatic patients present innate trastuzumab-resistance, and most patients develop acquired resistance within the first year of treatment^{4,5}. Therefore, better knowledge of the molecular mechanisms of trastuzumab action is crucial for improving HER2+ breast cancer treatment in an attempt to overcome these resistance problems.

It has been previously described that trastuzumab blocks HER2 signaling by preventing HER2 homo/heterodimerization, thus inducing antibody-dependent cell-mediated cytotoxicity, promoting HER2 receptor internalization, and inhibiting its cleavage, which in turn blocks the MAPK, mTOR, and PI3K/Akt pathways⁶. However, there are no validated markers for resistance to HER2-targeted trastuzumab treatment. Little is known about the role of miRNAs in trastuzumab response, although some studies have identified miRNAs involved in carcinogenesis, malignancy, prognosis, and treatment response in different cancers⁷. For example, miR-34a and miR-221/222 are involved in the response to docetaxel and tamoxifen in breast cancer⁸, and miR-10b and miR-21

¹INCLIVA Biomedical Research Institute, 46010 Valencia, Spain. ²Pathology Department, IIS-Fundación Jiménez Díaz, 28040 Madrid, Spain. ³Cancer Research Program, IMIM (Hospital del Mar Research Institute), 08003 Barcelona, Spain. ⁴Medical Oncology Department, Hospital del Mar, 08003 Barcelona, Spain. ⁵Pompeu Fabra University, 08002 Barcelona, Spain. ⁶Oncology and Hematology Department, Hospital Clínico Universitario, 46010 Valencia, Spain.

*These authors contributed equally to this work. Correspondence and requests for materials should be addressed to P.E. (email: pilar.eroles@uv.es)

overexpression are associated with poor outcomes in breast cancer. Ichikawa and colleagues demonstrated that miR-26a and miR-30b are implicated in trastuzumab response by identifying them as trastuzumab-inducible miRNAs that are overexpressed in HER2 positive cell lines treated with trastuzumab⁹. However, the exact role of this overexpression in sensitivity or resistance to trastuzumab is still unexplained.

Following on from this work, our group focused on how miR-26a and miR-30b are implicated in this response to trastuzumab by examining the differences in miRNA expression in trastuzumab-sensitive, acquired-resistance, or innate-resistance HER2+ breast cancer cell lines. In addition, we also screened genes which may be targets of these miRNAs and which could also be involved in treatment response. We checked the regulation of *CCNE2* (Cyclin E2) by these miRNAs and its expression in different HER2+ breast cancer cell lines because this gene has been previously implicated in acquired trastuzumab resistance¹⁰ and is a possible miR-30b target⁹. We also describe possible miR-30b and miR-26a target genes related to the cell cycle and apoptosis that are likely implicated in trastuzumab response, and thus go on to suggest potential biomarkers which may be able to identify trastuzumab resistance in early treatment phases.

Materials and Methods

Cell lines and treatment. The HER2-positive breast cancer cell lines BT474 wt, BT474r, and HCC1954, were obtained from F.R.'s group at the *Fundación Jiménez Díaz* Hospital (Madrid) and the HER2-negative MDA-MB-231 breast cancer cell line was obtained from J.A.'s group at the IMIM, Hospital del Mar Research Institute (Barcelona). All the cell lines were grown at 37 °C with 5% CO₂. The MDA-MB-231, BT474 wt, and BT474r cell lines with acquired resistance, generated by culture with 15 µg/ml trastuzumab for 6–12 months¹¹, were grown in Dulbecco Modified Eagle medium nutrient mixture F-12 (DMEM/F12) with 2.5 mM L-Glutamine and 15 mM HEPES (Gibco), supplemented with 10% FBS and 1% penicillin-streptomycin. HCC1954 was cultured in RPMI 1640 medium (Gibco) containing 10% FBS, 1% penicillin-streptomycin, and 1% L-glutamine. Trastuzumab (Herceptin®, Roche) was used at 15 µg/ml.

RNA isolation. Total RNA containing small RNAs was isolated using mirVana miRNA isolation kit (Ambion, Austin, Texas) according to the manufacturer's protocol. The concentration and purity of the RNA obtained was measured as the OD260/280 ratio using a GeneQuant pro spectrophotometer (GE, Healthcare).

Transfection. Cell lines were transfected either with 50 nM hsa-miR-26a-5p or hsa-miR-30b-5p mirVana mimic or inhibitor miRNAs (Ambion), and, where relevant, with 50 nM or 100 nM *CCNE2* siRNA (#AM16708, Thermofisher), using the TransIT-X2 Dynamic Delivery System reagent (Mirus), following the manufacturer's instructions.

