Activation of the CRABPII/RAR pathway by curcumin induces retinoic acid mediated apoptosis in retinoic acid resistant breast cancer cells

PADMAMALINI THULASIRAMAN, GALEN GARRIGA, VEENA DANTHULURI, DANIEL J. McANDREWS and IMRAN Q. MOHIUDDIN

Department of Biomedical Sciences, College of Allied Health, University of South Alabama, Mobile, AL 36688, USA

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Abstract. Due to the anti-proliferative and anti-apoptotic effects of retinoic acid (RA), this hormone has emerged as a target for several diseases, including cancer. However, development of retinoid resistance is a critical issue and efforts to understand the retinoid signaling pathway may identify useful biomarkers for future clinical trials. Apoptotic responses of RA are exhibited through the cellular RA-binding protein II (CRABPII)/retinoic acid receptor (RAR) signaling cascade. Delivery of RA to RAR by CRABPII enhances the transcriotional activity of genes involved in cell death and cell cycle arrest. The purpose of this study was to investigate the role of curcumin in sensitizing RA-resistant triple-negative breast cancer (TNBC) cells to RA-mediated apoptosis. We provide evidence that curcumin upregulates the expression of CRABPII, RARβ and RARγ in two different TNBC cell lines. Co-treatment of the cells with curcumin and RA results in increased apoptosis as demonstrated by elevated cleavage of poly(ADP-ribose) polymerase and cleaved caspase-9. Additionally, silencing CRABPII reverses curcumin sensitization of TNBC cells to the apoptotic inducing effects of RA. These findings provide mechanistic insights into sensitizing TNBC cells to RA-mediated cell death by curcumin-induced upregulation of the CRABPII/RAR pathway.

Introduction

With an estimated 1.7 million cases of breast cancer diagnosed worldwide in 2012, this type of cancer remains the most common in women. Among the patients with breast cancer, triple-negative breast cancer (TNBC) accounts for 10-20% of the invasive breast cancer which is defined as estrogen receptor (ER)-negative, progesterone receptor (PR)-negative and human epidermal growth factor receptor 2 (HER2)-negative (1). Due to the lack of markers, it carries poor prognosis and presents an emerging need to understand the biology of this subtype of breast cancer and develop alternative therapeutic options.

Retinoids are structurally related to the hormone of vitamin A with all-trans-retinoic acid (ATRA) being the active metabolite. Despite the toxicity associated with retinoids, it is in general considered to be well tolerated pharmacological agent. Based on the notion that retinoic acid (RA) promotes cell differentiation, regulates proliferation and apoptosis, it has been combined with anthracycline-based chemotherapy to successfully treat acute promyelocytic leukemia with a success rate of 80% (2). Depending on its interacting partner, RA has distinct biological functions. The ligand binding domain of the retinoic acid receptor (RAR) which includes RARα, RARβ and RARγ and retinoic X receptors RXRα, RXRβ and RXRγ can interact with RA (3,4) and activate genes that are involved in anti-proliferation, apoptosis, differentiation and cell cycle arrest (5-9). However, binding of RA to the alternative nuclear receptor, peroxisome proliferator-activated receptor β/δ (PPARβ/δ) transduces signals which facilitate cell growth, promote cell survival and protect cells against apoptosis (10-13). Delivery of RA to RARs is facilitated by cellular RA-binding protein II (CRABPII), which sequesters RA, RARβ and RARγ and retinoic X receptors RXRα, RXRβ and RXRγ can interact with RA (3,4) and activate genes that are involved in anti-proliferation, apoptosis, differentiation and cell cycle arrest (5-9). However, binding of RA to the alternative nuclear receptor, peroxisome proliferator-activated receptor β/δ (PPARβ/δ) transduces signals which facilitate cell growth, promote cell survival and protect cells against apoptosis (10-13). Delivery of RA to RARs is facilitated by cellular RA-binding protein II (CRABPII), which sequesters RA, translocates to the nucleus, channels RA to RAR and enhances the transcriptional activity of RAR target genes (6,8,14). On the other hand, fatty acid-binding protein 5 (FABP5) can transport RA to its cognate receptor, PPARβ/δ which targets genes that