Cell viability. BT474 wt, BT474r, HCC1954, and MDA-MB-231 cells were cultured in 96-well plates and treated with 15 µg/ml trastuzumab for 7 days with or without prior transfection with miRNA mimics/inhibitors or siRNA. Cell viability was measured using a MTT-based Cell Growth Determination Kit (#GDC1; Sigma). MTT was added to each well under sterile conditions (to a final concentration of 10% of the total volume) and the plates were incubated for 4 h at 37 °C. Formazan crystals were dissolved in the solubilization solution at a 1:1 ratio and, starting from yellow MTT, formed purple crystals via succinate dehydrogenase only in viable cells. Absorbance was measured at 570 nm in a microplate reader with background correction at 690 nm. The significance of any differences were assessed using the Student t test.

Sensitivity or resistance to trastuzumab was assessed using the following equations: $\text{Log}(N_t) = (\text{Log}2/\text{DT}) \times T + \text{Log}(N_0)$ and fold change in growth rate = $\text{DT with drug} / \text{DT without drug}$, where DT = doubling time, N_t = number of cells after treatment, T = time of treatment, and N_0 = number of cells before treatment. Cell lines with more than a 20% decrease in cell number in response to trastuzumab treatment were considered to be trastuzumab-sensitive¹².

Flow cytometry. BT474 wt and BT474r cell lines were cultured in 6-well plates and treated with 15 µg/ml trastuzumab for 7 days with or without prior transfection with hsa-miR-30b-5p mirVana miRNA mimic. The cells were labelled with 0.5 mg/ml annexin-V conjugated to the FITC fluorochrome and 0.5 mg/ml of propidium iodide (PI) using the FITC Annexin V Apoptosis Detection Kit I (BD Pharmingen™). The cells were incubated for 15 minutes in the dark at room temperature, followed by the addition of 400 µl of binding buffer. Annexin V binds apoptotic cells because of phosphatidylserine membrane residue translocation, and PI is incorporated because of loss of plasma membrane integrity. The amount of PI and annexin V labelling was analyzed by flow cytometry using a FACSVerse™ flow cytometer (BD) to determine cell viability by measuring the percentage of apoptotic cells present.

For cell cycle analysis, aliquots of 2×10^6 cells were kept on ice prior to fixation. The cells were washed twice with cold PBS, decanted, and the pellet was resuspended before the addition of ethanol. The cells were then fixed by slowly pipetting 1 ml of 80% ice-cold ethanol into the cell suspension while vortexing; the cells were left in ethanol at −20 °C for 2 hours and were then washed twice with cold PBS. Finally they were resuspended in 1 ml freshly-made DAPI/TX-100 solution and incubated for 30 minutes at room temperature before analysis by flow cytometry (excitation at 405 nm and emission at 450 nm).

Gene and miRNA expression by qRT-PCR. The expression of *CCNE2* (cyclin E2), *CASP3* (caspase 3), and *APAF1* (apoptotic peptidase activating factor 1) mRNA and levels of miR-26a and miR-30b were determined by real-time qPCR. First, 1000 ng RNA was retro-transcribed to cDNA using a High-Capacity cDNA Reverse Transcription kit (Applied Biosystems) and a TaqMan® MiRNA Reverse Transcription kit; cDNA was then synthesized at 25 °C for 10 min and 37 °C for 2 hours for mRNA retro-transcription and 16 °C for 30 min, 42 °C for 30 min, and 85 °C for 5 min for miRNA retro-transcription, and was then stored at −20 °C until use.

Figure 1. Cell viability after trastuzumab treatment. BT474 wt, BT474r, HCC1954, and MDA-MB-231 cells were exposed to 15 μ g/ml of trastuzumab for 7 days and their viability was assessed using MTT assays. The percentage of viable cells was calculated with respect non-treated cells. * $p < 0.05$.

TaqMan[®] primers for housekeeping miRNA *RNU43* as well as miR-26a and miR-30b were obtained from Applied Biosystems. Transcript levels were detected using a 9700HT Fast Real-Time PCR system (Applied Biosystems) and PCR reactions were performed with a TaqMan[®] Universal Master Mix (Applied Biosystems) and TaqMan[®] 20 $\times$ assay following the manufacturer's protocol. PCR conditions were: 50 $^{\circ}$ C for 2 min, 95 $^{\circ}$ C for 10 min, 40 cycles of 95 $^{\circ}$ C for 15 sec, and 60 $^{\circ}$ C for 1 min for gene expression, or 95 $^{\circ}$ C for 10 min, followed by 40 cycles of 95 $^{\circ}$ C for 15 sec, and 60 $^{\circ}$ C for 1 min for miRNA expression.

Results were normalized according to the expression of housekeeping *GAPDH* mRNA or *RNU43* miRNA. The threshold cycle value (C_T) was determined for each measurement and mRNA or miRNA expression was calculated relative to the control using the comparative critical threshold ($2^{-\Delta\Delta C_T}$) method where: ΔC_T = target C_T minus the housekeeping control C_T , $\Delta\Delta C_T$ = sample ΔC_T minus the baseline (non-treated cells) ΔC_T , and the relative expression level = $2^{-\Delta\Delta C_T}$. Each experiment was performed in technical and biological triplicate.