Correspondence to: Dr Padmamalini Thulasiraman, Department of Biomedical Sciences, College of Allied Health, University of South Alabama, Mobile, AL 36688, USA
E-mail: pthulasiraman@southalabama.edu

Abbreviations: RA, retinoic acid; CRABPII, cellular retinoic acid-binding protein II; RAR, retinoic acid receptor; TNBC, triple-negative breast cancer; PARP, poly(ADP-ribose) polymerase; ER, estrogen receptor; PR, progesterone receptor; HER2, human epidermal growth factor receptor 2; ATRA, all-trans-retinoic acid; PPARβ/δ, peroxisome proliferator-activated receptor β/δ; FABP5, fatty acid-binding protein 5; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; DMEM, Dulbecco's modified Eagle's medium; FBS, fetal bovine serum; qRT-PCR, quantitative real-time polymerase chain reaction; DMSO, dimethyl sulfoxide

Key words: cellular retinoic acid-binding protein II, retinoic acid receptor, breast cancer, curcumin, retinoic acid
are involved in proliferation and cell growth (10-13). Due to the dual and opposing function of RA, the growth inhibitory effects of RA are determined by the expression of the regulatory factors, CRABPII, RARs, FABP5 and PPARβ/δ. The differential effects of retinoids on anti-proliferation, cell cycle arrest and apoptosis is dependent on the concentration of RA and the time frame of RA treatment (15,16). Retinoids repress genes involved in cell division and cell proliferation, which is followed by differentiation, without affecting cell viability. While decrease in proliferation and cell cycle arrest of cancer cells occurs at an earlier time frame of within 2 days, the apoptotic effects of retinoids begin on day 4 of treatment, with increased apoptosis after day 5 or 6 of treatment (15,16).

Recent study has demonstrated the importance of CRABPII in mammary carcinoma tumor growth suppression through RA-dependent and RA-independent mechanism (17). The clinical significance of CRABPII has been highlighted in several types of cancer, including non-small lung cancer (18), non-myeloma skin cancer (19) and pancreatic cancer cells (20), pinpointing that restoring the CRABPII signaling pathway may serve as a therapeutic intervention to ameliorate the efficacy of RA or sensitize cancer cells to this hormone. While CRABPII acts to deliver RA to its RAR receptors, each of the different isoforms of RAR exhibits specific function with each receptor regulating a subset of distinct genes (21-26). RARβ has been implicated in inflammation and tumor suppression (21-23), while the loss of RARγ has been demonstrated in the progression of malignant squamous cell carcinoma (24,25). Selective activation of RARs by retinoids induces autophagy in ER-positive breast cancer cells (26). Though there may be alternative RA-resistant mechanisms in cancer, the loss of sensitivity to RA in cancers has been related to the lack of CRABPII and RAR expression with increased expression of FABP5 and PPARβ/δ (11,19,27,28).

Phytochemicals have been extensively studied for the treatment of diseases such as cancer. Curcumin, an active ingredient in the dietary spice turmeric (Curcuma Longa), possesses antioxidant, anti-inflammatory and anticancer properties (29,30). The apoptotic effects of curcumin have been observed in several cancers including breast (31), pancreatic (32), prostate (33) and lung (34), while having no cytotoxic effects on healthy cells (35). Despite its low solubility and bioavailability, the combination of curcumin with conventional chemotherapeutic agents has been demonstrated to be effective in cancer regression (36,37). Synthesis of nanocarriers has not only increased the solubility in aqueous solution but has also improved the bioavailability of curcumin towards cancer cells (38-40). Several studies have demonstrated the ability of curcumin to increase the sensitivity of cancer cells to chemotherapeutic drugs (41,42). Through multiple mechanisms, curcumin and its analogs sensitize cancer cells to chemotherapeutic agents, thus overcoming drug resistance and improving susceptibility to growth suppression by conventional drug treatments (43-47).

Previously, we reported that 30 µM curcumin sensitizes TNBC cells to RA-mediated growth suppression by altering the expression level of FABP5/PPARβ/δ pathway and targeting PPARβ/δ target genes (45). However, we did not explore the role of curcumin on the second arm of the retinoid pathway, namely CRABPII/RAR, and in this study we evaluated the hypothesis that curcumin mediated upregulation of the CRABPII/RAR pathway in TNBC cells promotes sensitivity to RA-mediated apoptosis. The overall aim of this study is to investigate the effect of curcumin on the CRABPII/RAR pathway and assess the contribution of this pathway in sensitizing TNBC to RA-mediated growth suppression by triggering cell death. In order to achieve this aim, we examine whether regulation of CRABPII/RAR by curcumin is dose-dependent and the functional consequence of this pathway in increasing sensitivity to RA. Furthermore, we explored the impact of silencing CRABPII on apoptosis facilitated by curcumin and/or RA. In the present study, we demonstrate that while 30 µM curcumin induces CRABPII, it does not affect RARs in TNBC cells. However, lower doses of curcumin (5 and 10 µM) upregulate CRABPII, RARβ and RARγ expression in these cells. We also provide evidence that induction of the CRABPII/RAR pathway by curcumin sensitizes RA-resistant TNBC cells to RA-mediated apoptosis, and knockdown of CRABPII in TNBC cell lines reverses the sensitization of the apoptotic effects of RA by curcumin.

Materials and methods

Reagents. Antibodies for RARα, RARβ, RARγ and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) were obtained from Abcam (Cambridge, MA, USA). Anti-CRABPII was purchased from R&D Systems (Minneapolis, MN, USA). Caspase-9 and poly(ADP-ribose) polymerase (PARP) antibodies were obtained from Cell Signaling Technology, Inc. (Danvers, MA, USA). Anti-mouse and anti-rabbit immunoglobulin was from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA). Curcumin (C-1386) and ATRA (R-2625) were purchased from Sigma-Aldrich (St. Louis, MO, USA). 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenylenetrazolium bromide (MTT) reagent was purchased from Sigma-Aldrich. CRABPII and control siRNA were obtained from Santa Cruz Biotechnology, Inc.

Cell lines. MDA-MB-231 and MDA-MB-468 cells were maintained in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum (FBS) with antibiotics. MDA-MB-231 cells were a kind gift from Dr Ming Tan (Mitchell Cancer Institute, Mobile, AL, USA). MDA-MB-468 was purchased from American Type Culture Collection (ATCC; Manassas, VA, USA).

Western blot analyses. Cells were cultured in 100-mm plates and treated with curcumin and/or ATRA in media containing 10% charcoal treated FBS for the indicated time. Cells were lysed in a buffer containing 150 mM NaCl, 10 mM Tris, pH 7.2, 0.1% SDS, 1% Triton X-100, 1% deoxycholate, 5 mM EDTA and 1 mM PMSF for 1 h. The concentration of the whole cell protein was determined using the Bradford assay. Cell lysate was resolved by SDS-PAGE and transferred onto a nitrocellulose membrane. The membrane was blocked in 10% bovine serum albumin (BSA) in Tris-buffered saline containing 0.05% Tween-20 for 1 h at room temperature. The membrane was then probed using the appropriate antibody, CRABPII,
RARα, RARβ, RARγ, PARP or caspase-9 at a dilution of 1:1,000 overnight at 4˚C. After a 30 min wash, the appropriate secondary antibody was added for 1 h at room temperature prior to exposure. Anti-GAPDH (1:2,000) was used as a loading control and was incubated with the membrane for 1 h at 4˚C. The antigen-antibody complex was visualized with SuperSignal West Pico Chemiluminescent Substrate (Thermo Fisher Scientific, Inc., Hanover Park, IL, USA).