Western immunoblotting. Cells were seeded and then exposed to the indicated concentration of trastuzumab. After treatment and/or transfection experiments, cells were washed twice with cold PBS, and the monolayer was scraped into 1 ml of Pierce RIPA buffer (Thermo #89900). The lysates were transferred to a clean microfuge tube, placed on ice for 15 min, and centrifuged for 10 min at 14,000 rpm. The supernatant was transferred to a clean microfuge tube, and the protein concentration was determined. Protein extracts (40 μ g) were boiled in Laemmli buffer and resolved on a 10% SDS-polyacrylamide gel, before transfer onto a nitrocellulose membrane. Membranes were blocked in 5% BSA for 1 hour and then incubated with antibodies to APAF1 (Abcam, #32372), CCNE2 (Abcam, #40890), and CASP3 (Abcam, #32351) overnight at 4 $^{\circ}$ C. The membranes were subsequently washed and then incubated for 1 hour with an anti-rabbit IgG horseradish peroxidase-linked secondary antibody (Cell signaling, #7074). The membranes were then washed and briefly incubated using with an Amersham ECL Western Blotting detection reagent (GE Healthcare, #RPN2209).

Statistical analysis. The sample and control groups were compared using a two-tailed Student t-test. All data presented includes the standard deviation (SD). P -values of less than 0.05 were considered to be statistically significant.

Results

The effect of trastuzumab on cell viability. First we examined the effect of trastuzumab on cell viability in our HER2-positive (BT474 wt, BT474r, and HCC1954) and HER2-negative (MDA-MB-231) breast cancer cell lines. After seven days of trastuzumab treatment we observed a 40% reduction in BT474 wt proliferation ($p = 0.01$) but no changes in viability in the BT474r, HCC1954, or MDA-MB-231 cells (Fig. 1). Annexin-V assays showed that BT474 wt cell viability was reduced from 97.7% to 22.6% after trastuzumab treatment while BT474r cells were not affected by the same treatment.

Both miR-30b and miR-26a are trastuzumab-responsive miRNAs. Similar to previous studies, the three HER2-positive and one HER2-negative breast cancer cell lines were exposed to trastuzumab⁹ and then miR-30b and miR-26a expression were analyzed in these cells by real-time qPCR analysis. As shown in Fig. 2, the fold-expression of miR-30b and miR-26a (relative to endogenous *RNU43*) increased after trastuzumab treatment in BT474 wt cells (~ 3 fold, $p = 0.002$ and $p = 0.003$ for miR-30b and miR-26a, respectively), while no significant changes in expression were observed in the other cell lines, which were all resistant to trastuzumab treatment.

MiR-30b and miR-26a overexpression sensitize cells to trastuzumab treatment. Transfection with miR-30b or miR-26a mimics resulted in a decrease in BT474 wt cell viability ($\sim 20\%$ reduction, $p = 0.036$ and $p = 0.022$ for miR-30b and miR-26a, respectively) in normal growth conditions, but no change in the other cell lines. However, when trastuzumab was added alongside these miRNA mimics, viability considerably reduced in both BT474 wt ($\sim 60\%$ reduction, $p = 0.0012$ and $p = 0.004$ for miR-30b and miR-26a, respectively) and BT474r ($\sim 40\%$ reduction, $p = 0.01$ and $p = 0.009$ for miR-30b and miR-26a, respectively) cells (Fig. 3). No changes were

Figure 2. Expression levels of miR-30b and miR-26a. The levels of miR-30b and miR-26a were measured with and without trastuzumab treatment in BT474 wt (trastuzumab-sensitive) and BT474r (trastuzumab-resistant) cells. Fold-expression was calculated in relation to endogenous RNU43. ** $p < 0.01$.

Figure 3. The effect of miR-30b and miR-26a overexpression or downregulation on cell viability. BT474 wt, BT474r, and HCC1954 cell lines were transfected with miR-30b and miR-26a mimetics or inhibitors and the viability of the cells was assayed after 7 days after trastuzumab treatment and under control conditions. The experiments were performed in triplicate and the values are calculated in relation to the negative control transfections with scramble miRNAs. * $p < 0.05$, ** $p < 0.01$.

observed in HCC1954 and MDA-MB-231 (data not shown). In contrast, transfection with miR-30b or miR-26a inhibitors did not affect the viability of any of the cell lines either in normal conditions or with trastuzumab treatment.

The flow cytometry viability assays showed that the percentage of viable BT474 wt cells reduced from 97.7% to 4.11% when miR-30b mimic was transfected and to 2.66% when this mimic was combined with trastuzumab treatment. MiR-30b overexpression in BT474r reduced cell viability from 97.6% to 44.8%, however this result did not differ between cohorts transfected with miR-30b mimic alone or combined with trastuzumab.