Quantitative real-time polymerase chain reaction (qRT-PCR). Cells were treated with curcumin for 4 h, and RNA was extracted using TRIzol (Life Technologies, Grand Island, NY, USA). As outlined in the protocol for the high-capacity RNA to cDNA kit from Applied Biosystems (Foster City, CA, USA), 2 µg total RNA was reverse transcribed into cDNA. To determine expression of CRABPII, RARα, RARβ and RARγ, qRT-PCR was carried out by using commercially available TaqMan chemistry and assay on demand probes (Applied Biosystems). GAPDH was used for normalization. Detection and data analysis were carried out on the ABI StepOnePlus Real-Time PCR system. Relative quantity of gene expression was performed using 2-ΔΔCt method (48).

Cell viability assay. MDA-MB-231 and MDA-MB-468 cells (5,000 cells/well) were cultured in media containing 10% charcoal treated FBS and plated in a 96-well plate overnight. Cells were then treated with curcumin in the presence or absence of ATRA for 72 h. Controls used were DMSO and/or ethanol depending on the drug treatment. After 72 h, 5 µg/ml of MTT reagent was added directly to the cells for 3 h and allowed to incubate at 37˚C. The media was removed from the plate and the intact cells were resuspended in 150 µl of 0.04 M HCl in isopropanol. The cells were placed in the incubator for 15 min to solubilize the cells, and were then mixed completely. Absorbance was read at 570 nm to determine cell proliferation.

Cell transfection. MDA-MB-231 and MDA-MB-468 cells were transfected with CRABPII siRNA according to the protocol from Invitrogen Life Technologies (Grand Island, NY, USA). Briefly, 30 pmol of control or CRABPII siRNA was mixed with 500 µl Gibco Opti-MEM® I medium without serum by Life Technologies in a 6-well plate. Lipofectamine™ RNAiMAX (5 µl) was added to the diluted RNAi molecules for 20 min at room temperature. MDA-MB-231 and MDA-MB-468 (250,000 cells) were diluted in 2.5 ml of antibiotics-free media and added to the plates containing the RNAi duplex/Lipofectamine™ mixture labeled with control or CRABPII siRNA. After 24 h, the cells were treated with 10 µM curcumin in the presence or absence of 1 µM ATRA for 96 h. ATRA was added every 2 days. Cells were then lysed, protein was extracted, quantitated and protein extract was loaded onto a gel for SDS-PAGE. The blot was probed with the appropriate antibodies.

Statistical analysis. Statistical significance of differences between treatments was determined using two-tailed Student's t-test and P-values were noted. Differences between groups were considered statistically significant at p<0.05.

Results
Concentration-dependent impact of curcumin on CRABPII and RAR expression in TNBC cell lines. Knowing that TNBC cells are resistant to RA due to the expression patterns of CRABPII and FABP5 (11,20,49), our previous study demonstrated that 30 µM curcumin reduced the expression of FABP5 and its cognitive receptor, PPARβ/δ (45). In this study, we sought to examine the effect of curcumin on the CRABPII/RAR pathway in TNBC cells. As shown in Fig. 1A, 30 µM curcumin enhanced the CRABPII mRNA expression level compared to control (p=0.016) in MDA-MB-231 cells, however, had no effect on the RAR isoforms (data not shown).
Since CRABPII delivers RA to RAR isoforms, we tested whether the induction of CRABPII was sufficient to sensitize RA-resistant MDA-MB-231 cells to RA-mediated apoptosis. To investigate cell death, we tested PARP, an indicator of apoptosis and RAR target, caspase-9. As shown in Fig. 1B, treatment of MDA-MB-231 cells with 30 µm curcumin for 48 h completely converted full length PARP to cleaved PARP, and thus there was no effect with the combination of curcumin and RA. Examination of the caspase-9 showed there was no difference in the expression of cleaved caspase-9 in cells treated with curcumin compared to the co-treatment with curcumin and ATRA (Fig. 1B). Despite upregulation of CRABPII by 30 µm curcumin, this dose of curcumin induced apoptosis in 48 h, independent of the CRABPII pathway.

Because 30 µm curcumin is a high-dose which alone induces apoptosis within 48 h, we examined the expression level of CRABPII and RAR isoforms at lower doses of curcumin (5 and 10 µM). Compared to control, 5 and 10 µM curcumin induced CRABPII mRNA expression, with statistical significance (p<0.05), in MDA-MB-231 cells by ~3-3.5-fold (Fig. 2A). Concomitantly, we examined the effect of lower doses of curcumin on CRABPII protein expression. Consistent with the upregulation of CRABPII mRNA expression by 5 and 10 µM curcumin, we also observed that at both of these doses, curcumin induced CRABPII protein expression in MDA-MB-231 cells (Fig. 2B). Because curcumin affects CRABPII expression in RA-resistant MDA-MB-231 cells, we further tested whether curcumin regulates CRABPII mRNA and protein expression in the TNBC cell line, MDA-MB-468. As shown in Fig. 2C, both of the lower doses of curcumin, in comparison to control, upregulated the expression of the CRABPII mRNA (p<0.05). Correspondingly, curcumin induced CRABPII protein expression level in MDA-MB-468 (Fig. 2D).

We next investigated whether 5 and 10 µm curcumin had an effect on the cognate receptors of CRABPII, RAR isoforms α, β and γ. At both of these doses, RARβ and RARγ protein expression were upregulated in MDA-MB-231 cells compared to their respective controls (Fig. 3A and B). To determine whether regulation of RARβ and RARγ by curcumin is a global effect among TNBC cells, treatment of MDA-MB-468 cells with 5 and 10 µM curcumin induced both RARβ and RARγ protein expression (Fig. 3C and D). Interestingly, we observed that curcumin did not affect the expression of RARα in either of the TNBC cell lines, MDA-MB-231 and MDA-MB-468 (data not shown). These results suggest that lower doses of 5 and 10 µM curcumin enhance RARβ and RARγ protein expression in TNBC cells.

Lower doses of curcumin sensitize TNBC cells to RA-mediated apoptosis. Suppression of cell growth by RA is followed by RA-induced apoptosis. Previously, we showed that treatment
of TNBC cells with 30 µM curcumin for 48 h sensitized these cells to RA-mediated growth suppression by reducing the expression level of the FABP5/PPARβ/δ pathway (45). In this study, we observed that lower doses of curcumin (5 and 10 µM) induced the expression of CRABPII, RARβ and RARγ (Figs. 2 and 3). To determine the effect of lower doses of curcumin on sensitizing TNBC cells to RA-mediated apoptosis, we treated MDA-MB-231 and MDA-MB-468 cells with 5 and 10 µM curcumin in the presence or absence of 1 µM ATRA for 72 h. As expected, RA did not inhibit cell proliferation, but it sensitized cells to RA-mediated apoptosis (Fig. 4).
growth in MDA-MB-231 cells (Fig. 4A). While the lower concentration of curcumin (5 µM) did not suppress growth of MDA-MB-231 cells after 72 h, 10 µM curcumin did reduce growth compared to control, with statistical significance (p=0.03) by ~20% (Fig. 4A). Combining either 5 or 10 µM curcumin with ATRA sensitized MDA-MB-231 cells to RA-mediated growth suppression, with a robust effect with 10 µM curcumin and ATRA (Fig. 4A). Similar findings were concluded in MDA-MB-468 cells. RA treatment had marginal growth inhibitory effects on MDA-MB-468 cells (p<0.05). In comparison to control treated cells, lower doses of curcumin (5 and 10 µM) statistically reduced growth of MDA-MB-468 cells (p<0.05) but the combination of curcumin with RA sensitized MDA-MB-468 cells to RA-induced growth suppression, with a more pronounced effect with 10 µM curcumin (Fig. 4B).