The effect of trastuzumab on the cell cycle. Analysis of the cell cycle phases by flow cytometry (Fig. 4) showed that trastuzumab induces G1 arrest (4.7% increase compared to control) in BT474 wt cells. Transfection with miR-30b mimic induced stronger G1 arrest in BT474 wt cells (6.8% increase), which also increased when miR-30b overexpression was combined with trastuzumab treatment (6.7% increase). Proliferative (S + G2) BT474 wt cells decreased from 31.3% to 29% when trastuzumab was added further decreased to 26.87% when miR-30b mimic was also transfected, and reduced to 22.08% when combined with trastuzumab. Neither trastuzumab treatment nor miR-30b mimic transfection caused G1 cell cycle arrest in BT474r cells and BT474r

Figure 4. Distribution of the cell cycle phases after trastuzumab treatment. The percentage of cells in G1, S, and G2 phases was evaluated in BT474 wt and BT474r cells, with or without miR-30b overexpression, and with or without trastuzumab treatment.

Figure 5. The effect of trastuzumab on CASP3, CCNE2, and APAF1 expression. The levels of CASP3, CCNE2, and APAF1 genes with or without trastuzumab treatment were analyzed in BT474 wt and BT474r cells. The experiments were performed in triplicate and the values were related to endogenous GAPDH expression. * $p < 0.05$, ** $p < 0.01$.

proliferative cells remained the same following trastuzumab treatment, but the addition of miR-30b mimic decreased the number of BT474r proliferative (S + G2) cells by 5.13% both with and without the presence of trastuzumab.

The impact of miR-30b and miR-26a on cell cycle genes. Because both miR-30b and miR-26a seemed to be implicated in regulating cell viability, we checked how these miRNAs modulate the cell cycle and cell proliferation. First we measured the expression of the cell cycle regulator gene *CCNE2* and the two pro-apoptotic genes *APAF1* and *CASP3*, in normal conditions and after trastuzumab treatment. *CASP3* and *APAF1* showed increased expression after this treatment in BT474 wt cells (~3 fold, $p = 0.0012$ and $p = 0.0017$, respectively) but no significant differences were observed in *CCNE2* expression. In contrast, only *CCNE2* expression significantly increased ($p = 0.032$) after trastuzumab treatment in BT474r cells. Basal expression of *CASP3* and *CCNE2* was lower in resistant BT474r cells compared to wild-type BT474 wt cells ($p = 0.018$ and $p = 0.029$, respectively), whereas basal *APAF1* expression was similar in both cell lines (Fig. 5). HCC1954 cells did not show changes in *APAF*, *CASP3* or *CCNE2* expression (data not shown).

Western blot analysis showed that APAF1 and CASP3 proteins were highly expressed in BT474 wt cells after trastuzumab treatment, whereas CCNE2 expression did not change. In contrast, the opposite was seen in BT474r cells: APAF1 and CASP3 did not change but CCNE2 was strongly expressed. There were no changes in protein expression after treatment in the HCC1954 cell line (Fig. 6A).

We also checked gene expression levels in these cell lines after miR-30b or miR-26a mimic or inhibitor transfection (Fig. 7). In normal conditions, transfection with either miR-30b or miR-26a mimic resulted in a significant increase in the expression of the pro-apoptotic *CASP3* and *APAF1* genes in BT474 wt (*CASP3*: $p = 0.026$ and $p = 0.032$, *APAF1*: $p = 0.035$ and $p = 0.042$, for miR30b and miR-26a respectively) and BT474r cells (*CASP3*: $p = 0.034$ and $p = 0.019$, *APAF1*: $p = 0.041$ and $p = 0.034$, for miR30b and miR-26a respectively), but not in HCC1954 cells. In addition, *CCNE2* expression was also significantly downregulated in BT474 wt and BT474r cell lines after miR-30b or miR-26a mimic transfection (BT474 wt: $p = 0.042$ and $p = 0.035$, BT474r: $p = 0.032$ and $p = 0.041$, for miR30b and miR-26a respectively), but not in HCC1954. In contrast, these results were reversed when the cells were transfected with miR-30b or miR-26a inhibitors: the *CASP3* and *APAF1* genes were downregulated while *CCNE2* was upregulated (BT474 wt: $p = 0.002$ and $p = 0.021$, for miR-30b and miR-26a respectively; BT474r: $p = 0.034$ for miR-30b and not significant changes for miR-26a). In the same way, transfection

Figure 6. The effect of trastuzumab on CASP3, CCNE2, and APAF1 protein expression by western blot analysis. (A) Levels of APAF1, CCNE2, and CASP3 proteins were measured in BT474 wt, BT474r, and HCC1954 cells exposed to trastuzumab (15 µg/ml for 7 days). Actin was used as loading control. (B) Levels of CCNE2 protein in BT474 wt, BT474r, and HCC1954 cells transfected with miR-30b mimetic.

with miR-30b mimic produced a decrease in CCNE2 protein expression in both BT474 wt and BT474r cell lines compared to controls (Fig. 6B).

When trastuzumab was combined with the miR-30b or miR-26a mimics in BT474 wt and BT474r cells, CASP3 and APAF1 expression significantly increased and CCNE2 expression significantly decreased in comparison to control cells reaching similar levels of the mimics without trastuzumab (Fig. 7). However, this expression pattern appeared to be attenuated both in sensitive (BT474 wt) and resistant (BT474r) cell lines when trastuzumab treatment was combined with miR-30b or miR-26a inhibitors.

CCNE2 is involved in cell viability. CCNE2 silencing by siRNA (100 nM) transfection led to a decrease in cell proliferation in all of the breast cancer cell lines to ~40% viability ($p = 0.001$) in BT474 wt, ~25% viability ($p = 0.05$) in BT474r, ~40% viability ($p = 0.0005$) in HCC1954, and ~20% viability ($p = 0.003$) in MDA-MB-231 cells compared to their respective controls (Fig. 8).

CCNE2 expression is involved in acquired trastuzumab resistance. We compared cell viability in trastuzumab-treated and non-treated cell lines transfected with siRNA using MTT assays. As shown in Fig. 8, BT474 wt cells were initially sensitive to trastuzumab (~50% viability vs. control, $p = 0.05$) even when also transfected with 50 nM siRNA (~50% viability), but cell viability reduced to 25% when 100 nM siRNA was transfected in combination with trastuzumab ($p = 0.02$ vs. non-treated sample). BT474r was initially resistant to trastuzumab (~100% viability) but viability reduced to ~25% ($p = 0.03$) and ~10%, ($p = 0.01$) respectively, after transfection with 50 nM or 100 nM siRNA, respectively in combination with trastuzumab, showing an overall 50% decrease in viability ($p = 0.03$ vs. non-treated samples). Thus, according to the criteria set out in the Materials and Methods, this cell line was considered to be trastuzumab-sensitive after transfection with 50 nM or 100 nM CCNE2 siRNA. HCC1954 and MDA-MB-231 cells were resistant to trastuzumab and remained so after CCNE2 silencing (~40% and ~25% final viability, respectively), representing a non-significant change compared to non-treated samples in both cases.

Discussion

HER2+ breast tumor treatment has been greatly improved by anti-HER2 agents like trastuzumab; however, primary and secondary resistance is a substantial problem¹³. To investigate the possible mechanisms behind this resistance we must strive to understand the pathways and mechanisms modified by trastuzumab treatment. Altered miRNA profiles have been identified in several cancers in relation to malignancy and prognosis, and in metastatic breast cancer tumors miR-21 and miR-10b have been shown to be upregulated¹⁴. Some miRNAs, such as miR-34a or miR-221, are related to treatment response and are involved in docetaxel and tamoxifen response^{15,16}, while miR-30b and miR-26a are upregulated in HER2+ BT474 wt cells when treated with trastuzumab⁹. Following this line of investigation, we demonstrated that this upregulation occurs in BT474 wt cells but not in acquired-resistance BT474r cells generated by our group or in the innately-resistant HCC1954 cell line. Thus, upregulation of these miRNAs may be a marker for sensitivity to trastuzumab treatment.

In addition to validating the miR-26a and miR-30b-mediated reduction in cell viability in BT474 wt cells first shown by Ichikawa *et al.*, we also investigated the effect of miR-26a and miR-30b mimics and inhibitors, either alone or combined with trastuzumab, in cell lines that are either sensitive or resistant to this drug. Our results showed that while miR-26a and miR-30b mimics alone only had an effect in BT474 wt cells, when combined with trastuzumab they decreased cell proliferation in both the BT474 wt and BT474r cell lines, thus suggesting that these miRNAs are related to the acquisition of trastuzumab resistance. In addition, cell viability and cell cycle analyses confirmed previous results which showed that trastuzumab¹⁷ and miR-30b mimics⁹ decrease BT474 wt cell viability and produce G1 arrest, while trastuzumab does not affect the cell cycle in BT474r cells¹⁰. We also revealed that miR-30b mimics, alone or in combination with trastuzumab, decrease the number of proliferative BT474r cells.

Given the aforementioned effects of miR-30b and miR-26a (which are upregulated in response to trastuzumab) we tried to identify some of their cell cycle or apoptosis-related target genes. Previous studies linked the response to EGFR tyrosine kinase treatment to pro-apoptotic genes such as caspases and APAF1 in breast¹⁸

Figure 7. Regulation of CASP3, CCNE2, and APAF1 expression by miR-30b and miR-26a. The BT474 wt, BT474r, and HCC1954 cell lines were transfected with miR-30b and miR-26a mimetics or inhibitors and the expression levels of CASP3, CCNE2, and APAF1 were evaluated by quantitative PCR after trastuzumab treatment or under control conditions. The experiments were performed in triplicate and the values were normalized to endogenous GAPDH gene expression. * $p < 0.05$, ** $p < 0.01$.