Knowing that 10 µM curcumin had a greater impact on sensitization of TNBC cells to RA-mediated growth suppression at 72 h (Fig. 4A and B), and the fact that 10 µM curcumin induced CRABPII, RARβ and RARγ expression in these cells (Figs. 2 and 3), we tested whether this dose of curcumin could sensitize TNBC cells to RA-mediated apoptosis. Because the apoptotic effects of RA occur subsequent to growth suppression, we examined cell death by curcumin and RA at a later time-point. Cells were treated with 10 µM curcumin in the presence or absence of 1 µM ATRA for 96 h and examined for the protein expression of cleaved caspase-9. MDA-MB-231 transfected with control siRNA was also treated with curcumin and as expected, curcumin induced apoptosis by activation of caspase-9 (37 kDa), which was further enhanced in the presence of RA (Fig. 5A). However, when the cells were transfected with CRABPII siRNA, the combination of curcumin and RA abolished the active 37 kDa form of caspase-9 (Fig. 5A). These results suggest that curcumin mediated upregulation of CRABPII sensitizes TNBC cells to RA-mediated apoptosis by the induction of caspase-9, one of the RAR target genes. We also silenced CRABPII in MDA-MB-468 cells and combinatorial treatment of the cells with curcumin and RA enhanced expression of cleaved/active caspase-9 (37 and 35 kDa), in comparison to curcumin alone (Fig. 5B). However, knockdown of CRABPII reduced active caspase-9 (35 kDa) and abolished the expression of the cleaved/active 37 kDa caspase-9 (Fig. 5B).

Discussion

Sensitivity of cancer cells to RA is determined by two distinct pathway, CRABPII/RAR and FABP5/PPARβ/δ (10,11,14, 20,49). While the delivery of RA to the FABP5/PPARβ/δ pathway results in increased proliferation, transporting RA by CRABPII to RAR inhibits proliferation and promotes apoptosis (10,11,14). Reducing the ratio of FABP5/CRABPII can overcome the resistance of cancer cells to RA by shifting the delivery of RA to RAR by CRABPII instead of activating the FABP5/PPARβ/δ pathway (10,11,14).

Phytochemicals such as curcumin exhibit chemosensitizing properties which may circumvent toxicity issues faced with traditional chemotherapeutic agents (50). While they are less toxic and have no side effects, combinatorial treatment...
of phytochemicals with chemotherapeutic agents can be an alternative method to reduce the dosage of traditional chemotherapeutic regimens and lower cardiac toxicity associated with them (50). Owing to drug resistance in cancer, curcumin has been shown to be an effective adjuvant in reversing chemoresistance and sensitizing cancer cells to chemotherapeutic drugs (41-47,51-54).

Our previous study reported that the potential therapeutic activity of curcumin to sensitize RA-resistant TNBC cells to RA-mediated growth suppression was exhibited through the inhibition of the FABP5/PPARβ/δ pathway (45). The present study focused on the effect of curcumin on the CRABPII/RAR pathway to reverse resistance of TNBC cells to RA by activation of apoptosis. Results of the present study demonstrate that low concentrations of 5 and 10 µM curcumin increase the expression level of CRABPII, while concomitantly induces expression of its cognate receptor, RARβ and RARγ. Similarly, a recent study demonstrated that curcumin upregulated the expression of RARβ in several cancer cells, including TNBC MDA-MB-231 cells and activated RARβ via epigenetic regulation (22). Though the RAR subtypes have different functions and regulate distinct RARE targeted genes, RAR signaling plays a critical role in cancer progression. While RARβ is epigenetically silenced in tumor cells and its loss has been associated with lymph node metastasis (21,55,56), RARγ has been implicated as a tumor suppressor and restoration of this retinoid receptor reverses the tumorigenic potential of mouse keratinocytes (57,58). While curcumin induces the expression of several regulatory genes involved in apoptosis (59), we present evidence in this study that curcumin upregulates genes involved in the retinoid signaling pathway, namely CRABPII, RARβ, and RARγ in mammary carcinoma MDA-MB-231 and MDA-MB-468 cells.