Figure 8. CCNE2 silencing sensitizes BT474r cells to trastuzumab treatment. CCNE2 gene was silenced using siRNA in BT474 wt (A), BT474r (B), HCC1954 (C), and negative control MDA-MB-231 (D), and cell viability was assessed by MTT with and without trastuzumab treatment (15 μ g/ml for 7 days) * $p < 0.05$, ** $p < 0.01$.

and other cancers¹⁹. Thus, we studied the effect of trastuzumab on pro-apoptotic genes and showed that CASP3 and APAF1 were upregulated at the gene and protein level in BT474 wt cells when treated with trastuzumab, but that the same effect was not observed in trastuzumab-resistant cell lines. This finding suggests that the apoptotic processes induced by trastuzumab may be mediated in part by the APAF1 apoptosis pathway and that cell lines which do not overexpress this gene after trastuzumab treatment have higher proliferation and survival rates¹⁹.

APAF1, activated via CARD (caspase recruitment domain), initiates the activation of caspases such as caspase-3^{20,21}. Given that trastuzumab induces miR-30b and miR-26a upregulation, as well as CASP3 and APAF1 upregulation in BT474 wt cells, we tried to identify any relationships between these miRNAs and genes.

Figure 9. Schematic representation of trastuzumab response mechanism proposed for BT474 wt (sensitive) and BT474r (resistant) cell lines.

Transfection experiments demonstrated that upregulation of miR-30b and miR-26a with mimics induced *CASP3* and *APAF1* overexpression in BT474 wt and BT474r cells, while inhibitors of these miRNAs decreased *CASP3* and *APAF1* expression independently of the presence of trastuzumab. Thus, these results suggest that apoptotic *CASP3* and *APAF1* genes may be indirectly regulated by miR-30b and miR-26a through a previously undescribed intermediate mechanism (Fig. 9).

It is known that PI3K/Akt activation is essential for HER2-mediated apoptosis suppression^{22,23}; activation of the PI3K/Akt pathway by different mechanisms has been described both in HER2+ breast cancer patients with trastuzumab resistance²⁴ and in BT474r cells¹². Indeed, trastuzumab treatment inhibits the PI3K/Akt pathway by Akt dephosphorylation²⁵. Given that activated Akt pathways inhibit *APAF1* expression and its binding with *CASP9* (which in turn regulates *CASP3* expression), it seems logical that inactivation of the Akt pathway by trastuzumab in BT474 wt cells results in increased *APAF1*, *CASP3*, and *CASP9* expression which does not occur in resistant cell lines with active Akt²⁶. Indeed, we confirmed that *CASP9* is also significantly expressed in BT474 wt cells, but not in BT474r cells, after trastuzumab treatment (Supplementary Figure 1).

Previous studies have proposed *CCNE2* as a miR-30b target gene⁹, and have shown the role of *CCNE2* in trastuzumab-acquired resistance¹⁰. Here, we not only demonstrated that *CCNE2* is a miR-30b target gene (because both the gene and its protein were underexpressed when BT474 wt and BT474r cells were transfected with miR-30b mimic and increased when they were transfected with its inhibitor), but also that *CCNE2* is a miR-26a target. In addition, *CCNE2* expression is related to trastuzumab response¹⁰: while *CCNE2* expression reduced in BT474 wt cell lines treated with trastuzumab, it increased in BT474r cells, indicating that *CCNE2* overexpression could be a mechanism of trastuzumab resistance. However, given that the HCC1954 cell line with innate trastuzumab resistance did not overexpress *CCNE2*, we propose that it is a mechanism of acquired-resistance only.

Our findings agree with those by Scaltriti and colleagues (2012), who transfected *CCNE2* siRNA into the trastuzumab-resistant BT474 cell line generated in-house, and showed that resistance to this drug depends on *CCNE2* overexpression in these cells. We confirmed these results in our resistant cell line, BT474r, and also demonstrated that once *CCNE2* was silenced, the viability of these cells reduced by more than 20% which, according to O'Brien *et al.*, should be considered drug-sensitive¹². Although *CCNE2* could be part responsible for acquired resistance in BT474r cells, it seems that it is not related to the innate resistance displayed by HCC1954 cell lines because transfection with *CCNE2* siRNA decreased proliferation but did not affect trastuzumab response.

Given the relevance of the Akt pathway and *CASP3* and *APAF1* genes in trastuzumab response, and the implication of miR-26a and miR-30b in treatment sensitivity, it is possible that these apoptotic genes are trans-regulated by a direct miR-26a or miR-30b target gene. If activated Akt could be used as a trastuzumab-treatment response marker in HER2+ breast cancer^{26–28}, *CASP3* and *APAF1* overexpression may also represent possible treatment-sensitivity markers. Thus, it would be interesting to go deeper into the relationship between the PI3K/AKT pathway and the miRNAs and genes relevant in this context in order to validate this proposed mechanism. Similarly, evaluating genes and proteins which are altered in samples from trastuzumab-resistant breast cancer patients would also provide valuable information. Despite these limitations, we propose that *CCNE2* overexpression may represent a potential biomarker of acquired trastuzumab resistance in patients with HER2+ breast cancer. Finally, new therapies combining trastuzumab with *CCNE2* inhibitors might represent a new strategy for overcoming acquired resistance to this drug.

References

1. Torre, L. A. *et al.* Global cancer statistics, 2012. *CA Cancer J Clin* **65**, 87–108, doi: 10.3322/caac.21262 (2015).
2. Eroles, P., Bosch, A., Perez-Fidalgo, J. A. & Lluch, A. Molecular biology in breast cancer: intrinsic subtypes and signaling pathways. *Cancer Treat Rev* **38**, 698–707, doi: 10.1016/j.ctrv.2011.11.005 (2012).
3. Nguyen, P. L. *et al.* Breast cancer subtype approximated by estrogen receptor, progesterone receptor, and HER-2 is associated with local and distant recurrence after breast-conserving therapy. *J Clin Oncol* **26**, 2373–2378, doi: 10.1200/JCO.2007.14.4287 (2008).
4. Marty, M. *et al.* Randomized phase II trial of the efficacy and safety of trastuzumab combined with docetaxel in patients with human epidermal growth factor receptor 2-positive metastatic breast cancer administered as first-line treatment: the M77001 study group. *J Clin Oncol* **23**, 4265–4274, doi: 10.1200/JCO.2005.04.173 (2005).
5. Brufsky, A. Trastuzumab-based therapy for patients with HER2-positive breast cancer: from early scientific development to foundation of care. *Am J Clin Oncol* **33**, 186–195, doi: 10.1097/COC.0b013e318191bfb0 (2010).
6. Molina, M. A. *et al.* Trastuzumab (herceptin), a humanized anti-Her2 receptor monoclonal antibody, inhibits basal and activated Her2 ectodomain cleavage in breast cancer cells. *Cancer Res* **61**, 4744–4749 (2001).