Although 30 µM of curcumin induces the mRNA expression level of CRABPII, it did not affect the expression of RARs. However, lower doses of curcumin (5 and 10 µM) induces the protein expression level of not only CRABPII but also RARβ and RARγ in two TNBC cell lines suggesting that this is a global effect among TNBC cells. Alterations in gene expression associated with different doses of curcumin are not unusual and have been well documented with other genes (60). The differential effects of curcumin have been observed with several apoptotic genes that are upregulated or downregulated by only higher dose of curcumin, while the reverse has also been observed for some genes that are upregulated only at lower doses of curcumin (60). As noted in this study, the differences in concentration of curcumin to induce the CRABPII/RAR pathway may be an important determinant in identifying the required optimum dosage to sensitize TNBC cells to RA for future in vivo studies.

The concentration-dependent regulation of CRABPII and RARs by curcumin determined the outcome on the activation of apoptotic proteins, PARP and caspase-9. Although 30 µM curcumin upregulates the mRNA level of CRABPII, this dose of curcumin completely activates PARP and induces activation of caspase-9 in 48 h. However, the fact that 30 µM curcumin does not regulate the RARs indicates that this concentration of curcumin induces apoptosis independent of the CRABPII/RAR and 30 µM curcumin does not sensitize MDA-MB-231 cells to RA-induced apoptosis.

Curcumin has a differential effect on gene regulation and cell death initiated by this agent dose- and time-dependently (60,61). The present data demonstrates that curcumin can re-activate the CRABPII/RAR pathway in TNBC cells and cause RA to initiate apoptosis by activation of PARP and caspase-9. Such doses of curcumin (5 and 10 µM) upregulates RARβ and RARγ, as well as CRABPII in TNBC cells. Combination of 10 µM curcumin with RA for 96 h sensitizes TNBC cells to apoptosis mediated by RA as evidenced by increased PARP cleavage. Because 10 µM curcumin induces RARβ and RARγ in TNBC cells, curcumin at this dose sensitizes the cells to RA-mediated apoptosis through RAR-dependent activation of caspase-9. Initiation of cell death by RA itself is not sufficient to regulate apoptosis by RA, and hence shutting of RA from the cytosol to the nucleus by CRABPII facilitates binding of RA to RARs and enhances the transcriptional activation of genes such as caspase-9 involved in the retinoid signaling pathway. To extend these studies and gain a mechanistic understanding on the role of curcumin on the CRABPII/RAR pathway, our results also provide evidence that silencing CRABPII prevents curcumin from sensitizing TNBC cells to RA-induced activation of caspase-9. Taken together, our data suggest that in order to activate cell death by RA in RA-resistant TNBC cells, CRABPII and RAR pathway have to be upregulated by lower concentrations of curcumin and these two proteins work in concert to sensitize cells to RA-mediated apoptosis.

In conclusion, the present study revealed that reversing the resistance of TNBC to RA-induced apoptosis is dependent on the dose of curcumin and length of treatment. Accordingly, lower concentrations of curcumin induce CRABPII, RARβ and RARγ, and thus upregulation of CRABPII/RAR pathway contributes to the sensitization of TNBC cells to apoptosis by RA. As such this study highlights a novel mechanism by which RA-resistant mammary carcinoma cells can be resensitized to RA-mediated apoptosis by curcumin. The effectiveness in the combination of curcumin with RA warrants further consideration for its use in RA-resistant TNBC cells. Overall, this study provides mechanistic insights on the role of curcumin to reverse RA resistance in breast cancer cells through the regulation of the CRABPII/RAR pathway, and highlights the potential of using curcumin as a therapeutic adjuvant in RA resistant cancers.

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