7. Asensio, P. E., Martin, E. T., Merlo, B. P., Armas, E. E. & Hernandez, A. L. MicroRNAs in Breast Cancer: One More Turn in Regulation. *Curr Drug Targets* **17**, 1083–1100 (2016).
8. Zhao, J. J. *et al.* MicroRNA-221/222 negatively regulates estrogen receptor alpha and is associated with tamoxifen resistance in breast cancer. *J Biol Chem* **283**, 31079–31086, doi: 10.1074/jbc.M806041200 (2008).
9. Ichikawa, T. *et al.* Trastuzumab produces therapeutic actions by upregulating miR-26a and miR-30b in breast cancer cells. *PLoS One* **7**, e31422, doi: 10.1371/journal.pone.0031422 (2012).
10. Scaltriti, M. *et al.* Cyclin E amplification/overexpression is a mechanism of trastuzumab resistance in HER2+ breast cancer patients. *Proc Natl Acad Sci USA* **108**, 3761–3766, doi: 10.1073/pnas.1014835108 (2011).
11. Zazo, S. *et al.* Generation, characterization, and maintenance of trastuzumab-resistant HER2⁺ breast cancer cell lines. *Am J Cancer Res.* **6**(11), 2661–2678, eCollection (2016).
12. O'Brien, N. A. *et al.* Activated phosphoinositide 3-kinase/AKT signaling confers resistance to trastuzumab but not lapatinib. *Mol Cancer Ther* **9**, 1489–1502, doi: 10.1158/1535-7163.MCT-09-1171 (2010).
13. Rexer, B. N. & Arteaga, C. L. Intrinsic and acquired resistance to HER2-targeted therapies in HER2 gene-amplified breast cancer: mechanisms and clinical implications. *Crit Rev Oncog* **17**, 1–16 (2012).
14. Sato, F. *et al.* MicroRNA profile predicts recurrence after resection in patients with hepatocellular carcinoma within the Milan Criteria. *PLoS One* **6**, e16435, doi: 10.1371/journal.pone.0016435 (2011).
15. Kastl, L., Brown, I. & Schofield, A. C. miRNA-34a is associated with docetaxel resistance in human breast cancer cells. *Breast Cancer Res Treat* **131**, 445–454, doi: 10.1007/s10549-011-1424-3 (2012).
16. Qian, B. *et al.* High miR-21 expression in breast cancer associated with poor disease-free survival in early stage disease and high TGF-beta1. *Breast Cancer Res Treat* **117**, 131–140, doi: 10.1007/s10549-008-0219-7 (2009).
17. Spector, N. L. & Blackwell, K. L. Understanding the mechanisms behind trastuzumab therapy for human epidermal growth factor receptor 2-positive breast cancer. *J Clin Oncol* **27**, 5838–5847, doi: 10.1200/JCO.2009.22.1507 (2009).
18. Way, T. D., Kao, M. C. & Lin, J. K. Degradation of HER2/neu by apigenin induces apoptosis through cytochrome c release and caspase-3 activation in HER2/neu-overexpressing breast cancer cells. *FEBS Lett* **579**, 145–152, doi: 10.1016/j.febslet.2004.11.061 (2005).
19. Deng, W. G., Kwon, J., Ekmekcioglu, S., Poindexter, N. J. & Grimm, E. A. IL-24 gene transfer sensitizes melanoma cells to erlotinib through modulation of the Apaf-1 and Akt signaling pathways. *Melanoma Res* **21**, 44–56, doi: 10.1097/CMR.0b013e3283382155 (2011).
20. Hofmann, K., Bucher, P. & Tschopp, J. The CARD domain: a new apoptotic signalling motif. *Trends Biochem Sci* **22**, 155–156 (1997).
21. Li, P. *et al.* Cytochrome c and dATP-dependent formation of Apaf-1/caspase-9 complex initiates an apoptotic protease cascade. *Cell* **91**, 479–489 (1997).
22. Albanell, J., Codony, J., Rovira, A., Mellado, B. & Gascon, P. Mechanism of action of anti-HER2 monoclonal antibodies: scientific update on trastuzumab and 2C4. *Adv Exp Med Biol* **532**, 253–268 (2003).
23. Dean-Colomb, W. & Esteva, F. J. Her2-positive breast cancer: herceptin and beyond. *Eur J Cancer* **44**, 2806–2812, doi: 10.1016/j.ejca.2008.09.013 (2008).
24. Jensen, J. D. *et al.* PIK3CA mutations, PTEN, and pHER2 expression and impact on outcome in HER2-positive early-stage breast cancer patients treated with adjuvant chemotherapy and trastuzumab. *Ann Oncol* **23**, 2034–2042, doi: 10.1093/annonc/mdr546 (2012).
25. Nagata, Y. *et al.* PTEN activation contributes to tumor inhibition by trastuzumab, and loss of PTEN predicts trastuzumab resistance in patients. *Cancer Cell* **6**, 117–127, doi: 10.1016/j.ccr.2004.06.022 (2004).
26. Chandarlapaty, S. *et al.* Frequent mutational activation of the PI3K-AKT pathway in trastuzumab-resistant breast cancer. *Clin Cancer Res* **18**, 6784–6791, doi: 10.1158/1078-0432.CCR-12-1785 (2012).
27. Berns, K. *et al.* A functional genetic approach identifies the PI3K pathway as a major determinant of trastuzumab resistance in breast cancer. *Cancer Cell* **12**, 395–402, doi: 10.1016/j.ccr.2007.08.030 (2007).
28. Gayle, S. S., Arnold, S. L., O'Regan, R. M. & Nahta, R. Pharmacologic inhibition of mTOR improves lapatinib sensitivity in HER2-overexpressing breast cancer cells with primary trastuzumab resistance. *Anticancer Agents Med Chem* **12**, 151–162 (2012).

Acknowledgements

The present work was supported by grants from the Spanish Ministry of Economy and Competitiveness (MINECO) and co-financed by FEDER funds (AES Program, grants PI12/00680, PI15/00146 and PIE15/00008 to J.A., PI12/01552 and PI15/00934 to F.R., PI12/01421, PI15/01617 and CB16/12/00481 to A.L.) and the Cancer Network (RD12/0036/0051 to J.A. and RD12/0036/0070 to A.L.). E.T. and A.A.-A. were supported by grants from (Institute of Health Carlos III, RETICS), RD12/0036/0070 and PI15/01617. B.P. is supported by the grant of Conselleria D'Educació, Investigació (Valencia) APOTIP/2016/A/008 and P.G.-A is supported by Fundación Conchita Rábago de Jiménez Díaz.

Author Contributions

Conception and design: A.L., P.E. Development of the methodology: E.T., A.A.-A., S.B., B.P., S.Z., P.G.-A. Acquisition of data: E.T., A.A.-A., S.B. Analysis and interpretation of data: E.T., A.A.-A., J.A., A.R., F.R., A.L., P.E. Writing, review and/or revision of the manuscript: A.L., P.E. Administrative, technical, or material support: S.Z., P.G.-A., B.P. Study supervision: J.A., A.R., F.R., A.L., P.E.

Additional Information

Supplementary information accompanies this paper at <http://www.nature.com/srep>

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Tormo, E. *et al.* The role of miR-26a and miR-30b in HER2+ breast cancer trastuzumab resistance and regulation of the CCNE2 gene. *Sci. Rep.* **7**, 41309; doi: 10.1038/srep41309 (2017).

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>